

Overview of Global Warming, Ozone Depletion, and Air Quality

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>




<https://www.videoblocks.com/video/earth-sunset-spacewalk-view-from-space-station-r7dydlcsgjd23vml0>

Lecture 2

1 February 2022

Announcements

1) AOSC Weekly Seminar Feb 3: 3:30 pm (Thursday)




DEPARTMENT OF
ATMOSPHERIC &
OCEANIC SCIENCE

DR. RYAN STAUFFER

INSTITUTION: NASA/GSFC CHEMISTRY AND ATMOSPHERIC DYNAMICS
LABORATORY

**TITLE: 'SOUTHERN HEMISPHERE ADDITIONAL OZONESONDES
(SHADOZ): ACCOMPLISHMENTS FROM 25 YEARS OF TROPICAL AND
SUBTROPICAL OZONE PROFILING'**

Abstract: Ozone (O₃) is a vital trace gas in Earth's atmosphere. It shields life from harmful UV radiation, but is a strong oxidant and harmful pollutant when found in high quantities near the surface. Ozone is also a greenhouse gas in the troposphere, so characterizing the vertical distribution of O₃ is critical to determining its role in climate and effects on air quality. Satellite observations give us excellent spatial coverage of the total column O₃ in the atmosphere, but are unable to resolve the O₃ vertical profile with accuracy, especially below the stratosphere. Balloon-borne ozonesondes bridge this gap by providing high vertical resolution measurements from the surface to over 30 km altitude. Before the late 1990s, ozonesonde measurements in the tropics were sparse, so the Southern Hemisphere Additional Ozonesondes (SHADOZ) strategic network organized over a dozen stations to frequently launch ozonesondes for pollution and climate studies. This talk features the history and accomplishments of SHADOZ from 1998-2021 including work on satellite and model validation, biomass burning, weather and climate links to O₃, and O₃ trends, as well as data reprocessing and quality assurance activities with global partners.



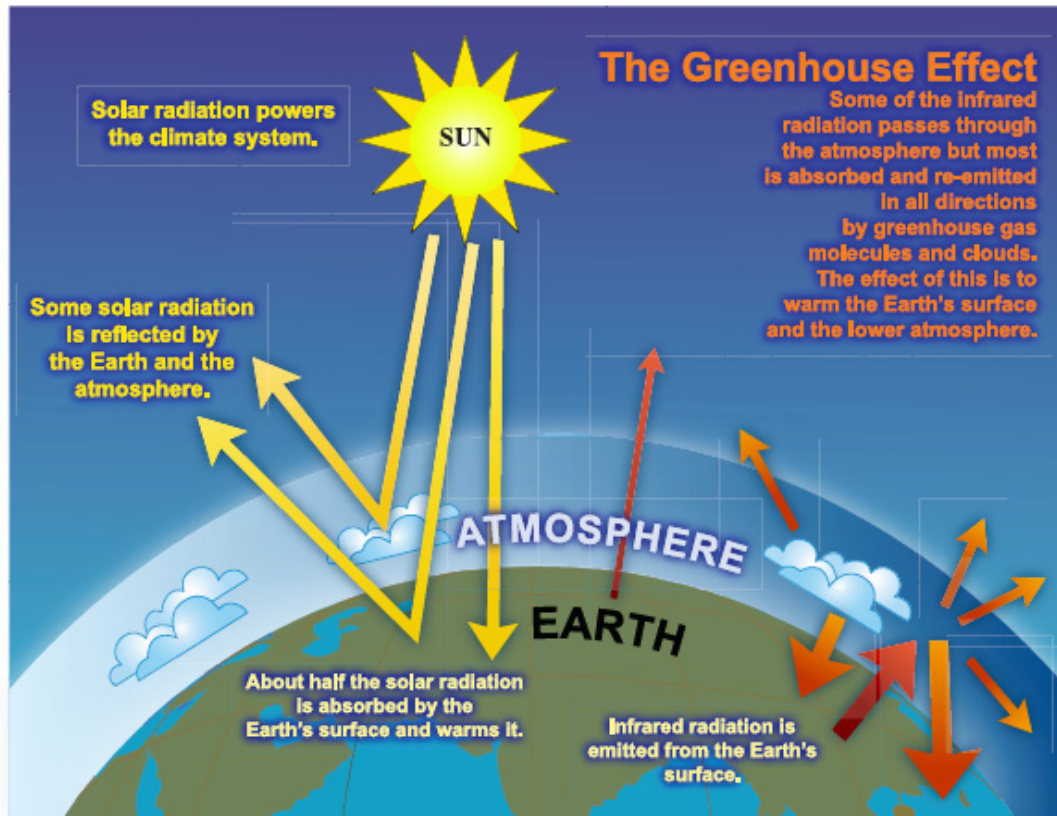
**3:30 PM EST
THURSDAY, FEBRUARY 3, 2022**

PRESENTED IN HYBRID VIRTUAL/IN-PERSON FORMAT
SEMINAR LOCATION: ATLANTIC BUILDING, ROOM 2400
CONTACT: JACOB WENEGRAT (WENEGRAT@UMD.EDU) OR JOSEPH KNISELY (JKNISELY@UMD.EDU)

<https://aosc.umd.edu/seminars>

2) Please have a calculator available for class on Thursday

Greenhouse Effect



FAQ 1.3, Figure 1. An idealised model of the natural greenhouse effect. See text for explanation.

Question 1.3, IPCC, 2007

Radiative Forcing of Climate, 1750 to 2019

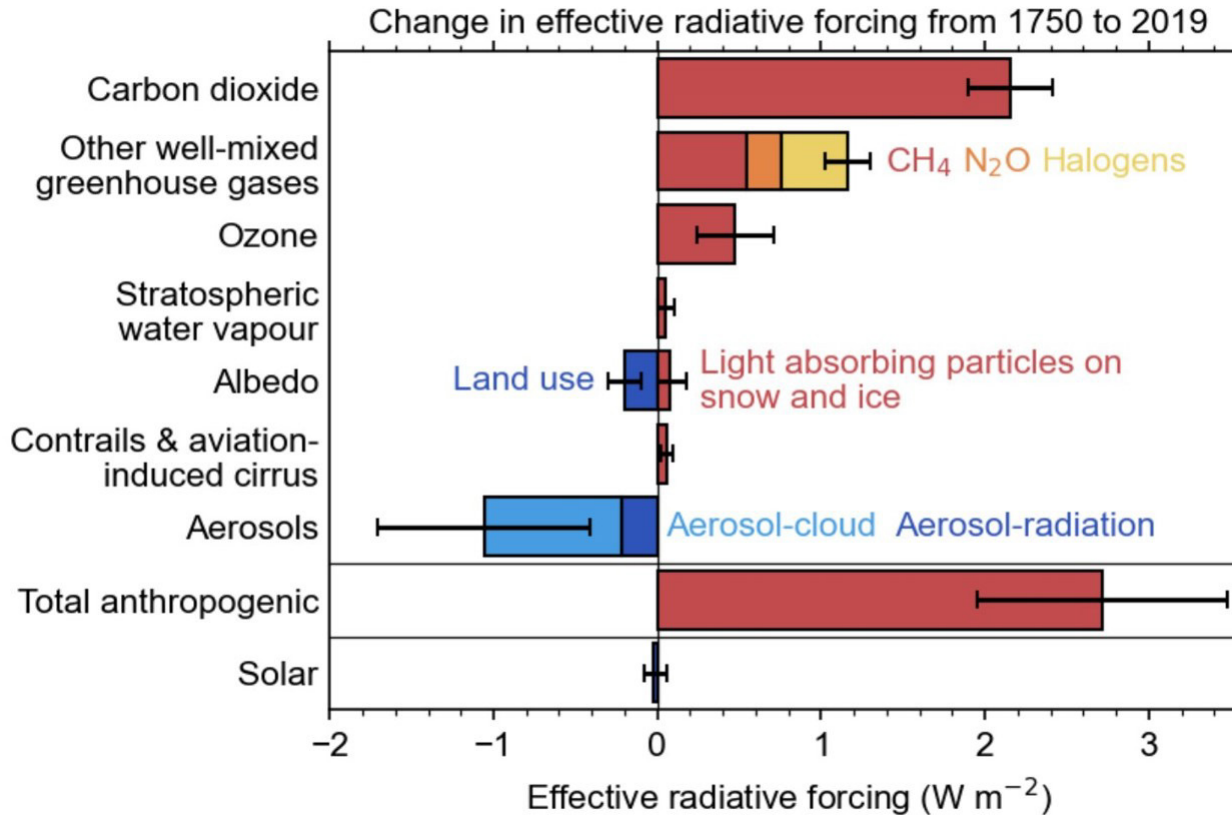
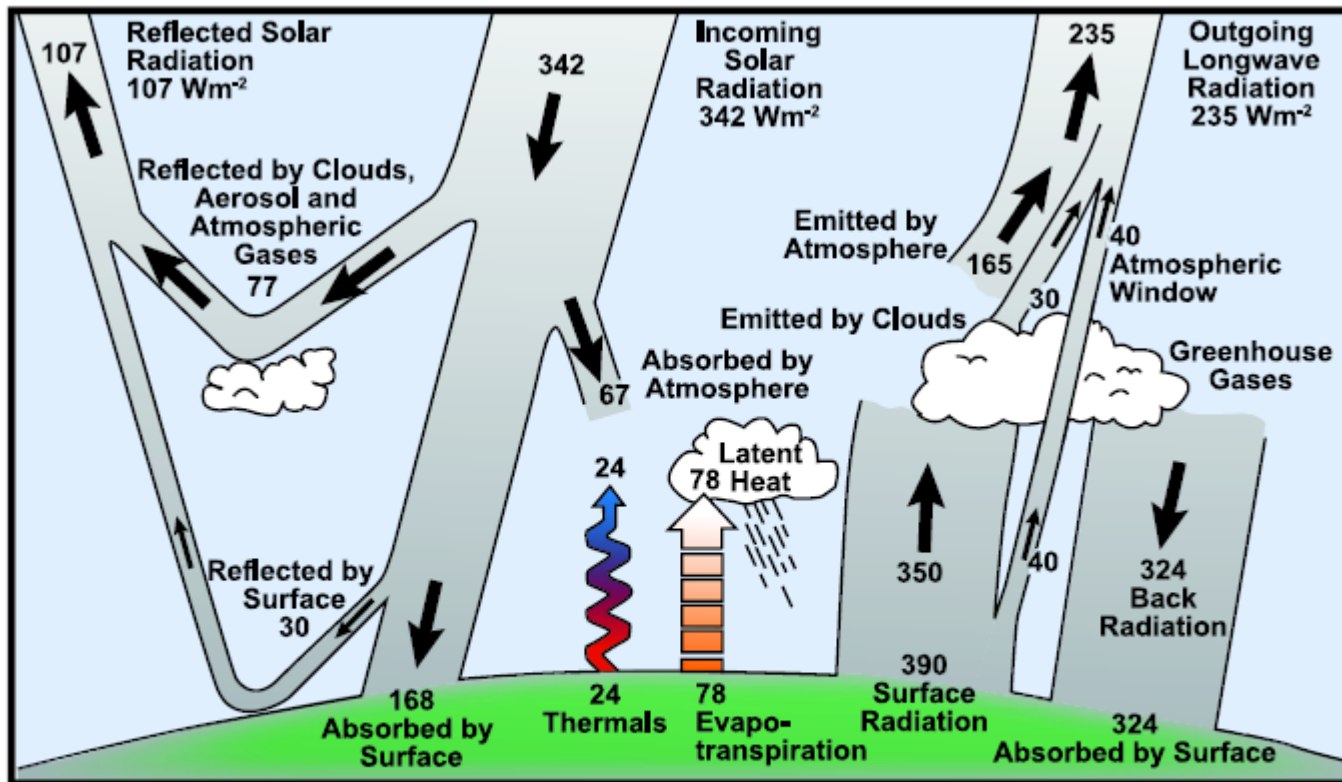


Figure 7.6, IPCC (2021)

https://www.ipcc.ch/report/ar6/wg1/downloads/report/IPCC_AR6_WGI_TS.pdf

Radiative Forcing



FAQ 1.1, Figure 1. Estimate of the Earth's annual and global mean energy balance. Over the long term, the amount of incoming solar radiation absorbed by the Earth and atmosphere is balanced by the Earth and atmosphere releasing the same amount of outgoing longwave radiation. About half of the incoming solar radiation is absorbed by the Earth's surface. This energy is transferred to the atmosphere by warming the air in contact with the surface (thermals), by evapotranspiration and by longwave radiation that is absorbed by clouds and greenhouse gases. The atmosphere in turn radiates longwave energy back to Earth as well as out to space. Source: Kiehl and Trenberth (1997).

