

# Chemistry-Climate Interactions

Presented by

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Geophysical Fluid Dynamics Laboratory Review

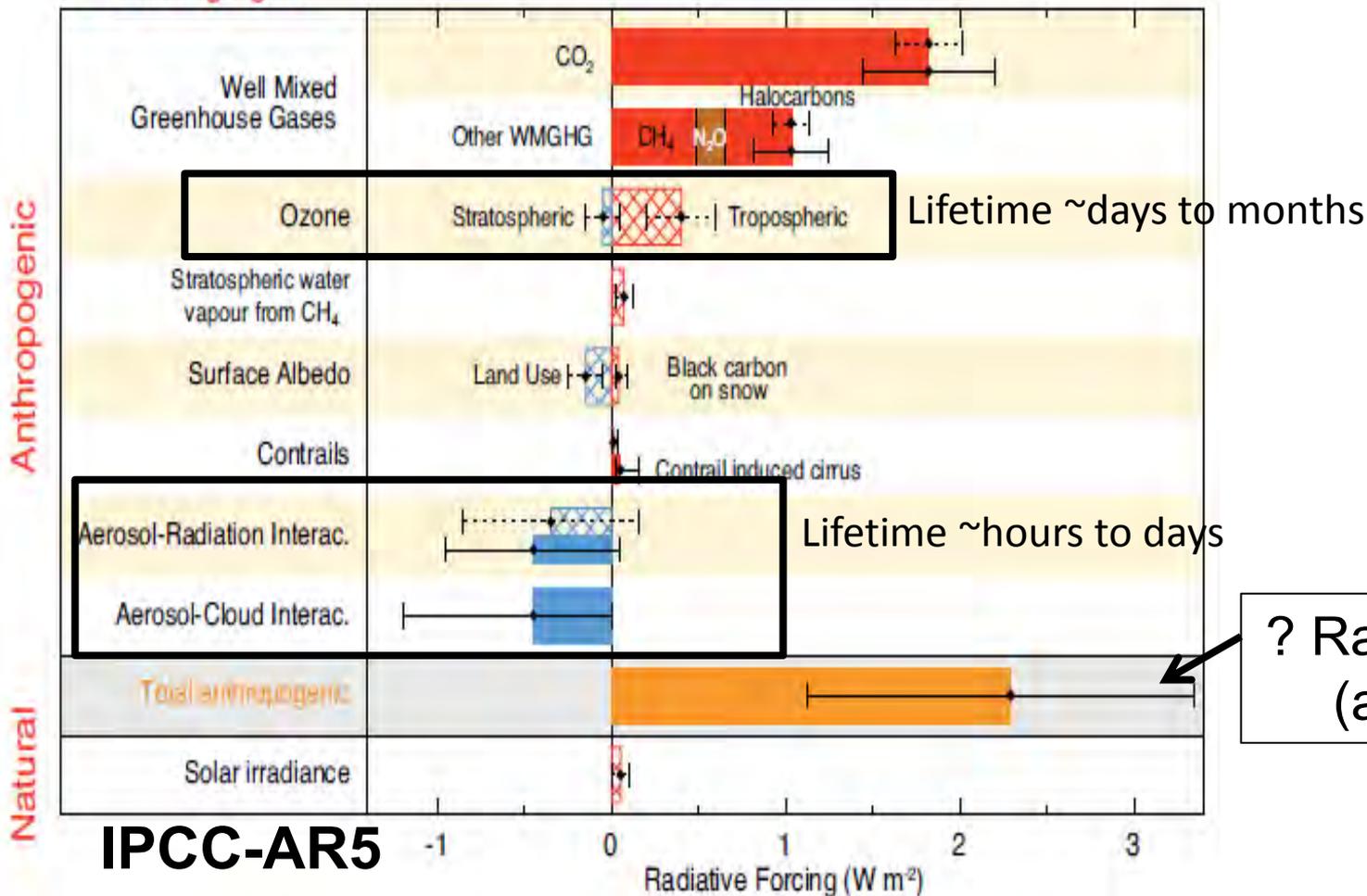
May 20 – May 22, 2014



# Atmospheric Chemistry plays a key role in the Climate System

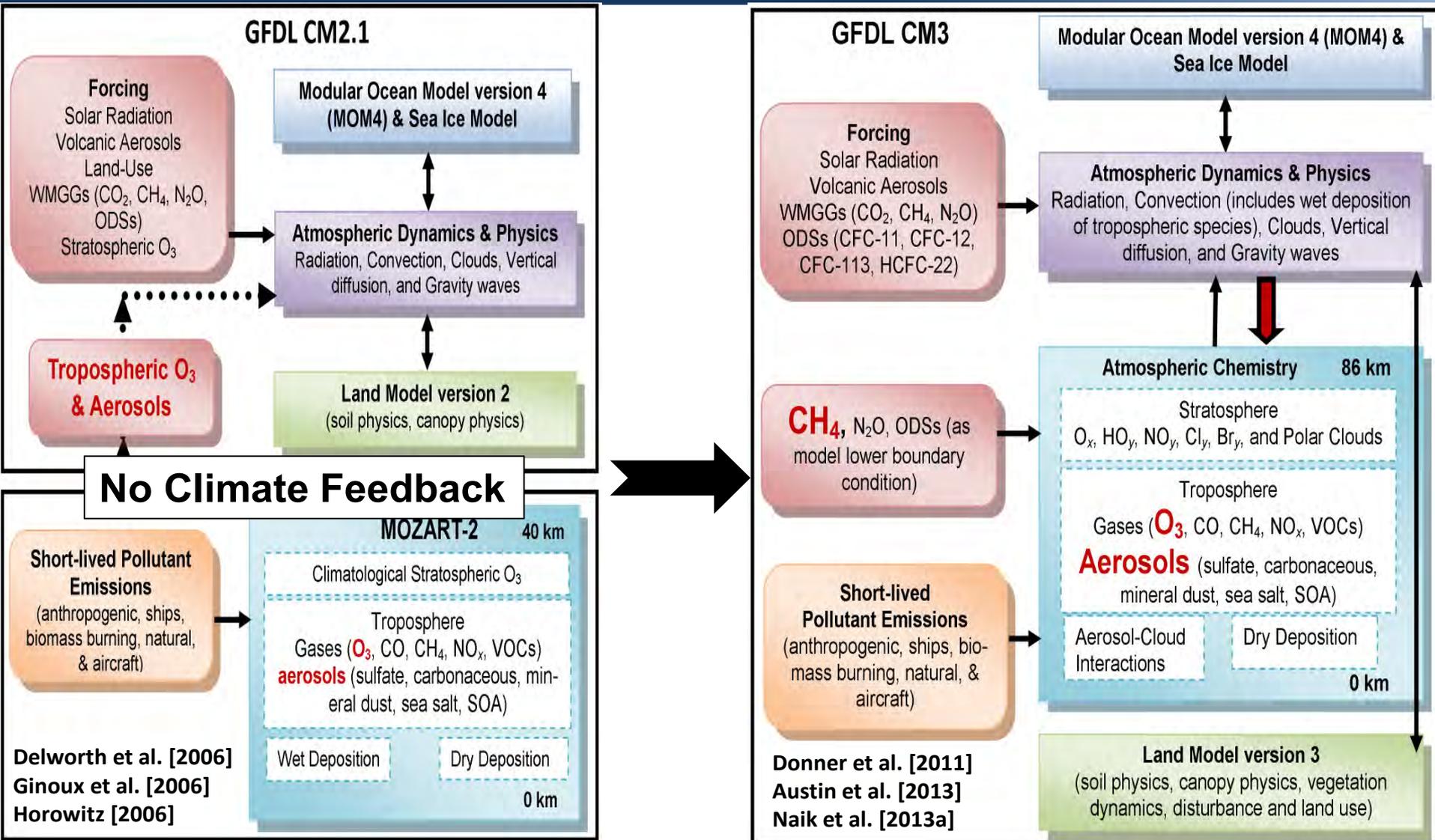
