

Radiative Forcing

AOSC / CHEM 433 & AOSC / CHEM 633

Ross Salawitch

Class Web Sites:

<http://www2.atmos.umd.edu/~rjs/class/fall2020>

<https://myelms.umd.edu/courses/1291919>

Goals:

- Understanding interaction between gases and IR radiation
- Radiative forcing of greenhouse gases
- Radiative forcing of aerosols

Wavenumber = 1 / Wavelength

$$1 \text{ } \mu\text{m} \text{ (micron)} = 10^{-6} \text{ m}$$

$$1 \text{ nm} \text{ (nanometer)} = 10^{-9} \text{ m}$$

$$\text{Therefore, } 1 \text{ } \mu\text{m} = 1000 \text{ nm}$$

Lecture 7

24 September 2020

Announcements

- Problem Set #2 due Fri, 2 Oct, 5 pm
 - Has been posted
 - Slightly different assignment for students enrolled in 433 & students enrolled in 633

09/24	Radiative Forcing	Chemistry in Context, Sec 2.4, 2.5, 2.6, 3.3 & 3.4 (14 pages) Paris Beacon of Hope Sec 1.2 (intro), 1.2.1 (please review), & 1.2.3.6 (8 pages)	AT 7	Lecture 7 2020 Zoom Video		Green Chemistry, Chapter 3.4 (Sections 3.4.4.1 to 3.4.4.4 provide a nice mathematical complement to the lecture material) Myhre et al., GRL, 1998 Bera et al., JPC, 2009	Quiz 7
09/29	Modeling Earth's Climate: Water Vapor, Aerosol, Cloud, & Albedo Feedbacks	Chemistry in Context, Sec 3.9 (6 pages) Houghton, pg. 105-116	AT 8	Lecture 8 2020 Zoom Video		Bony et al., 2006	Quiz 8
10/01	Consequences of Climate Change	Chemistry in Context, Sec 3.10 (5 pages) Forbes Article	No AT	Lecture 9 2020 Zoom Video		Union of Concerned Scientists Climate Reality Project Climate Change and Disease NY Times, Bangladesh NY Times, Kiribati	No Quiz
10/02					Problem Set 2 due today at 5 pm: 433 Students 633 Students		

<http://www2.atmos.umd.edu/~rjs/class/fall2020>

Announcements: Outside of Class

1) Thurs, 24 Sept : AOSC Weekly Seminar (today @ 3:30 pm)

Professor Nicole Riemer, University of Illinois

Aerosol Mixing State: Synthesizing Measurements and Models

Atmospheric aerosols are complex mixtures of different chemical species, and individual particles exist in many different shapes and morphologies. These characteristics contribute to the aerosol mixing state, which continuously evolves in the atmosphere.

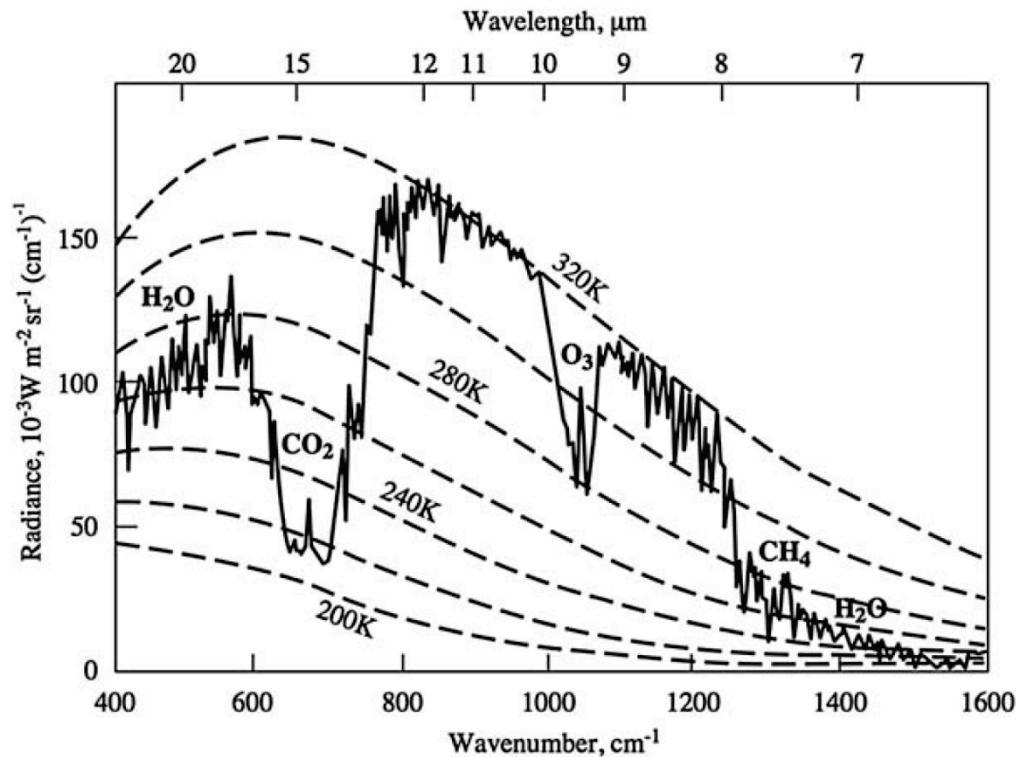
Atmospheric models have become sufficiently complex to incorporate aspects of aerosol mixing state in their predictions. Concurrently, sophisticated measurement techniques have been developed to probe various physicochemical properties of particles. However, there is no single instrument that can characterize all aspects of aerosol mixing state, and it has proven challenging to quantitatively compare measured mixing state attributes with mixing state predictions.

This talk will present a framework to synthesize a picture of the ambient aerosol from models and observations and identify current gaps in this endeavor. We will focus on suitable metrics to quantify mixing state, sampling strategies to determine these metrics that are accessible for both models and observations, and modeling strategies to scale up detailed modeling results on the particle scale to global predictions of mixing state.

<https://aosc.umd.edu/seminars/department-seminar>

Email Joseph Knisely at jknisely@umd.edu for Zoom connection info

Overview



Hanel et al., JGR, 1972:

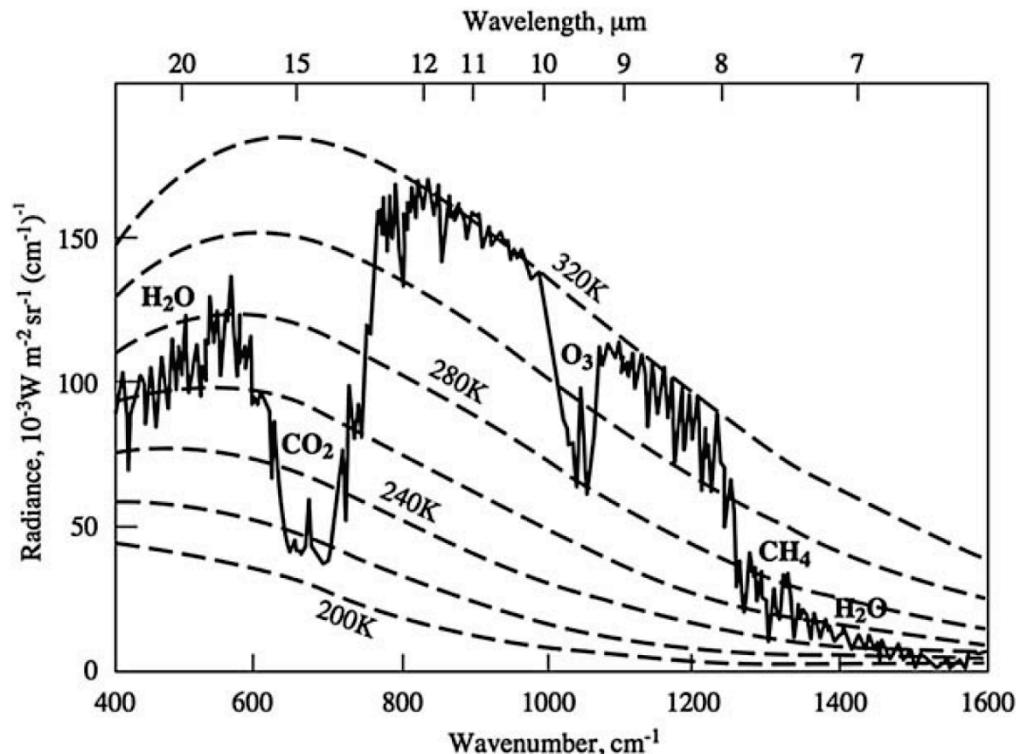
<https://agupubs.onlinelibrary.wiley.com/doi/abs/10.1029/JC077i015p02629>

Viewed from space and averaged over space and time, Earth emits $\sim 238 \text{ W/m}^2$ of thermal radiation between wavelengths of 5 and 50 μm .

The terrestrial emission spectrum matches that of a combination of blackbody spectra of temperatures between 220 and 320K.

The four most important gases that absorb terrestrial radiation (H_2O , CO_2 , CH_4 , O_3) are noted.

Overview



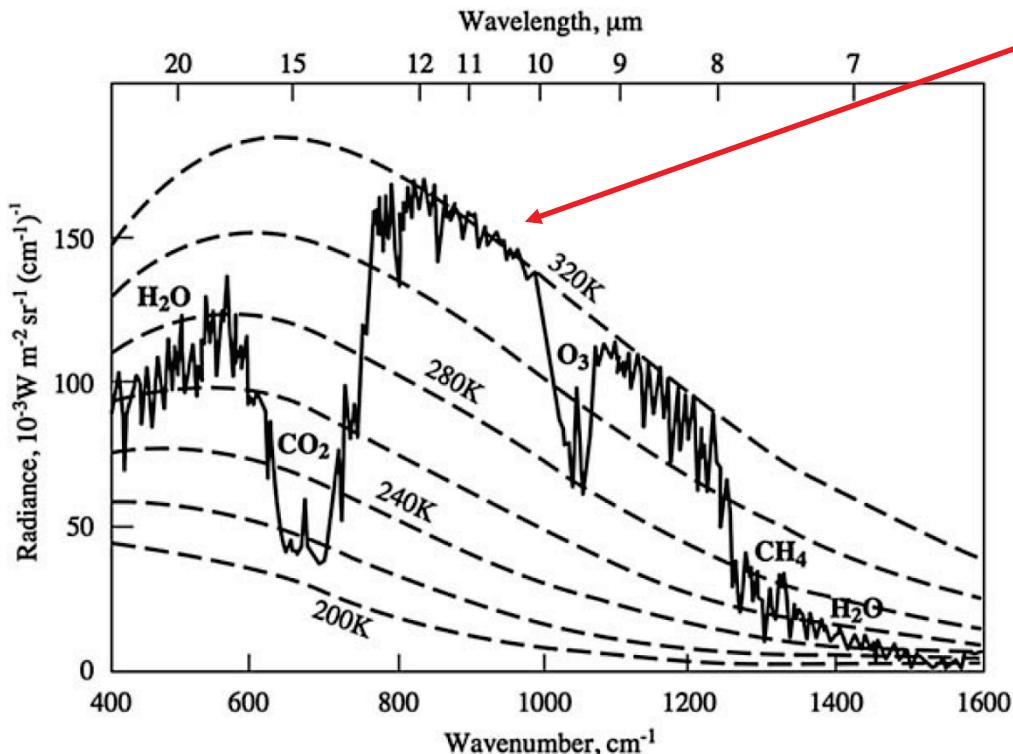
Hanel et al., JGR, 1972:

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Wavenumber = 1 / Wavelength

$$\begin{aligned}1 / 1600 \text{ cm}^{-1} &= 6.25 \times 10^{-4} \text{ cm} = 6.25 \times 10^{-6} \text{ m} = 6.25 \mu\text{m} \\1 / 400 \text{ cm}^{-1} &= 2.50 \times 10^{-3} \text{ cm} = 2.50 \times 10^{-5} \text{ m} = 25.0 \mu\text{m}\end{aligned}$$

Overview



320 K = 47°C = 116.6°F
where was this image acquired?

Hanel et al., JGR, 1972:

<https://agupubs.onlinelibrary.wiley.com/doi/abs/10.1029/JC077i015p02629>

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Overview

News & Blogs

CATEGORY 6™

NEW STORIES

INFOGRAPHICS

POSTERS

Africa's Hottest Reliably Measured Temperature on Record: 124.3°F on Thursday in Algeria

Dr. Jeff Masters · July 6, 2018, 8:08 AM EDT

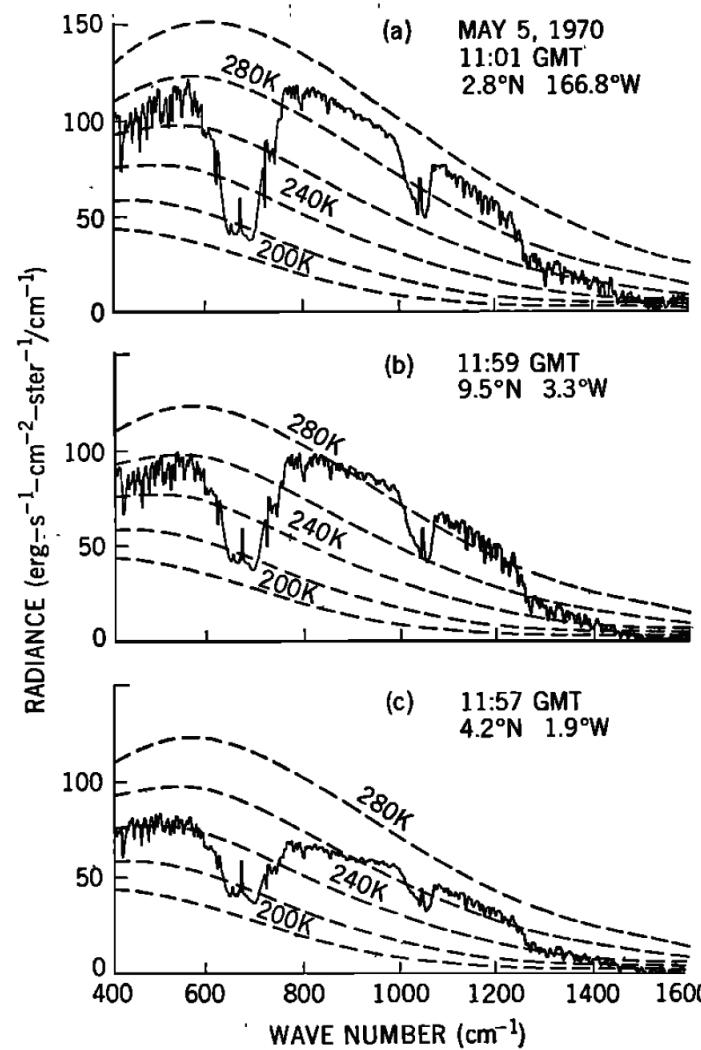
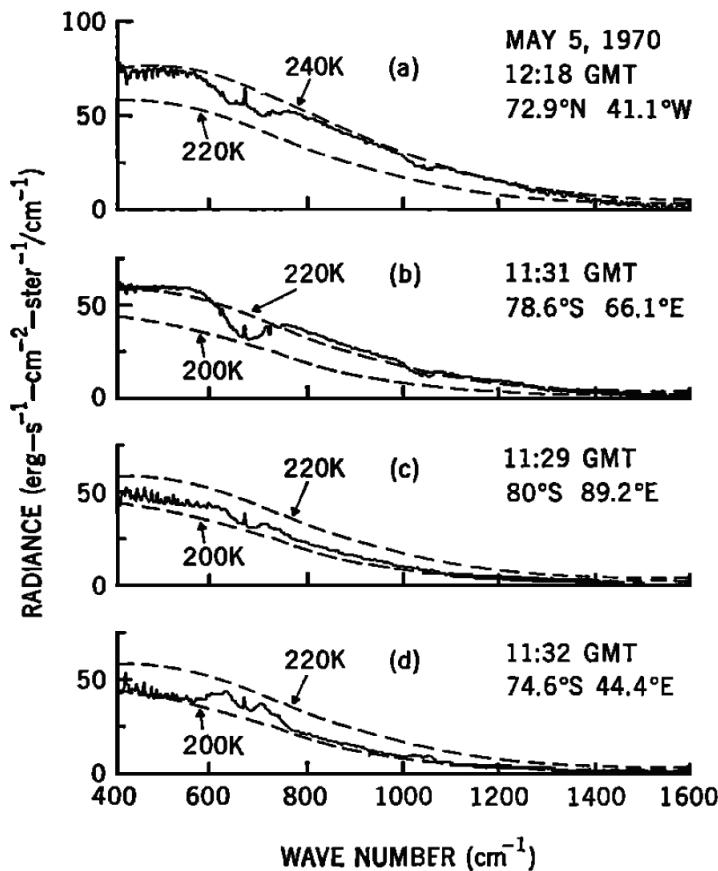
A historic heat wave in northern Africa on Thursday, July 5, brought Africa its hottest reliably measured temperature on record: 124.3°F (51.3°C), at [Ouargla, Algeria](#). Ouargla (population 190,000) is the capital city of Ouargla Province in the Algerian Sahara Desert, at an elevation of 719 feet (219 meters).



Location of Ouargla, Algeria, indicated by red marker. (Google)

<https://www.wunderground.com/cat6/Africas-Hottest-Reliably-Measured-Temperature-Record-1243F-Thursday-Algeria>

<https://www.washingtonpost.com/news/capital-weather-gang/wp/2018/07/06/africa-may-have-witnessed-its-all-time-hottest-temperature-thursday-124-degrees-in-algeria>



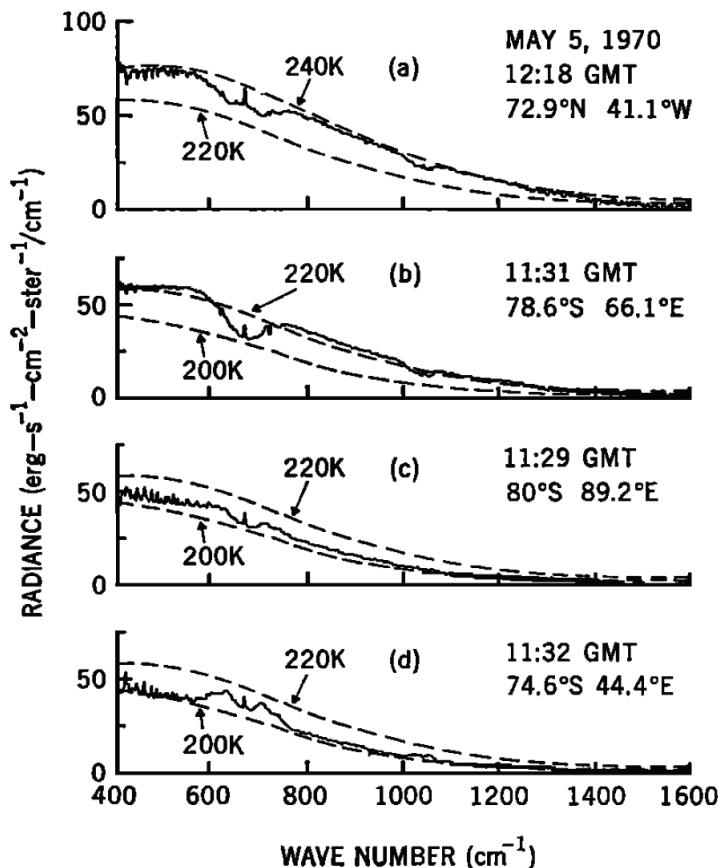
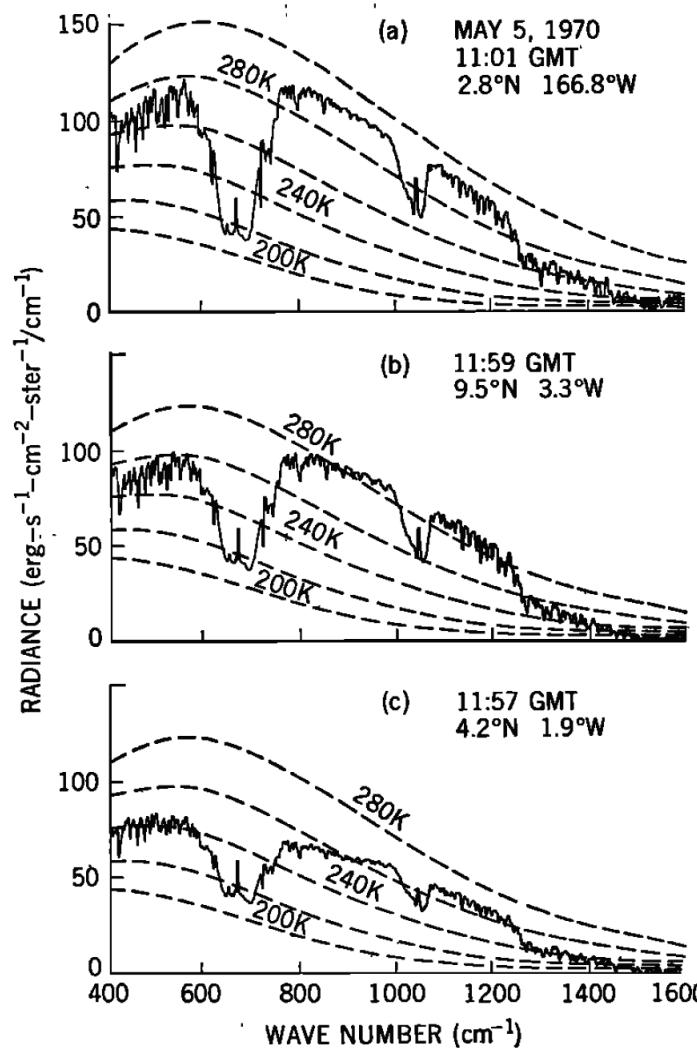


Fig. 12. Examples of polar spectra. The spectrum in *a* was obtained over Greenland; the spectra in *b*, *c*, and *d* were obtained over Antarctica.



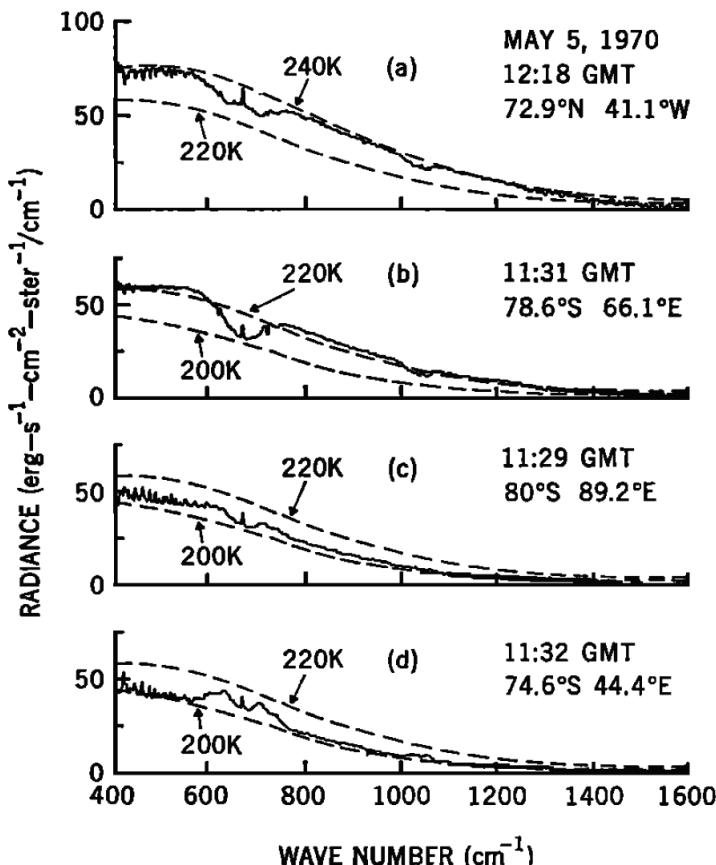


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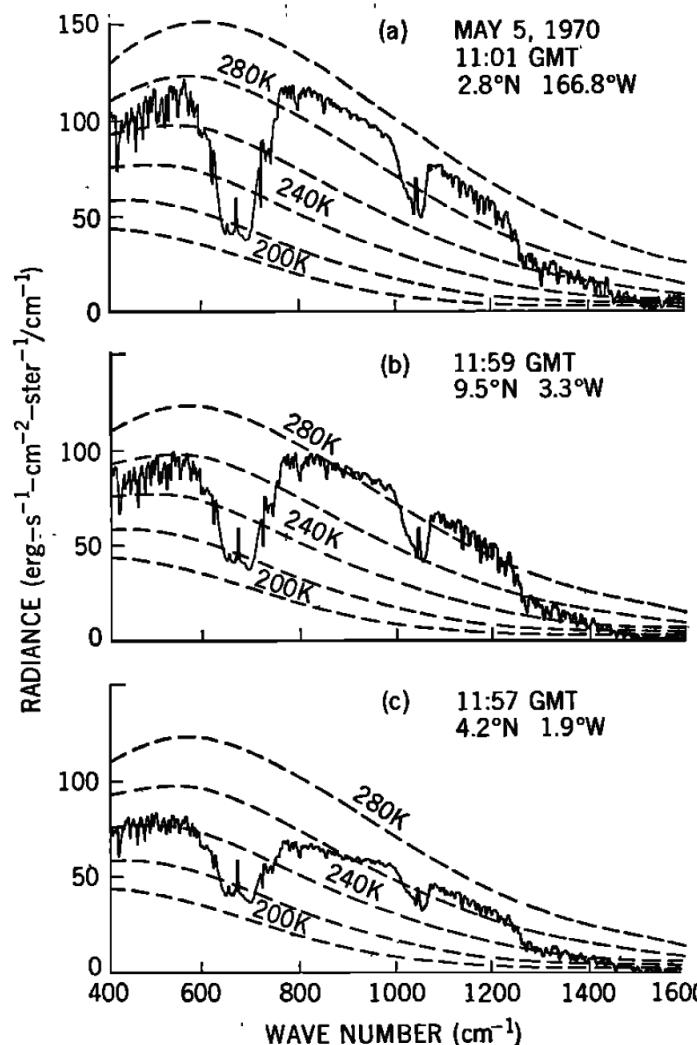


Fig. 13. Sample spectra illustrating effects of clouds in the window region of 800 to 1000 cm^{-1} . In this spectral region, a pronounced departure from a blackbody curve is noted in *b* and *c* in contrast to the spectra shown in Figure 14, for example.