Question 1.1, IPCC, 2007

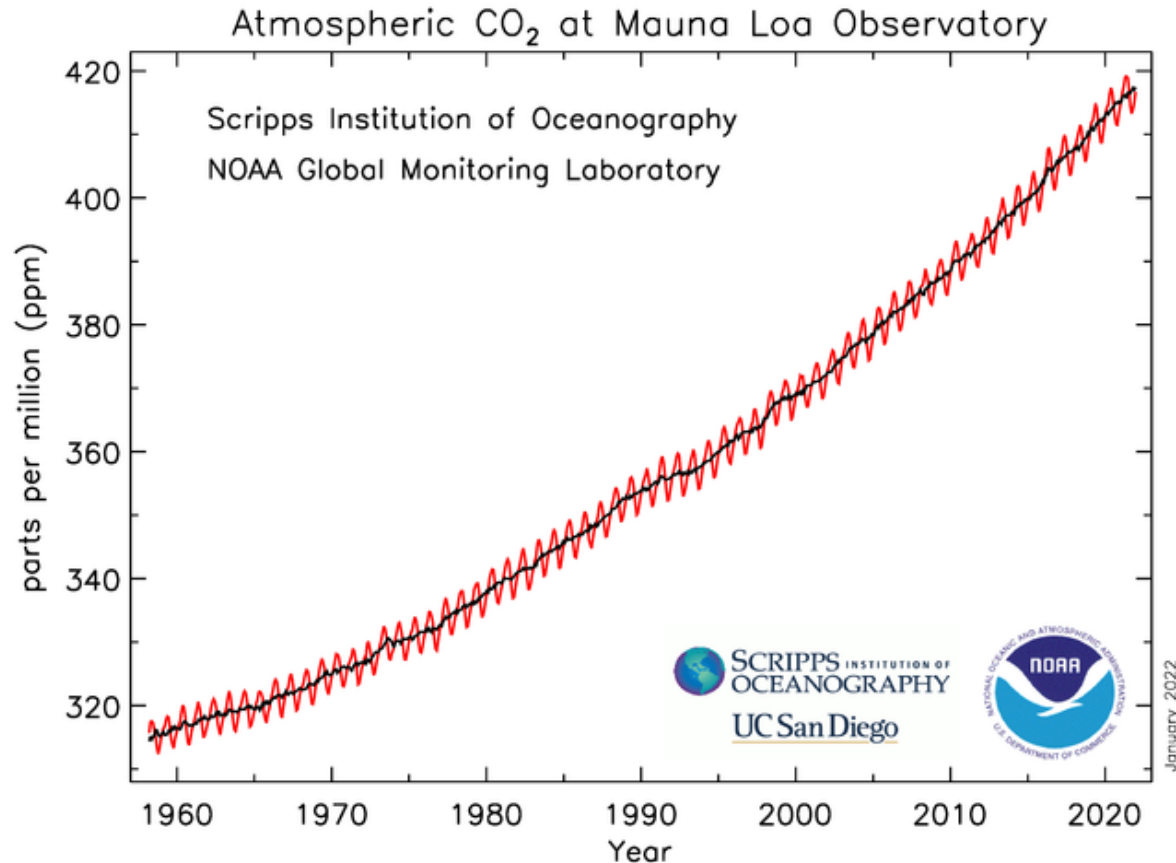
Radiative Forcing of Climate is Change in Energy
reaching the lower atmosphere (surface to tropopause) as GHGs rise.
“Back Radiation” is most important term.

Modern CO₂ Record

Global Mean on 30 Jan 2022: 419.41 parts per million (ppm)

30 Jan 2021: 416.19 parts per million (ppm)

Annual Rise about 3.2 ppm (0.77%)

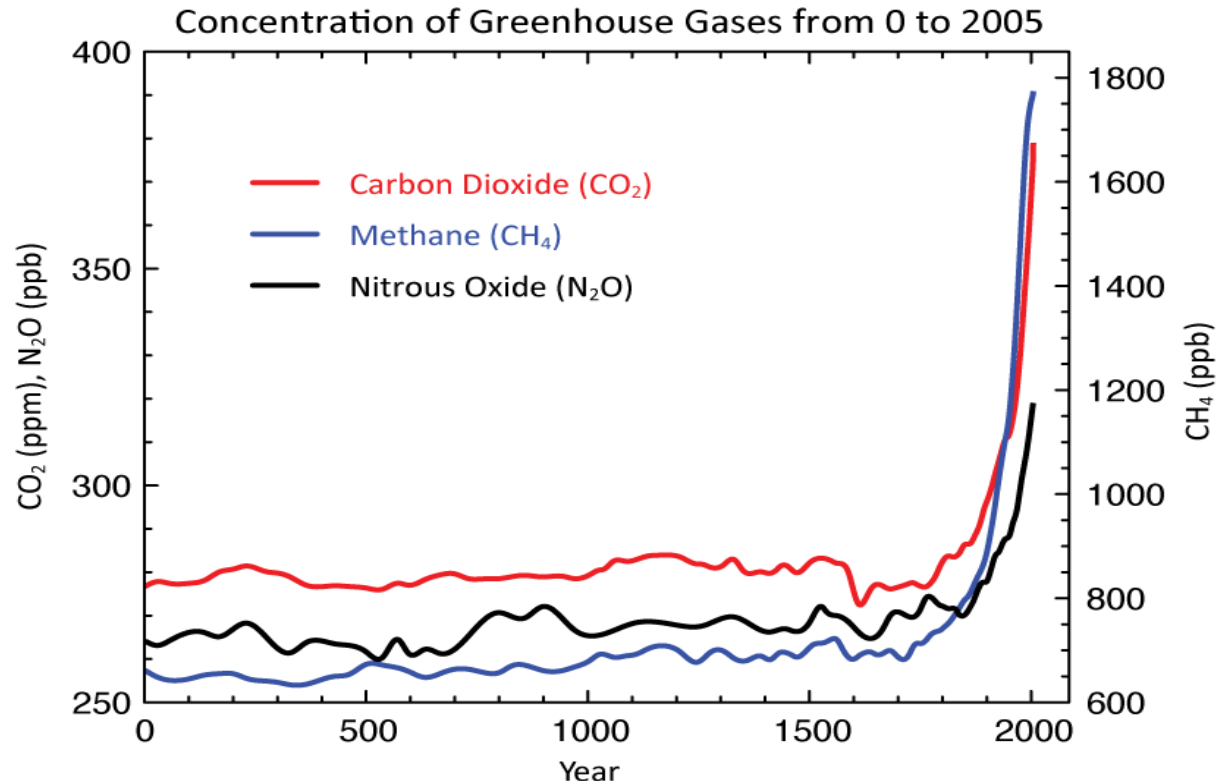


Legacy of Charles Keeling, Scripps Institution of Oceanography, La Jolla, CA

https://www.esrl.noaa.gov/gmd/webdata/ccgg/trends/co2_data_mlo.png

See also <https://www.co2.earth/daily-co2>

GHG Record Over Last Several Millennia



FAQ 2.1, Figure 1 (Errata). *Revised figure showing atmospheric concentrations of important long-lived greenhouse gases over the last 2,000 years. Using the combined and simplified data from Chapters 6 and 2, the original figure displayed the CH₄ curve incorrectly. The revised figure shows the same data correctly plotted. For further details please refer to the original figure caption.*

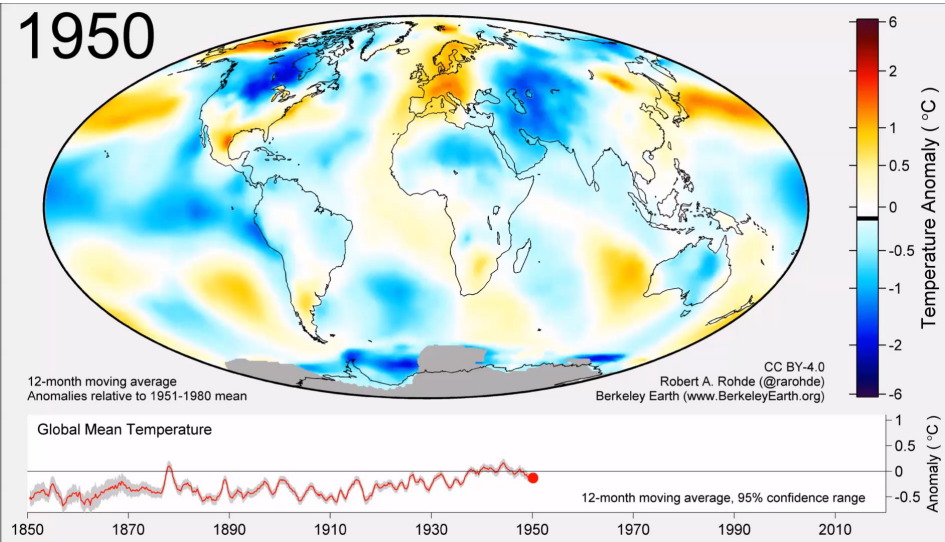
Question 2.1, IPCC, 2007 ... corrected

<https://www.ipcc.ch/site/assets/uploads/2018/05/ar4-wg1-errata.pdf>

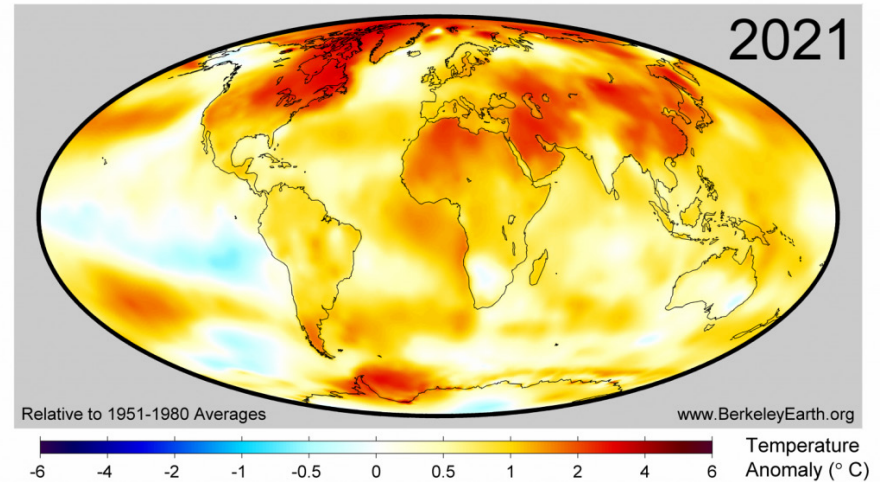
Correction issued upon realization the line for CH₄ had been plotted incorrectly

Berkeley Earth Animation of Global Warming

1950



2021



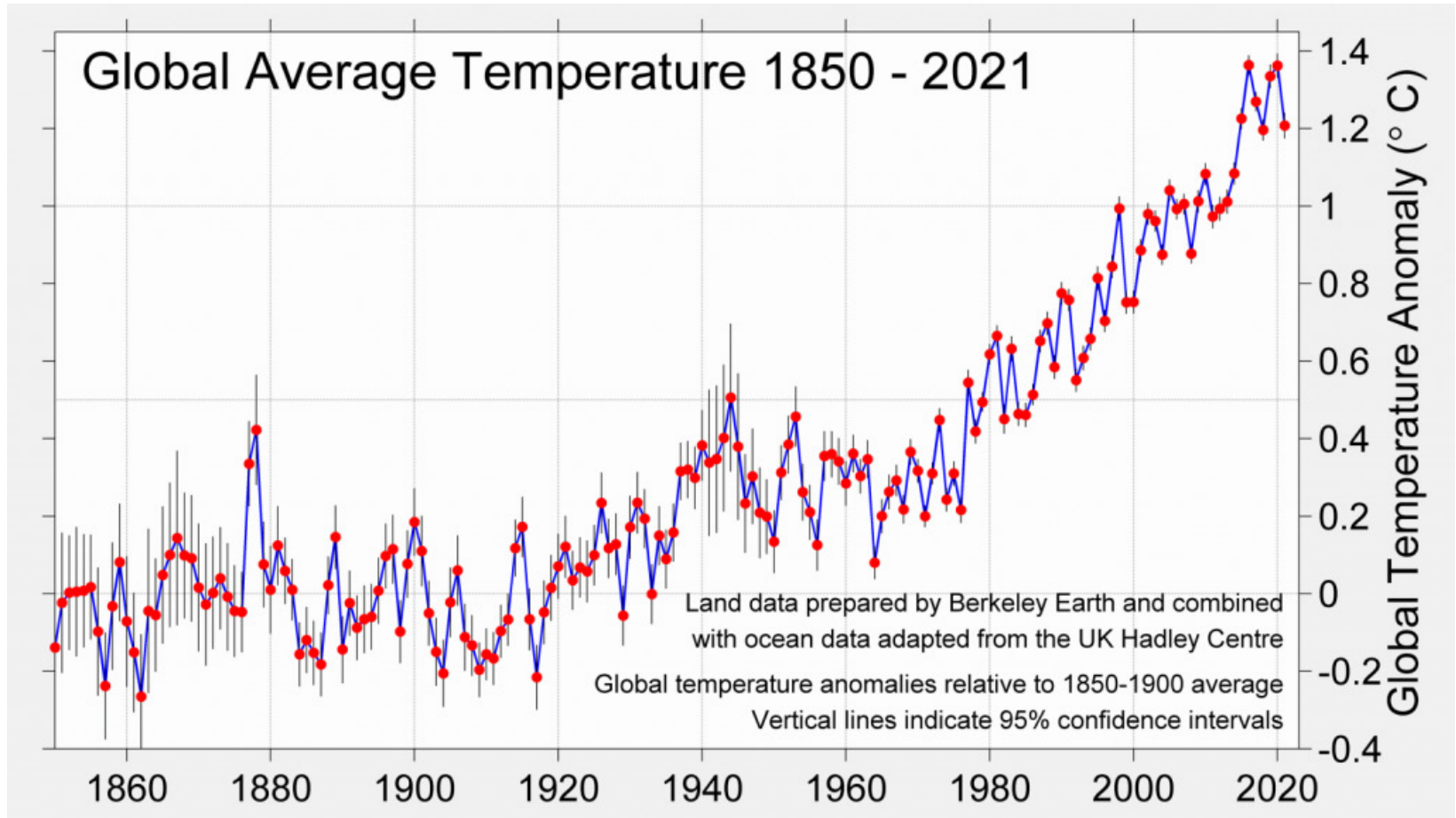
1°C (Celsius) warming is equivalent to 1.8°F (Fahrenheit) warming

Work of Robert Rohde and the Berkeley Earth Team

<http://berkeleyearth.org>

Animation at <https://twitter.com/RARohde/status/1217496115429494786>

Berkeley Earth Animation of Global Warming



The global mean temperature in 2021 was about 1.2°C above the average temperature from 1850-1900, a period often used as a pre-industrial baseline for global temperature targets. Nominally, 2021 was the 6th warmest year to have been directly observed, though 2015, 2018, and 2021 all cluster closely together relative to their uncertainty estimates.

Hence, the 5th, 6th, and 7th warmest years are all essentially tied.