Radiative forcing of climate between 1750 and 2011

Forcing agent



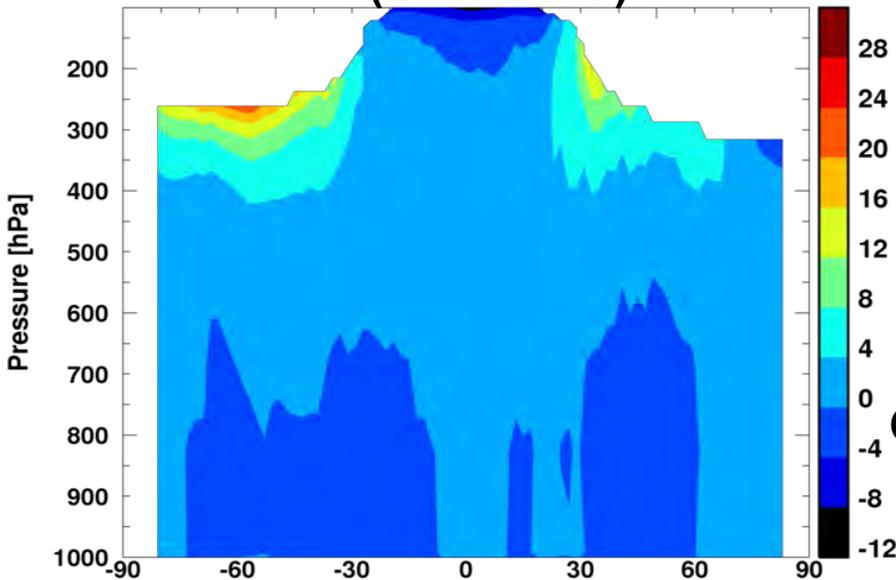
? Radiative Forcing (aerosols, O<sub>3</sub>)

# Seamless Atmospheric Chemistry in CM3



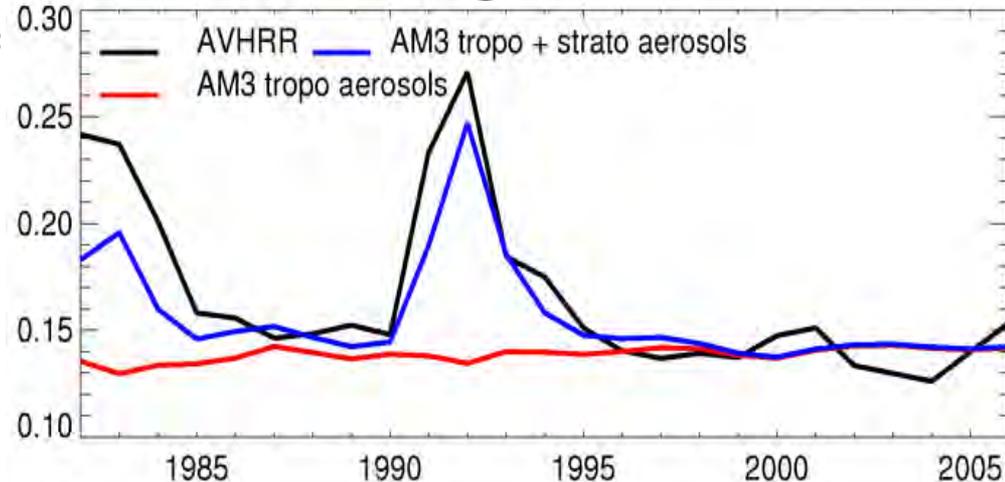
# AM3 (observed SST and Sea Ice) Captures Observations