Overview

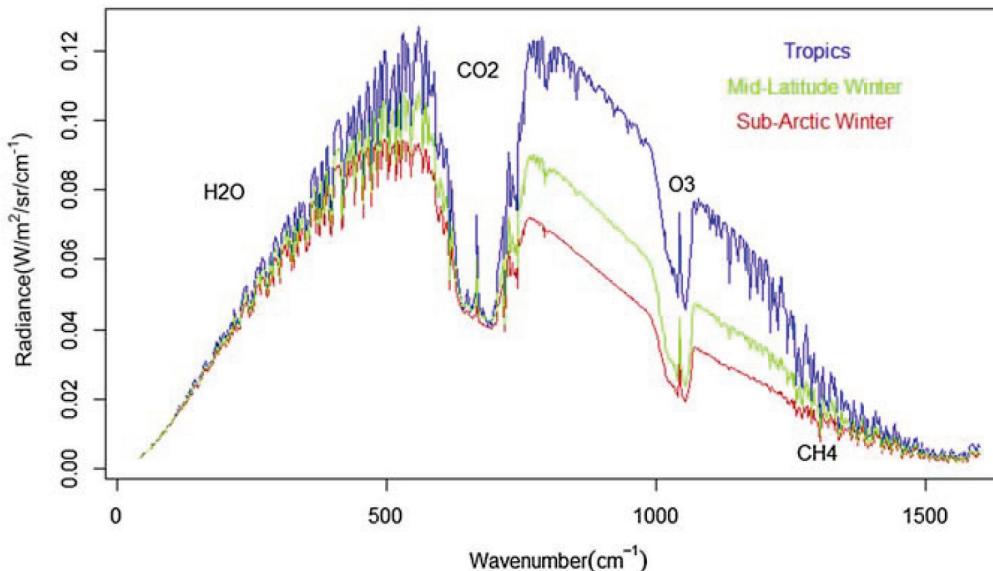


FIGURE 3.4.5 Overview of the earth's outgoing infrared radiation as a function of wave number (the inverse of wavelength) and latitude.⁴³ Radiances for this figure were calculated using Modtran and a web interface developed by David Archer available here: <http://climatedmodels.uchicago.edu/modtran/>.

Kirk-Davidoff, Chapter 3.4, *Green Chemistry: An Inclusive Approach*, 2018

- GHGs prevent outgoing energy emitted from the surface from being released back into space, thereby trapping this energy and releasing it in the form of heat.
- Averaged over space and time, the Earth radiates to space an amount of energy consistent with that of a black body at 255 K.
- Some spectral regions are nearly filled (i.e., 667 cm^{-1}) whereas many others exhibit negligible attenuation of outgoing radiation.
- A newly discovered “miracle compound” with a long atmospheric lifetime will be much more damaging to Earth’s climate system if it absorbs in a region that is _____, rather than a region that is _____.

Global Warming Potential

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Examples of Greenhouse Gases					
Name and Chemical Formula	Preindustrial Concentration (1750)	Concentration in 2008	Atmospheric Lifetime (years)	Anthropogenic Sources	Global Warming Potential
carbon dioxide CO ₂	270 ppm	388 ppm	50-200*	Fossil fuel combustion, deforestation, cement production	1
methane CH ₄	700 ppb	1760 ppb	12	Rice paddies, waste dumps, livestock	21
nitrous oxide N ₂ O	275 ppb	322 ppb	120	Fertilizers, industrial production, combustion	310
CFC-12 CCl ₂ F ₂	0	0.56 ppb	102	Liquid coolants, foams	8100

*A single value for the atmospheric lifetime of CO₂ is not possible. Removal mechanisms take place at different rates. The range given is an estimate based on several removal mechanisms.

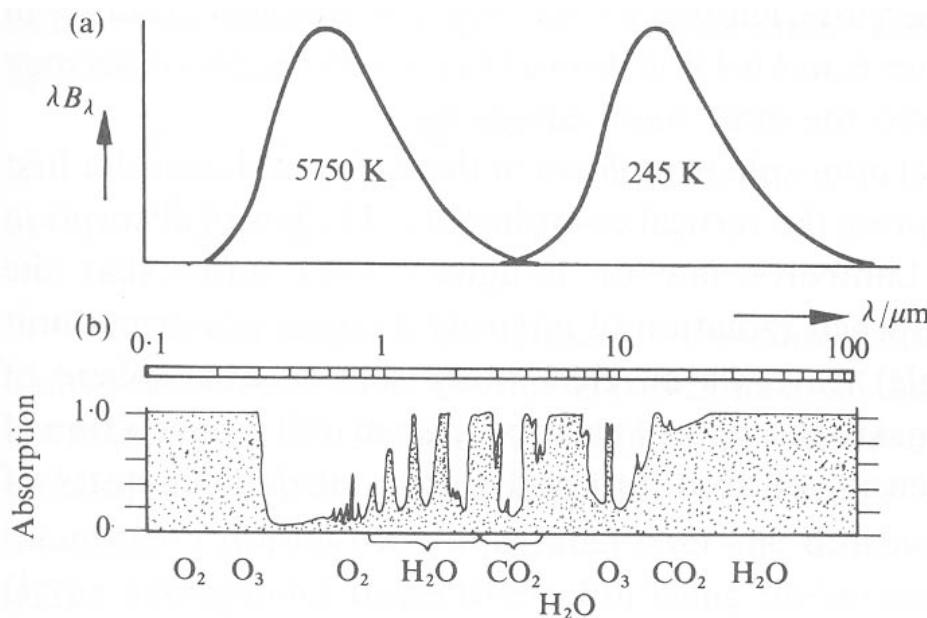
Chapter 3, *Chemistry in Context*

100 year time horizon

Some GHGs are much more effective than others, in terms of GWP (i.e., perturbation of RF per mass)

Atmospheric Radiation

- Solar irradiance (downwelling) at top of atmosphere occurs at wavelengths between ~200 and 2000 nm (~5750 K “black body” temperature)
- Thermal irradiance (upwelling) at top of the atmosphere occurs at wavelengths between ~5 and 50 μm (~245 K “black body” temperature for Earth’s atmosphere)



Panel (a): Curves of black-body energy versus wavelength for 5750 K (Sun’s approximate temperature) and for 245 K (Earth’s mean temperature). The curves are drawn with equal area since, integrated over the entire Earth at the top of the atmosphere, the solar (downwelling) and terrestrial (upwelling) fluxes must be equal.

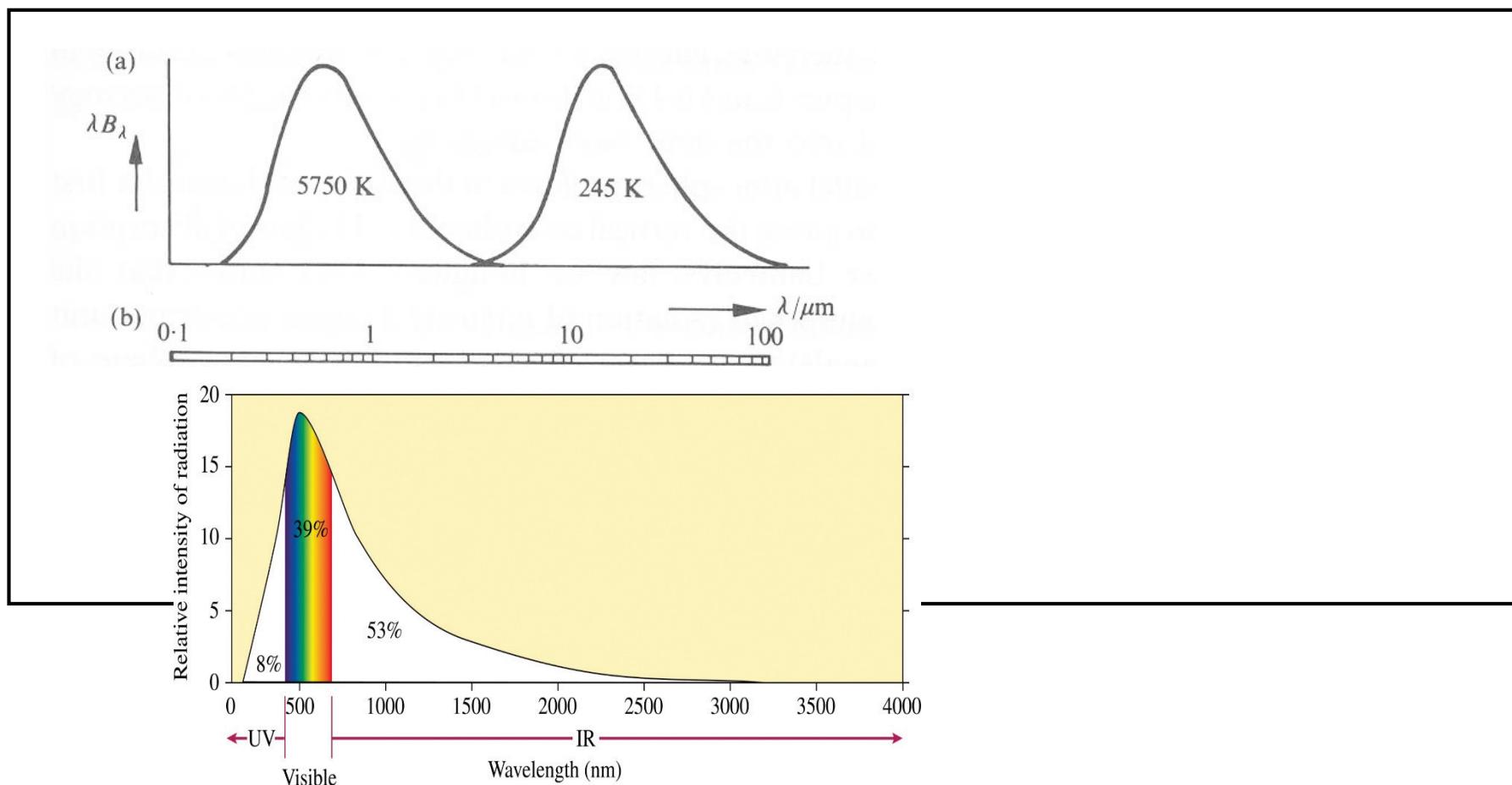
Panel (b): absorption by atmospheric gases for a clear vertical column of the atmosphere (1.0 represents complete absorption).

From Houghton, *Physics of Atmospheres*, 1991

- Absorption and photodissociation in the UV occurs due to changes in the electronic state (orbital configuration of electrons) of molecules
- Absorption and re-emission in the IR occurs due to changes in vibrational and rotational states of molecules with electric dipole moments

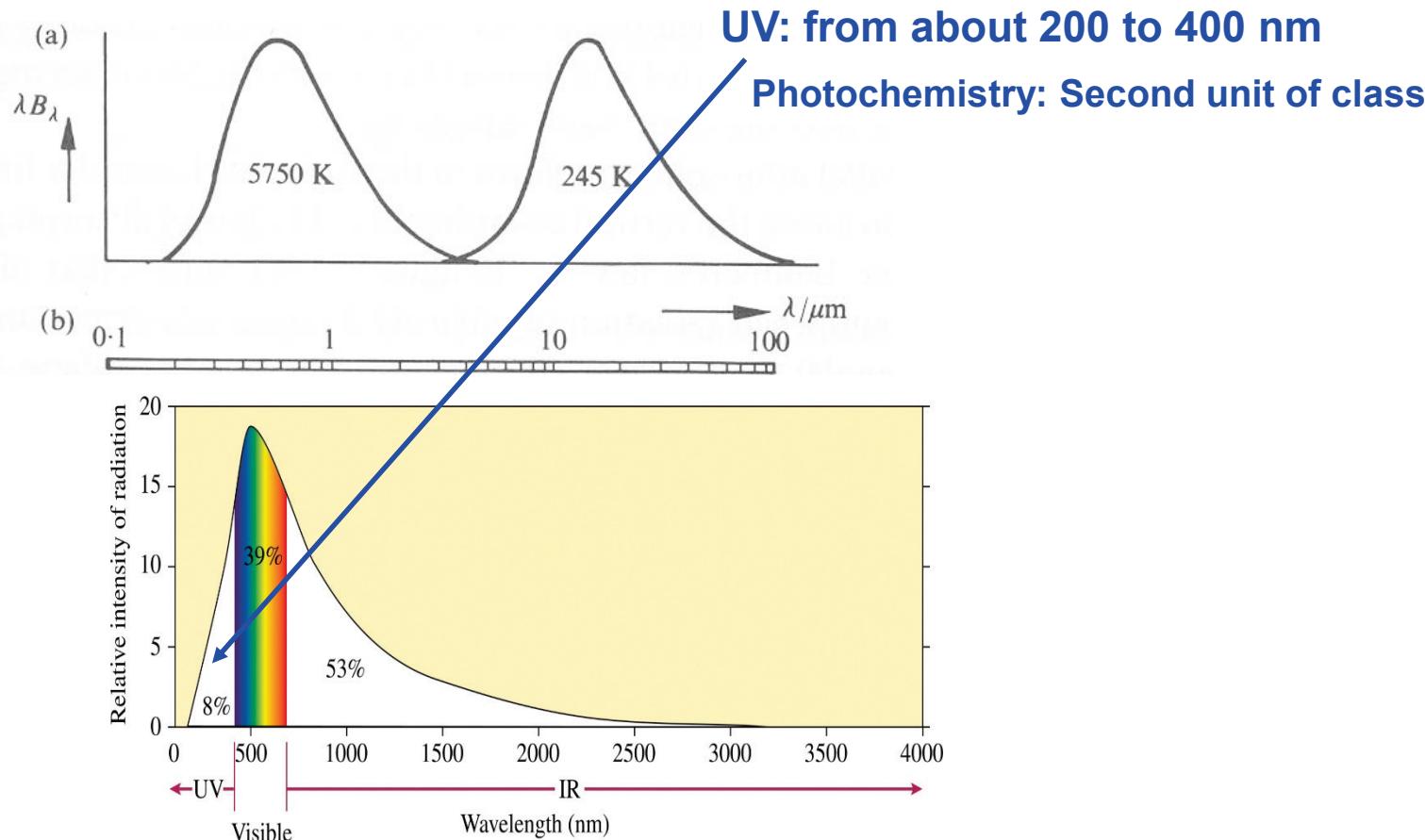
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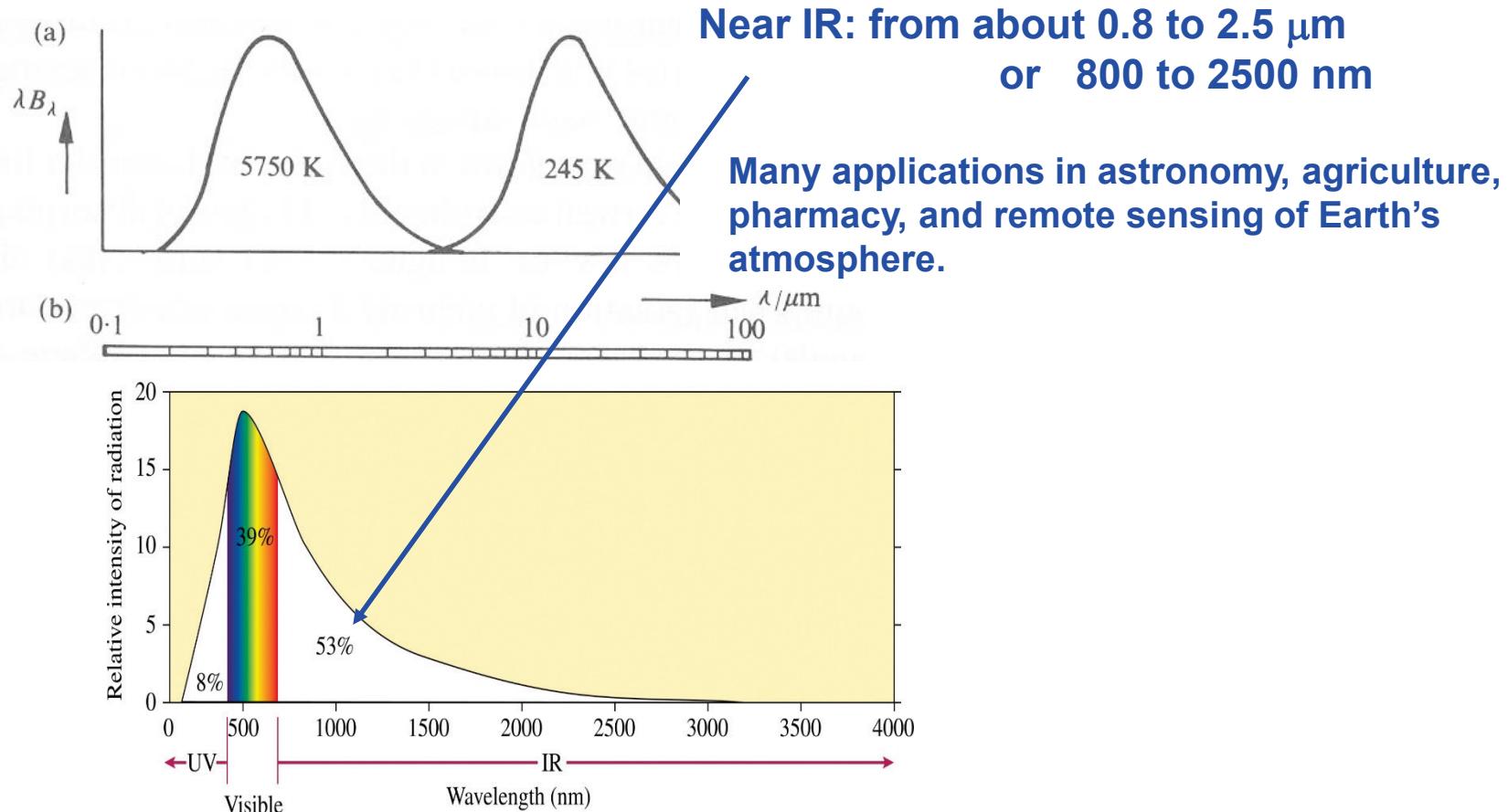
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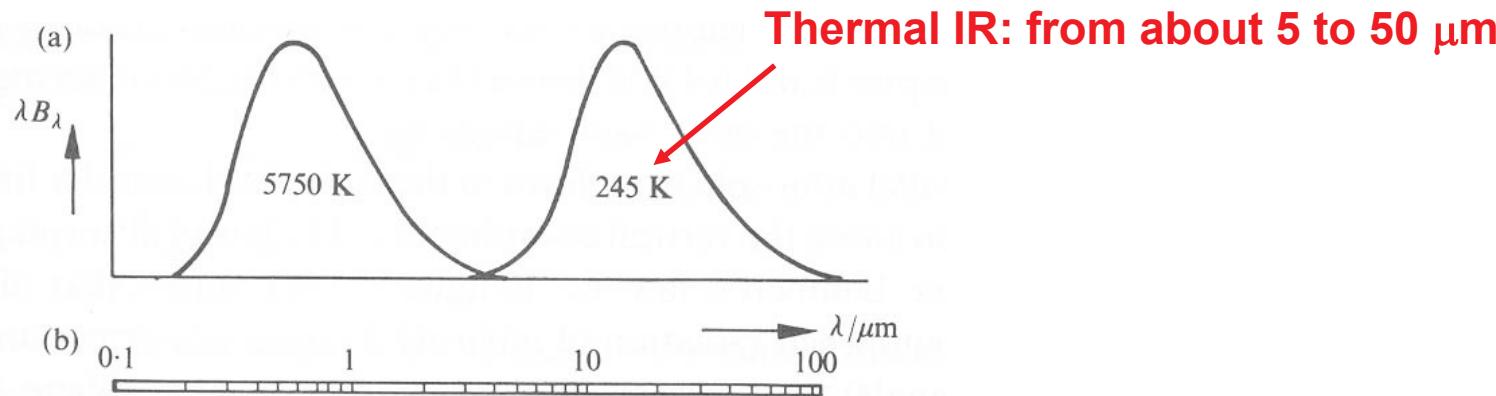
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Radiation & Molecules

Radiation can induce photo-dissociation (March 10 lecture), vibration, and rotation of molecules.

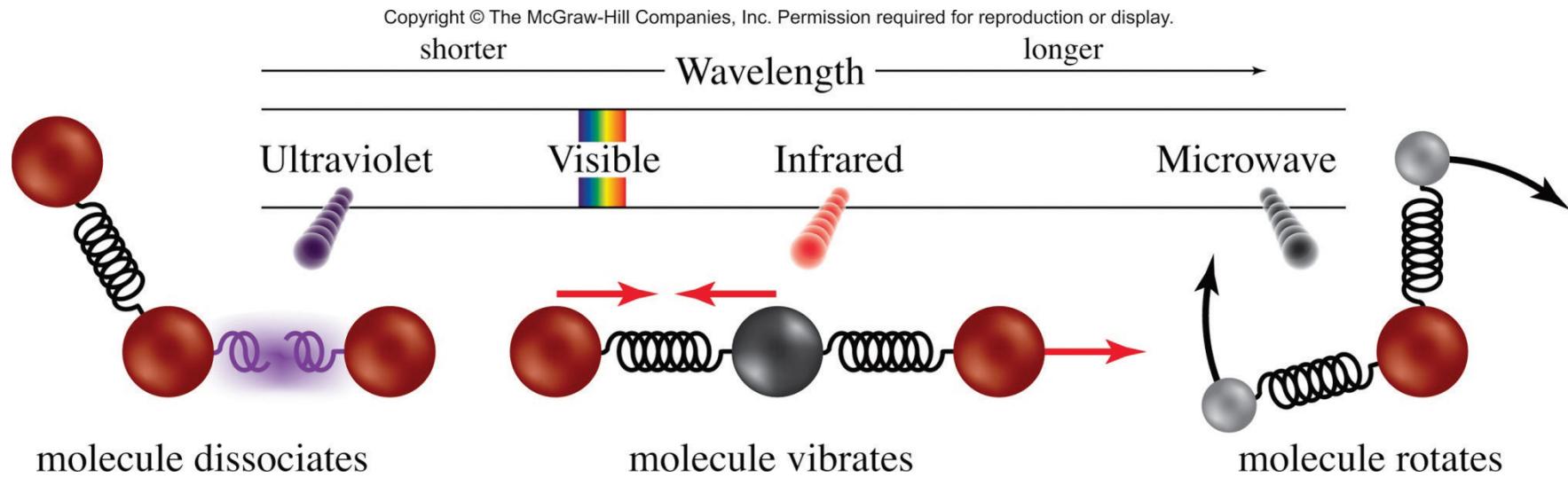


Fig 3.19, Chemistry in Context

Radiation & Molecules

Radiation can induce photo-dissociation (March 10 lecture), **vibration**, and rotation of molecules.

Thermal IR radiation is not energetic enough to break molecular bonds (i.e., photo-dissociate). Upon absorption, thermal IR will increase the vibrational energy of a molecule

CO₂ (linear molecule) has 4 vibrational modes (see below): for molecules vibrational frequencies are quantized. That is, only certain energies for the system are allowed. Most importantly, only photons with certain wavelengths (energies) will excite molecular vibrations.