Work of Robert Rohde and the Berkeley Earth Team

<http://berkeleyearth.org>

<http://berkeleyearth.org/global-temperature-report-for-2021/>

GWP – Global Warming Potential

$$\text{GWP}(\text{CH}_4) = \frac{\int_{\text{time initial}}^{\text{time final}} a_{\text{CH}_4} \times [\text{CH}_4(t)] dt}{\int_{\text{time initial}}^{\text{time final}} a_{\text{CO}_2} \times [\text{CO}_2(t)] dt}$$

where:

a_{CH_4} = Radiative Efficiency ($\text{W m}^{-2} \text{ kg}^{-1}$) due to an increase in CH_4

a_{CO_2} = Radiative Efficiency ($\text{W m}^{-2} \text{ kg}^{-1}$) due to an increase in CO_2

$\text{CH}_4(t)$ = time-dependent response to an instantaneous release of a pulse of CH_4

$\text{CO}_2(t)$ = time-dependent response to an instantaneous release of a pulse of CO_2

$$\text{GWP}(\text{N}_2\text{O}) = \frac{\int_{\text{time initial}}^{\text{time final}} a_{\text{N}_2\text{O}} \times [\text{N}_2\text{O}(t)] dt}{\int_{\text{time initial}}^{\text{time final}} a_{\text{CO}_2} \times [\text{CO}_2(t)] dt}$$

GWP – Global Warming Potential

Table TS.2. Lifetimes, radiative efficiencies and direct (except for CH₄) global warming potentials (GWP) relative to CO₂. {Table 2.14}

Industrial Designation or Common Name (years)	Chemical Formula	Lifetime (years)	Radiative Efficiency (W m ⁻² ppb ⁻¹)	Global Warming Potential for Given Time Horizon			
				SAR† (100-yr)	20-yr	100-yr	500-yr
Carbon dioxide	CO ₂	See below ^a	^b 1.4x10 ⁻⁵	1	1	1	1
Methane ^c	CH ₄	12 ^c	3.7x10 ⁻⁴	21	72	25	7.6
Nitrous oxide	N ₂ O	114	3.03x10 ⁻³	310	289	298	153

Notes:

† SAR refers to the IPCC Second Assessment Report (1995) used for reporting under the UNFCCC.

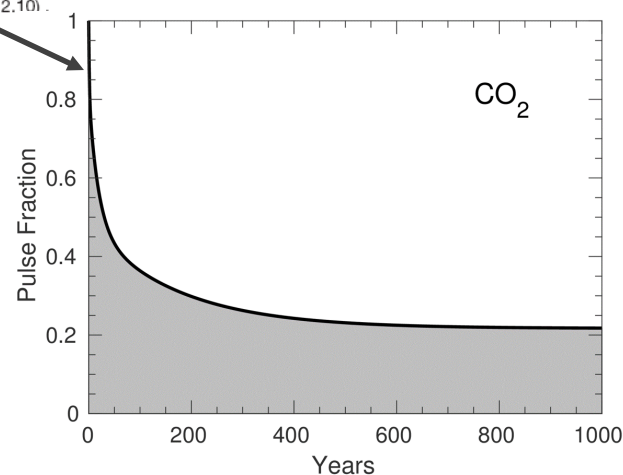
^a The CO₂ response function used in this report is based on the revised version of the Bern Carbon cycle model used in Chapter 10 of this report (Bern2.5CC; Joos et al. 2001) using a background CO₂ concentration value of 378 ppm. The decay of a pulse of CO₂ with time *t* is given by

$$a_0 + \sum_{i=1}^3 a_i \cdot e^{-t/\tau_i} \quad \text{where } a_0 = 0.217, a_1 = 0.259, a_2 = 0.338, a_3 = 0.186, \tau_1 = 172.9 \text{ years}, \tau_2 = 18.51 \text{ years}, \text{ and } \tau_3 = 1.186 \text{ years, for } t < 1,000 \text{ years.}$$

^b The radiative efficiency of CO₂ is calculated using the IPCC (1990) simplified expression as revised in the TAR, with an updated background concentration value of 378 ppm and a perturbation of +1 ppm (see Section 2.10.2).

^c The perturbation lifetime for CH₄ is 12 years as in the TAR (see also Section 7.4). The GWP for CH₄ includes indirect effects from enhancements of ozone and stratospheric water vapour (see Section 2.10).

from IPCC 2007 “Physical Science Basis”



$$CO_2(t) = 0.217 + 0.186 \times CO_2(t=0) e^{-t/1.286} + 0.338 \times CO_2(t=0) e^{-t/18.59} + 0.249 \times CO_2(t=0) e^{-t/172.9}$$

where all times are given in units of year

GWPs Now Come In **Two Flavors** (20 yr & 100 yr time horizons) And Are *Slight Moving Targets*

Global Warming Potentials (dimensionless)					
GHG	IPCC (1995) SAR	IPCC (2001) AR3	IPCC (2007) AR4	IPCC (2013) AR5	IPCC (2021) AR6
<i>100 Year Time Horizon</i>					
CH ₄	21	23	25	28	27.2 ^a or 29.8 ^b
N ₂ O	310	296	298	265	273
<i>20 Year Time Horizon</i>					
CH ₄	56	62	72	84	80.8 ^a or 82.5 ^b
N ₂ O	280	275	289	264	273

In part because best estimate of atmospheric lifetimes has evolved:

^aCH₄ from non-fossil fuel sources such as agriculture

^bCH₄ from fossil fuel sources

Atmospheric Lifetime (year)					
GHG	IPCC (1995) SAR	IPCC (2001) AR3	IPCC (2007) AR4	IPCC (2013) AR5	IPCC (2021) AR6
CH ₄	12	12	12	12.4	11.8
N ₂ O	114	114	114	121	109

IPCC : Intergovernmental Panel on Climate Change
SAR : Second Assessment Report; AR3: Third Assessment Report; AR4: Fourth Assessment Report
AR5: Fifth Assessment Report; AR6: Sixth Assessment Report

100 yr time horizon GWPs from latest IPCC report supposed to be used for International Book-Keeping

IPCC Sixth Assessment Report Global Warming Potentials

Greenhouse Gas	100 Year Time Period			20 Year Time Period		
	AR4 2007	AR5 2014	AR6 2021	AR4 2007	AR5 2014	AR6 2021
CO ₂	1	1	1	1	1	1
CH ₄ fossil origin	25	28	29.8	72	84	82.5
CH ₄ non fossil origin			27.2			80.8
N ₂ O	298	265	273	289	264	273

Does the new report mean your company has to update your calculations, and should you be adopting the 100-year or 20-year GWPs? The answer is not as simple as you might think and depends on regulation in your region and the purpose of your report. However, in the absence of any bespoke requirements, the best practice is to adopt the new AR6 100-year GWPs.

The Paris Rulebook states: ‘*Each Party shall use the 100-year time-horizon global warming potential (GWP) values from the IPCC Fifth Assessment Report, or 100-year time-horizon GWP values from a subsequent IPCC assessment report as agreed upon by the ‘Conference of the Parties serving as the meeting of the Parties to the Paris Agreement’ (CMA), to report aggregate emissions and removals of GHGs, expressed in CO₂-eq. Each Party may in addition also use other metrics (e.g., global temperature potential) to report supplemental information on aggregate emissions and removals of GHGs, expressed in CO₂-eq.*’.

<https://www.ercevolution.energy/ipcc-sixth-assessment-report>

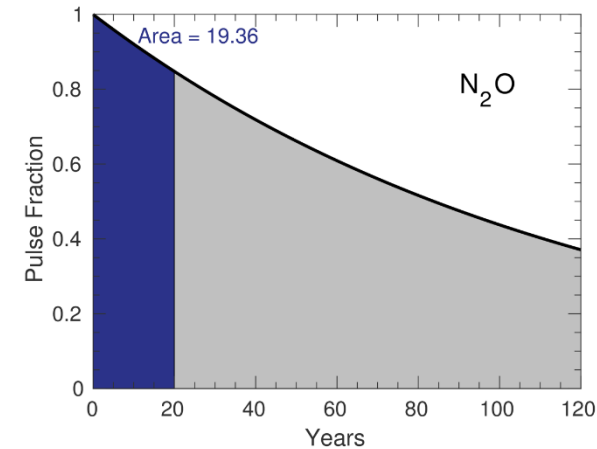
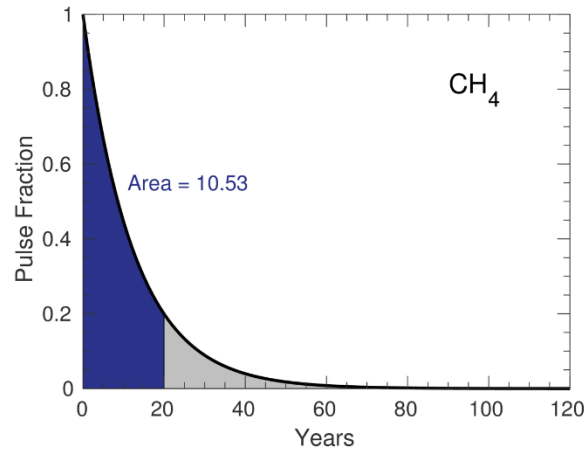
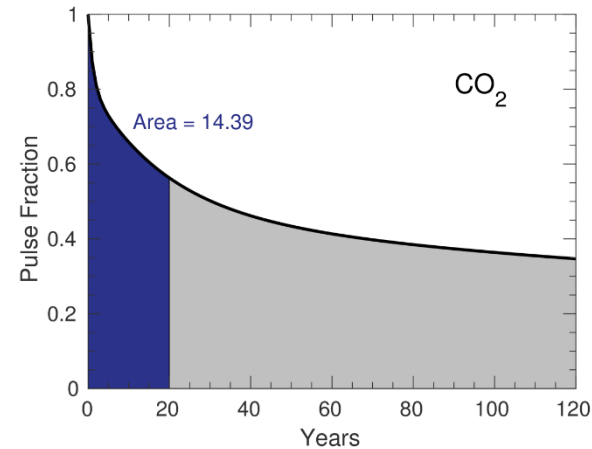
GWP – Global Warming Potential

$$\text{GWP}(\text{CH}_4) = \frac{\int_{\text{time initial}}^{\text{time final}} a_{\text{CH}_4} \times [\text{CH}_4(t)] dt}{\int_{\text{time initial}}^{\text{time final}} a_{\text{CO}_2} \times [\text{CO}_2(t)] dt}$$

$$\text{GWP}(\text{N}_2\text{O}) = \frac{\int_{\text{time initial}}^{\text{time final}} a_{\text{N}_2\text{O}} \times [\text{N}_2\text{O}(t)] dt}{\int_{\text{time initial}}^{\text{time final}} a_{\text{CO}_2} \times [\text{CO}_2(t)] dt}$$

Global Warming Potentials				
GHG	IPCC (1995)	IPCC (2001)	IPCC (2007)	IPCC (2013)
100 Year Time Horizon				
CH ₄	21	23	25	28
N ₂ O	310	296	298	265
20 Year Time Horizon				
CH ₄	56	62	72	84
N ₂ O	280	275	289	264

20 Year Time Horizon means time final = 20 years in these integrals



$$\text{CO}_2(t) = 0.217 + 0.186 \times \text{CO}_2(t=0) e^{-t/1.286} + 0.338 \times \text{CO}_2(t=0) e^{-t/18.59} + 0.249 \times \text{CO}_2(t=0) e^{-t/172.9}$$

$$\text{CH}_4(t) = \text{CH}_4(t=0) e^{-t/12.4}$$

$$\text{N}_2\text{O}(t) = \text{N}_2\text{O}(t=0) e^{-t/121.0}$$

where all times are given in units of year

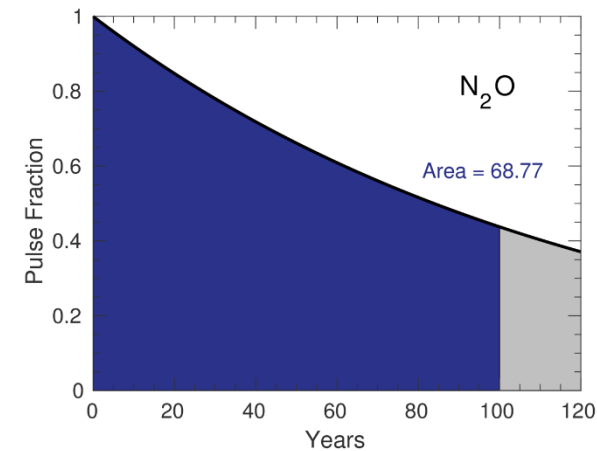
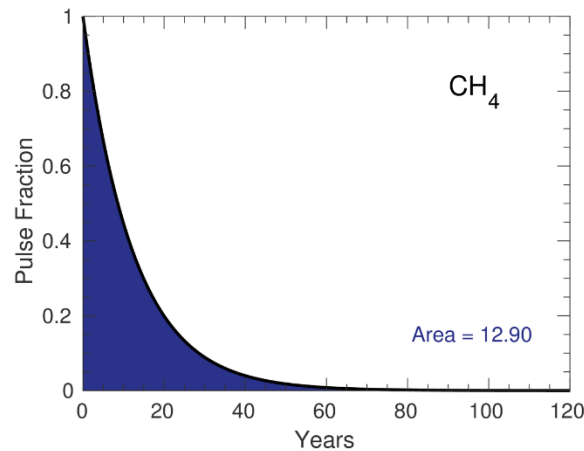
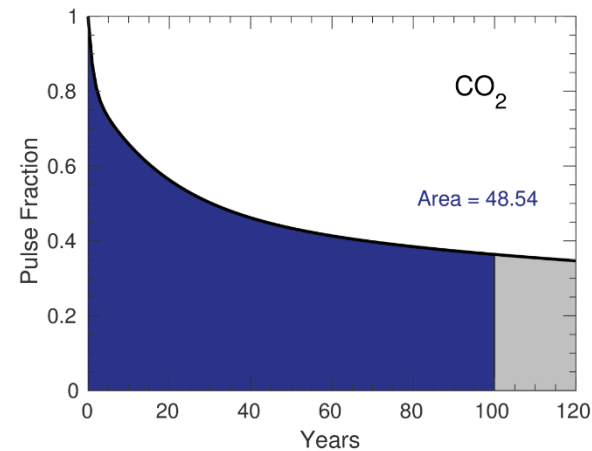
GWP – Global Warming Potential

$$\text{GWP}(\text{CH}_4) = \frac{\int_{\text{time initial}}^{\text{time final}} a_{\text{CH}_4} \times [\text{CH}_4(t)] dt}{\int_{\text{time initial}}^{\text{time final}} a_{\text{CO}_2} \times [\text{CO}_2(t)] dt}$$

$$\text{GWP}(\text{N}_2\text{O}) = \frac{\int_{\text{time initial}}^{\text{time final}} a_{\text{N}_2\text{O}} \times [\text{N}_2\text{O}(t)] dt}{\int_{\text{time initial}}^{\text{time final}} a_{\text{CO}_2} \times [\text{CO}_2(t)] dt}$$