## AM3 – TES Tropospheric O<sub>3</sub> (ppbv) (2005-2007)



AM3 captures the observed zonal mean O<sub>3</sub> to within  $\pm 4$  ppbv in much of the troposphere

## Global Mean Oceanic Aerosol Optical Depth (AOD) @ 550 nm

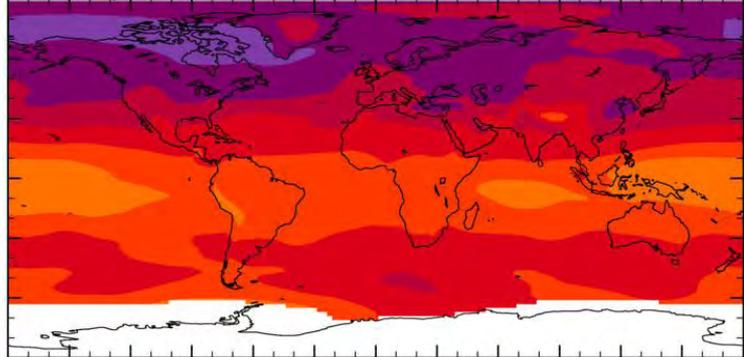


AM3 reproduces the observed evolution of total AOD over the ocean from 1982-2006

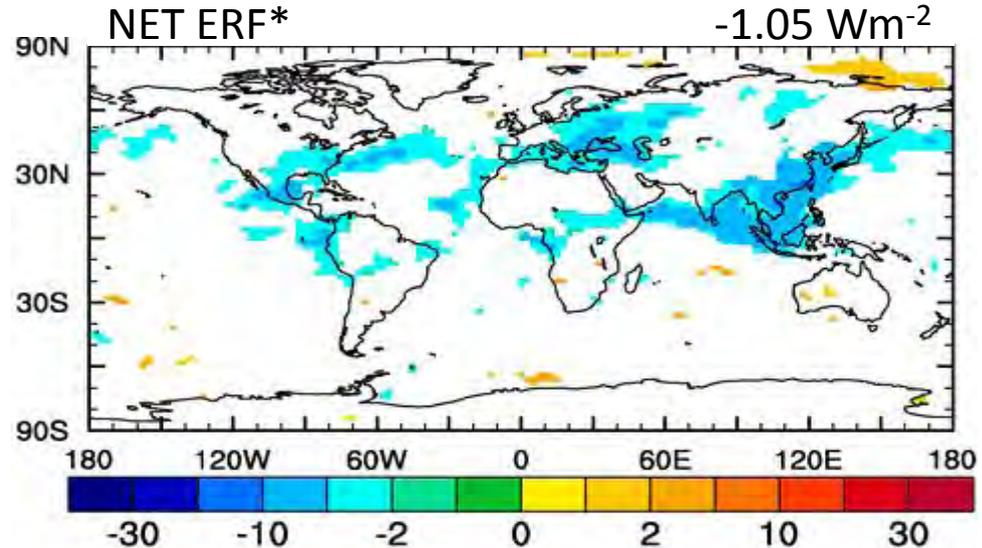
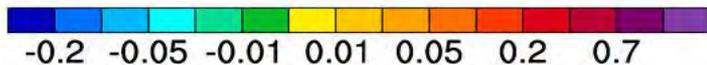
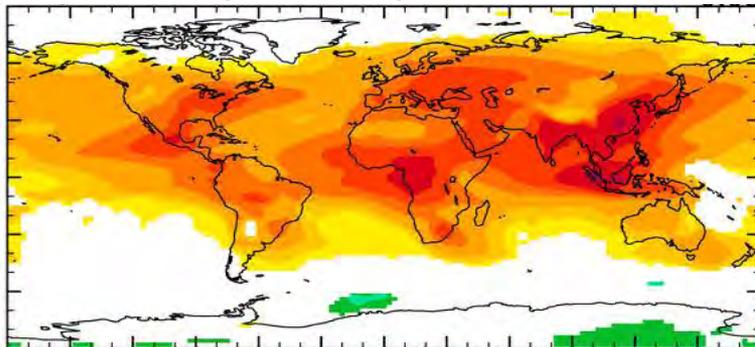
Naik et al. JGR [2013a]

# Net Negative Radiative Forcing (RF) due to Preindustrial to Present-day Increases in Tropospheric O<sub>3</sub> and Aerosols

$\Delta O_3$  (2000-1860) DU



$\Delta$ Aerosol Optical Depth (2000-1860)



Multi-model Mean [Shindell et al., 2013]