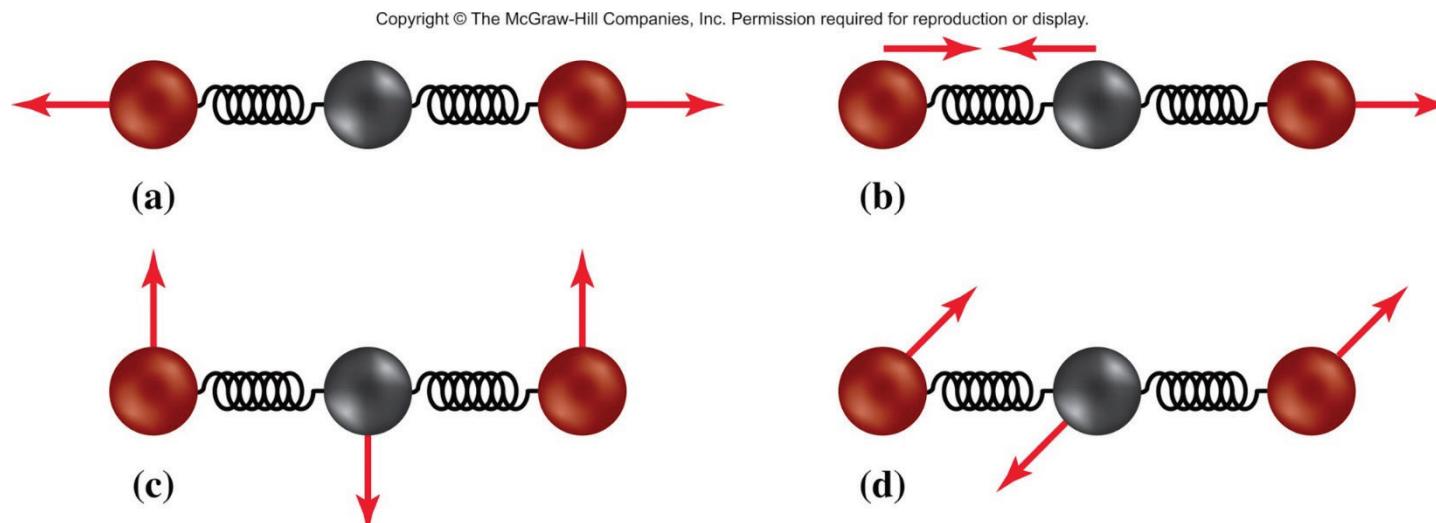


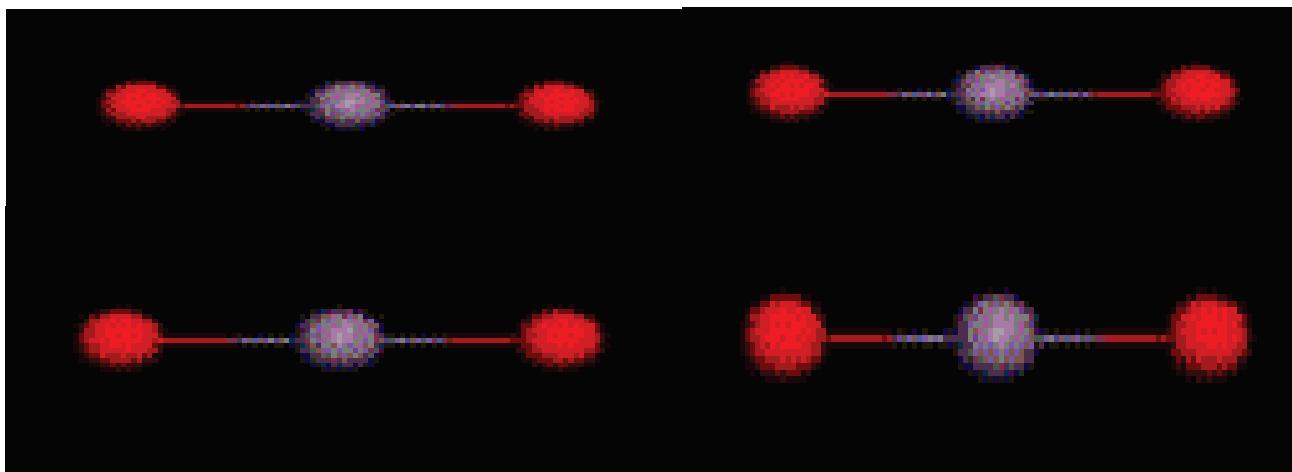
Fig 3.16, Chemistry in Context

Radiation & Molecules

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http://science.widener.edu/svb/ftir/ir_co2.html

Excitation of Molecules

A greenhouse gas must have either

- naturally occurring **dipole moment**
- exhibit a **dipole moment** during vibration

Dipole moment \Rightarrow product of magnitude of charges & distance of separation between charges:
i.e., a molecule is said to have a dipole moment if it has a non-zero spatial distribution of charge

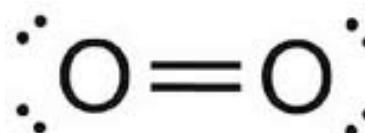
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No dipole moment, either naturally or during vibration:



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CO₂ has no natural dipole moment

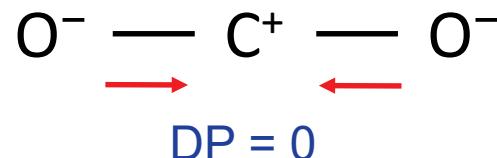


Fig 3.14, Chemistry in Context

Excitation of Molecules

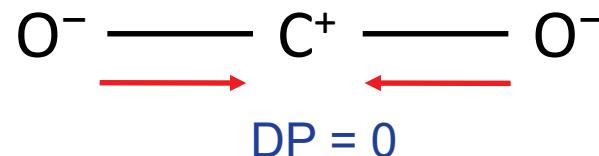
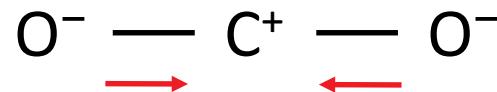
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Symmetric Stretch: no dipole moment

Symmetric stretch



Excitation of Molecules

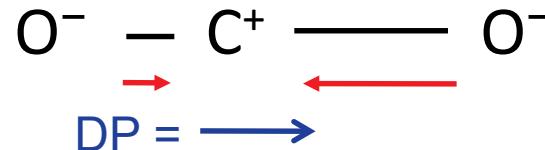
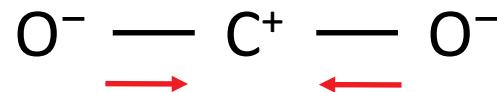
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Anti-symmetric Stretch: dipole moment

Anti-symmetric stretch

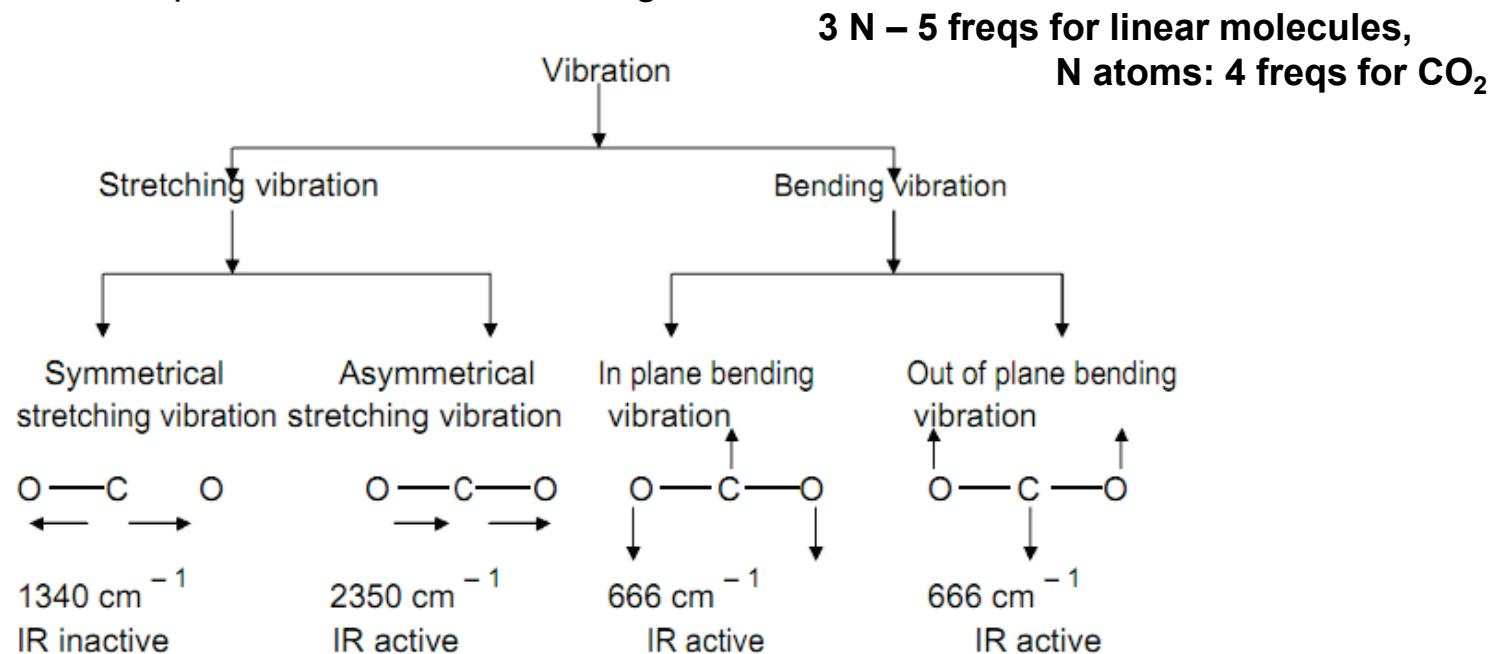


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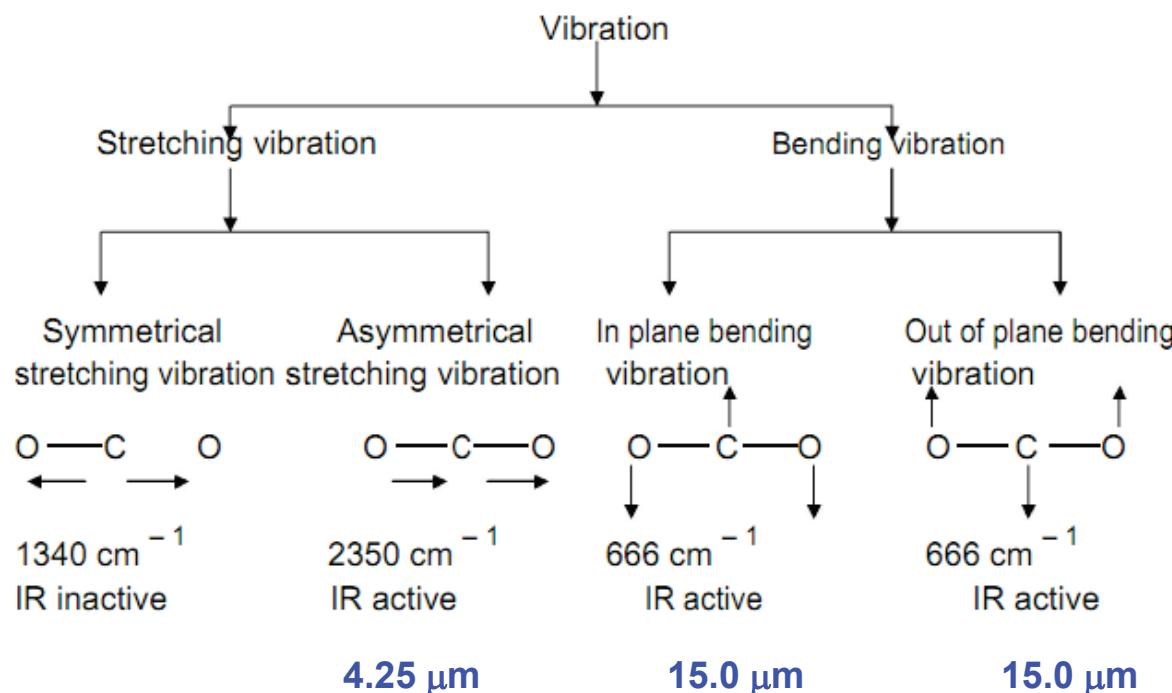


Excitation of Molecules

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$$1 / 666 \text{ cm}^{-1} = 1.50 \times 10^{-3} \text{ cm} = 15.0 \times 10^{-6} \text{ m} = 15.0 \mu\text{m}$$



<http://www.vidyarthiplus.in/2013/12/cy6151-engineering-chemistry-1.html#.VOUqai4RXIY>

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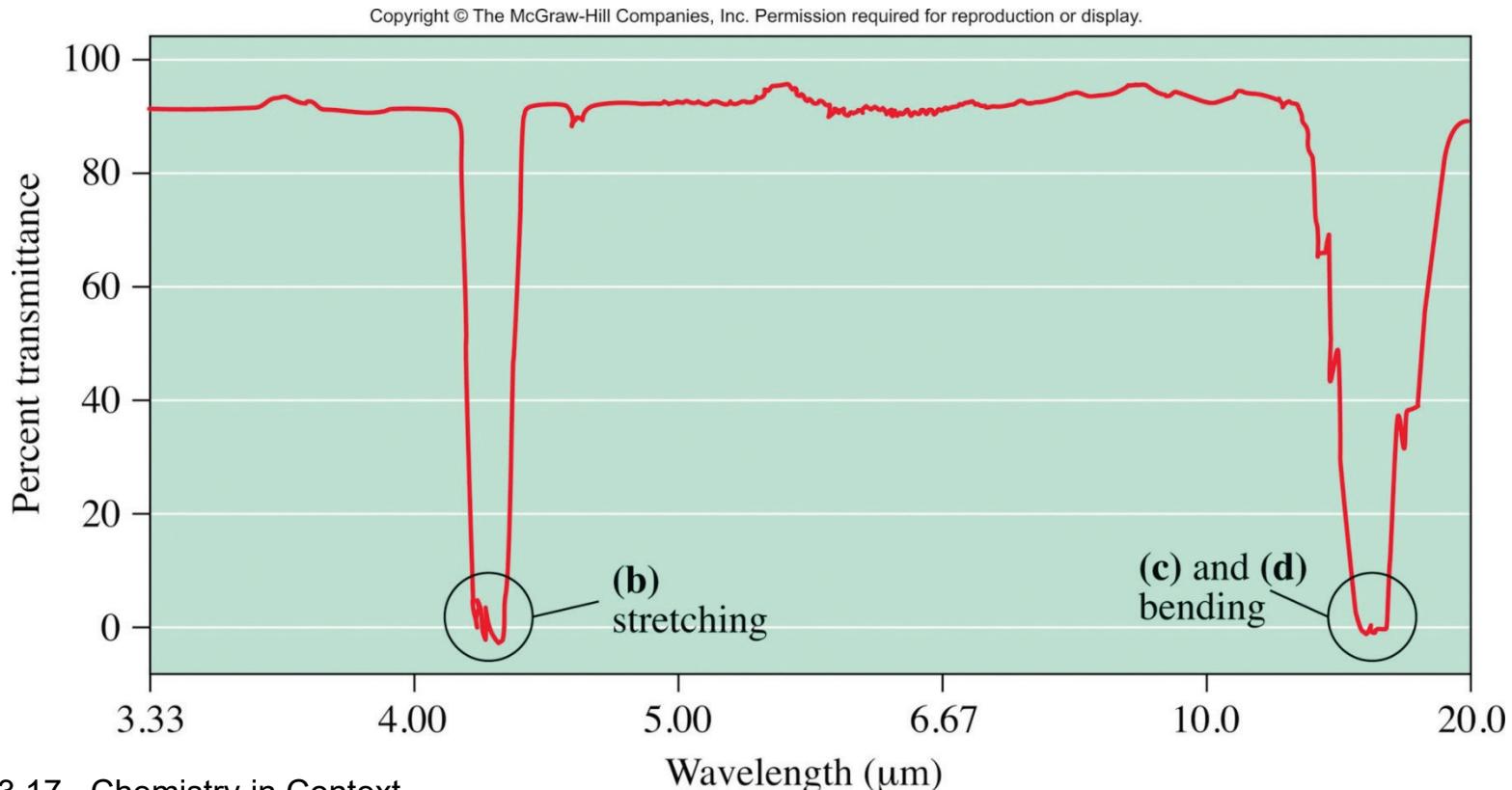


Fig 3.17, Chemistry in Context

Excitation of Molecules

Which transition involves more energy?

$$E = hc / \lambda$$

h (Planck's constant) = 6.62×10^{-27} erg sec; c (speed of light) = 3×10^8 cm/sec

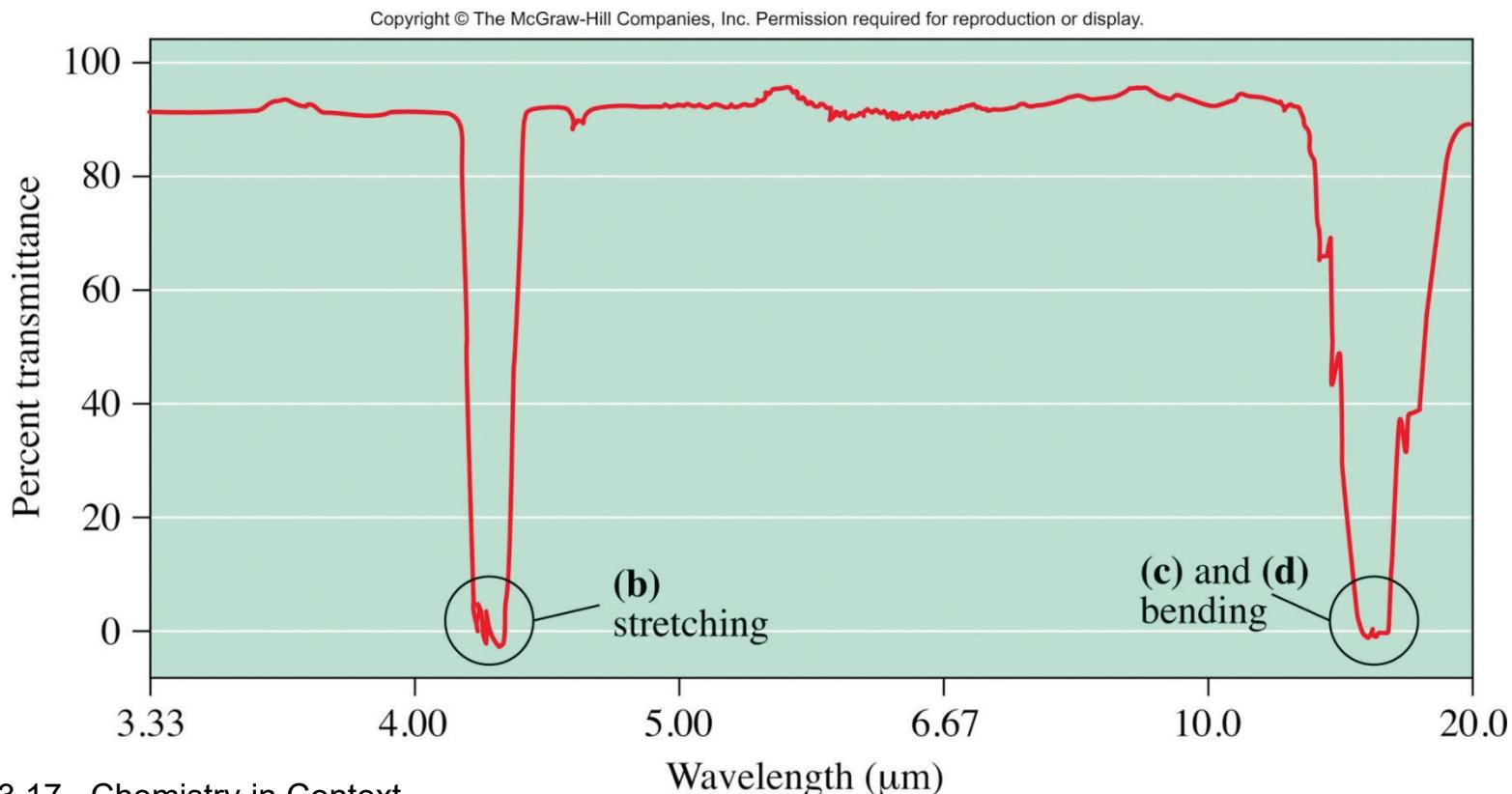


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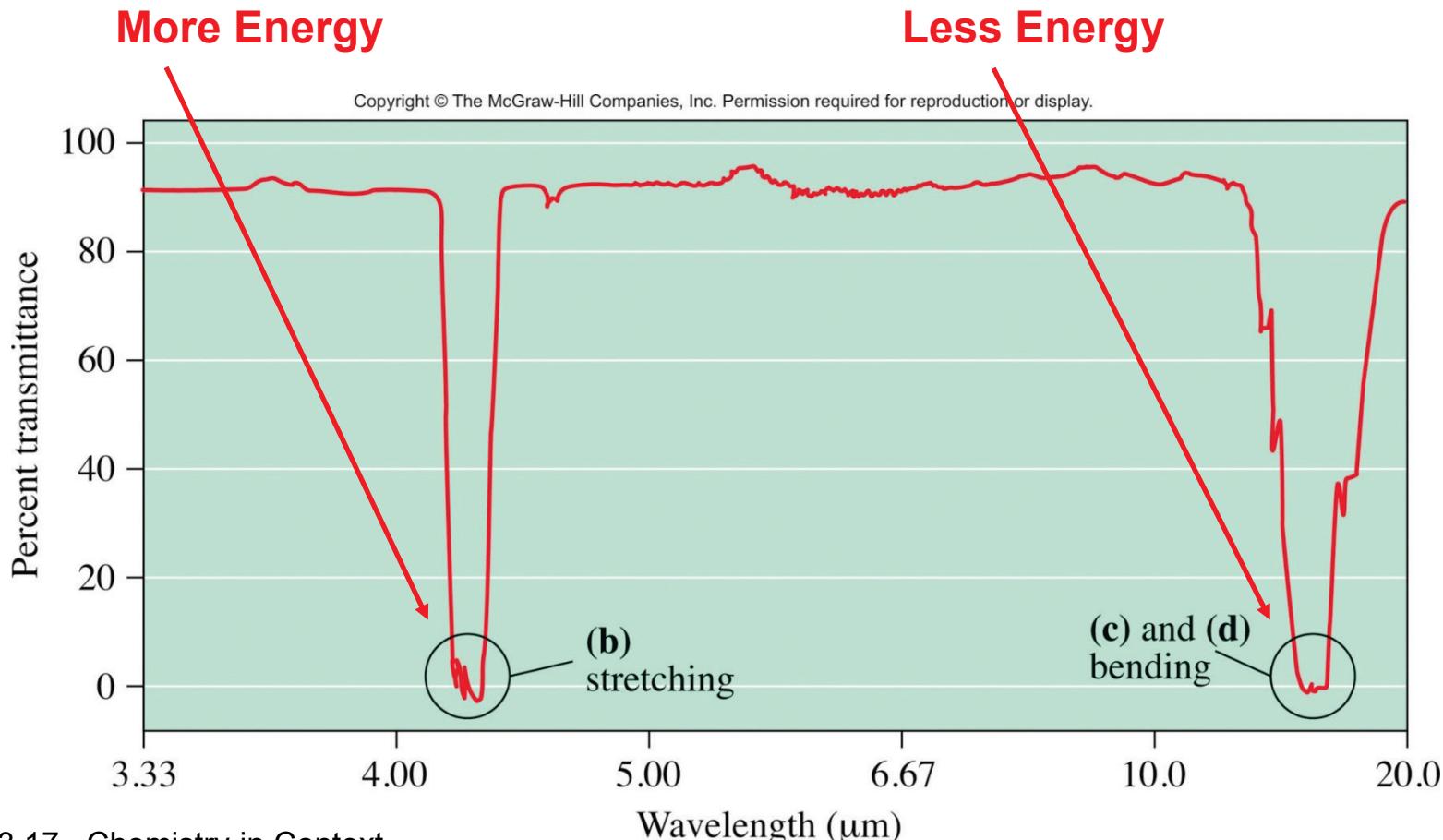


Fig 3.17, Chemistry in Context

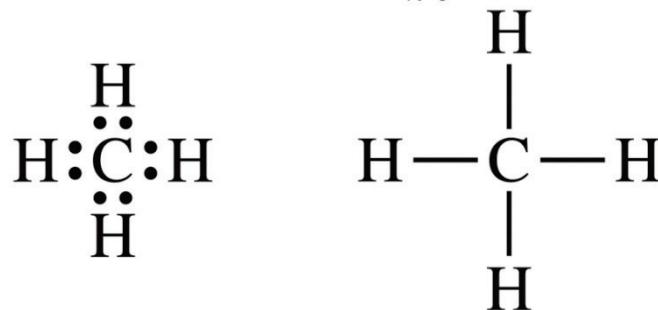
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CH₄ also has no natural dipole moment: charge is uniformly distributed



Figs 3.10 & 3.11, Chemistry in Context

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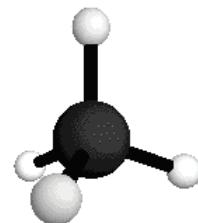
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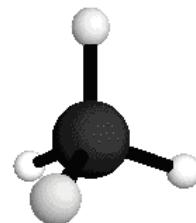
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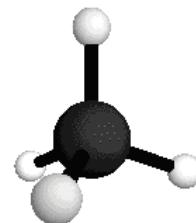
CH₄ has 4 unique vibrational modes, 2 of which interact with the IR field



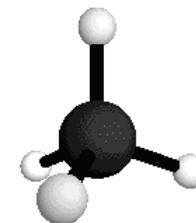
#1
3.3 μm



#2
6.3 μm



#3
3.2 μm



#4
7.6 μm

http://www2.ess.ucla.edu/~schauble/MoleculeHTML/CH4_html/CH4_page.html

Excitation of Molecules

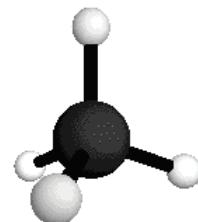
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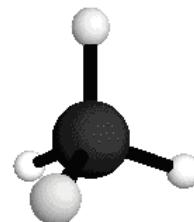
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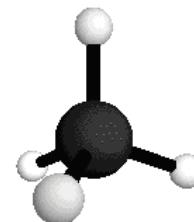
Only modes for which the C and H atoms both move are radiatively active



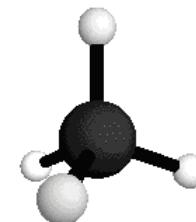
#1
3.3 μm



#2
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- naturally occurring **dipole moment**
- exhibit a **dipole moment** during vibration

Dipole moment \Rightarrow product of magnitude of charges & distance of separation between charges:
i.e., a molecule is said to have a dipole moment if it has a non-zero spatial distribution of charge

H₂O has a natural dipole moment (bent molecule) and absorbs in three spectral regions:

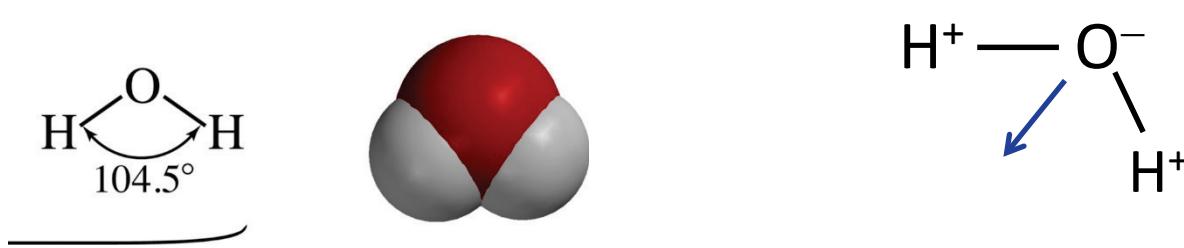


Fig 3.13, Chemistry in Context

Excitation of Molecules

A greenhouse gas must have either

- naturally occurring **dipole moment**
- exhibit a **dipole moment** during vibration