Global Warming Potentials				
GHG	IPCC (1995)	IPCC (2001)	IPCC (2007)	IPCC (2013)
100 Year Time Horizon				
CH ₄	21	23	25	28
N ₂ O	310	296	298	265
20 Year Time Horizon				
CH ₄	56	62	72	84
N ₂ O	280	275	289	264

100 Year Time Horizon means time final = 100 years in these integrals



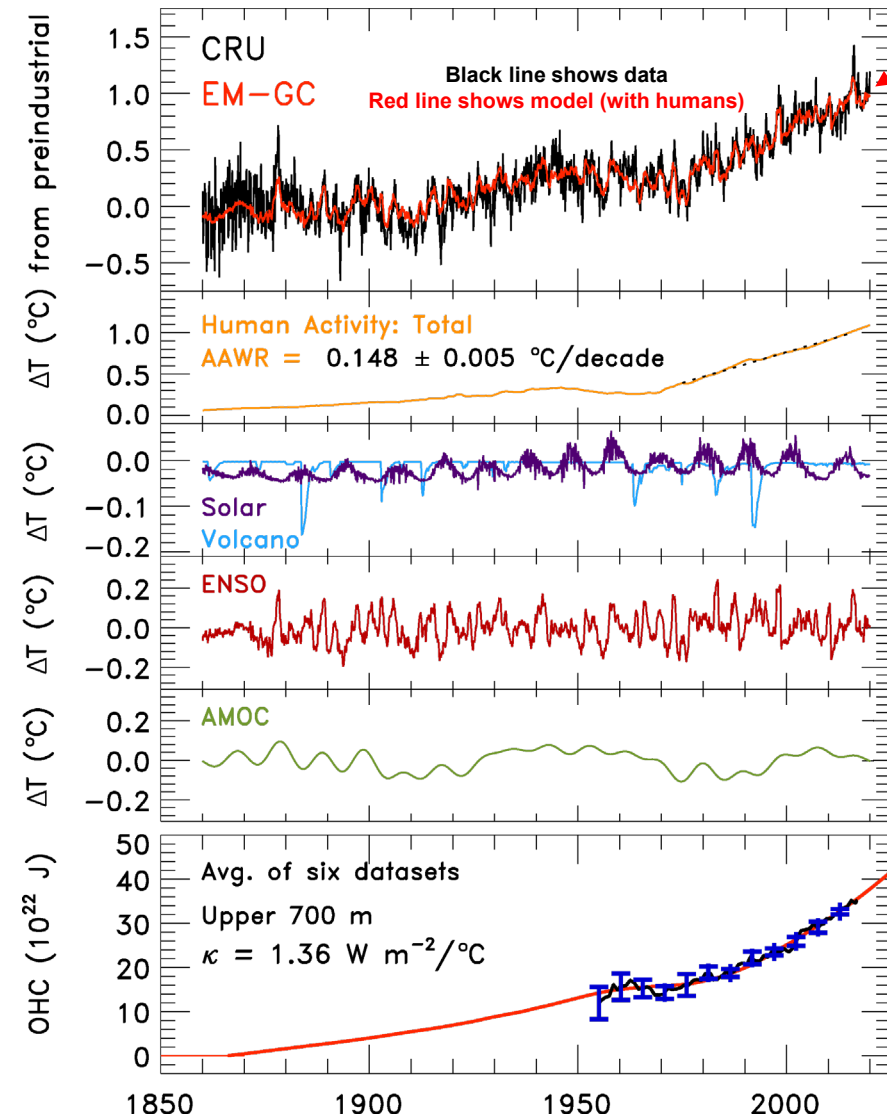
$$\text{CO}_2(t) = 0.217 + 0.186 \times \text{CO}_2(t=0) e^{-t/1.286} + 0.338 \times \text{CO}_2(t=0) e^{-t/18.59} + 0.249 \times \text{CO}_2(t=0) e^{-t/172.9}$$

$$\text{CH}_4(t) = \text{CH}_4(t=0) e^{-t/12.4}$$

$$\text{N}_2\text{O}(t) = \text{N}_2\text{O}(t=0) e^{-t/121.0}$$

where all times are given in units of year

Are humans responsible?



$$\Delta T_{MDL\ i} = (1 + \gamma) \left(\frac{GHG\ RF_i + LUC\ RF_i + Aerosol\ RF_i}{\lambda_p} \right) + C_0 + C_1 \times SOD_{i-6} + C_2 \times TSI_{i-1} + C_3 \times ENSO_{i-2} + C_4 \times AMOC_i - \left(\frac{Q_{OCEAN\ i}}{\lambda_p} \right)$$

where:

i denotes month

$$\lambda_p = 3.2 \text{ W m}^{-2} \text{ } ^{\circ}\text{C}^{-1}$$

$$1 + \gamma = \{1 - \lambda_{\Sigma}/\lambda_p\}^{-1}$$

GHG RF = RF due to all anthropogenic GHGs

LUC RF = RF due to Land Use Change

Aerosol RF = RF due to Tropospheric Aerosols

SOD = Stratospheric Optical Depth

TSI = Total Solar Irradiance

ENSO = El Niño Southern Oscillation

AMOC = Atlantic Meridional Overturning Circulation

Q_{OCEAN} = Ocean heat export =

$$\kappa(1 + \gamma)\{\Delta T_{MDL\ i} - \Delta T_{OCEAN\ SURFACE\ i}\}$$

CRU: Climate Research Unit of East Anglia, United Kingdom
EM-GC: Empirical Model of Global Climate, Univ of Maryland

Global warming is caused by CO₂, the greatest waste product of modern society, as well as CH₄, N₂O, and other GHGs.

Temperature will continue to rise until human emission of GHGs is curtailed

Canty *et al.*, 2013 <https://www.atmos-chem-phys.net/13/3997/2013/acp-13-3997-2013.html>

McBride *et al.*, 2021 <https://esd.copernicus.org/articles/12/545/2021>

Nicholls *et al.*, 2021 <https://agupubs.onlinelibrary.wiley.com/doi/full/10.1029/2020EF001900>

Figure provided by Laura McBride.

Are humans responsible?

Orbital variations: drive the ice ages but too small to drive modern warming

Volcanoes: no sustained forcing

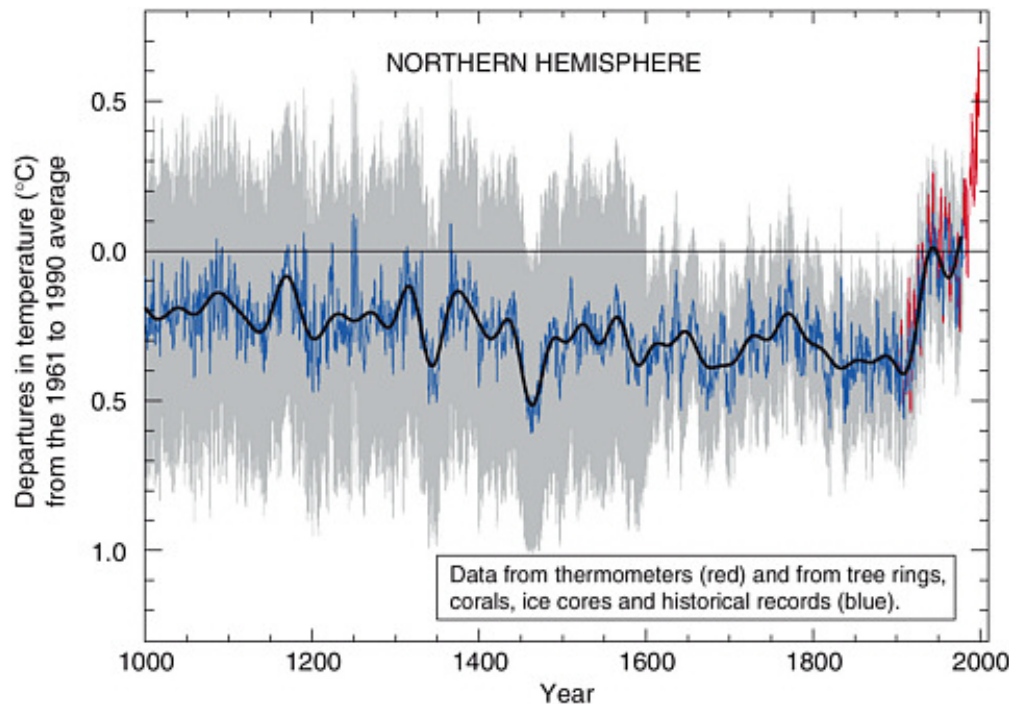
Solar variability:

Perhaps dominant forcing of Medieval Warming and Little Ice Age

Small effect since ~1860

Internal variability (eg, El Niño / La Niña) :

Climate record from 1000 to 1850 shows nothing like sustained, present rate of warming



Ozone in the Atmosphere

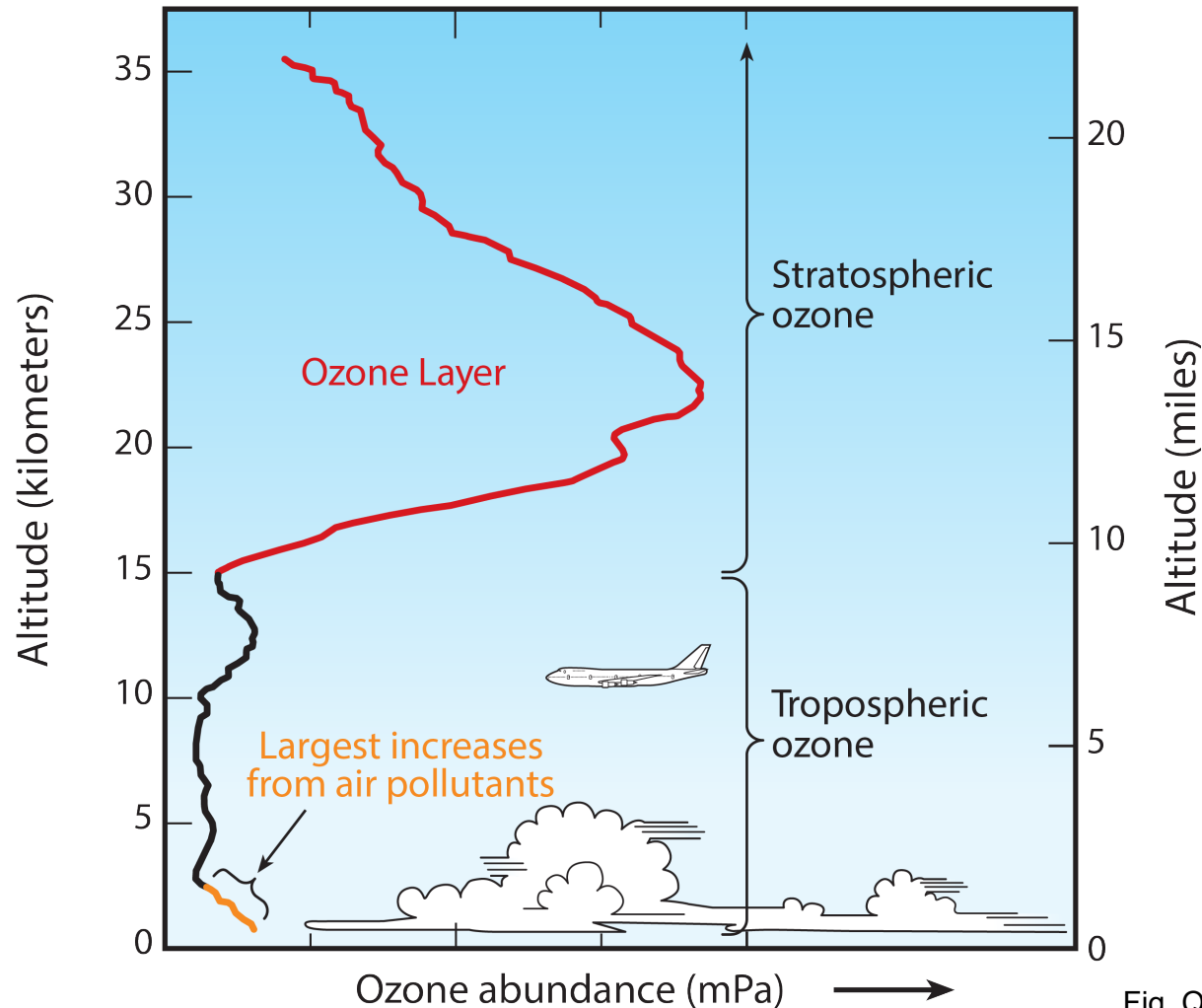


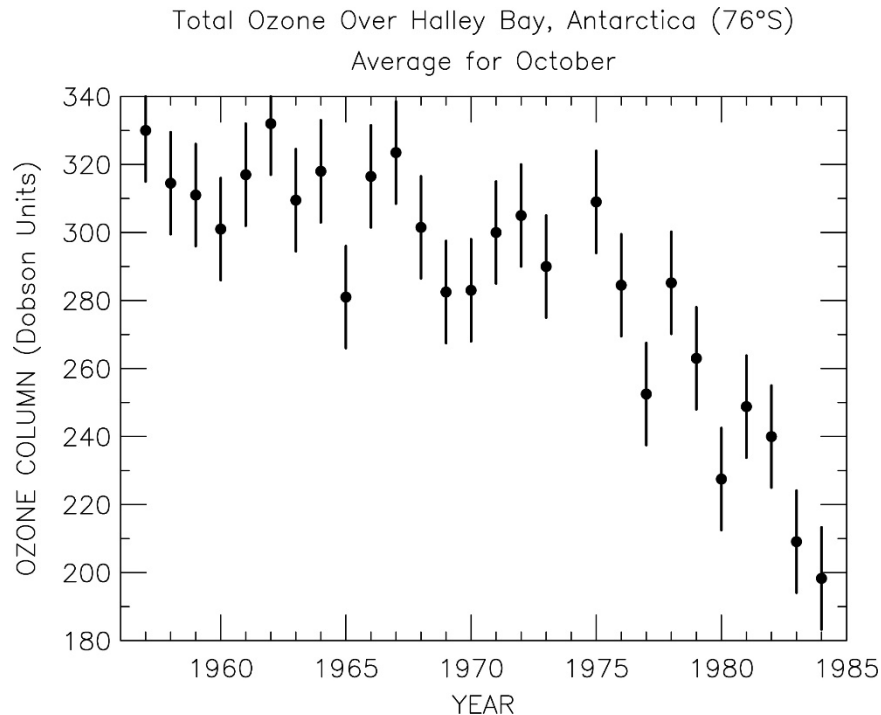
Fig. Q1-2
20 QAs about the Ozone Layer

**It is incredible that human activity
both destroys stratospheric ozone (so-called good ozone)
and produces tropospheric ozone (so-called bad ozone)**