$$O_3 \text{ RF} = 0.33 \pm 0.18$$

$$\text{Aerosol ERF}^* = -1.17 \pm 0.4$$

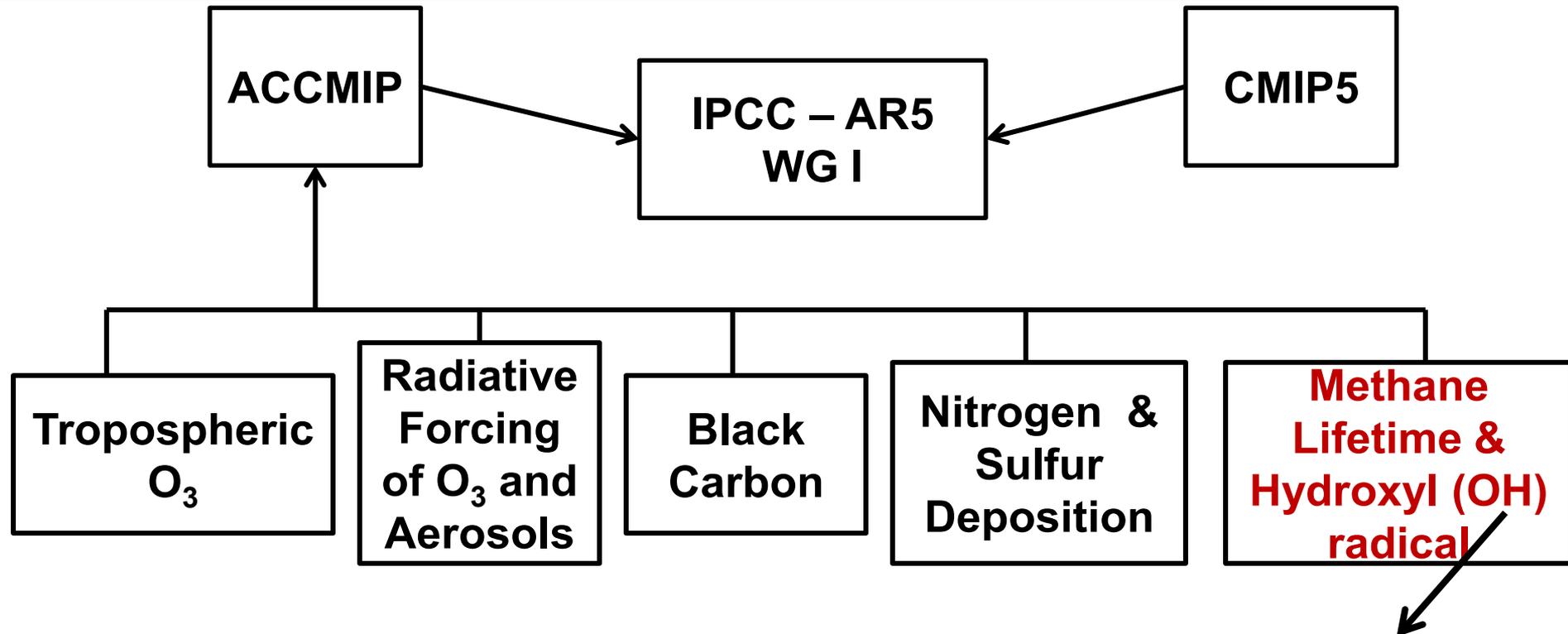
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$$\text{NET RF} = -0.84 \pm 0.50 \text{ W m}^{-2}$$

\*NET EffectiveRF: Change in TOA net radiative flux with fixed sea surface temperature

Naik et al. JGR [2013a]

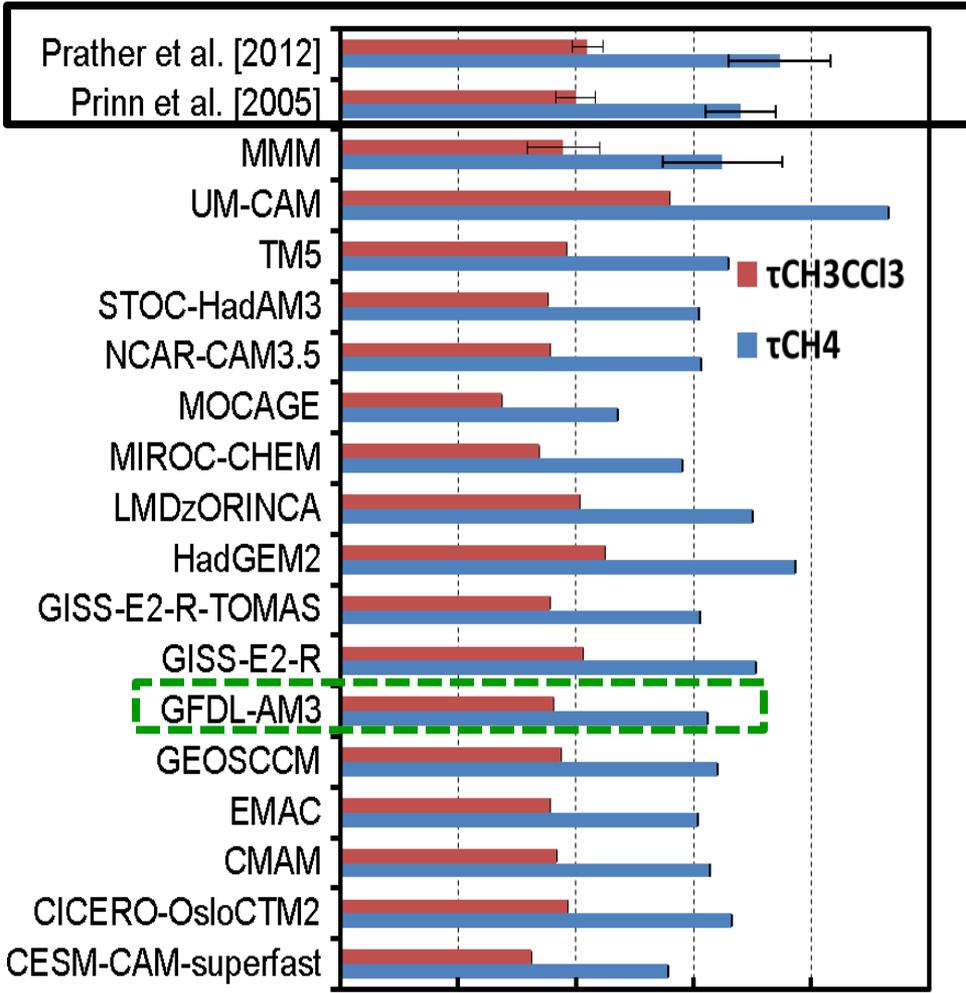
# Contribution to the Atmospheric Chemistry Climate Model Intercomparison Project (ACCMIP)