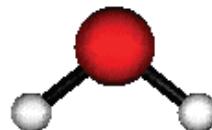
Dipole moment \Rightarrow product of magnitude of charges & distance of separation between charges:
i.e., a molecule is said to have a dipole moment if it has a non-zero spatial distribution of charge

H₂O has a natural dipole moment (bent molecule) and absorbs in three spectral regions:



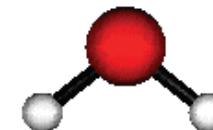
2.5 μm

Asymmetric
Stretch



2.6 μm

Symmetric
Stretch



6.1 μm

Bending
Mode

http://www2.ess.ucla.edu/~schauble/MoleculeHTML/H2O_html/H2O_page.html

Excitation of Molecules

A greenhouse gas must have either

- naturally occurring **dipole moment**
- exhibit a **dipole moment** during vibration

Dipole moment \Rightarrow product of magnitude of charges & distance of separation between charges:
i.e., a molecule is said to have a dipole moment if it has a non-zero spatial distribution of charge

N₂O also has a natural dipole moment (since it is an asymmetric molecule) and also absorbs in three spectral regions:



4.5 μm



7.8 μm



17.0 μm

http://www2.ess.ucla.edu/~schauble/MoleculeHTML/N2O_html/N2O_page.html

Absorption vs. Wavelength

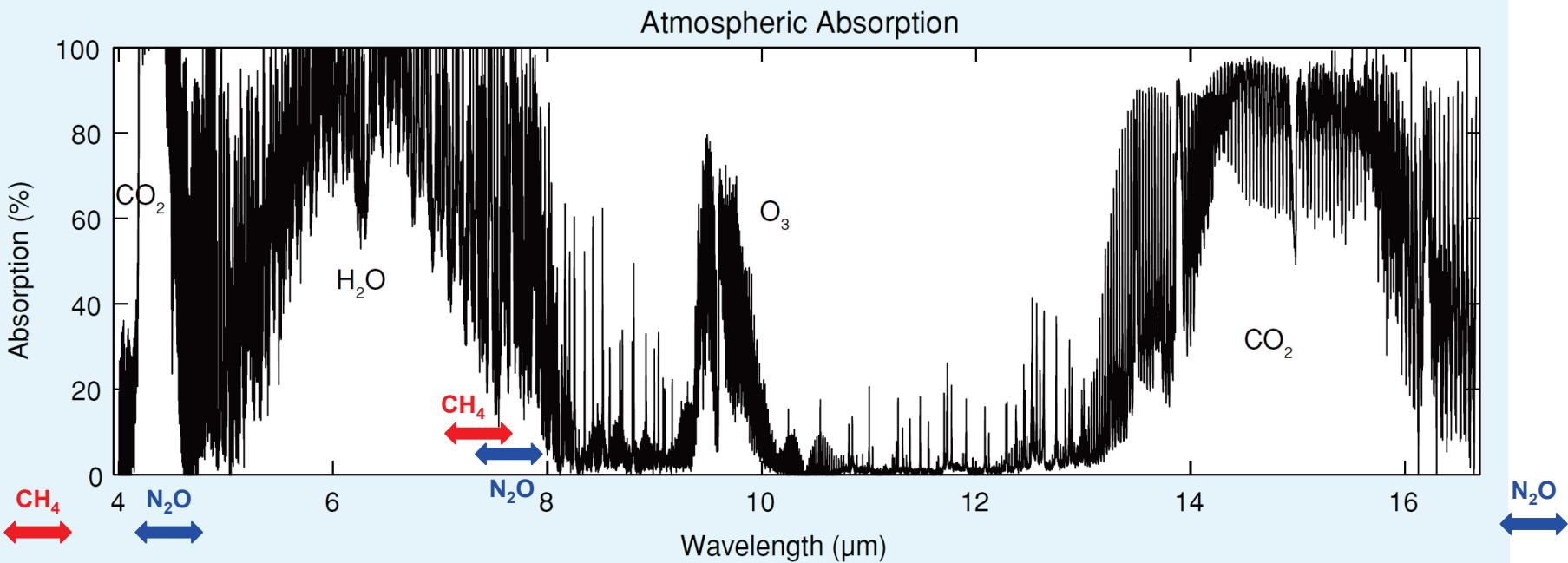


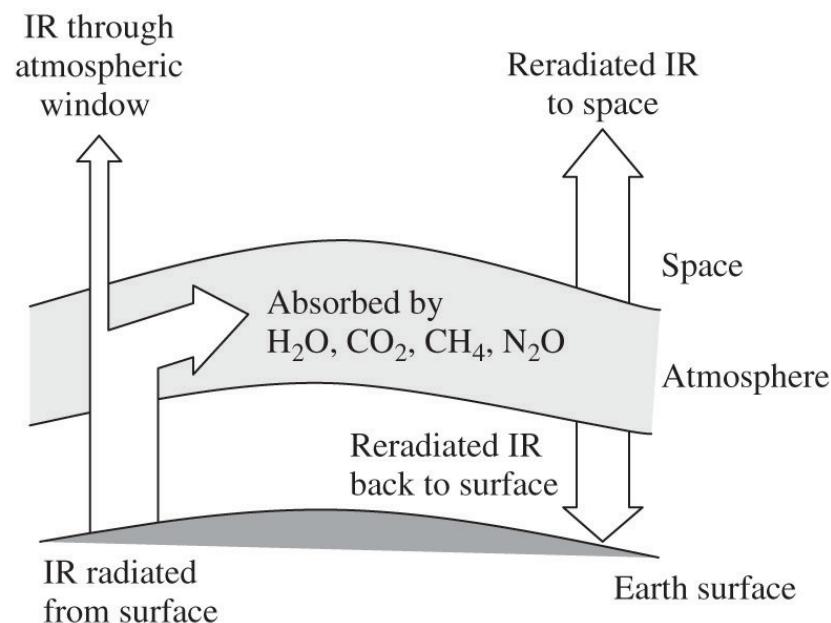
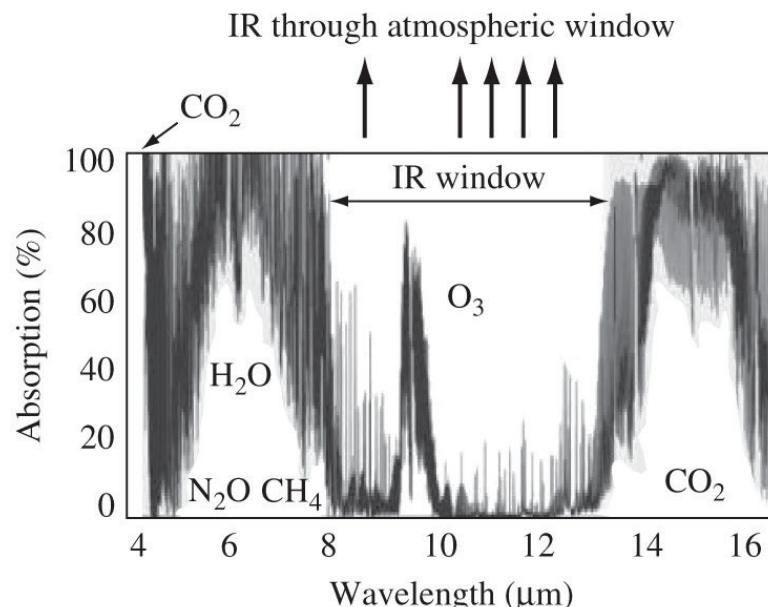
Fig 2.6, IPCC SROC (Special Report on Ozone layer and Climate), 2005
<https://www.ipcc.ch/site/assets/uploads/2018/03/sroc02-1.pdf>

The Greenhouse Effect

Molecules of that absorb specific wavelengths of IR energy experience different fates:

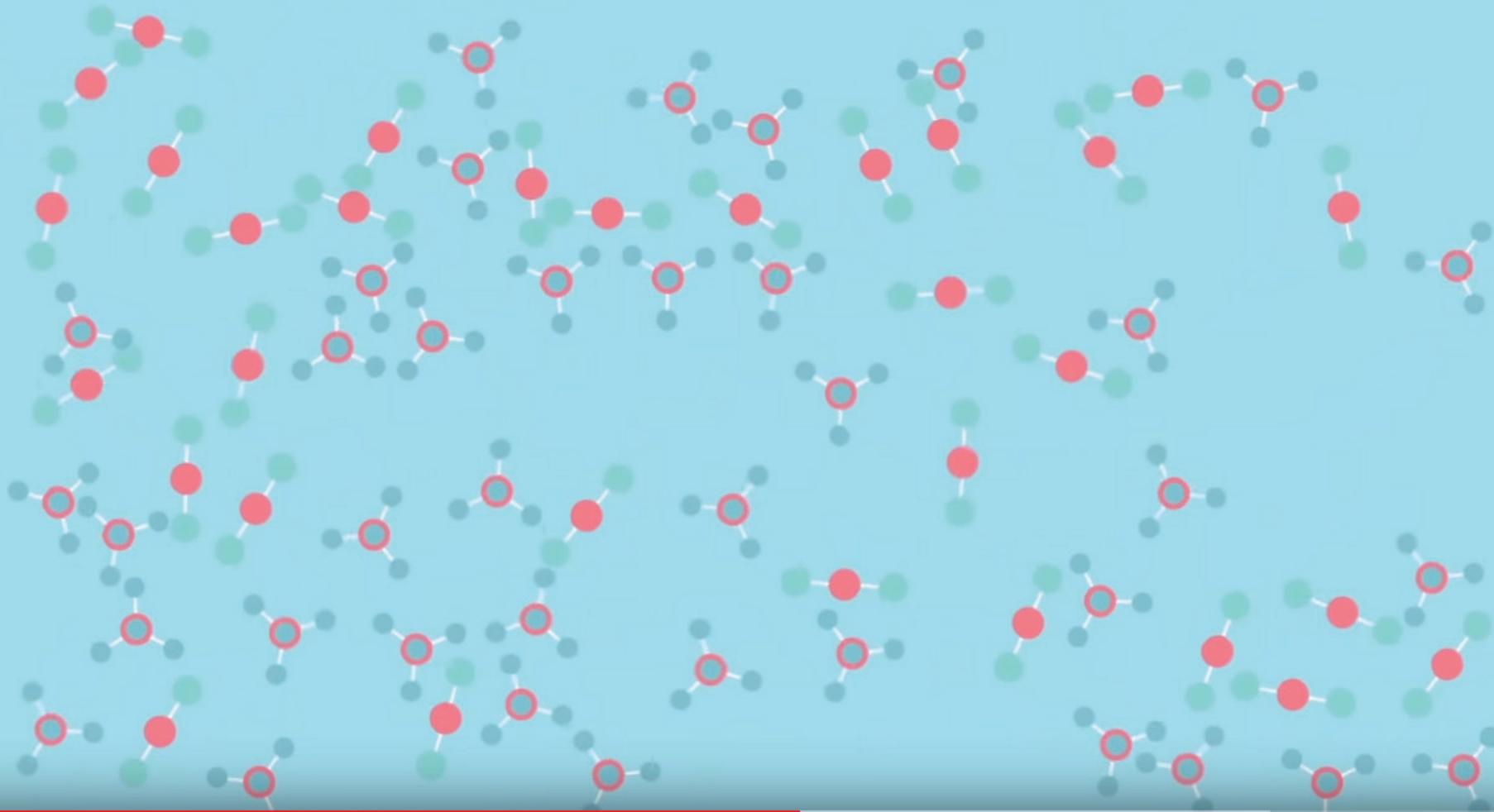
- Some hold that extra energy for a brief time, then re-emit it in all directions as heat.
- Others collide with atmospheric molecules such as N_2 and O_2 and transfer the absorbed energy to those molecules, as heat

Both processes “trap” radiation emitted by the Earth; this trapping of energy heats the lower atmosphere and surface



Masters, Intro. to Environmental Engineering and Science, 3d ed.

See Chapter 3.4 by Dan Kirk-Davidoff,
in *Green Chemistry: An Inclusive Approach*, 2018
in Additional Readings for a simple, differential equation description of the GHG effect
based on a so-called two layer model.



▶ ▶ 🔍 1:41 / 3:08

CC HD □ ☰

How Do Greenhouse Gases Actually Work?

1,312,336 views • May 26, 2015

like 25K

dislike 550

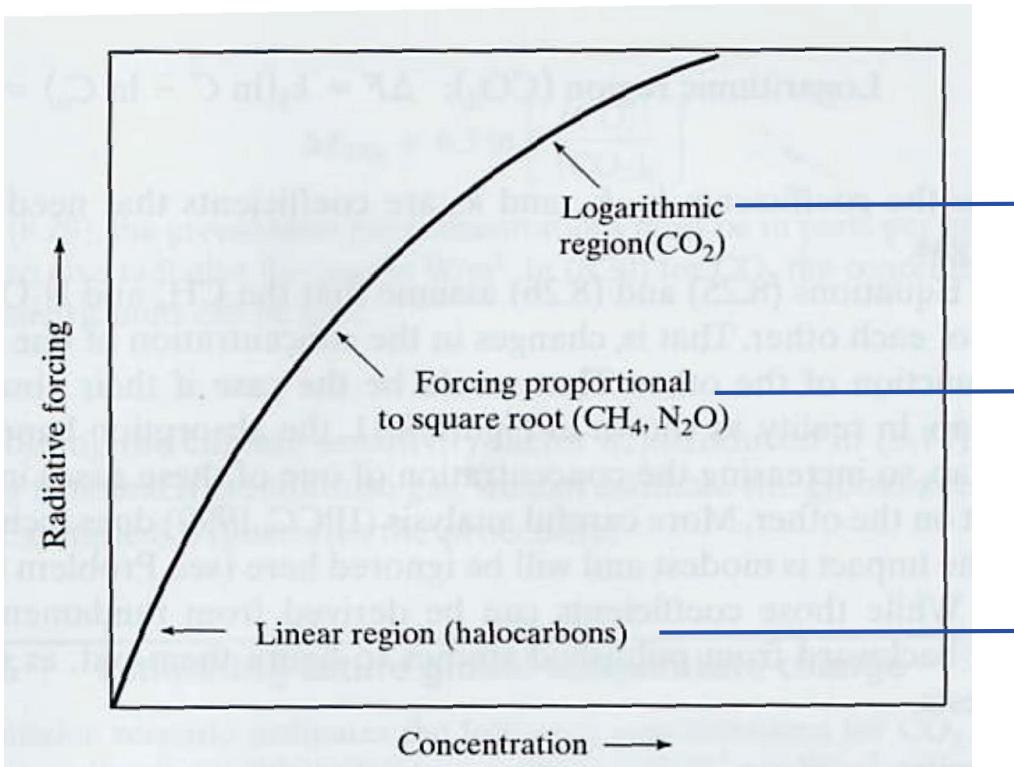
SHARE

SAVE

...

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

How does RF change with concentration?



Wigley (1987)

$$\Delta RF = \alpha \ln \left(\frac{C}{C_o} \right)$$

$$\Delta RF = \alpha \left(\sqrt{C} - \sqrt{C_o} \right)$$

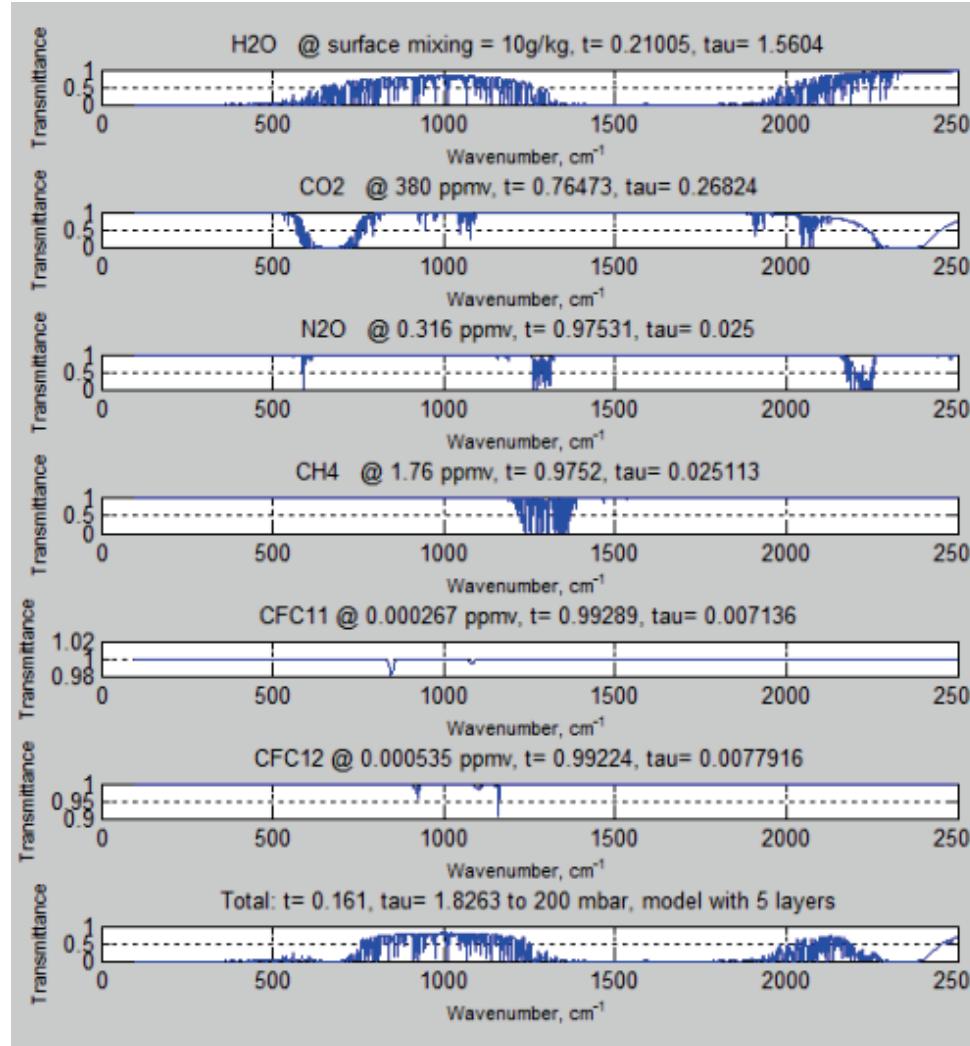
$$\Delta RF = \alpha (C - C_o)$$

Masters, Introduction to Environmental Engineering and Science, 1998

Effectiveness of a GHG depends on “saturation” of absorption band.

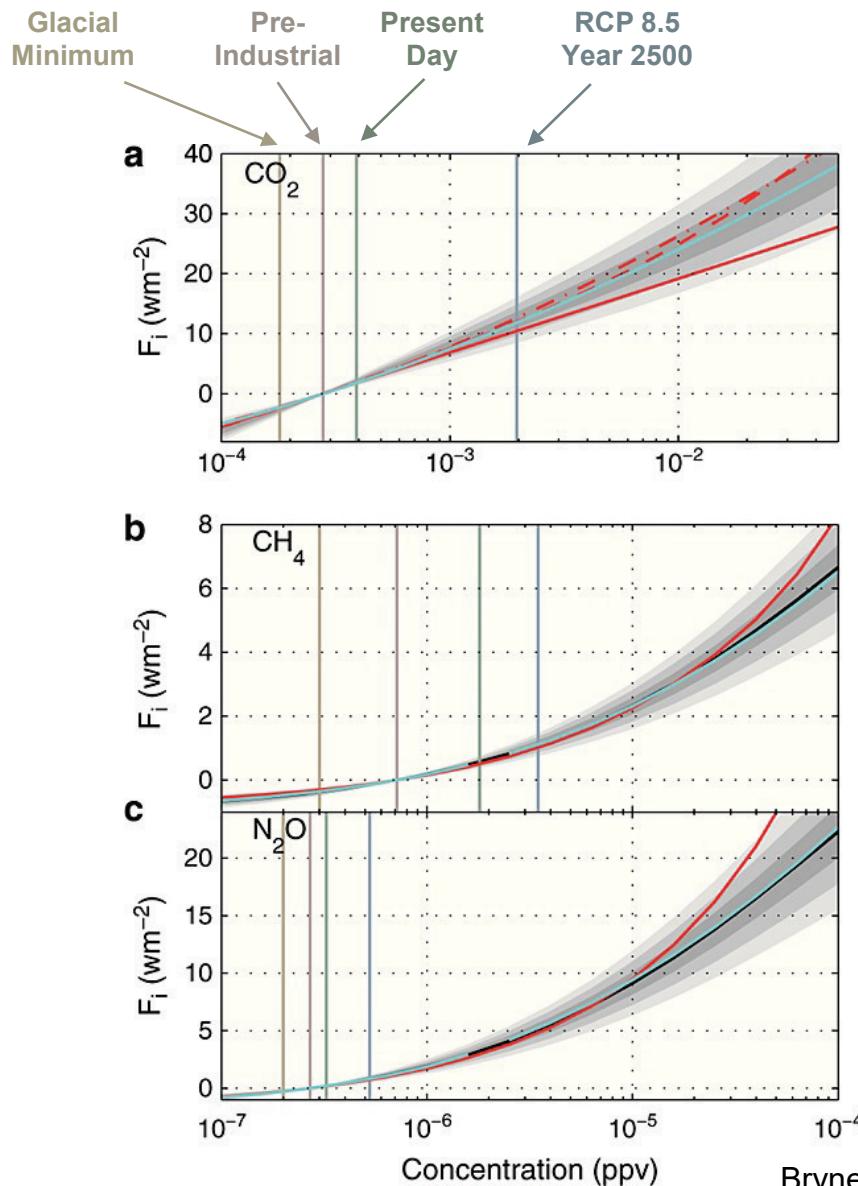
Highly saturated (most of the outgoing radiation is already absorbed) bands are less sensitive to increases in GHG concentration than partially or non saturated bands.

How does RF change with concentration?



<https://www.google.com/url?sa=i&url=https%3A%2F%2Fscienceofdoom.com%2F2011%2Fpage%2F4%2F&psig=AOvVaw0YxpuY4tcgapz8bhcYtC2&ust=1601056441281000&source=images&cd=vfe&ved=0CAIQjRxqFwoTCID7v5euguwCFQAAAAAdAAAABAD>

How does RF change with concentration?



- Black line is calculated RF using the Spectral Mapping for Atmospheric Radiative Transfer (SMART) radiative transfer code
- Light and dark grey show 1σ & 2σ uncertainties
- Cyan line is “fit” to the results
- Red lines are older fits from various IPCC and WMO/UNEP Ozone Depletion Reports

Bryne and Goldblatt, JGR, 2013

<https://agupubs.onlinelibrary.wiley.com/doi/full/10.1002/2013GL058456>

How does RF change with concentration?

Table 8.SM.1 | Supplementary for Table 8.3: RF formulae for CO₂, CH₄ and N₂O.

Gas	RF (in W m ⁻²)	Constant α
CO ₂	$\Delta F = \alpha \ln(C/C_0)$	5.35
CH ₄	$\Delta F = \alpha (\sqrt{M} - \sqrt{M_0}) - (f(M, N_0) - f(M_0, N_0))$	0.036
N ₂ O	$\Delta F = \alpha (\sqrt{N} - \sqrt{N_0}) - (f(M_0, N) - f(M_0, N_0))$	0.12

Notes:

$$f(M, N) = 0.47 \ln [1 + 2.01 \times 10^{-5} (MN)^{0.75} + 5.31 \times 10^{-15} M (MN)^{1.52}]$$

C is CO₂ in ppm.

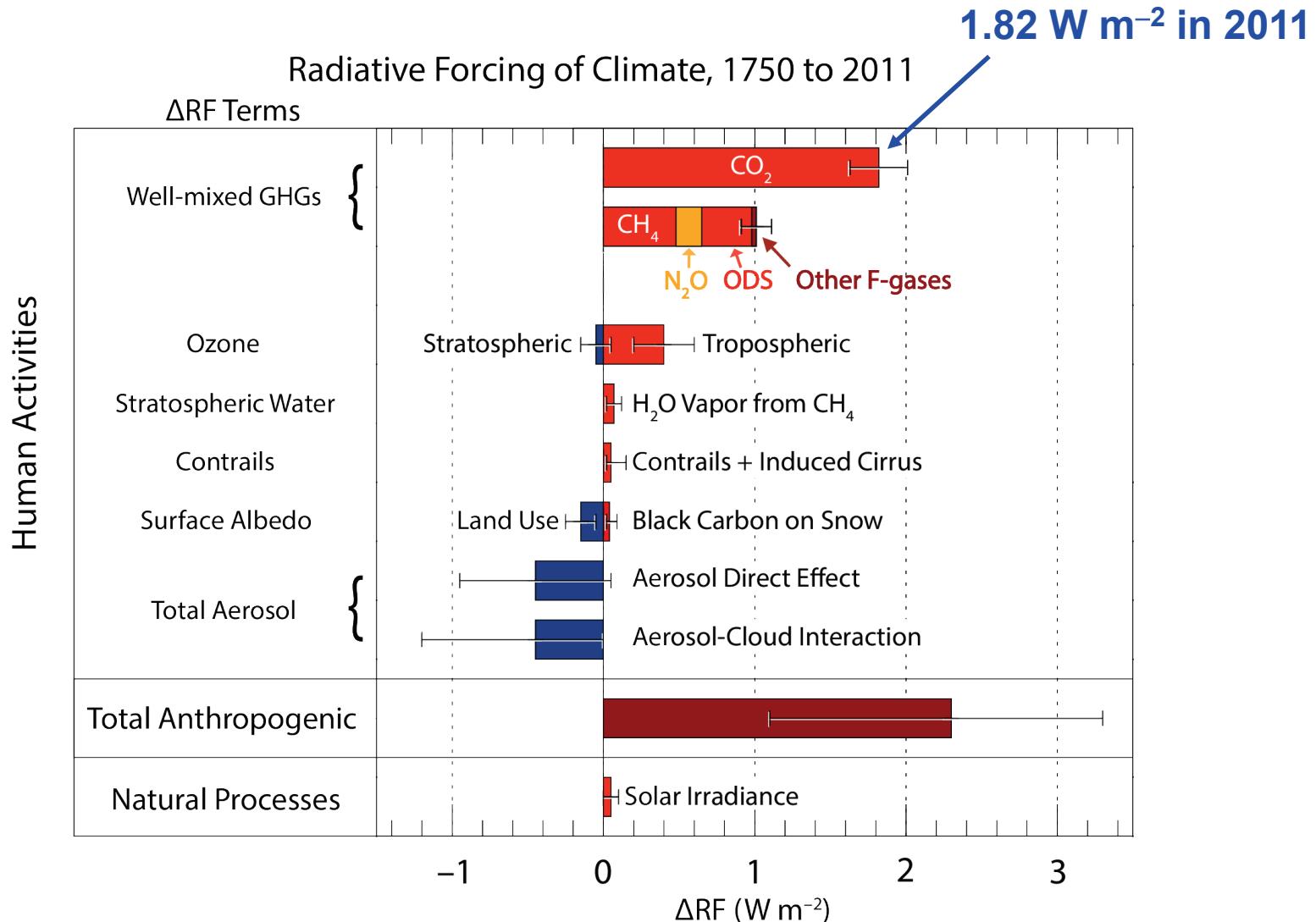
M is CH₄ in ppb.