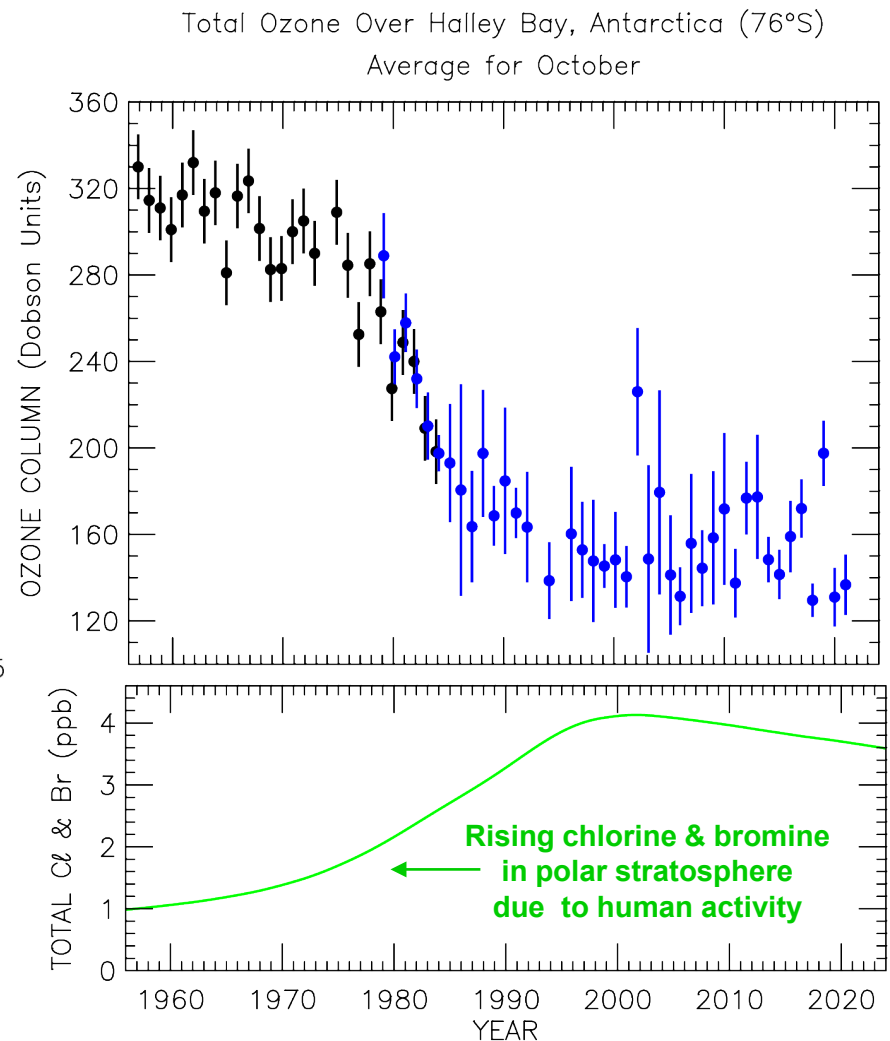
Earth's Atmosphere – Effect of Humans

Stratospheric Ozone – shields surface from solar UV radiation

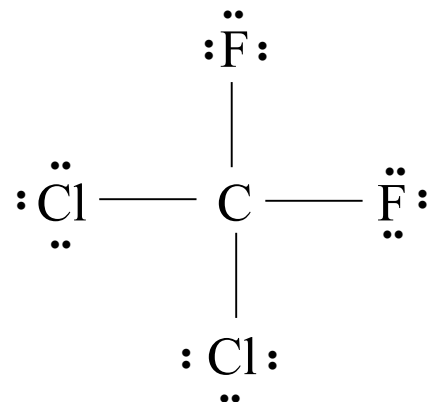
Update



After Farman *et al.*, Large losses of total ozone in Antarctica reveal Seasonal ClO_x/NO_x interaction, *Nature*, 315, 207, 1985.



CFC-12



How is it eventually removed from the atmosphere ?

What does it produce upon its removal ?

See pages 71 to 75, Ch 2, Chemistry in Context, for description of Lewis Dot Structures of atmospherically important species

**Note: you will not be tested on Lewis Dot Structures.
However, we want non-chemists to at least have been exposed to this concept for tracking the position of electrons, central for understanding atmospheric chemical reactions.**

Measurements of Reactive Chlorine From Space

Measurements of Chlorine Gases from Space

Annual mean 2006 (30°–70°N)

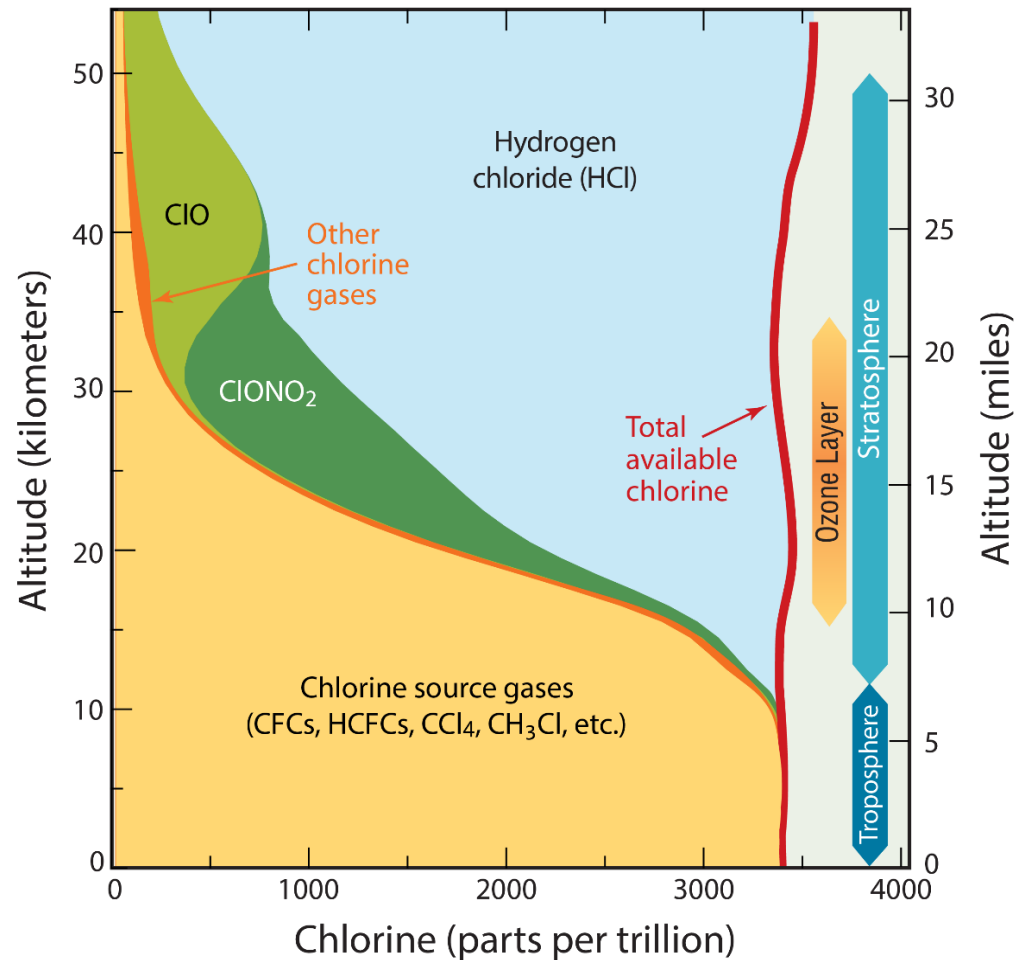
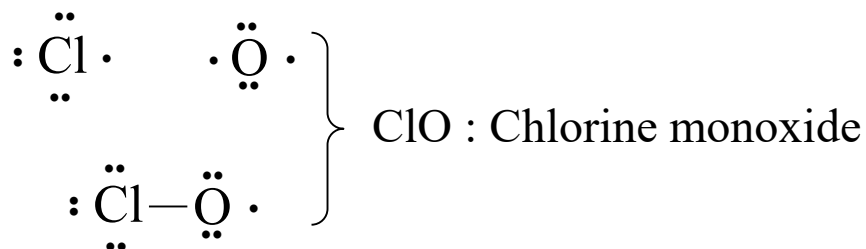


Fig. Q7-2, 20 QAs about the Ozone Layer

ClO (Chlorine Monoxide) is a Radical

Radicals

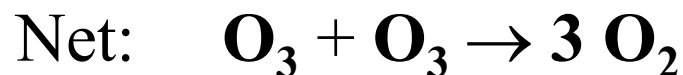
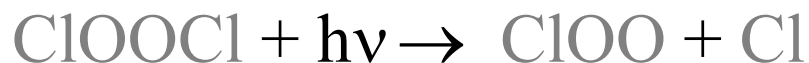
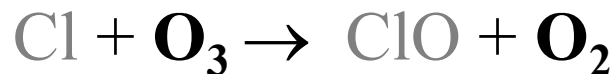
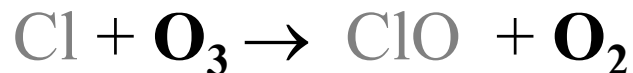
- Odd number of electrons - unpaired electron in outer valence shell
- Go to great lengths to pair off lone electron
- Exceptionally reactive



See pages 71 to 75, Ch 2, Chemistry in Context, for description of Lewis Dot Structures of atmospherically important species

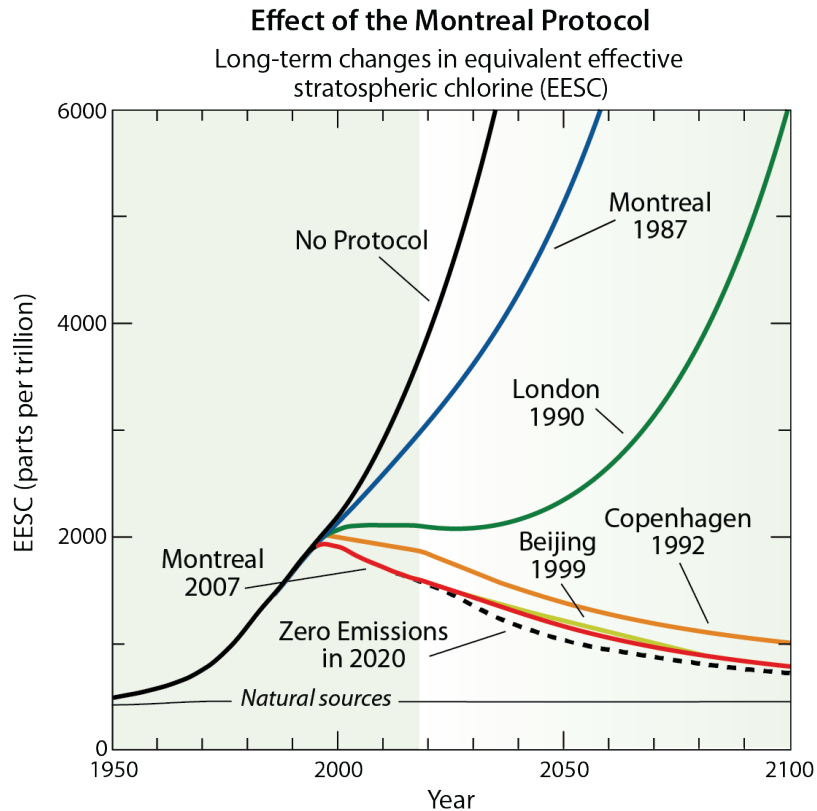
Note: you will not be tested on Lewis Dot Structures.
However, we want non-chemists to at least have been exposed to this concept for tracking the position of electrons, central for understanding atmospheric chemical reactions.

Chlorine Radicals Lead to Ozone Loss



Catalytic loss of ozone: this chemistry causes the Antarctic ozone hole

And Atmospheric Levels of these Pollutants are Declining



CFCs: Chlorofluorocarbons

Contain some combination of chlorine, fluorine, and at least one carbon. Freons are a trade name for CFCs.

Bromocarbons:

Contain bromine, perhaps chlorine, and at least one carbon. Halons are a trade name for bromocarbons.

HCFCs: Hydro-chlorofluorocarbons

Same as CFCs, except one or more hydrogen has replaced a chlorine.

HFCs: Hydrofluorocarbons

Contain some combination of hydrogen, fluorine, and carbon. **These gases do not contain any bromine or chlorine, and hence pose no damage to the ozone layer.** Some HFCs are potent GHGs.