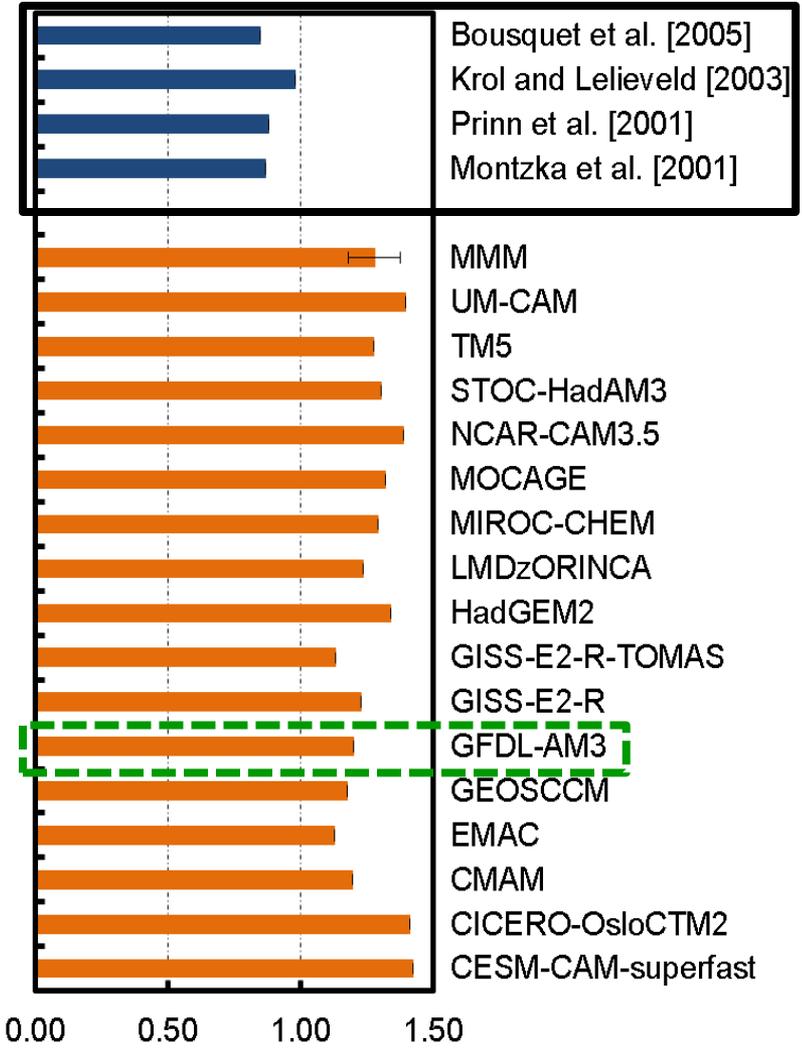
- **Dominant atmospheric oxidizing agent** → abundance and lifetime of radiatively active species (e.g.  $\text{CH}_4$ , ozone depleting substances)
- **Extremely short-lived (~1s)** → global measurements are hard to make, rely on proxy methods (e.g.  $\text{CH}_3\text{CCl}_3$  lifetime) or forward models to estimate global mean OH