N is N₂O in ppb.

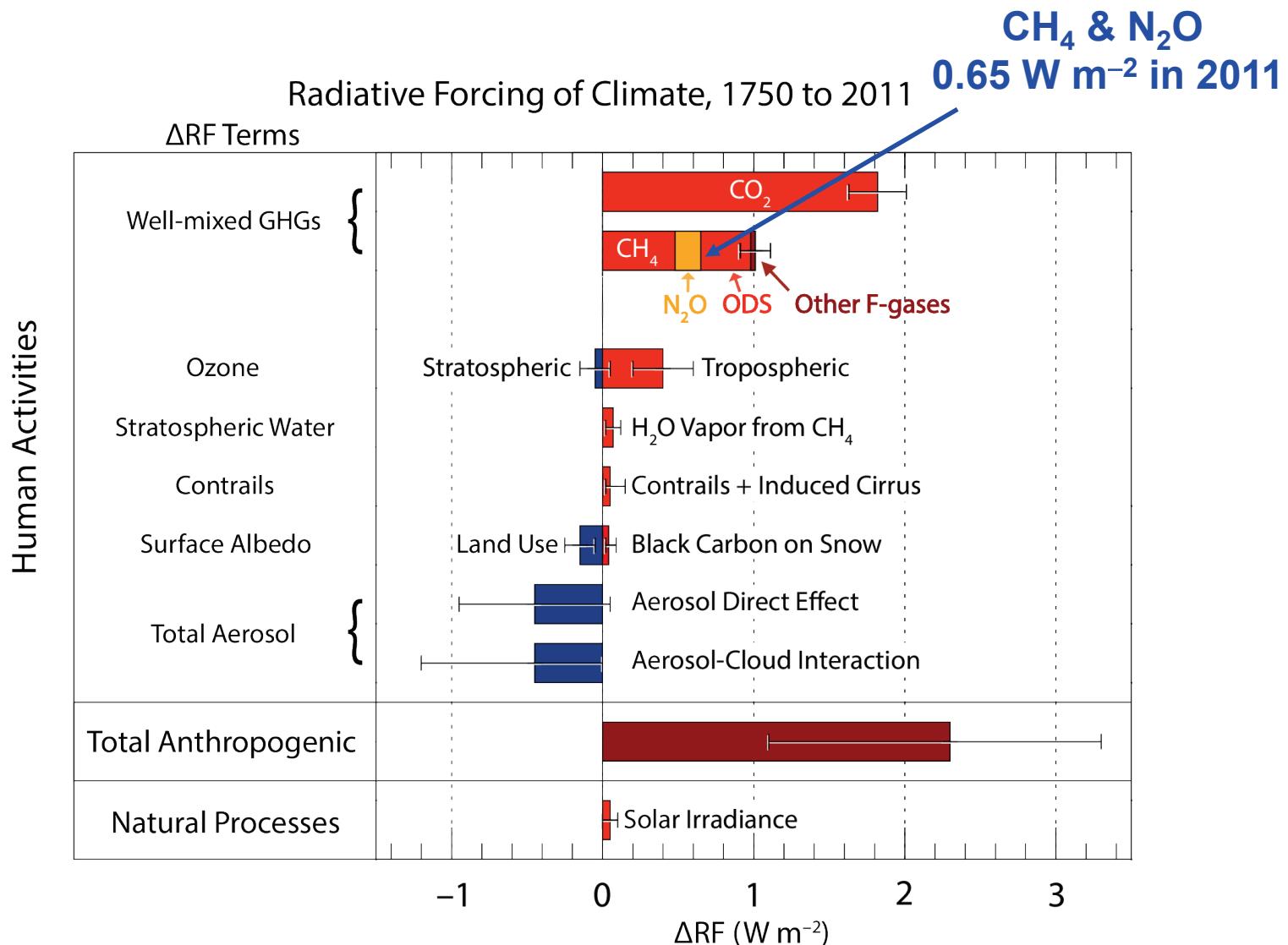
The subscript 0 denotes the unperturbed molar fraction for the species being evaluated. However, note that for the CH₄ forcing N₀ should refer to present-day N₂O, and for the N₂O forcing M₀ should refer to present-day CH₄.

IPCC Fifth Assessment Report, 2013

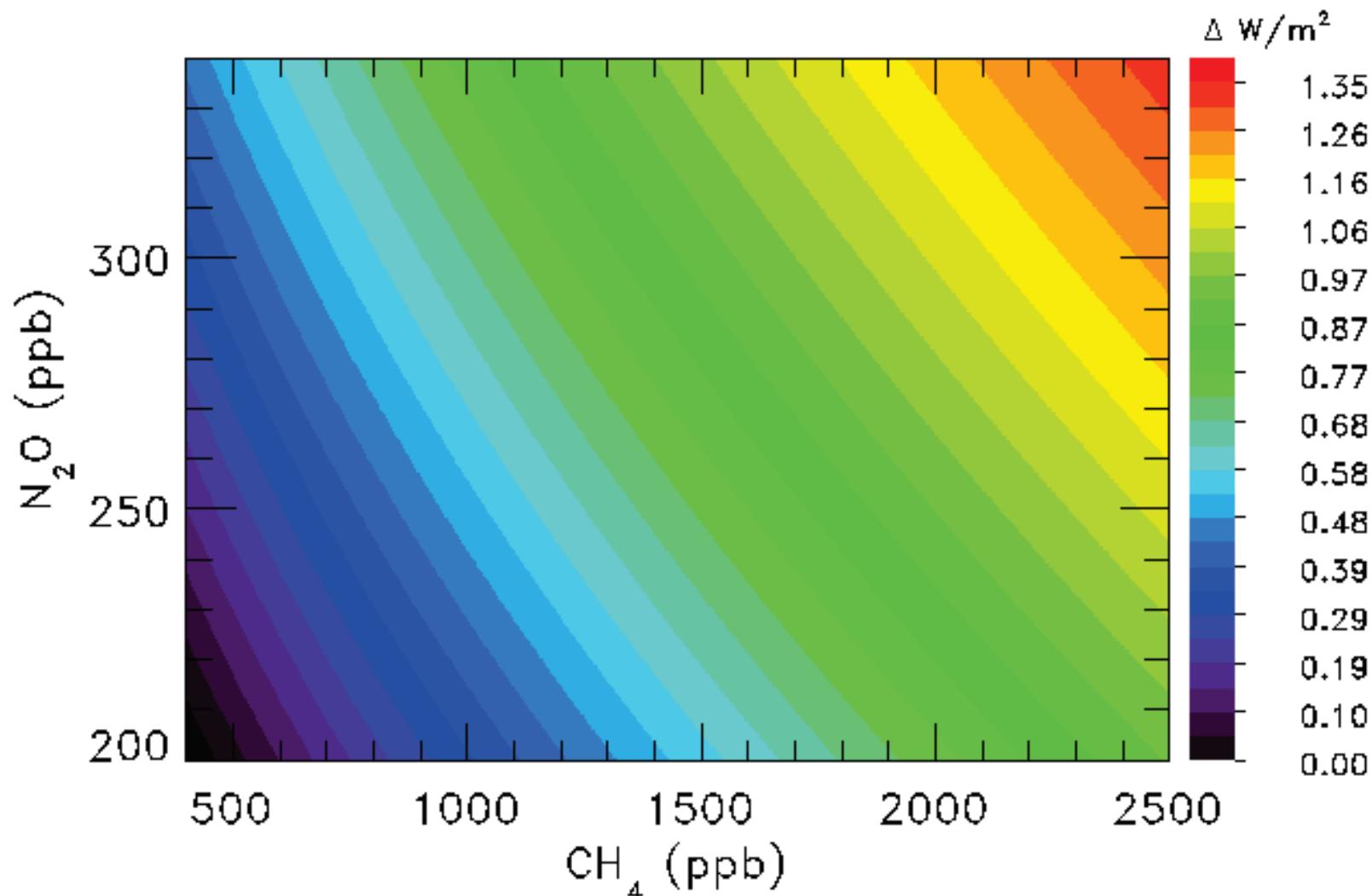
ΔRF of Climate



ΔRF of Climate

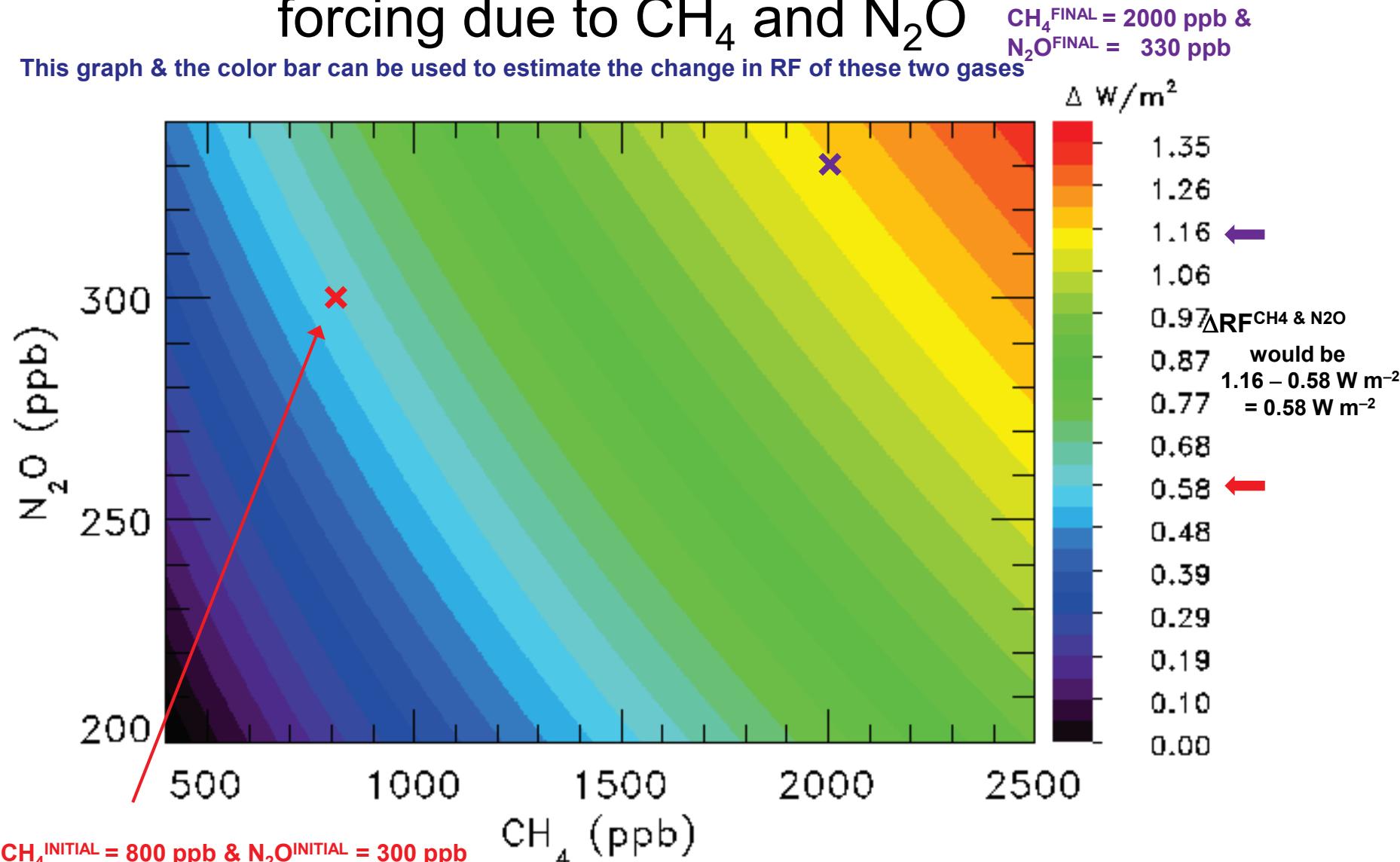


Graphical representation of surface radiative forcing due to CH₄ and N₂O



Graphical representation of surface radiative forcing due to CH_4 and N_2O

This graph & the color bar can be used to estimate the change in RF of these two gases



Absorption vs. Wavelength

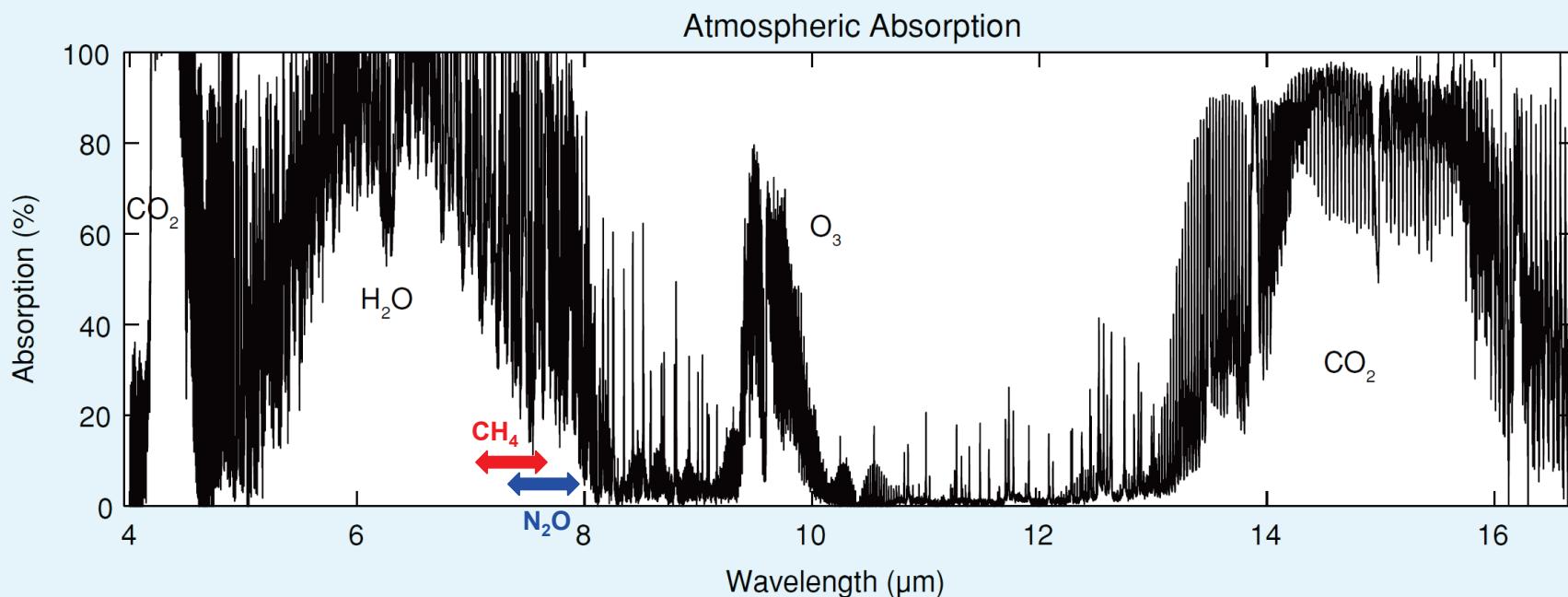
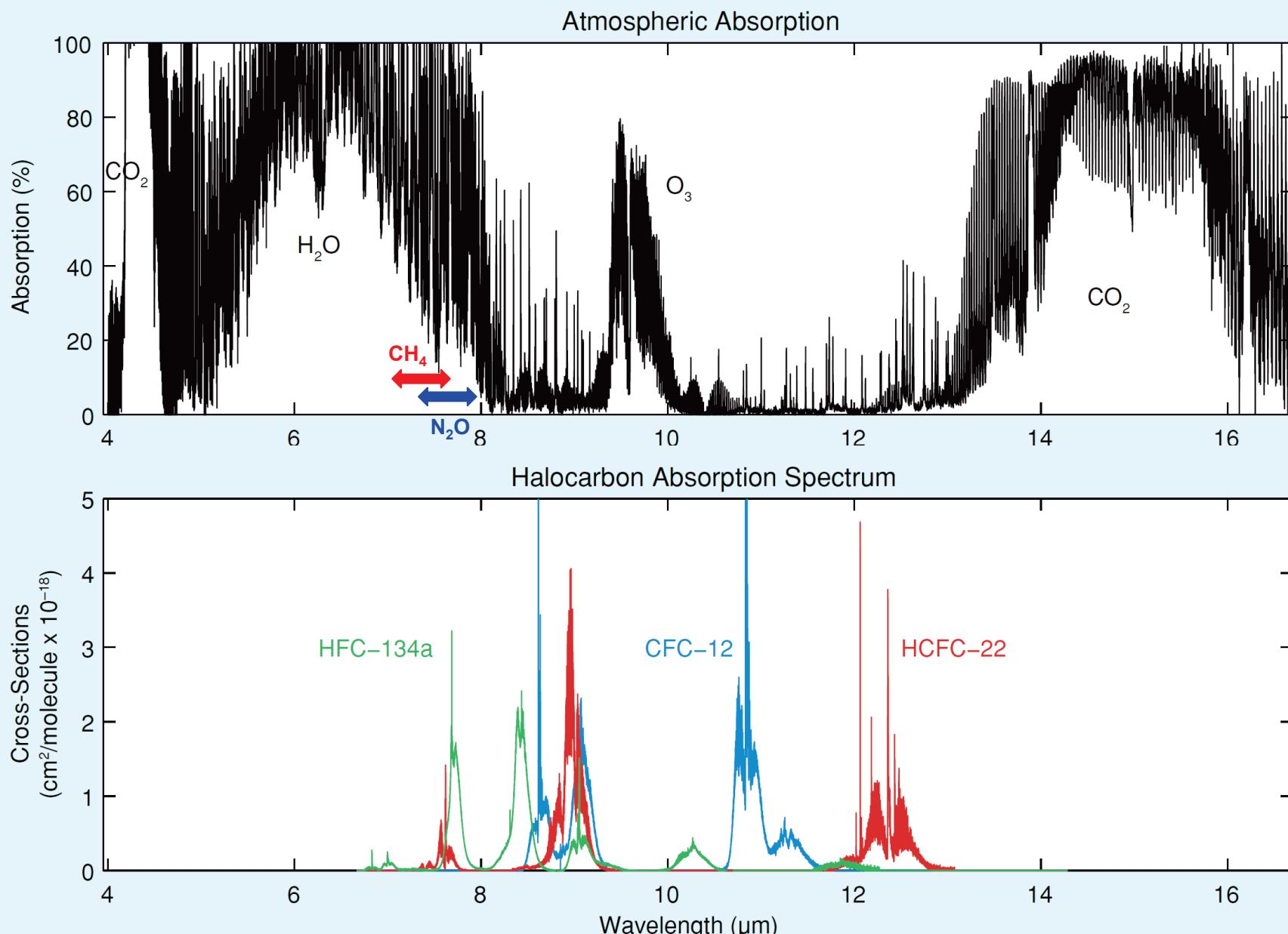


Fig 2.6, IPCC SROC (Special Report on Ozone layer and Climate), 2005
<https://www.ipcc.ch/site/assets/uploads/2018/03/sroc02-1.pdf>

Absorption vs. Wavelength



How does RF change with concentration?

Table 6.2: Simplified expressions for calculation of radiative forcing due to CO₂, CH₄, N₂O, and halocarbons.

The first row for CO₂ lists an expression with a form similar to IPCC (1990) but with newer values of the constants. The second row for CO₂ is a more complete and updated expression similar in form to that of Shi (1992). The third row expression for CO₂ is from WMO (1999), based in turn on Hansen *et al.* (1988).

Trace gas	Simplified expression Radiative forcing, ΔF (Wm ⁻²)	Constants
CO ₂	$\Delta F = \alpha \ln(C/C_0)$	$\alpha = 5.35$
CH ₄	Please see slide 42	
N ₂ O		
CFC-11a	$\Delta F = \alpha(X - X_0)$	$\alpha = 0.25$
CFC-12	$\Delta F = \alpha(X - X_0)$	$\alpha = 0.32$

C is CO₂ in ppm

M is CH₄ in ppb

N is N₂O in ppb

X is CFC in ppb

The constant in the simplified expression for CO₂ for the first row is based on radiative transfer calculations with three-dimensional climatological meteorological input data (Myhre *et al.*, 1998b). For the second and third rows, constants are derived with radiative transfer calculations using one-dimensional global average meteorological input data from Shi (1992) and Hansen *et al.* (1988), respectively.

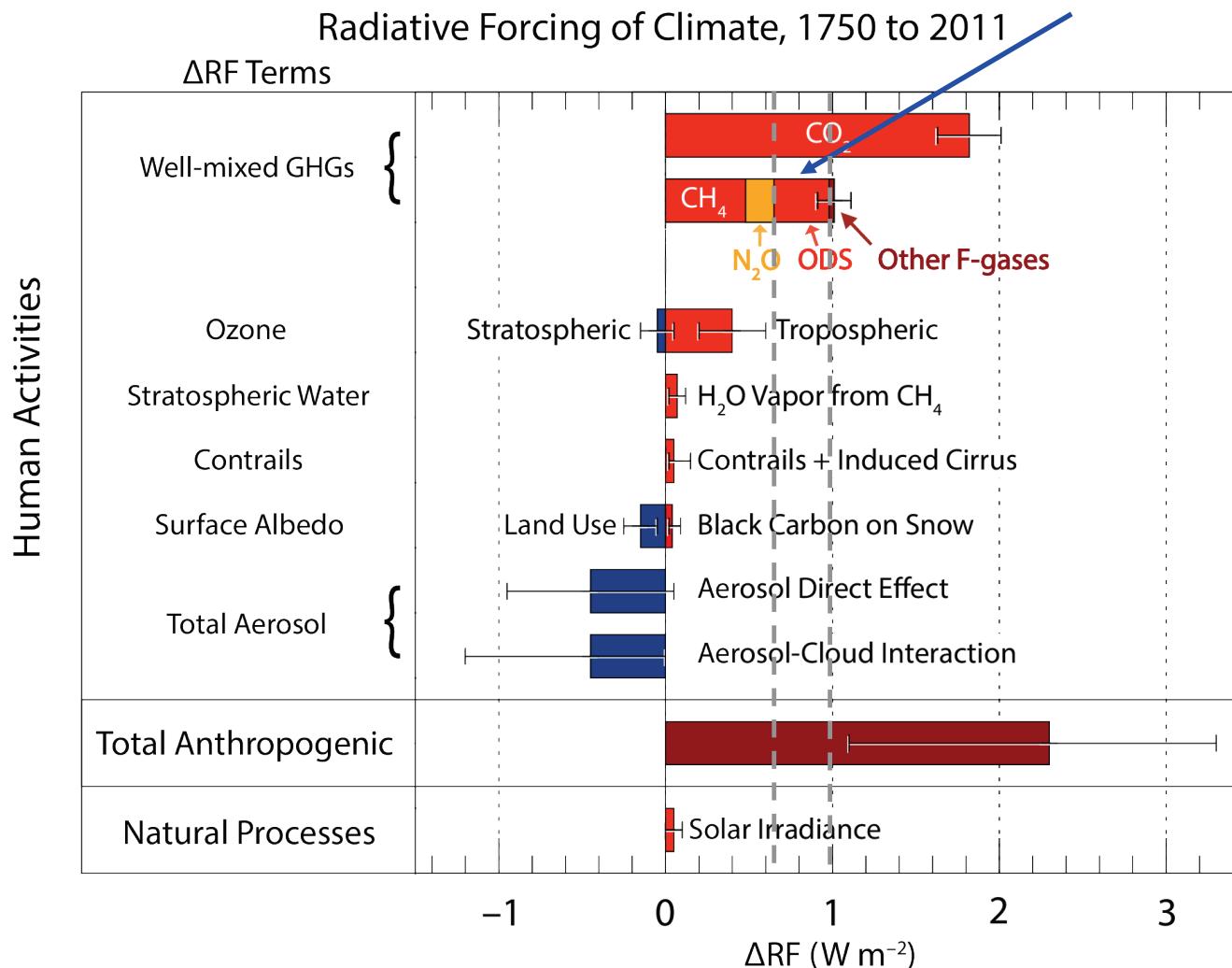
The subscript 0 denotes the unperturbed concentration.

^a The same expression is used for all CFCs and CFC replacements, but with different values for α (i.e., the radiative efficiencies in [Table 6.7](#)).