EESC: Equivalent, effective stratospheric chlorine. Reflects combined influence of chlorine and bromine on ozone, via a simple formula: $[\text{Chlorine}] + 60 \times [\text{Bromine}]$

Figure Q14-1, 20 QAs about the Ozone Layer

CFC Usage Prior to the Montreal Protocol

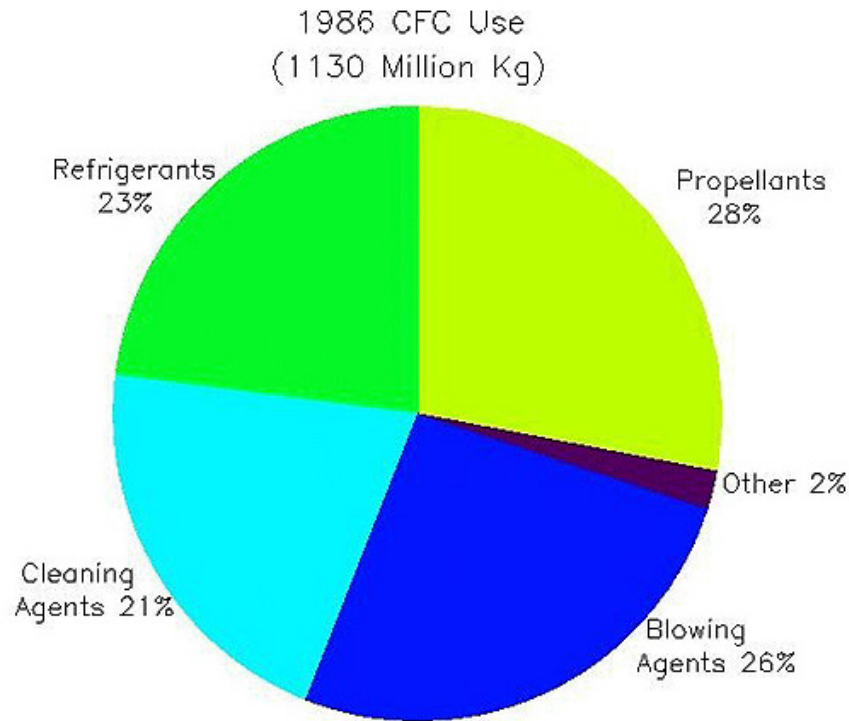
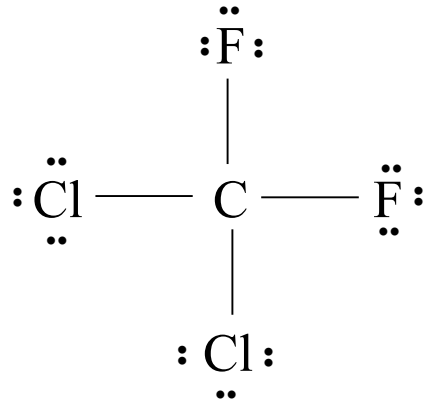


Figure 5b. Changing use patterns for CFCs (from Fisher and Midgley, 1994).

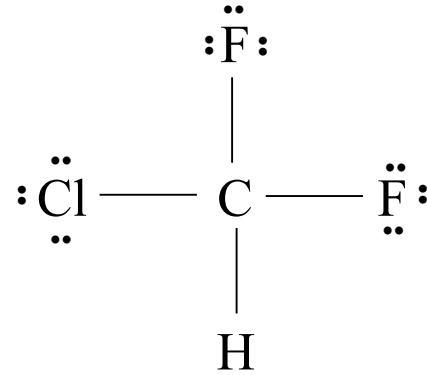
The uses of CFCs in various sectors before the 1987 Montreal Protocol, which required countries to phase out their production to protect the ozone layer.

From http://www.ccpo.odu.edu/SEES/ozone/class/Chap_10/index.htm
based upon <https://agupubs.onlinelibrary.wiley.com/doi/epdf/10.1029/94JD00738>

CFCs were replaced with HCFCs



CFC-12
or
Freon-12
or
R-12



HCFC-22
or
Freon-22
or
R-22

Phase out of CFCs and other Ozone Depleting Substances (ODSs)

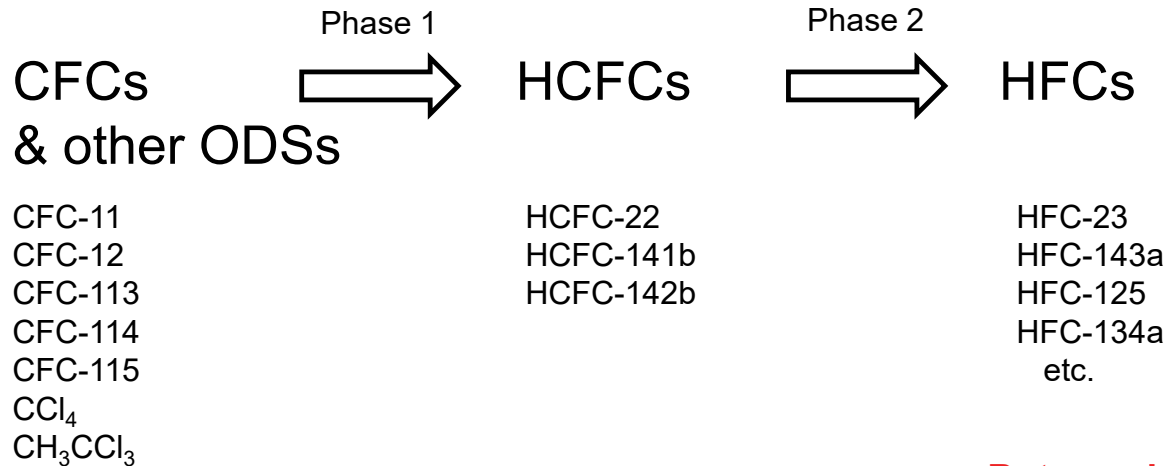
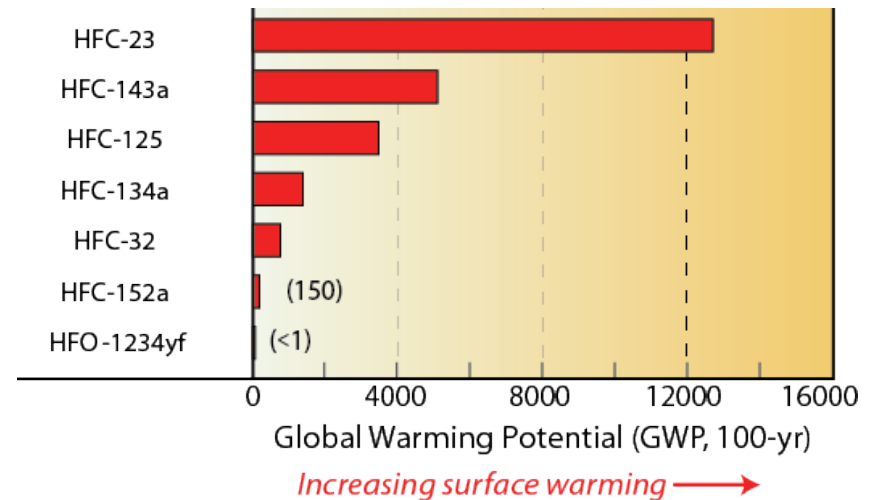


Figure Q17-3, 20 QAs about the Ozone L

**Harmful to
ozone layer**

**Less harmful to
ozone layer**

**But very harmful to
Earth's climate**



See http://www.atmos.umd.edu/~rjs/class/spr2020/supplemental_readings/Naming_Convention_for_CFCs_Halons.pdf
for a guide to CFC naming convention

Climate Benefit of the Kigali Amendment

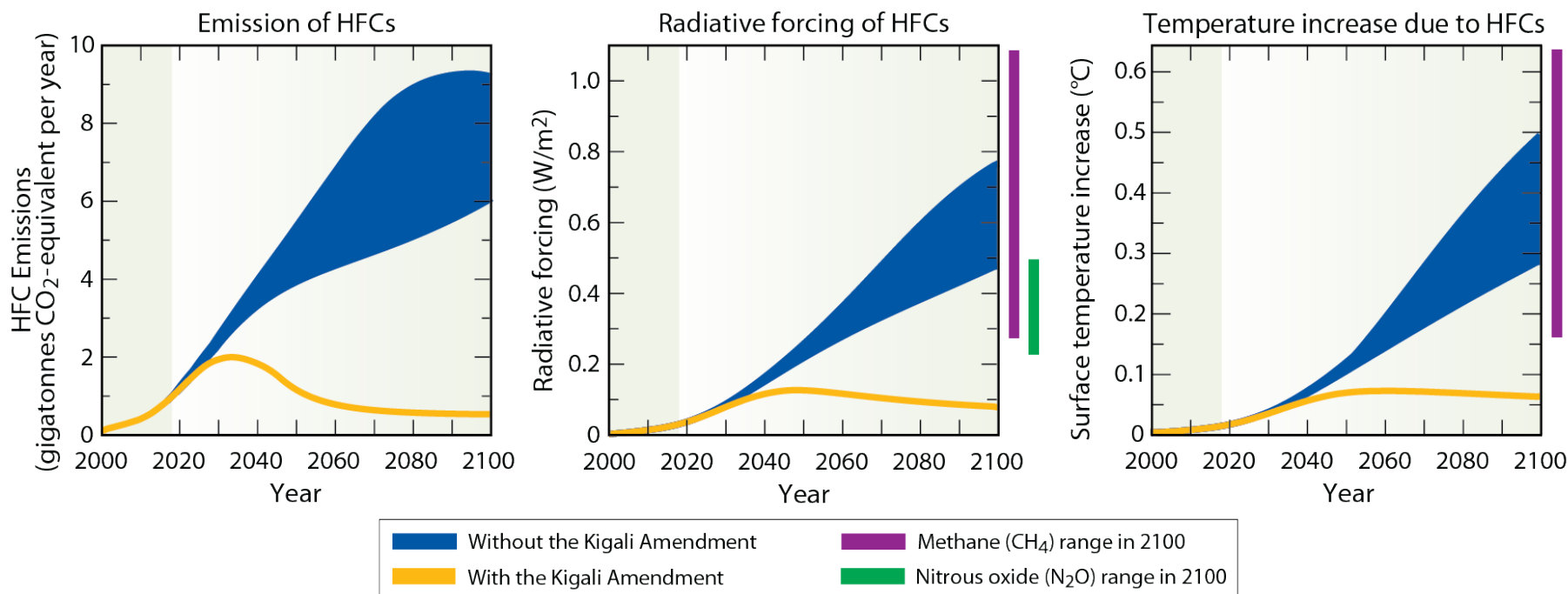


Figure Q19-2, 20 QAs about the Ozone Layer

Senate to Consider Kigali Amendment Ratification

November 16, 2021

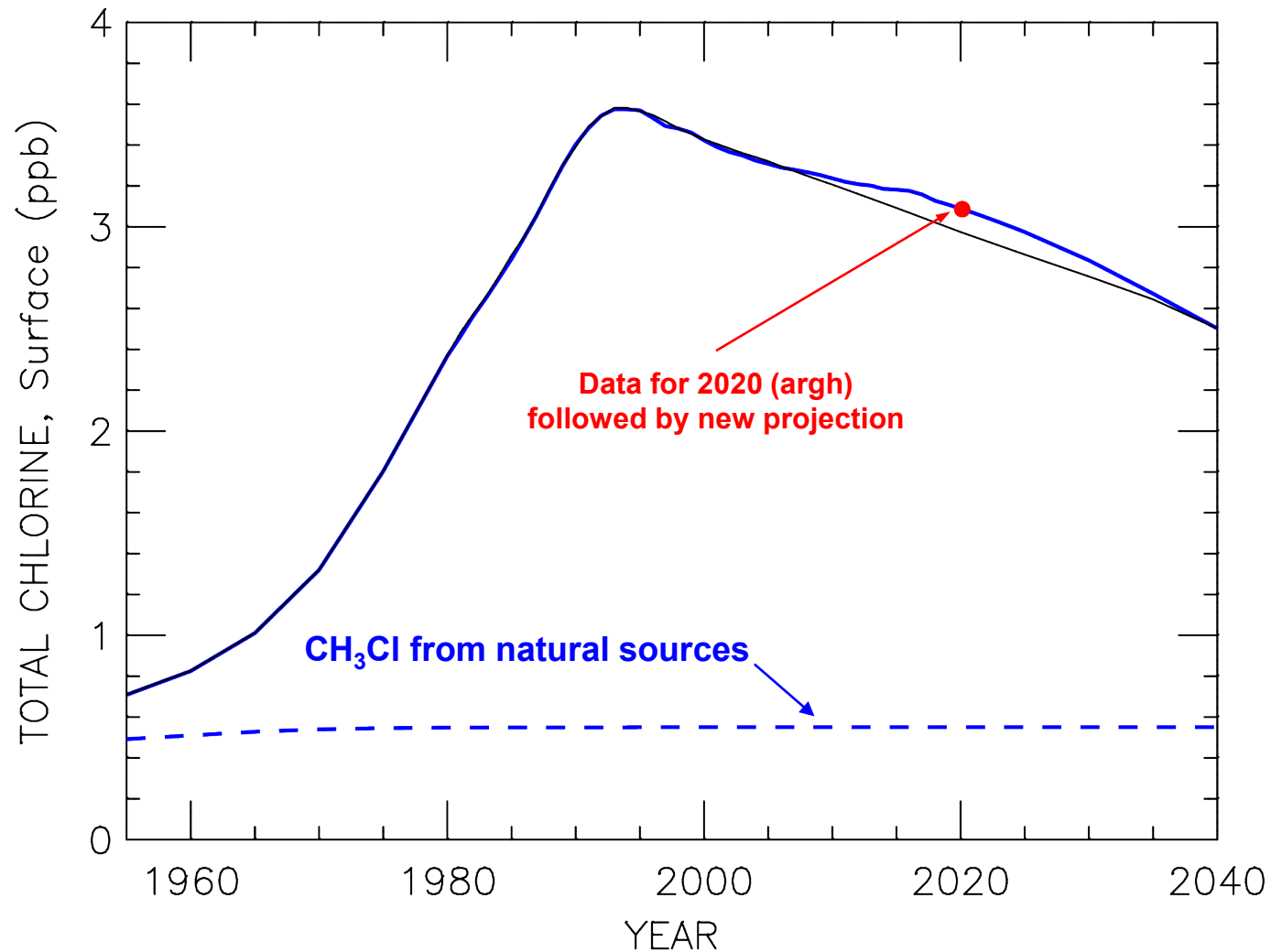
Christina Theodoridi

The White House today [sent](#) the Kigali Amendment to the Montreal Protocol, the treaty to phase down hydrofluorocarbons (HFCs), to the Senate for its advice and consent to U.S. ratification. This action follows up on President Biden's [executive order](#) earlier this year.