# Global Chemistry Climate Models (CCMs) overestimate mean OH relative to observational constraints

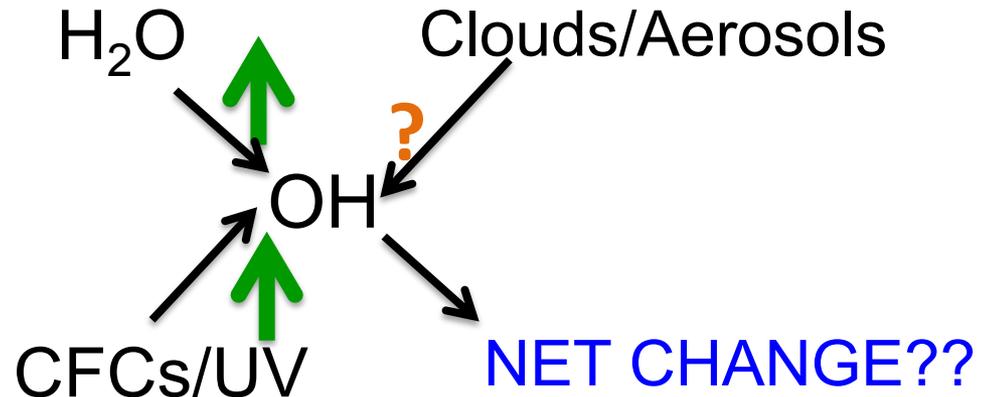
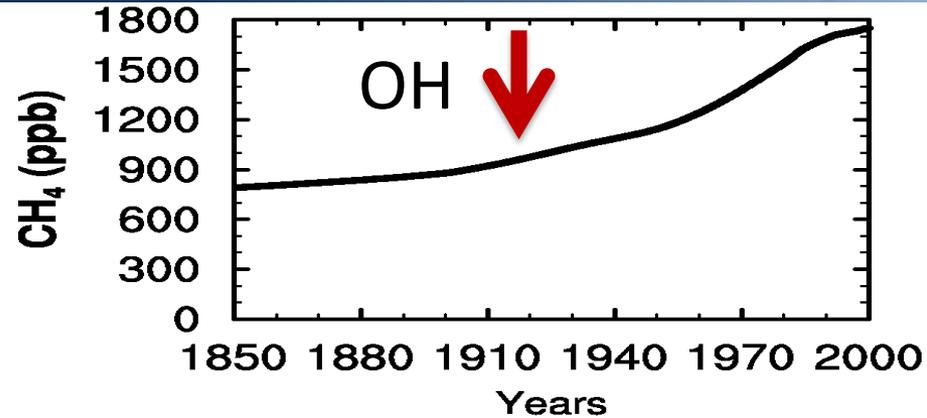
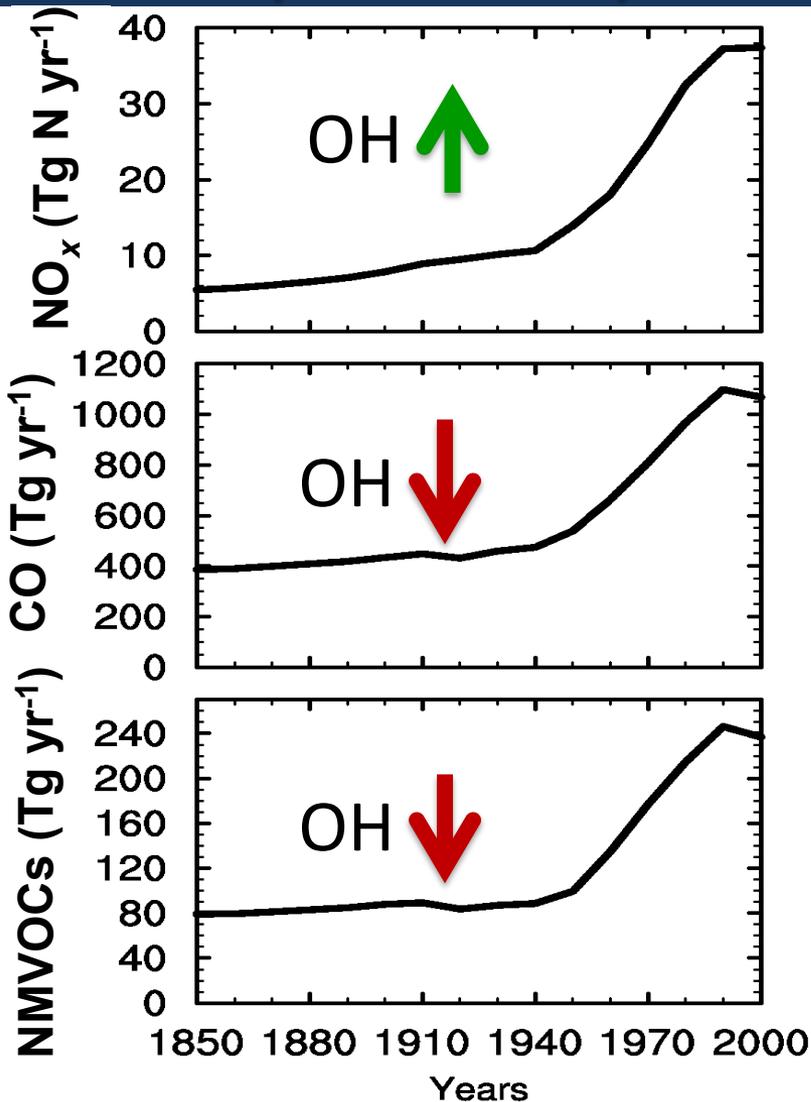
Years  
0.0 3.0 6.0 9.0 12.0 15.0



## OH North/South Ratio



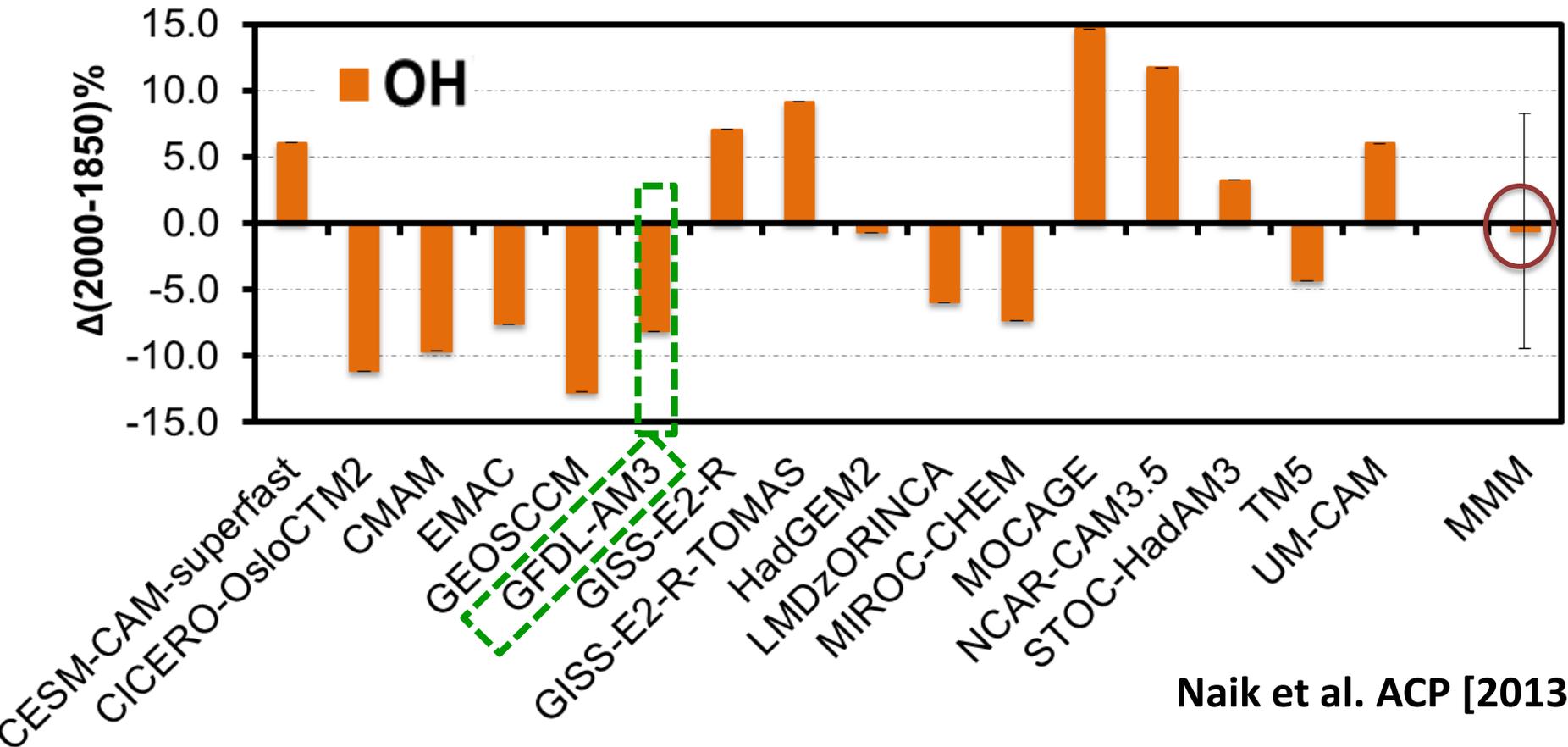
# How has global mean OH changed in response to historical (1850-2000) emission increases and climate change?