IPCC Third Assessment Report, 2001

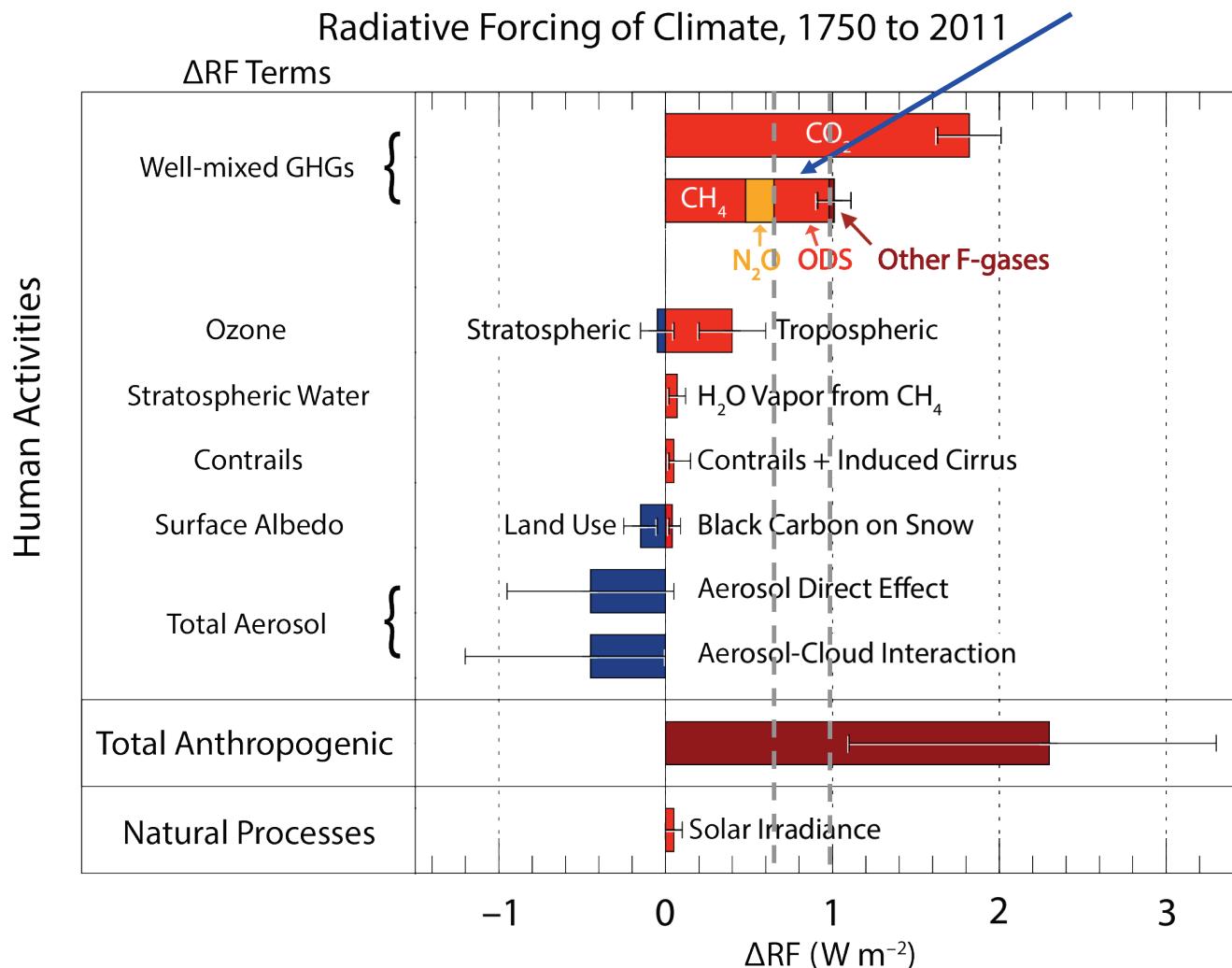
ΔRF of Climate

All Ozone Depleting Substances
 0.33 W m^{-2} in 2011
 CFC-11 & CFC-12 70% of total

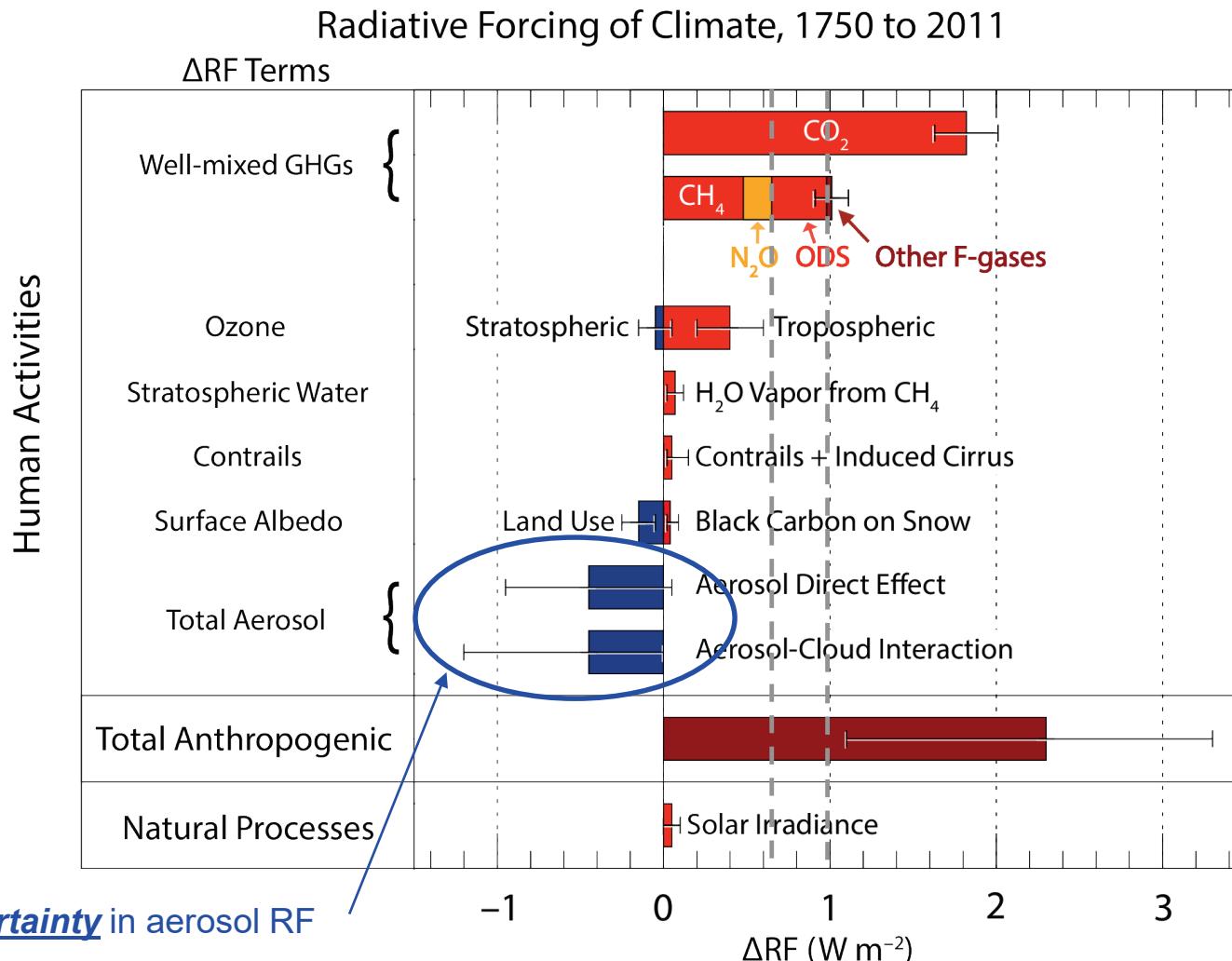


ΔRF of Climate

All Ozone Depleting Substances
 0.33 W m^{-2} in 2011
 CFC-11 & CFC-12 70% of total



ΔRF of Climate



- scatter and absorb radiation (**direct radiative forcing**)
- affect cloud formation (**indirect radiative forcing**)

RF Due to Tropospheric Aerosols: Direct Effect

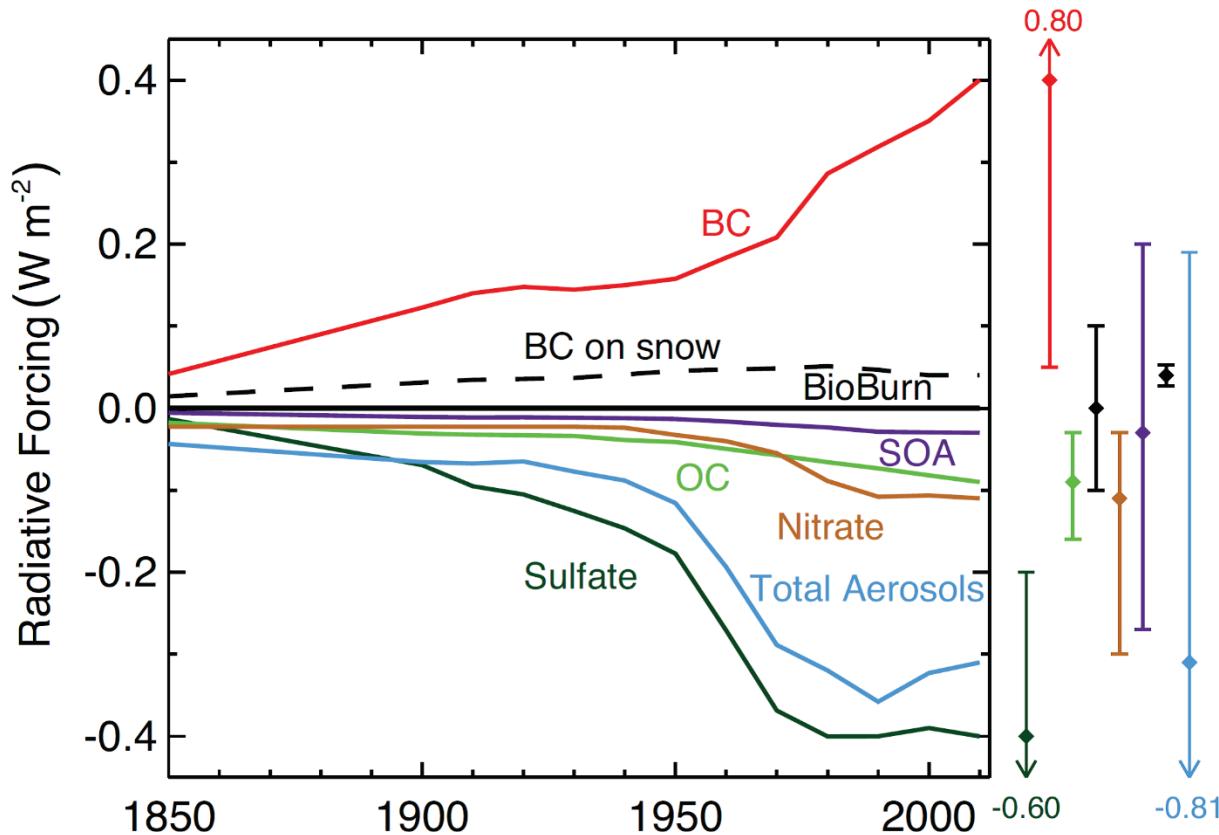


Figure 8.8 | Time evolution of RF due to aerosol–radiation interaction and BC on snow and ice.

Fig 8.8, IPCC 2013: Only Direct RF of aerosols considered here

RF Due to Tropospheric Aerosols: Indirect Effect

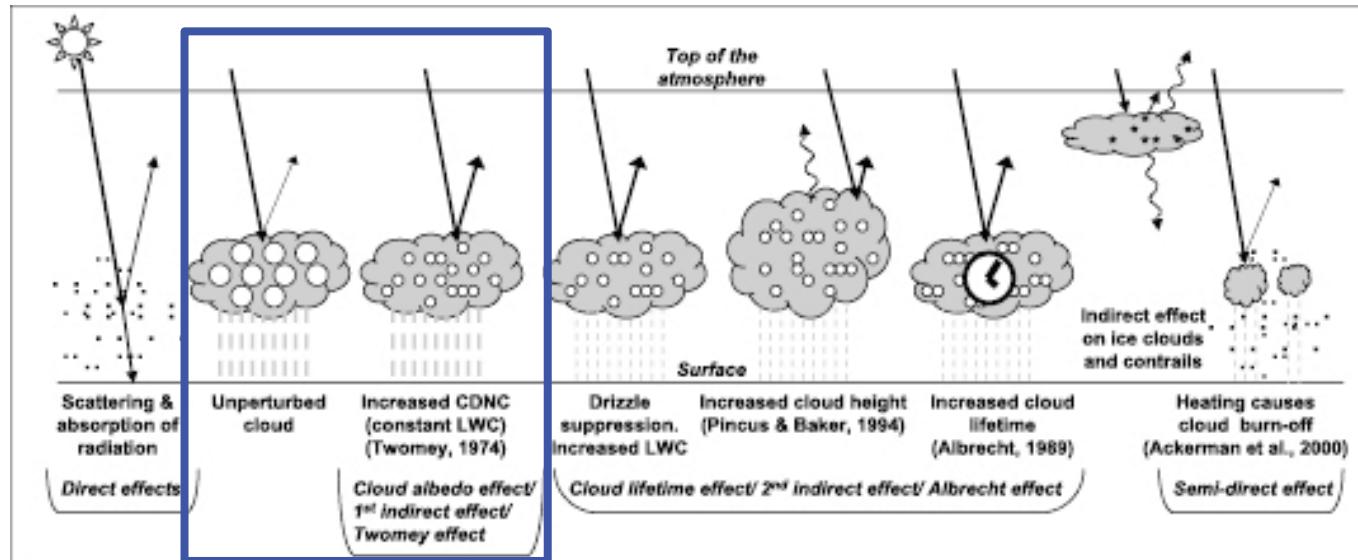
Indirect Effects of Aerosols on Clouds

Anthropogenic aerosols lead to more cloud condensation nuclei (CCN)

Resulting cloud particles consist of smaller droplets, promoted by more sites (CCN) for cloud nucleation

The cloud that is formed is therefore brighter (reflects more sunlight) \Rightarrow

Twomey effect, aka 1st Indirect Effect



Large uncertainty in aerosol RF

Fig 2-10, IPCC 2007

- scatter and absorb radiation (**direct radiative forcing**)
- affect cloud formation (**indirect radiative forcing**)

RF Due to Tropospheric Aerosols: Indirect Effect

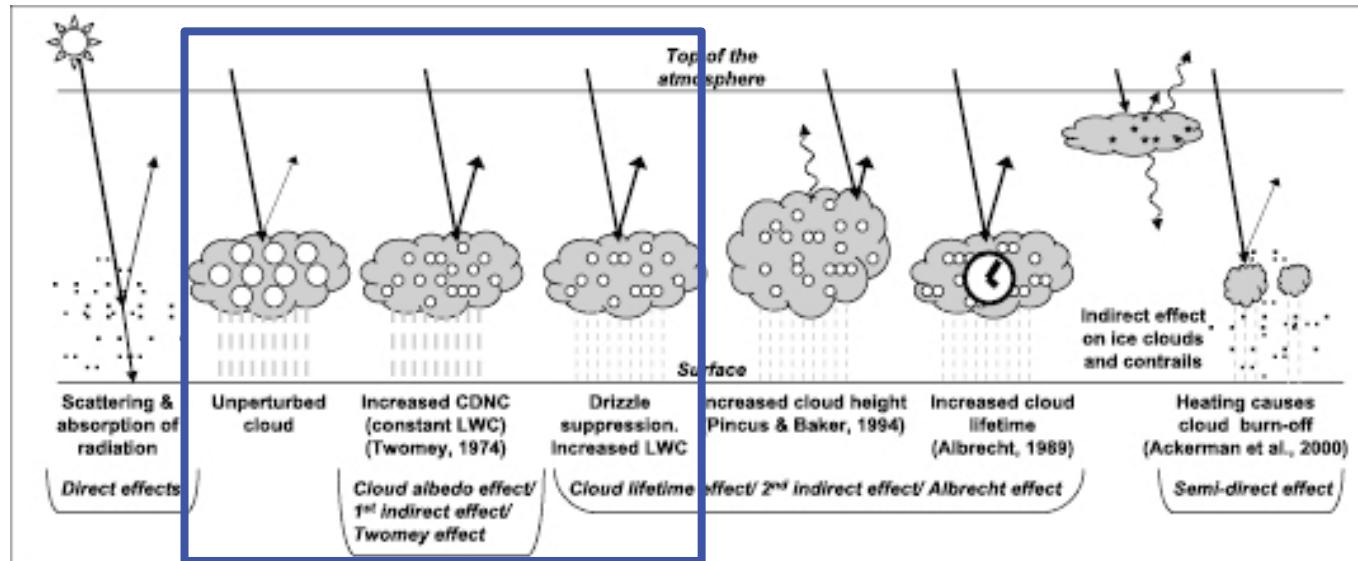
Indirect Effects of Aerosols on Clouds

Anthropogenic aerosols lead to more cloud condensation nuclei (CCN)

Resulting cloud particles consist of smaller droplets, promoted by more sites (CCN) for cloud nucleation

The cloud that is formed is therefore brighter (reflects more sunlight) and has less efficient precipitation, i.e. is longer lived) \Rightarrow

Albrecht effect, aka 2nd Indirect Effect

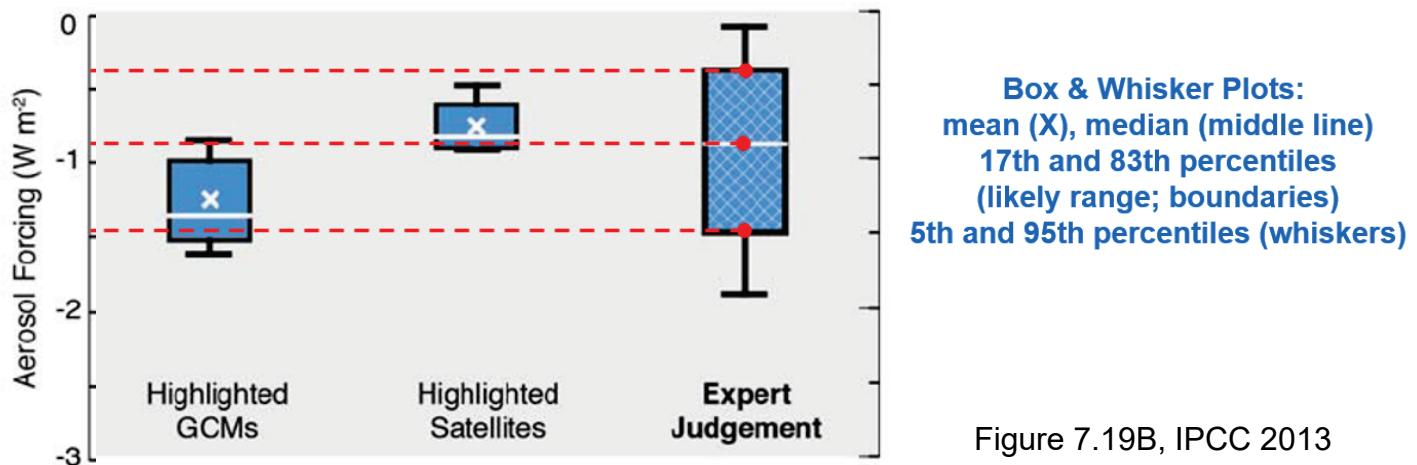


Large uncertainty in aerosol RF

Fig 2-10, IPCC 2007

- scatter and absorb radiation (**direct radiative forcing**)
- affect cloud formation (**indirect radiative forcing**)

Tropospheric Aerosol RF



ΔRF_{2011} GHGs $\approx 3.2 \text{ W m}^{-2}$ \Rightarrow climate change is complex but this quantity is well known

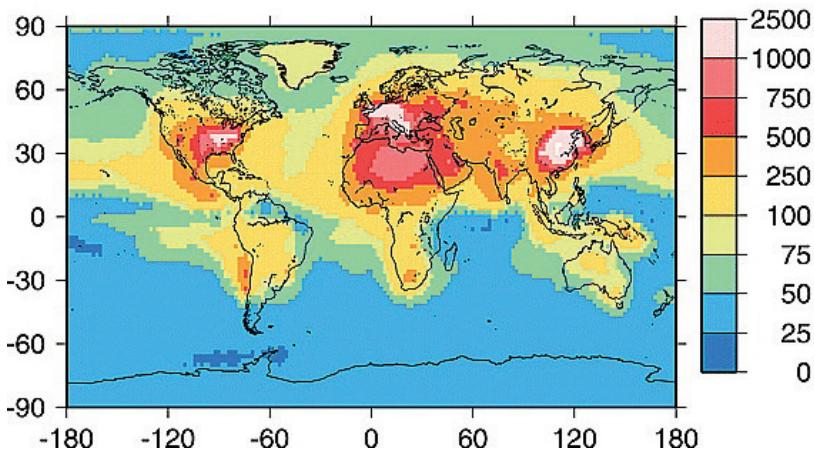
ΔRF_{2011} Aerosols: best estimate is -0.9 W m^{-2} , probably between -0.4 W m^{-2} and -1.5 W m^{-2} ;
could be between -0.1 W m^{-2} and -1.9 W m^{-2}

Large uncertainty in aerosol RF

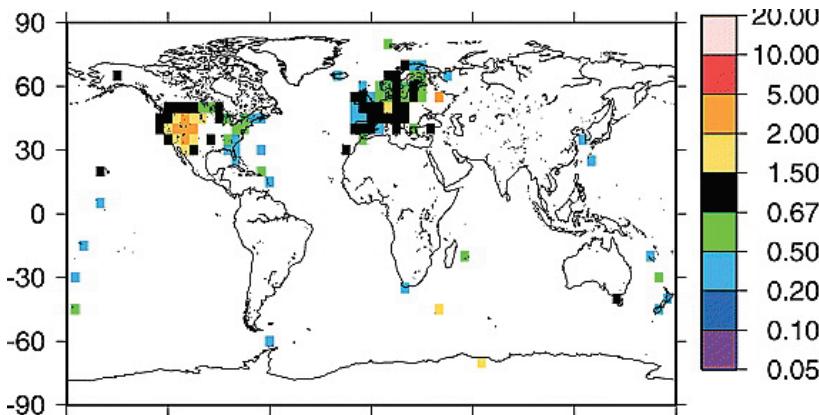
- scatter and absorb radiation (**direct radiative forcing**)
- affect cloud formation (**indirect radiative forcing**)

Tropospheric Sulfate Aerosols

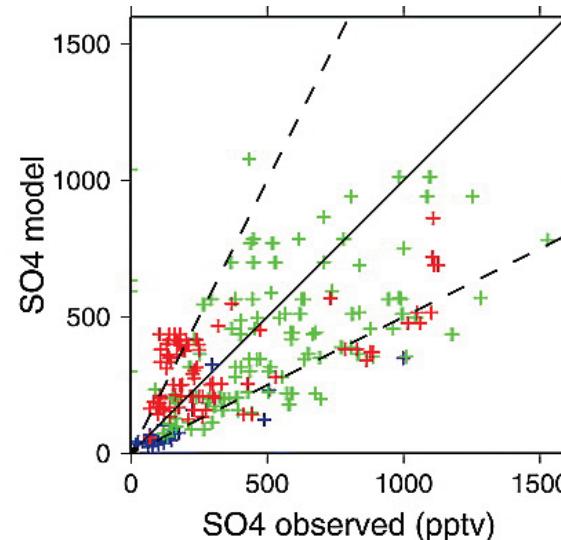
Modeled Sulfate (ppt)
Year 2000



Ratio of Modeled / Measured Sulfate



Modeled versus Measured Sulfate



Remote sites (blue), Europe (green), & United States (red).