<https://www.nrdc.org/experts/christina-theodoridi/senate-consider-kigali-amendment-ratification>

Montreal Protocol Has Banned Industrial Production of CFCs & Other ODS

Projections Based on 2018 World Meteorological Organization
Scientific Assessment of Ozone Depletion Report



Montreal Protocol Had Banned Most Industrial Production of CFCs & Other ODS

The New York Times

In a High-Stakes Environmental Whodunit, Many Clues Point to China

Interviews, documents and advertisements collected by The New York Times and independent investigators indicate that a major source — possibly the overwhelming one — is factories in China that have ignored a global ban and kept making or using the chemical, CFC-11, mostly to produce foam insulation for refrigerators and buildings.

“You had a choice: Choose the cheaper foam agent that’s not so good for the environment, or the expensive one that’s better for the environment,” said Zhang Wenbo, owner of a refrigerator factory here in Xingfu, in Shandong Province, where he and many other small-scale manufacturers said that until recently, they had used CFC-11 widely to make foam insulation.



Billboards in Xingfu, China, promoting locally made refrigerators. The city has around 1,700 businesses involved in the production of cooking and refrigeration equipment. Gillies Sabatini for The New York Times

<https://www.nytimes.com/2018/06/24/world/asia/china-ozone-cfc.html>

Air Quality Index

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.

Table 1.4 Levels for the Air Quality Index		
Air Quality Index (AQI) Values	Levels of Health Concern	Colors
<i>When the AQI is in this range:</i>	<i>...air quality conditions are:</i>	<i>...as symbolized by this color.</i>
0–50	Good	Green
51–100	Moderate	Yellow
101–150	Unhealthy for sensitive groups	Orange
151–200	Unhealthy	Red
201–300	Very unhealthy	Purple
301–500	Hazardous	Maroon

- Computed for each criteria pollutant even though many newspapers only give a single value (usually for worse index)
- In the U.S. health officials are generally concerned about elevated O₃, PM_{2.5}, and ultra-fine particles

Tropospheric Pollutants (The Air We Breathe)

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.

Table 1.2 U.S. National Ambient Air Quality Standards		
Pollutant	Standard (ppm)	Approximate Equivalent Concentration ($\mu\text{g}/\text{m}^3$)
Carbon monoxide		
8-hr average	9	10,000
1-hr average	35	40,000
Nitrogen dioxide		
Annual average	0.053	100
Ozone		
8-hr average	0.075	147
1-hr average	0.12	235
Particulates*		
PM ₁₀ , annual average	—	50
PM ₁₀ , 24-hr average	—	150
PM _{2.5} , annual average	—	15
PM _{2.5} , 24-hr average [†]	—	35
Sulfur dioxide		
Annual average	0.03	80
24-hr average	0.14	365
3-hr average	0.50	1,300

Note: A standard also exists for lead, but lead does not appear in this table since most of the U.S. is in compliance

*PM₁₀ refers to all airborne particles 10 μm in diameter or less. PM_{2.5} refers to particles 2.5 μm in diameter or less.

—The unit of ppm is not applicable to particulates.

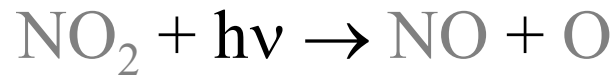
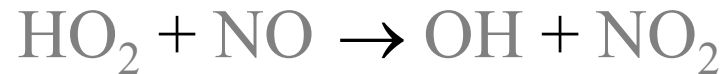
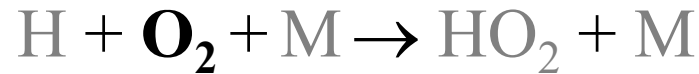
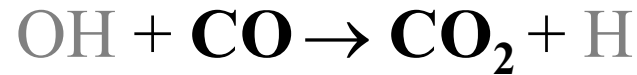
[†]PM_{2.5} standards are likely to be revised after 2011.

Chapter 1 Source: U.S. Environmental Protection Agency. Standards also exist for lead, but are not included here.

Chemistry in Context

Criteria pollutant: identified as being common-place and detrimental to human welfare (i.e., ubiquitous pollutant)

Tropospheric Ozone Production



Oxidation of **CO** in the presence of elevated **NO_x (NO + NO₂)**
leads to ***production*** of tropospheric ozone

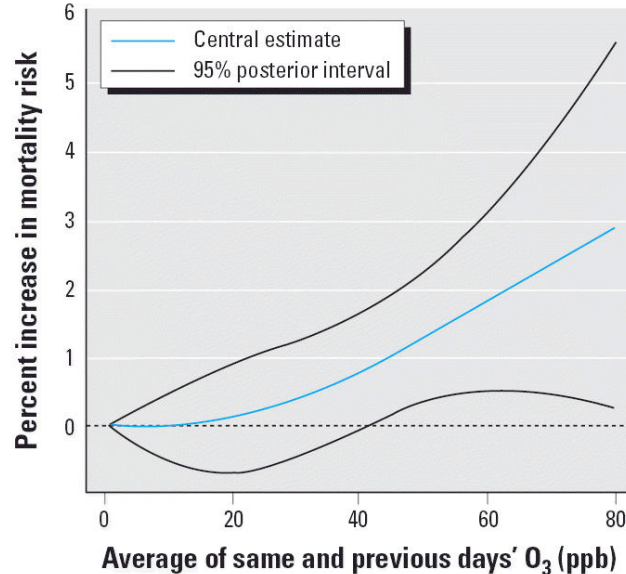
Air Quality Standards and Why We Care



For more information, have a look at:

<https://www.weforum.org/agenda/2018/11/deadly-air-pollution-shortens-lives-by-nearly-2-years-researchers>

<https://aqli.epic.uchicago.edu/pollution-facts>



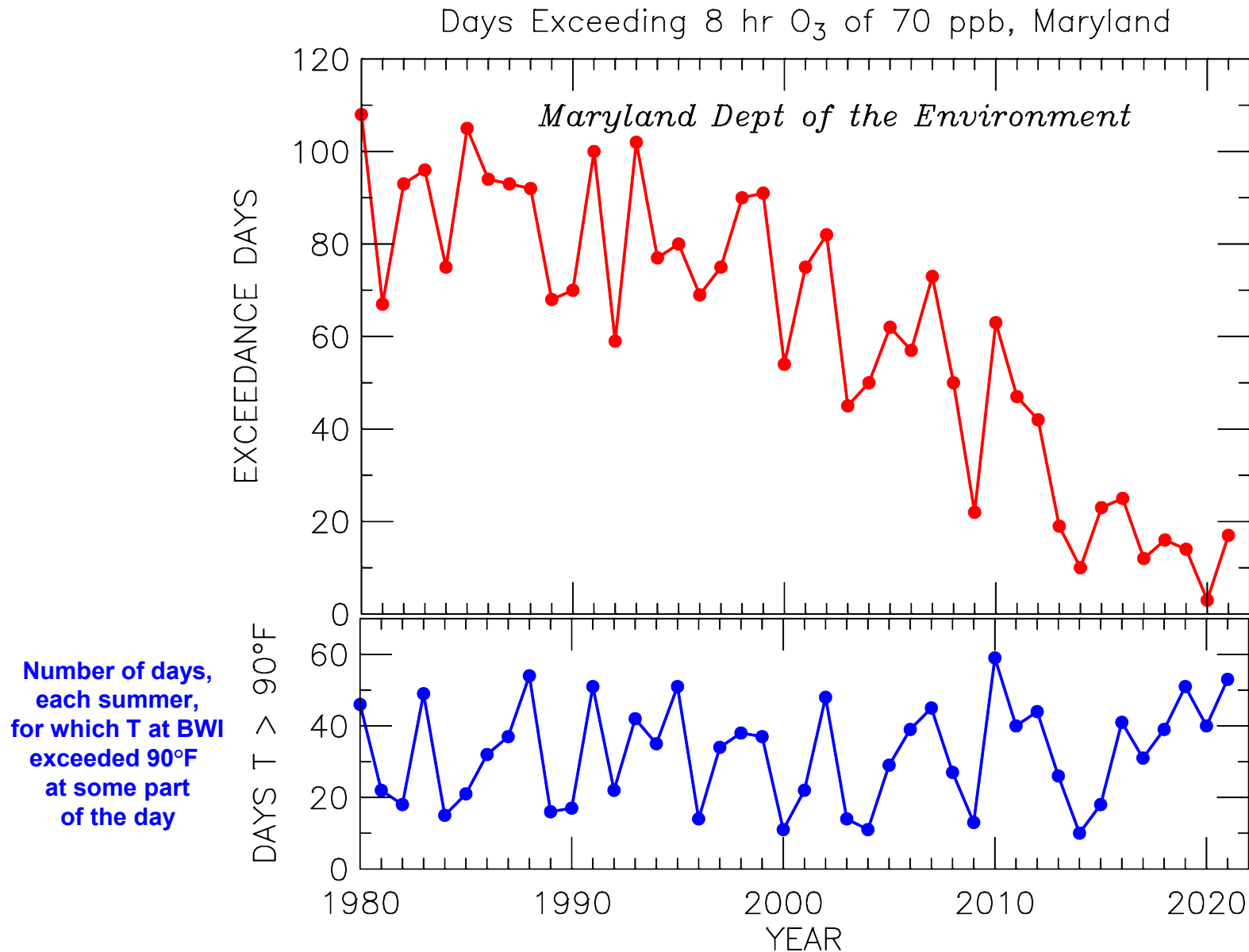
Increased risk of premature death (mortality) for all levels of surface O₃

Reductions in surface O₃ will benefit public health, regardless of present conditions

Bell *et al.*, 2006

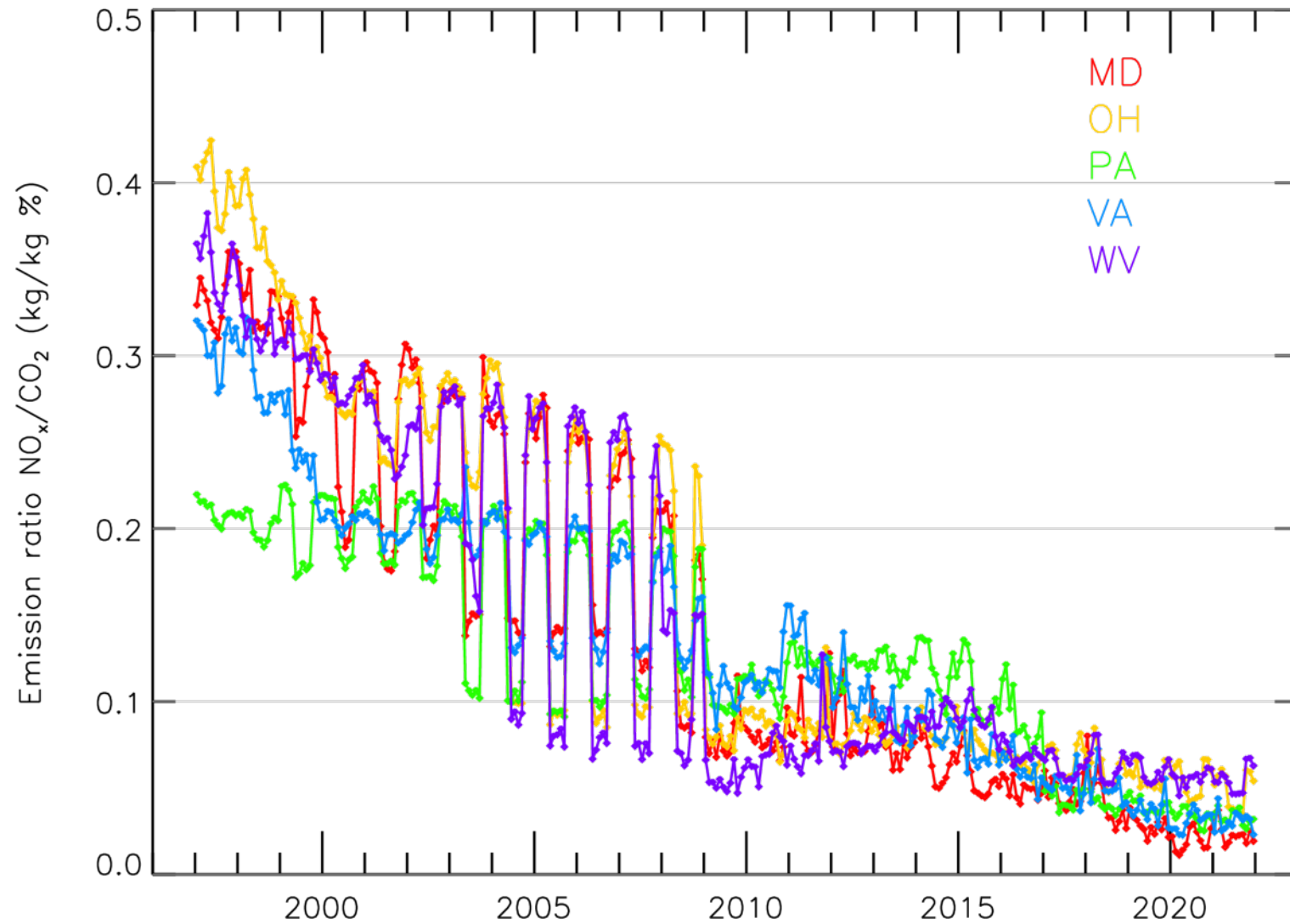
<http://www.ncbi.nlm.nih.gov/sites/ppmc/articles/PMC1440776>

Significant Improvements in Local Air Quality since early 1980s



<http://www.mde.state.md.us/programs/Air/AirQualityMonitoring/Pages/SeasonalReports.aspx>

Trends in power plant emissions of NO_x



Air Quality In The News

Biden Administration to Reinstate Mercury Pollution Rules Weakened Under Trump

The E.P.A. will resume enforcing limits on the release of mercury, a neurotoxin linked to developmental damage in children, from coal-burning power plants.