No consensus amongst prior modeling studies (1991-2012)

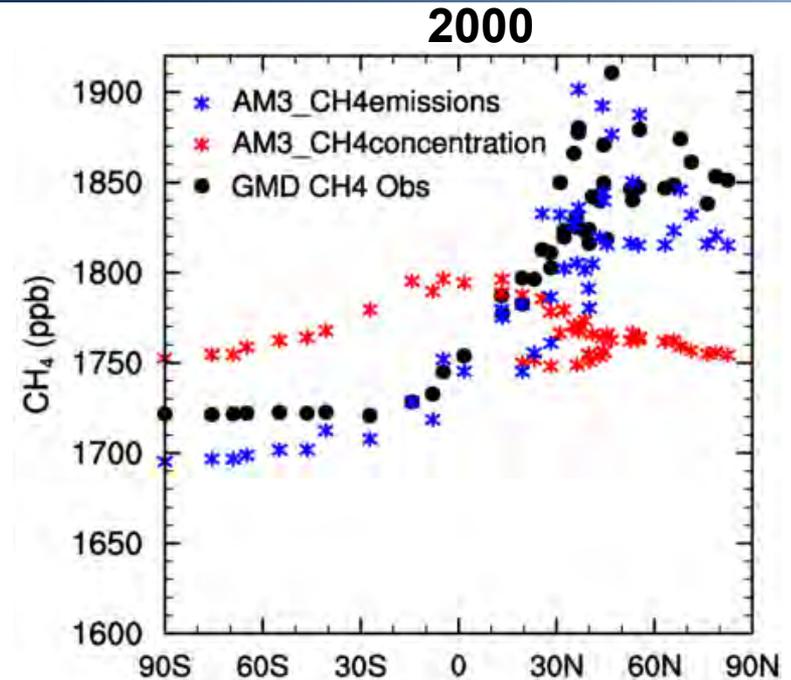
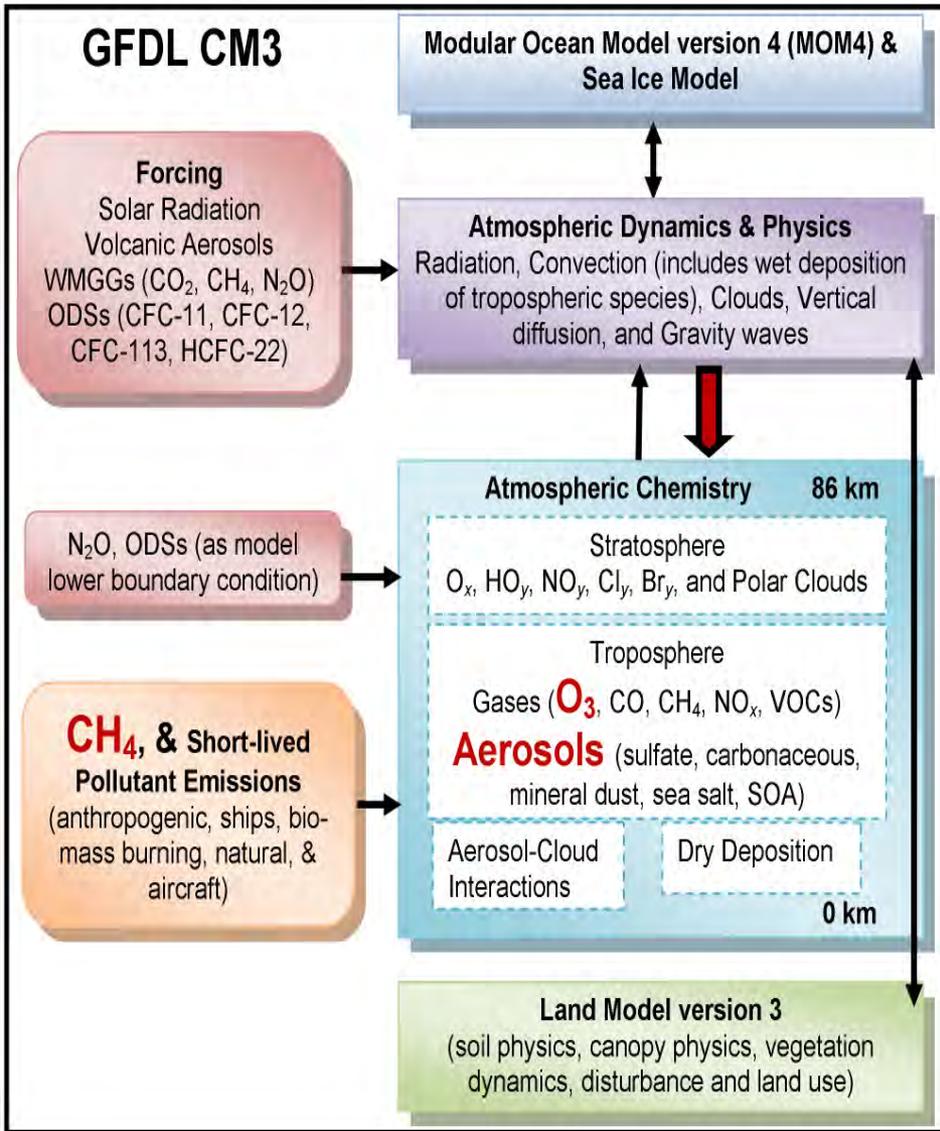
# Large intermodel diversity in the sign of OH change over the historical period with no clear trend