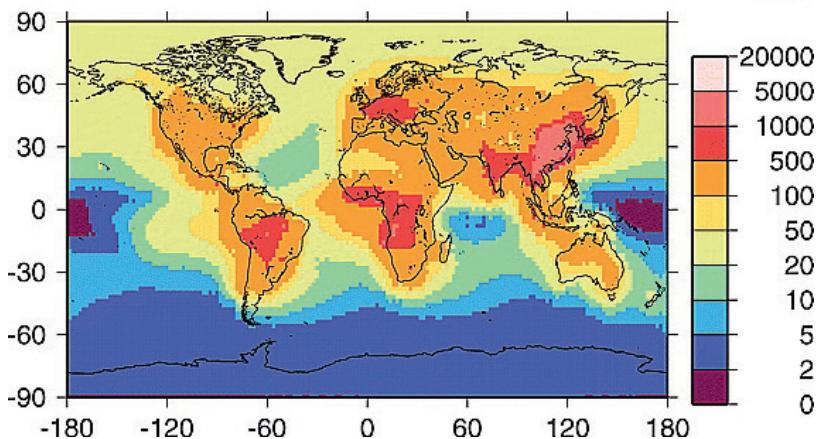
Koch et al., JGR, 2007

<https://agupubs.onlinelibrary.wiley.com/doi/full/10.1029/2005JD007024>

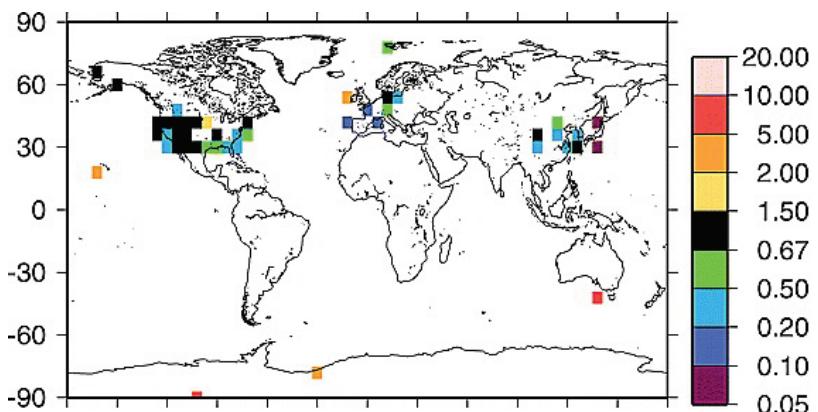
Tropospheric Sulfate Aerosols

Modeled Black Carbon (ng m^{-3})

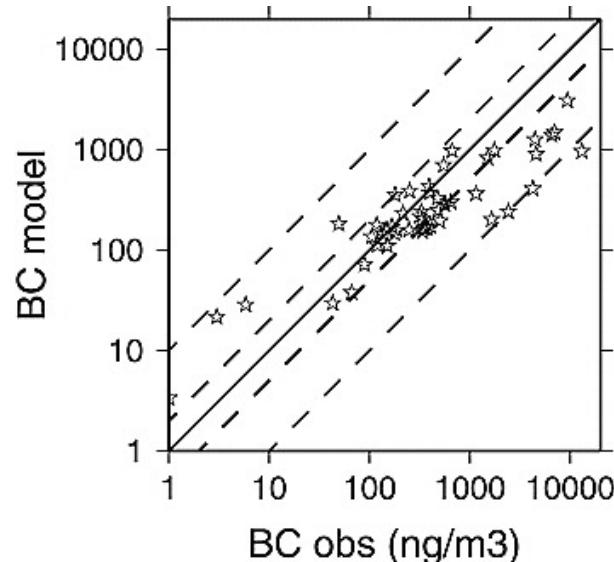
Year 2000



Ratio of Modeled / Measured Black Carbon



Modeled versus Measured Black Carbon

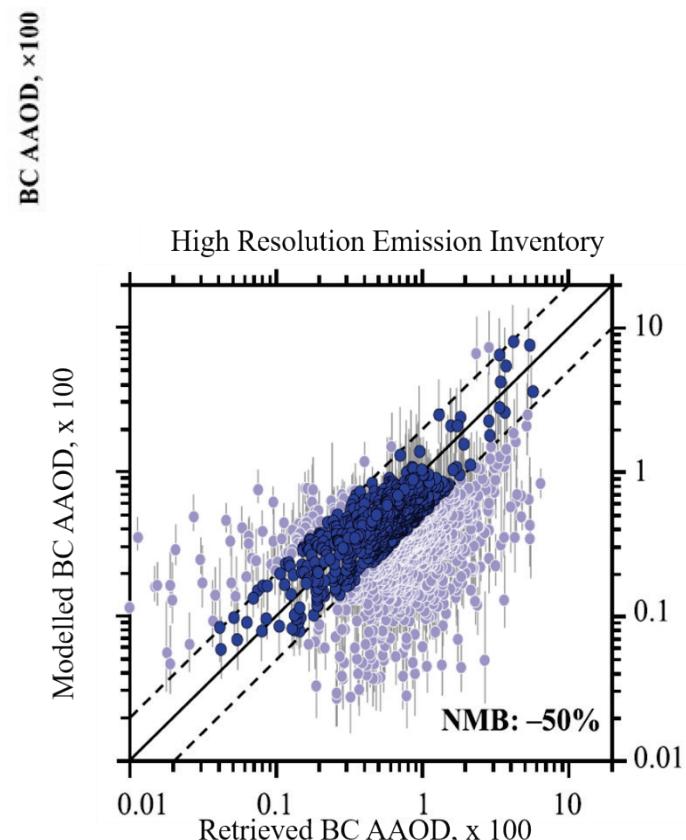
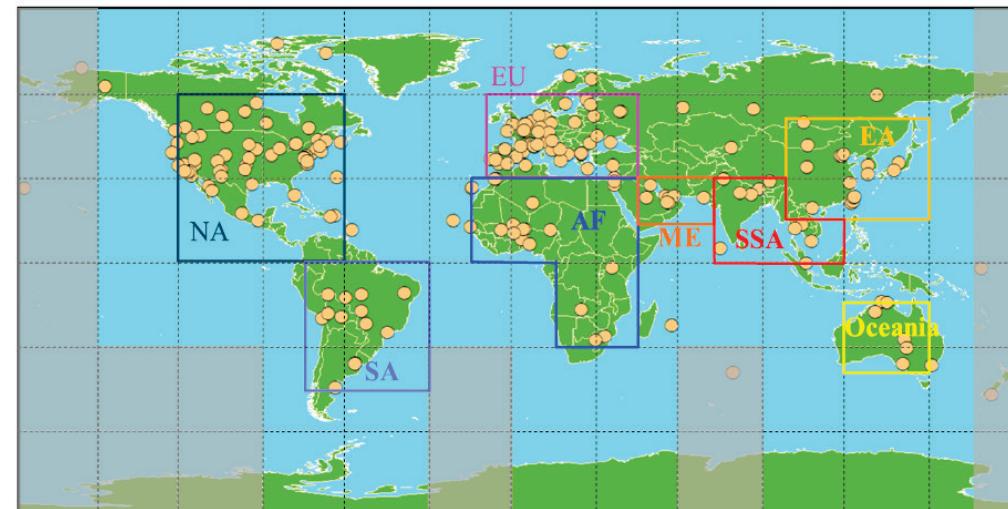
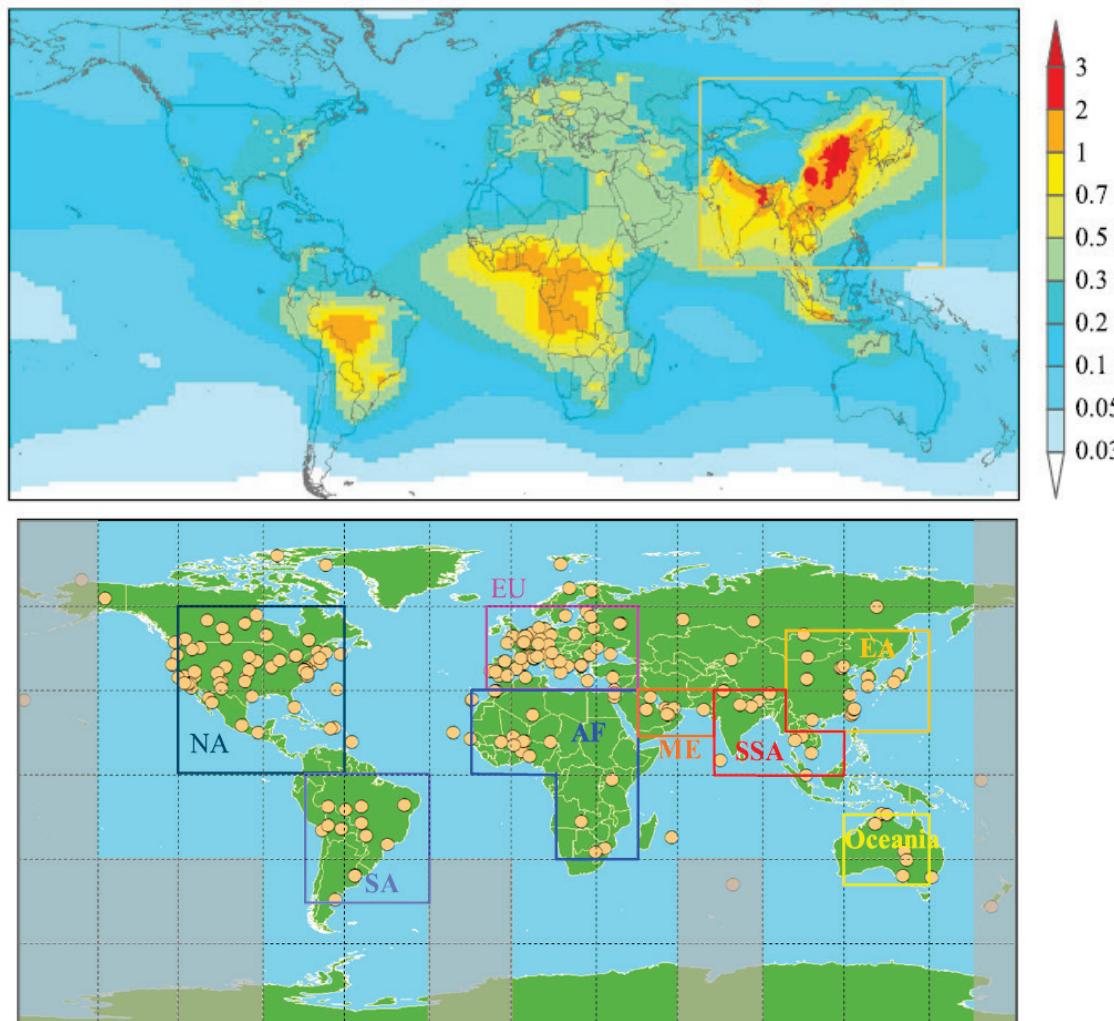


Koch et al., JGR, 2007

<https://agupubs.onlinelibrary.wiley.com/doi/full/10.1029/2005JD007024>

Black Carbon Aerosols

Simulated Black Carbon Aerosol Absorption Optical Depth (AAOD) at 900 nm for year 2007



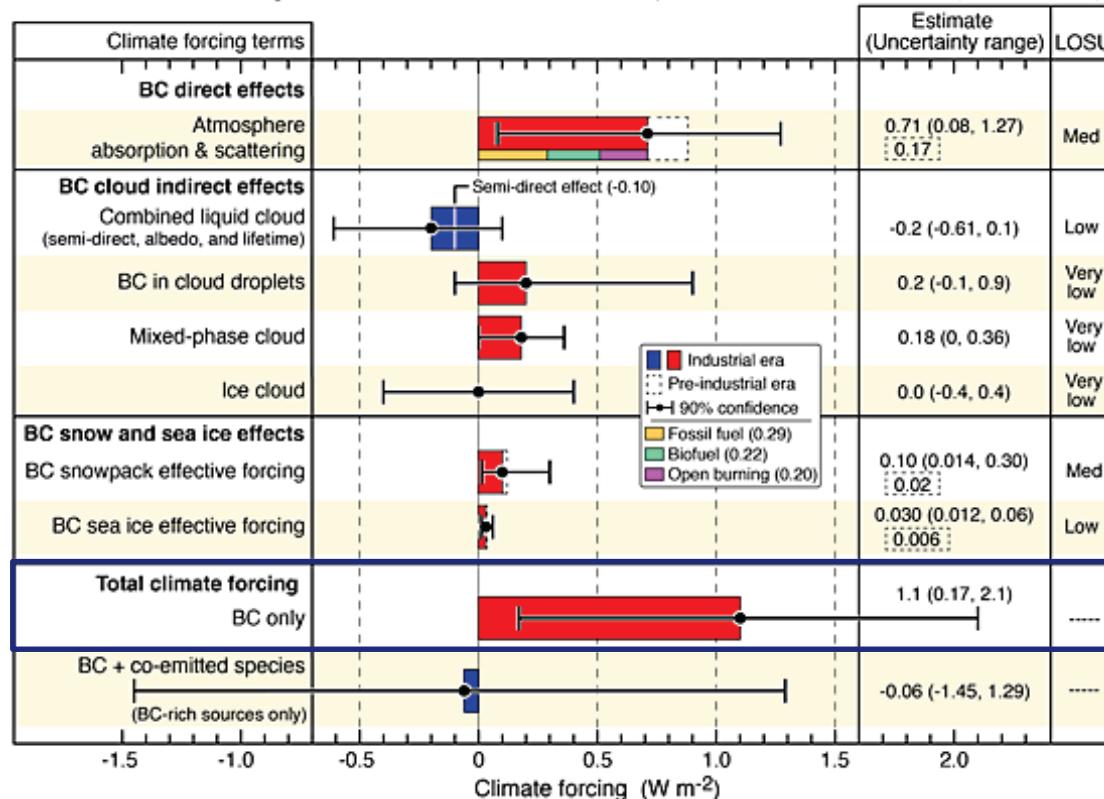
Wang et al., JGR, 2016

<https://agupubs.onlinelibrary.wiley.com/doi/full/10.1002/2015JD024326>

Black Carbon Aerosols

Bond *et al.*, Bounding the role of black carbon in the climate system: A scientific assessment, *JGR*, 2013

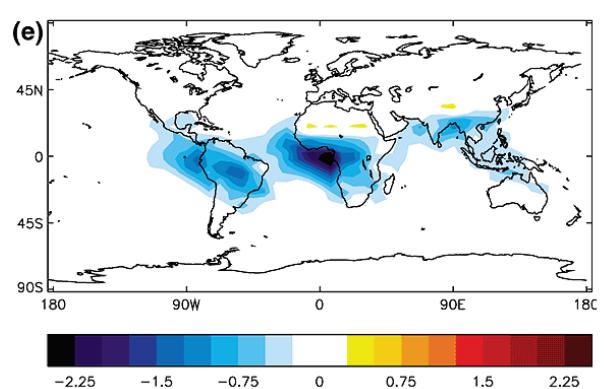
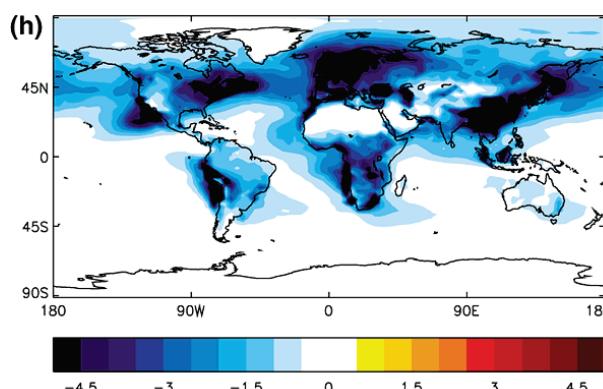
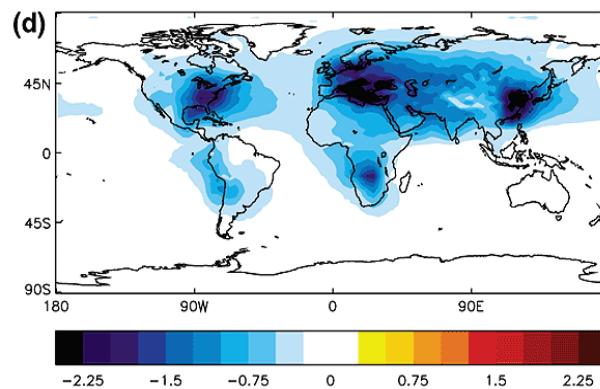
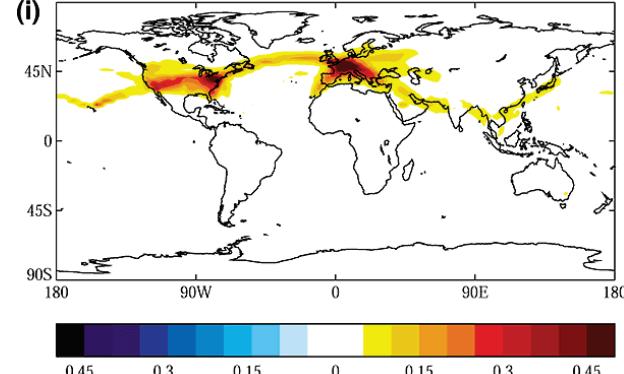
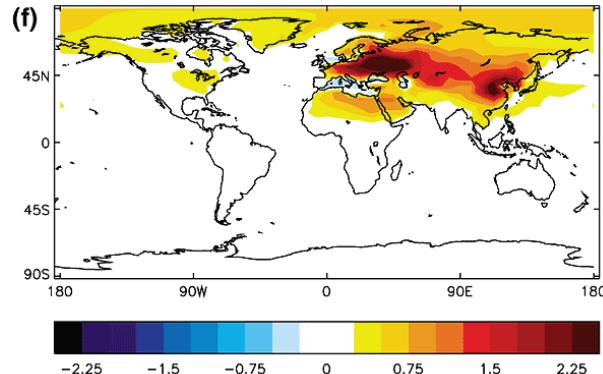
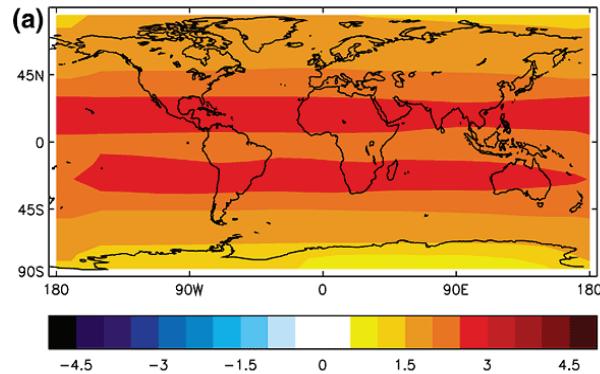
Global climate forcing of black carbon and co-emitted species in the industrial era (1750 - 2005)



Total Climate Forcing, Black Carbon Aerosols (W m⁻²)				
Report	IPCC (1995)	IPCC (2001)	IPCC (2007)	IPCC (2013)
ΔRF, BC	0.1 (0.03 to 0.3)	0.2 (0.1 to 0.4)	0.2 (0.05 to 0.35)	0.4 (0.05 to 0.80)

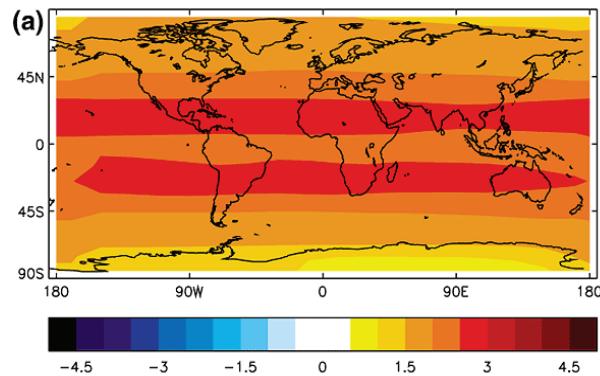
Global View

All forcings (1750-2000) are in Wm^{-2}

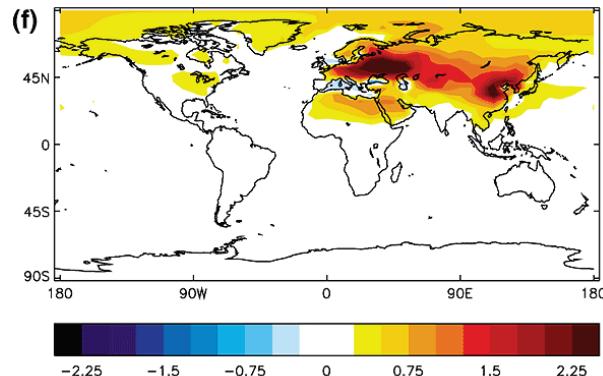


Global View

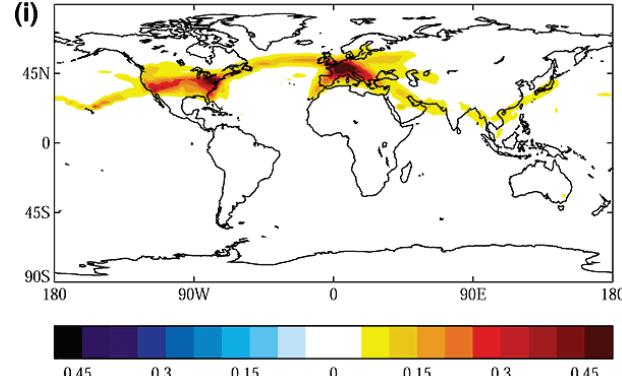
All forcings (1750-2000) are in Wm^{-2}



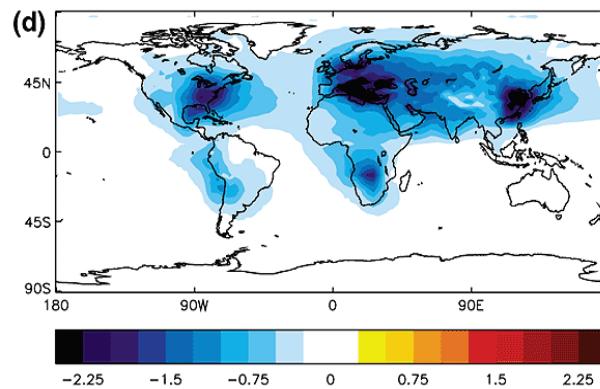
Greenhouse gases



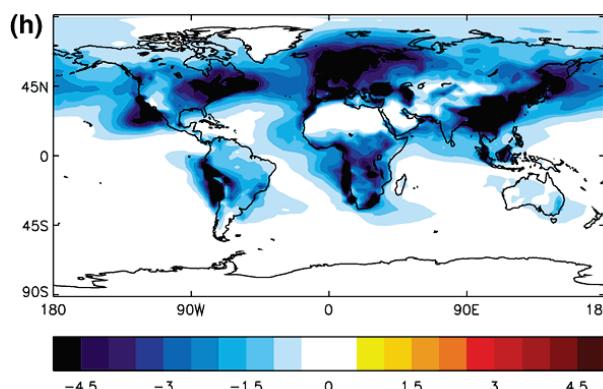
Organic and black carbon
from fossil fuel burning



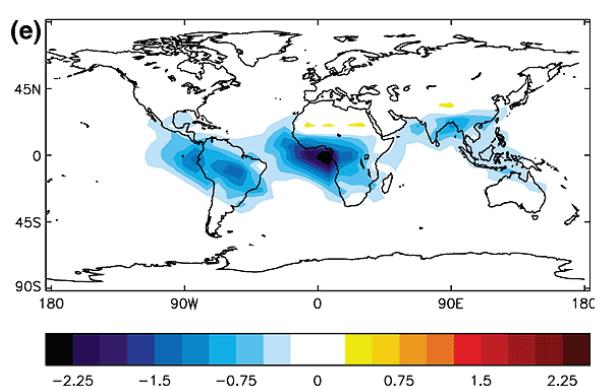
Aircraft contrails



Direct effect from
sulphate aerosols



Indirect effect from
sulphate aerosols



Organic and black carbon
from biomass burning

<https://www.ipcc.ch/report/ar3/wg1/chapter-6-radiative-forcing-of-climate-change/>