Brandon Shores Power Plant in Maryland, a plant that installed costly “scrubber” technology to reduce mercury emissions. Shannon Jensen for The New York Times



By Coral Davenport

Jan. 31, 2022 Updated 5:04 p.m. ET

WASHINGTON — The Biden administration on Monday reinstated a way of measuring the benefits of reducing air pollution, the first step in a plan that could tighten limits on the amount of mercury that can be discharged from coal-burning power plants.

Mercury is a neurotoxin that poses a particular danger to the brain development of children and fetuses.

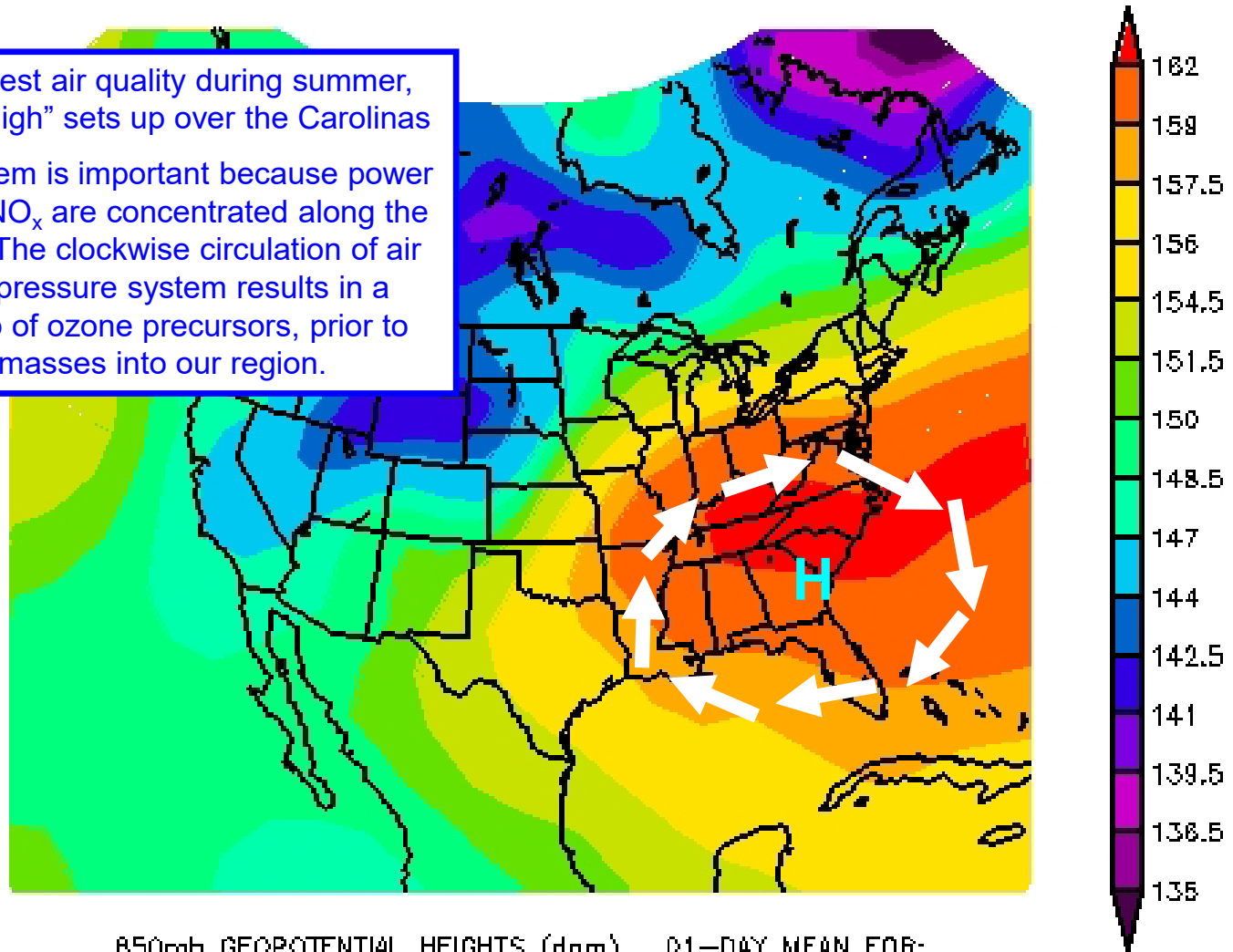
The mercury announcement is among several recent actions taken or planned by the Biden administration that are aimed at reducing pollution in air and water. After a first year in which President Biden tried to push ambitious climate legislation through Congress [only to see it stall](#), the administration is using its [regulatory machinery to try to curb pollution](#).

<https://www.nytimes.com/2022/01/31/climate/epa-mercury-pollution-coal.html>

Day-to-day meteorology (weather!) affects severity and duration of pollution episodes

Maryland has poorest air quality during summer, when a “Bermuda High” sets up over the Carolinas

This circulation system is important because power plant emissions of NO_x are concentrated along the Ohio River valley. The clockwise circulation of air around this high pressure system results in a significant build up of ozone precursors, prior to arrival of air masses into our region.



850mb GEOPOTENTIAL HEIGHTS (dam) 01-DAY MEAN FOR:
Sun JUL 04 1999

NCEP OPERATIONAL DATASET

<http://www.mde.state.md.us/assets/document/BJH%20-%20Basics%20on%20Ozone%20Transport.ppt>

Temperature Inversions and Air Quality

Temperature inversion: increase in temperature with height

Inversions important for Air Quality because they inhibit vertical mixing of air

Air pollutants can accumulate in cities ringed by mountains, such as Los Angeles, Mexico City, Denver, etc.

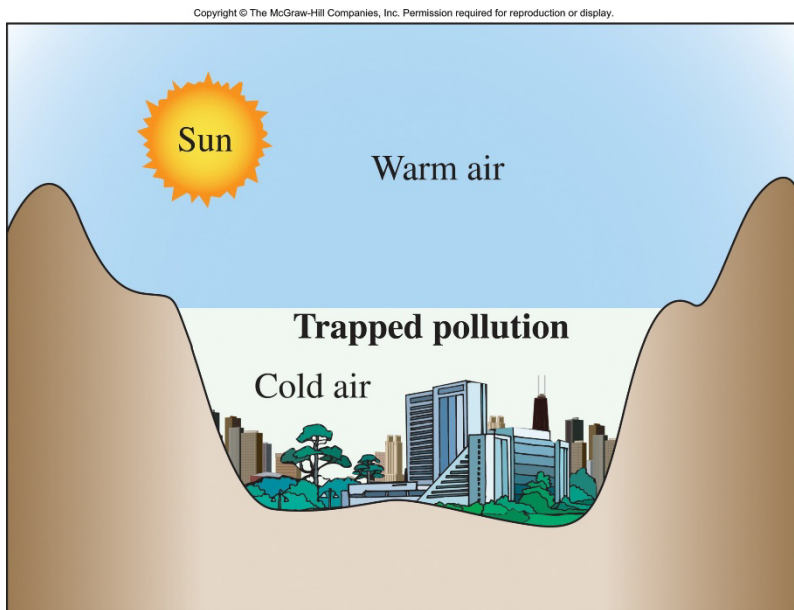
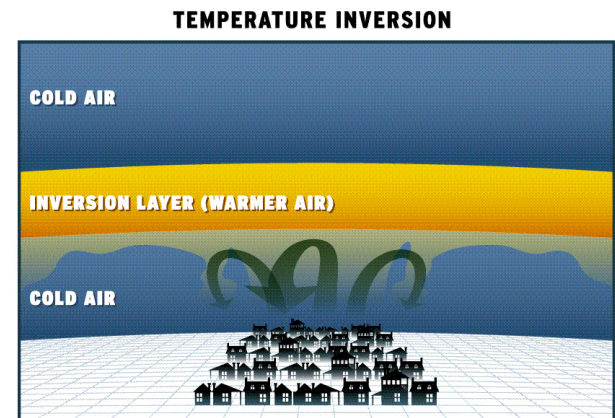
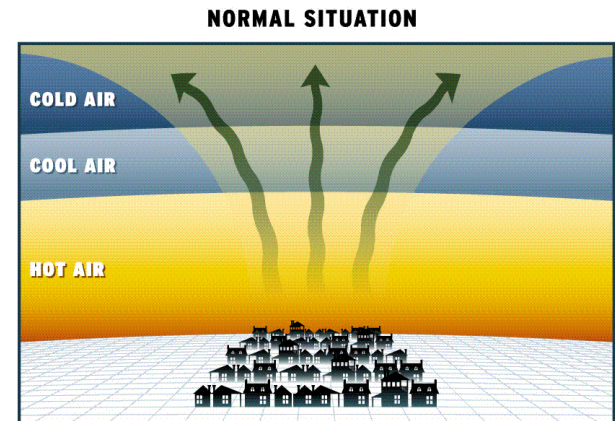
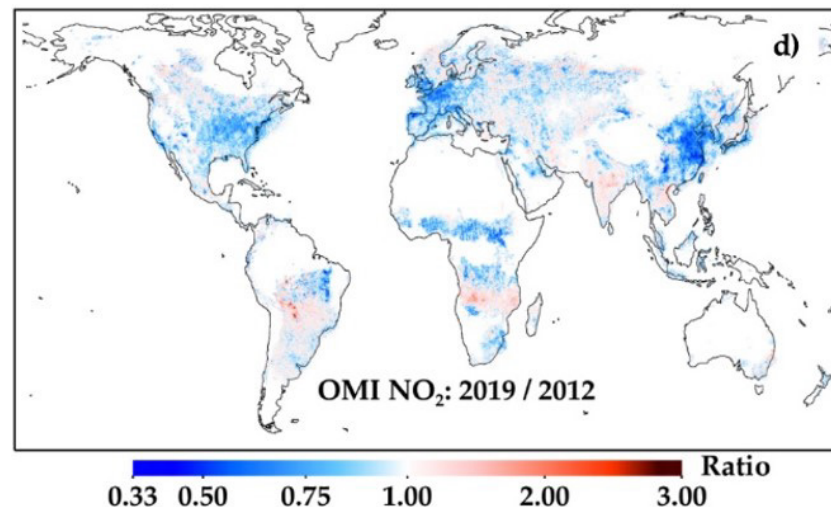
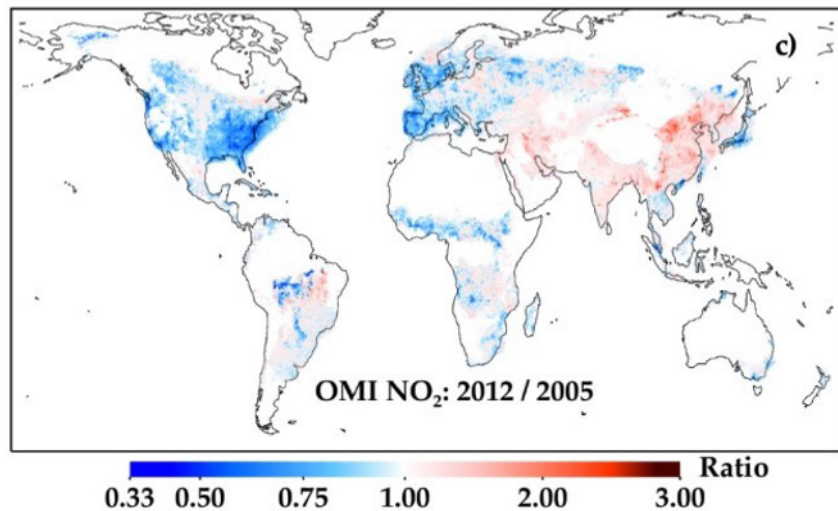
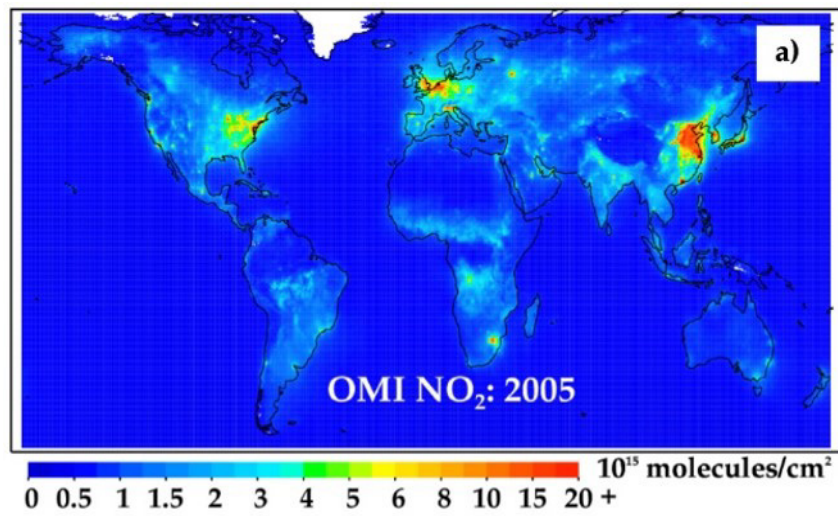


Figure 1.10, Chemistry in Context



<http://geographygems.blogspot.com/2011/09/smog.html>



Top: Tropospheric vertical column NO₂ (Trop NO₂) measured in 2005

Bottom Left: Ratio of Trop NO₂ measured in 2012 and 2005

Bottom Right: Ratio of Trop NO₂ measured in 2019 and 2012

All observations from the NASA OMI instrument

Goldberg *et al.*, *ERL*, 2021

<https://iopscience.iop.org/article/10.1088/1748-9326/ac2c34/meta>