Global mean OH is well-buffered against perturbations  
[Lelieveld et al. 2004, Montzka et al. 2011]



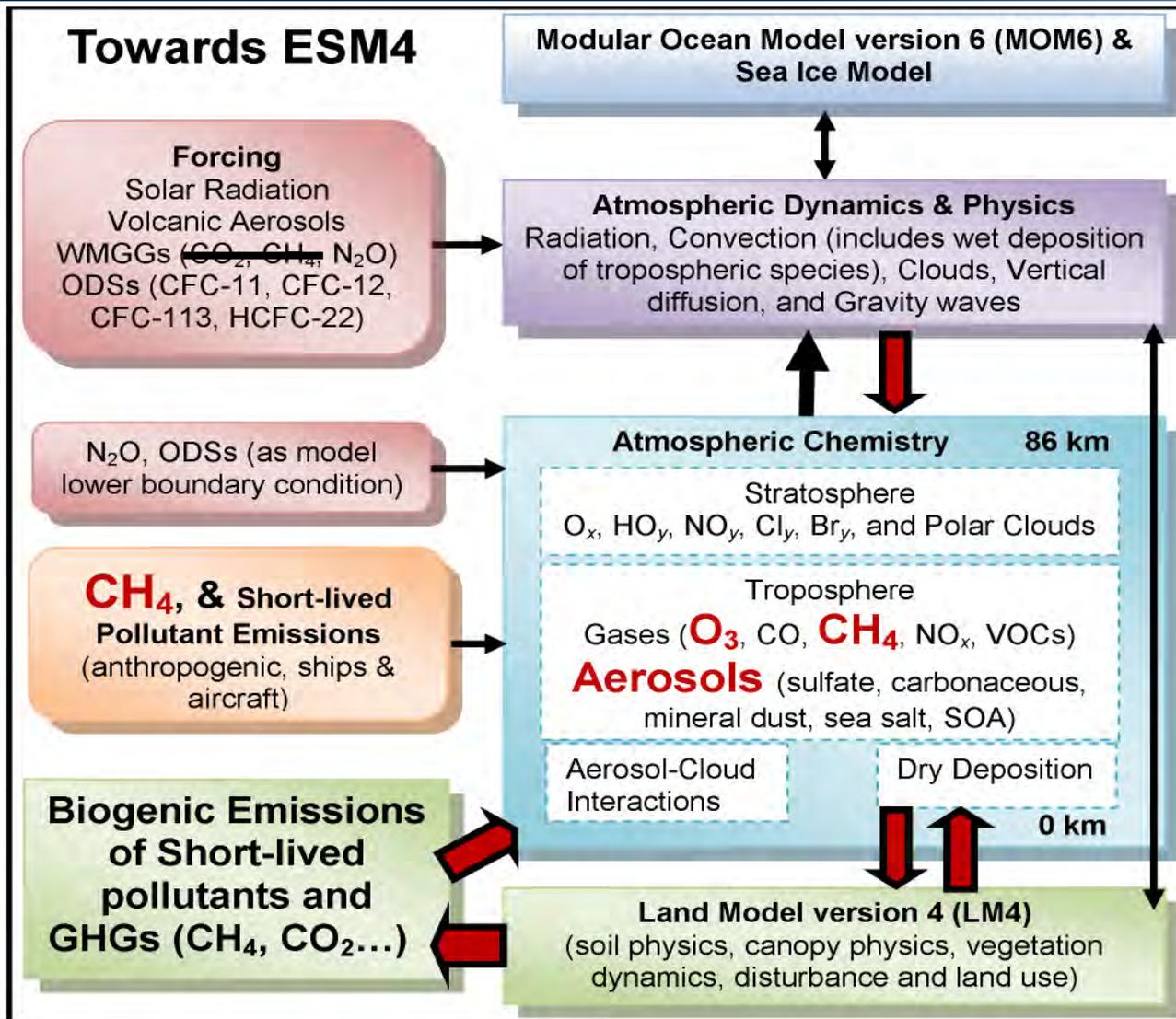
Naik et al. ACP [2013b]

# Future Directions: Realistic CH<sub>4</sub> Simulation



- Realistic simulation of CH<sub>4</sub> will allow us to investigate:
  - drivers of past and future changes in atmospheric CH<sub>4</sub>
  - impact on other chemical species, e.g., tropospheric O<sub>3</sub> and its RF

# Future Directions: Atmospheric Chemistry in the Earth System



- **A comprehensive treatment of atmospheric chemistry within GFDL's global climate model (CM3)** allows us to advance the scientific understanding of the effect of short-lived pollutants on climate
- **GFDL helped lead** a multi-model investigation of the global mean OH historical trend
  - Models do not give clear indication of global OH trend; multi-model mean change is consistent with recent observational inference
- **Integrating atmospheric chemistry within the Earth System Model** opens up the possibility of addressing whole new sets of questions within the broader **Earth System Science**

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