Combining RF GHGs & Aerosols

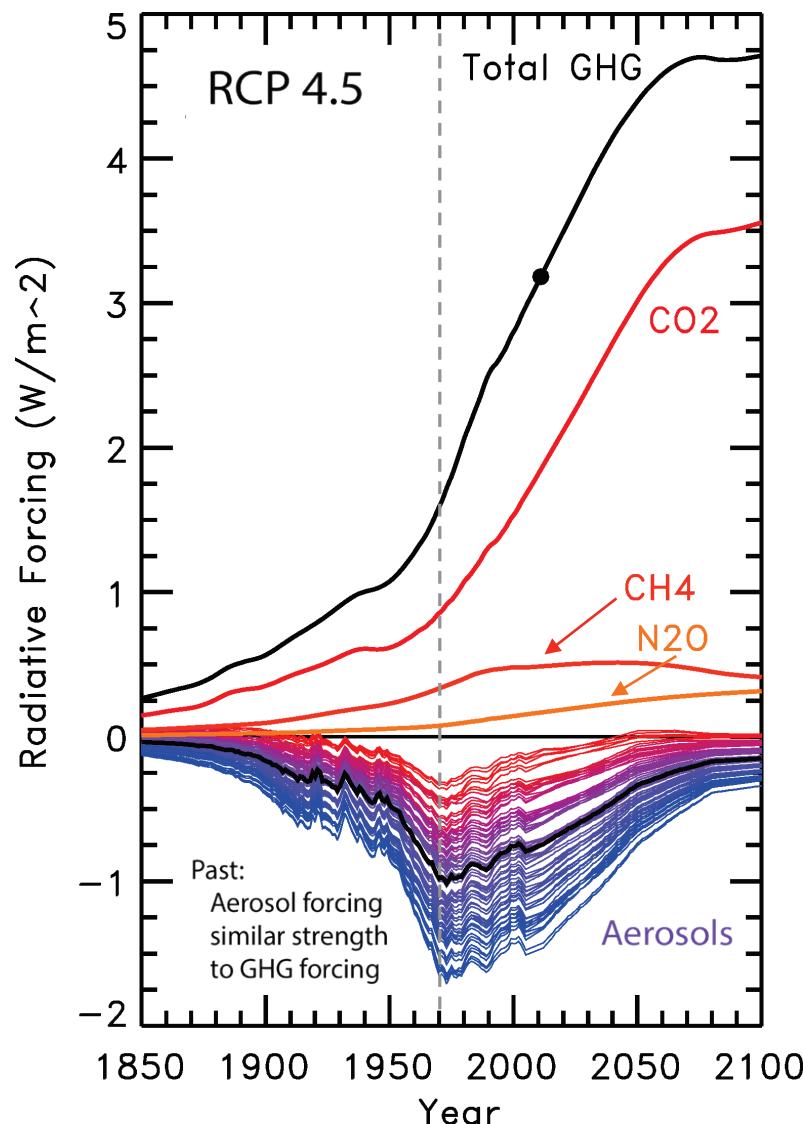


Fig 1.10, Paris, Beacon of Hope

Combining RF GHGs & Aerosols

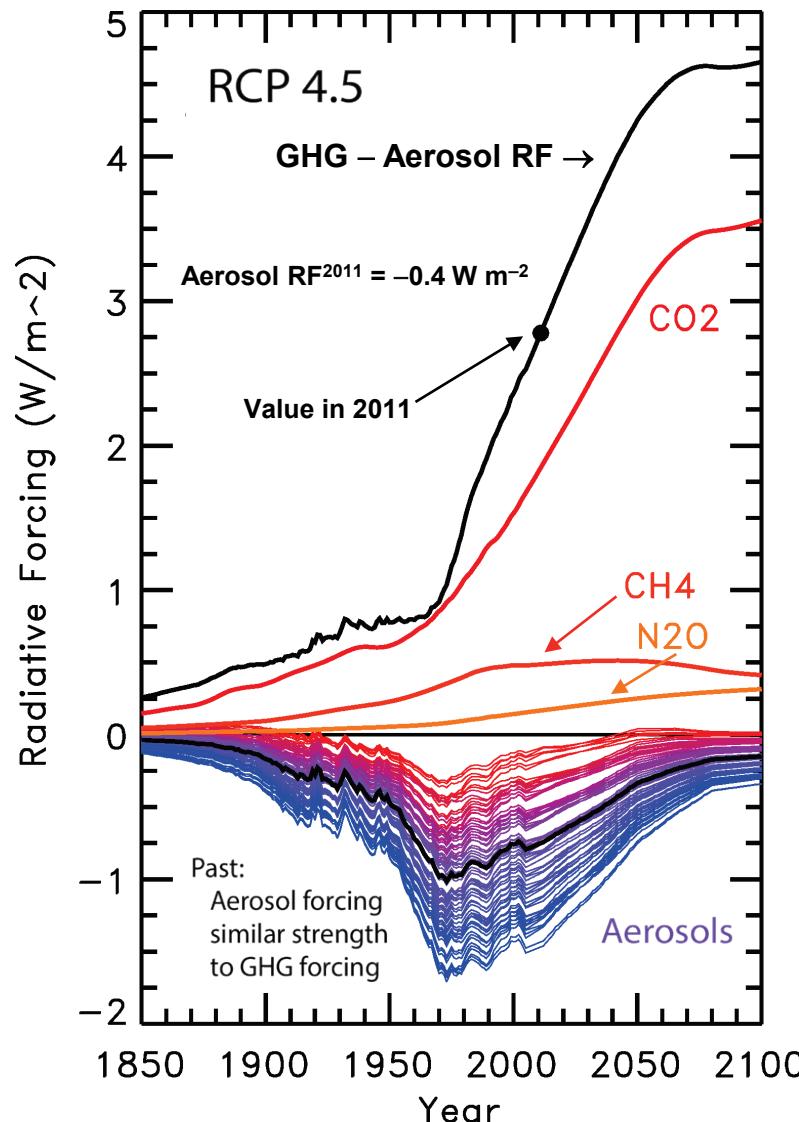


Fig 1.10, Paris, Beacon of Hope

Combining RF GHGs & Aerosols

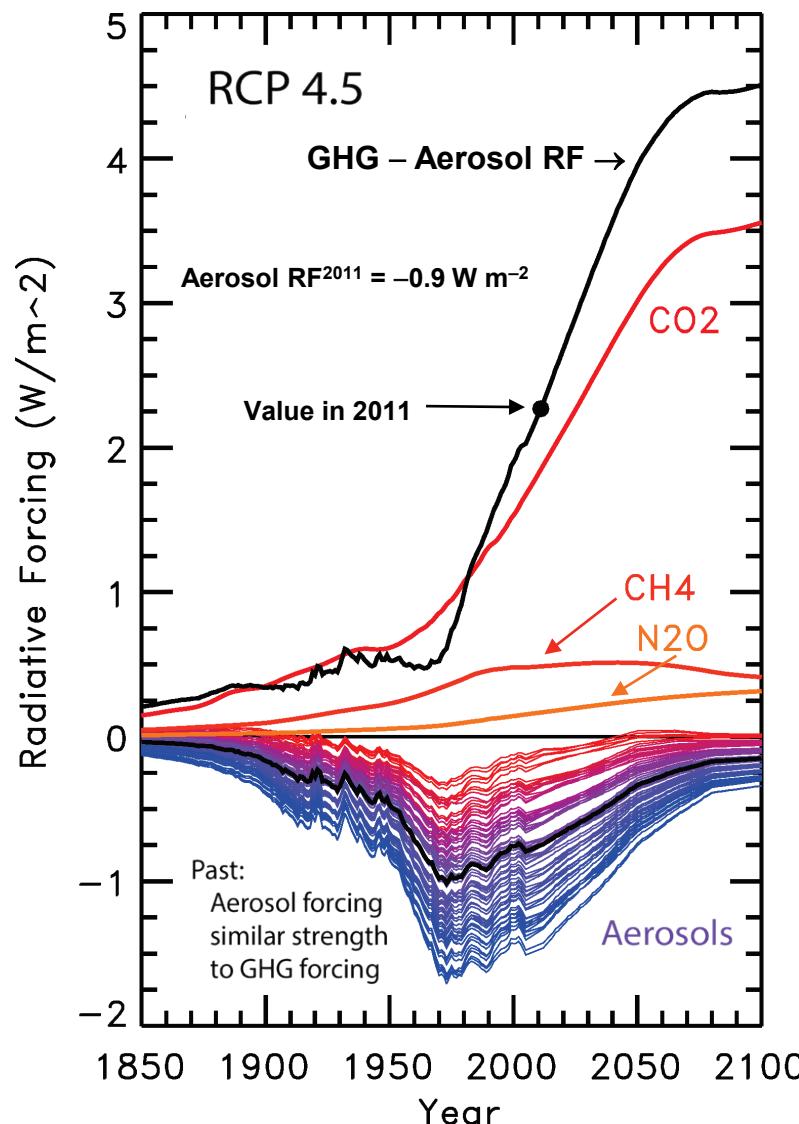


Fig 1.10, Paris, Beacon of Hope

Combining RF GHGs & Aerosols

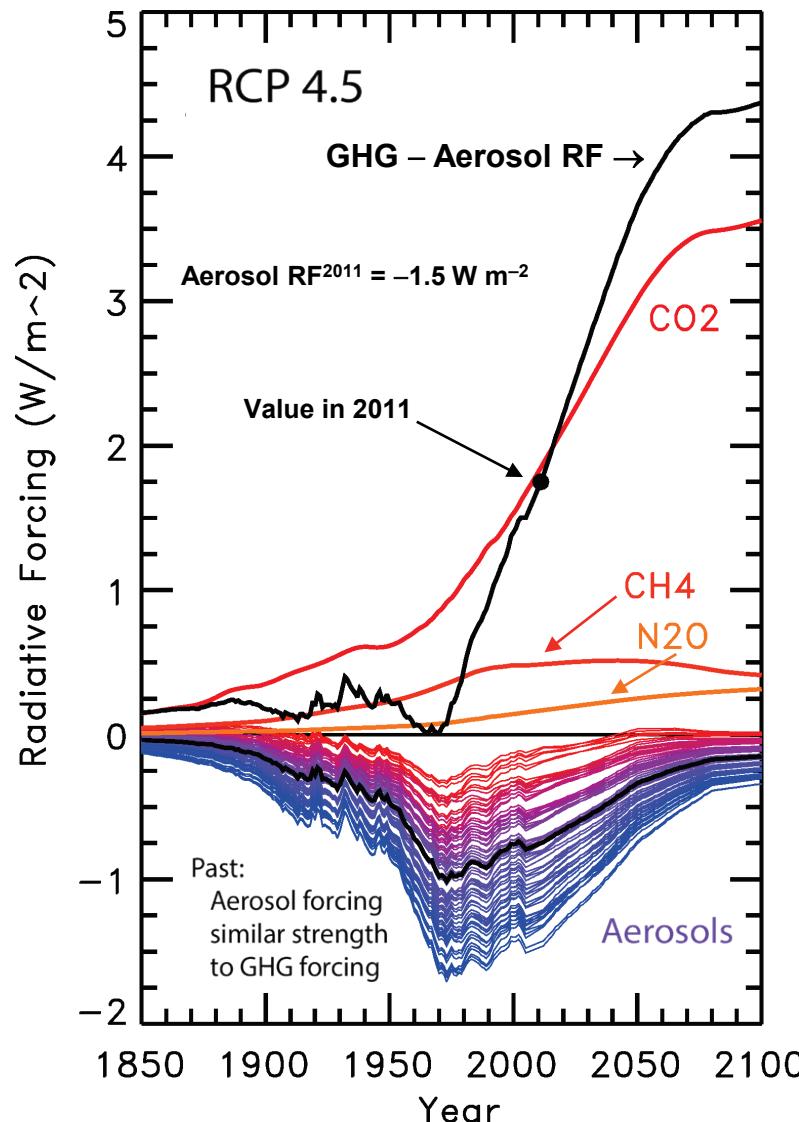
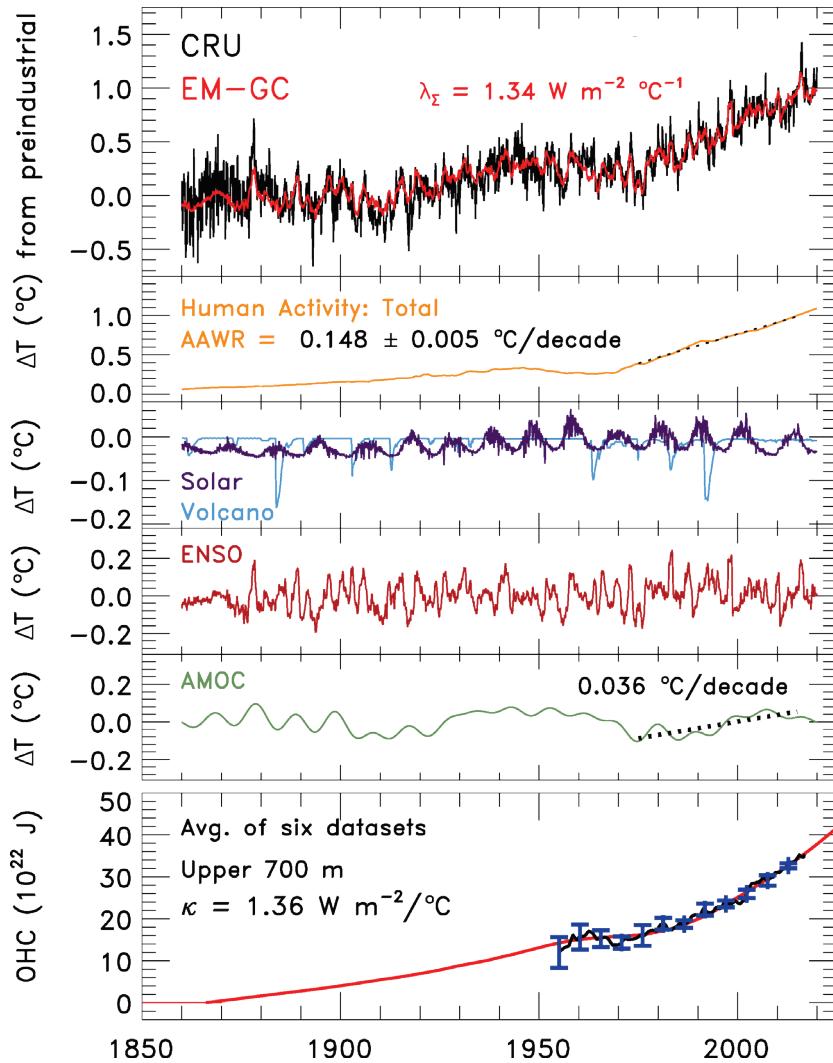


Fig 1.10, Paris, Beacon of Hope

Univ of Md Empirical Model of Global Climate



AAWR: Attributable Anthropogenic Warming Rate

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

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

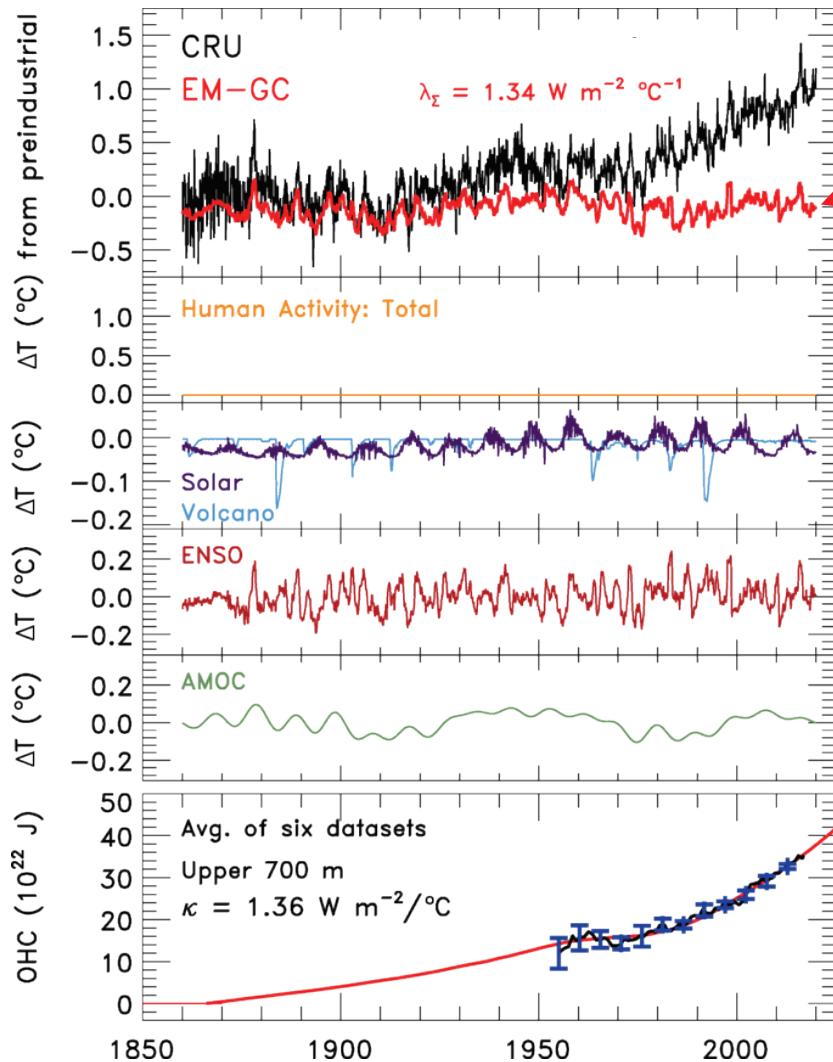
AMOC = Atlantic Meridional Overturning Circulation

Q_{OCEAN} = Ocean heat export =

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

Canty et al., ACP, 2013; Hope et al., Springer Climate, 2017, Hope et al., manuscript in preparation, 2020;
McBride et al., manuscript in preparation, 2020.

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

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

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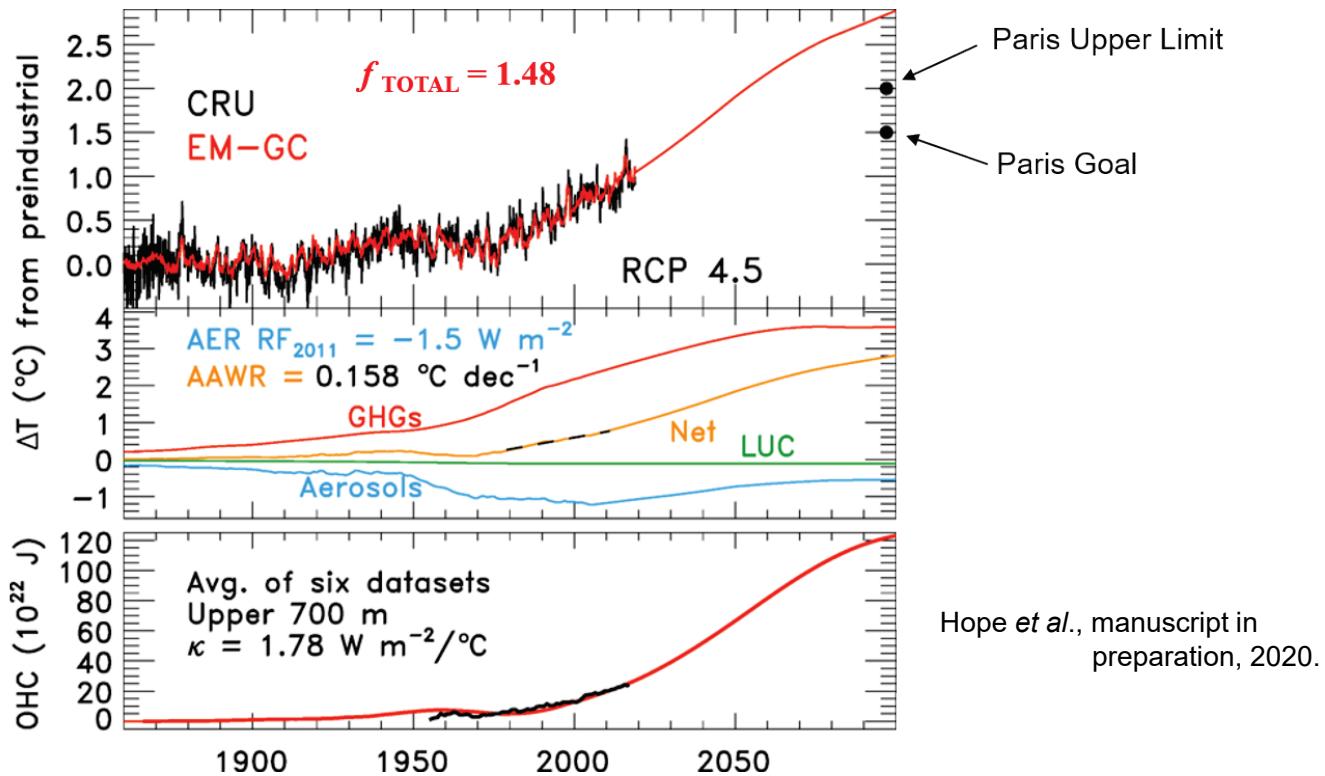
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Uncertainty in RF due to aerosols is a huge complication that places a fundamental uncertainty on how well future global warming can be forecast

$$\Delta T \approx \lambda_{\text{BB}} (1+f_{\text{TOTAL}}) \Delta RF - OHE$$

f_{TOTAL} : feedbacks due to water vapor, lapse rate, clouds, etc.

OHE : export of heat from atmosphere to world's oceans



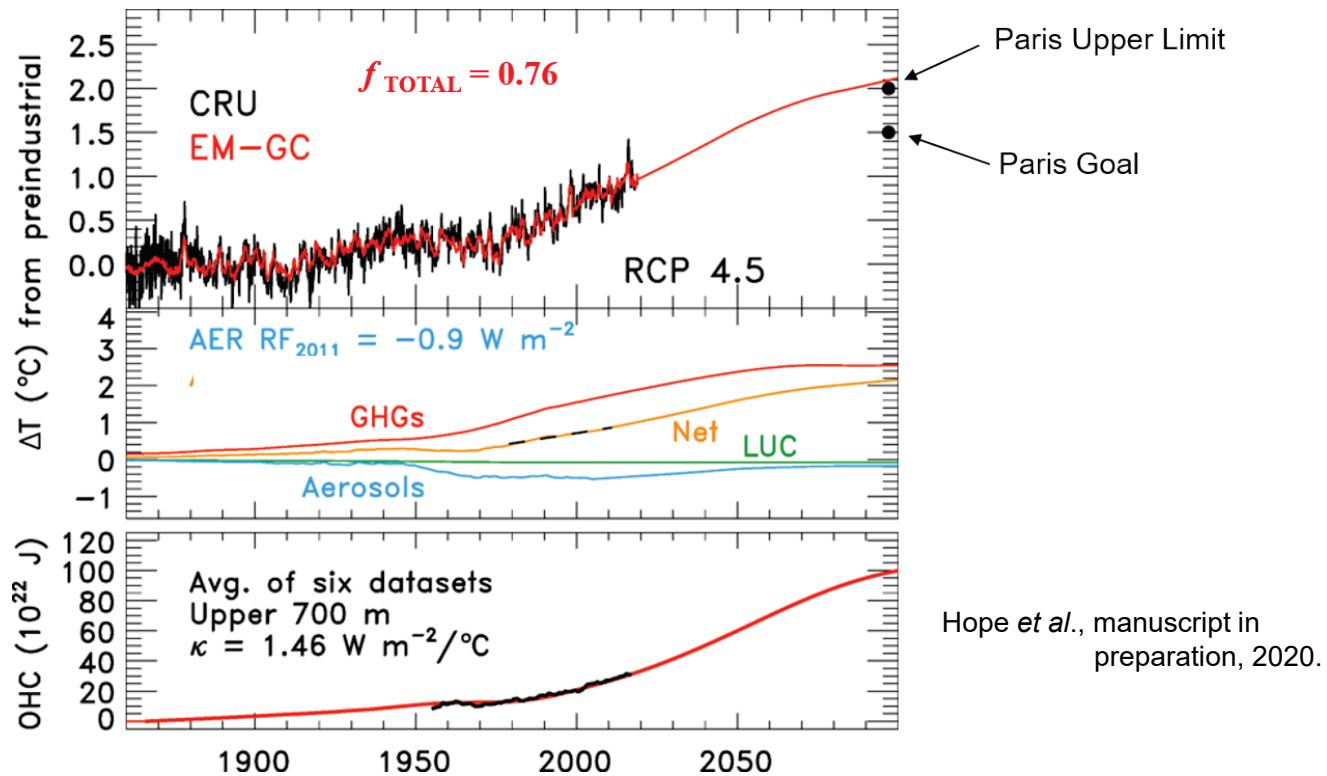
We assume that whatever value of climate feedback is inferred from the climate record will persist into the future. For Aerosol RF in 2011 of -1.5 W m^{-2} & assuming best estimate for H₂O and Lapse Rate feedback is correct, this simulation implies sum of other feedbacks (clouds, surface albedo) must be **strongly positive**.

Uncertainty in RF due to aerosols is a huge complication that places a fundamental uncertainty on how well future global warming can be forecast

$$\Delta T \approx \lambda_{\text{BB}} (1+f_{\text{TOTAL}}) \Delta \text{RF} - \text{OHE}$$

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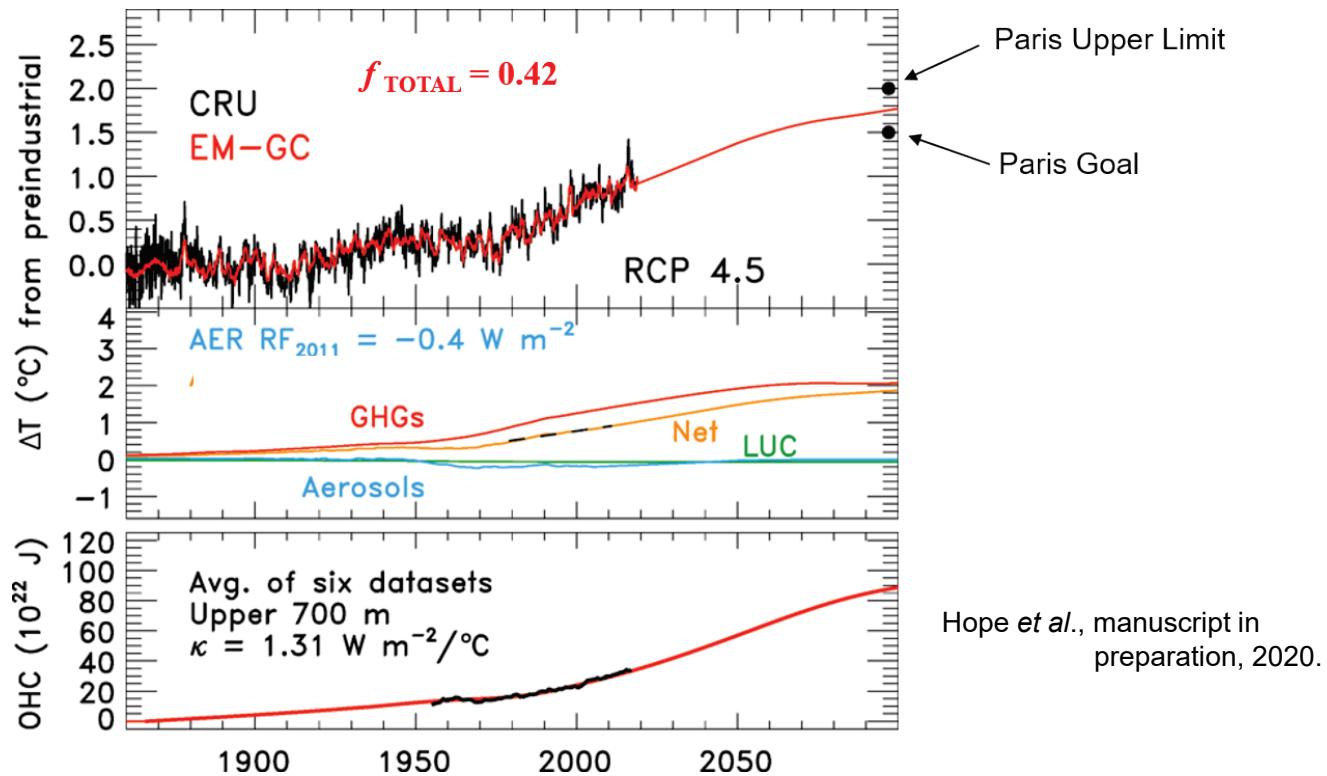
We assume that whatever value of climate feedback is inferred from the climate record will persist into the future. For Aerosol RF in 2011 of -0.9 W m^{-2} & assuming best estimate for H₂O and Lapse Rate feedback is correct, this simulation implies sum of other feedbacks (clouds, surface albedo) must be **moderately positive**.

Uncertainty in RF due to aerosols is a huge complication that places a fundamental uncertainty on how well future global warming can be forecast

$$\Delta T \approx \lambda_{\text{BB}} (1+f_{\text{TOTAL}}) \Delta RF - OHE$$

f_{TOTAL} : feedbacks due to water vapor, lapse rate, clouds, etc.

OHE : export of heat from atmosphere to world's oceans



We assume that whatever value of climate feedback is inferred from the climate record will persist into the future. For Aerosol RF in 2011 of -0.4 W m^{-2} & assuming best estimate for H₂O and Lapse Rate feedback is correct, this simulation implies sum of other feedbacks (clouds, surface albedo) must be **close to zero**.