

Ozone and methane changes and interactions

Regional to global scales

Ivar S.A. Isaksen

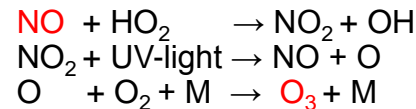
University of Oslo/CICERO

Tropospheric impact of NO_x and CO emissions on ozone and methane

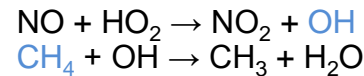
NO contribute to composition change/RF through:

Chemical formation of the secondary compound O₃
Reduction in CH₄ through reduced lifetime (enhanced OH)

Enhanced ozone production:



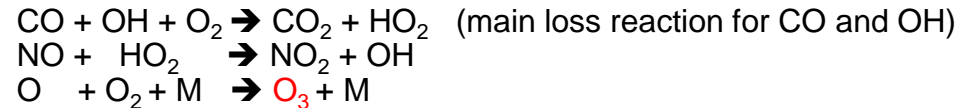
Enhanced methane loss:



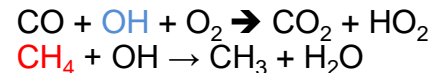
CO (and VOC) contribute to composition change/RF through:

Chemical formation of the secondary compound O₃
Enhanced CH₄ through increased lifetime (reduced OH)

Enhanced ozone production:



Enhanced methane production:

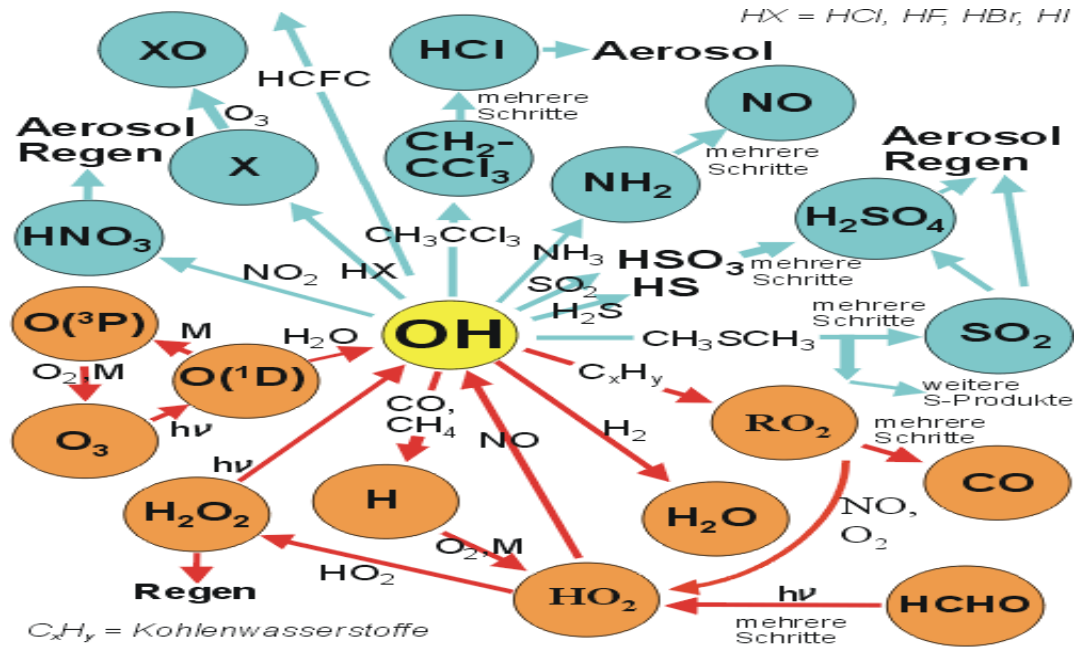


NO: **Increased** climate effect through O₃, **reduced** climate effect through methane

CO: **Increased** climate effect through O₃, **increased** climate effect methane methane

:

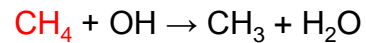
OH a key compound in atmospheric chemistry



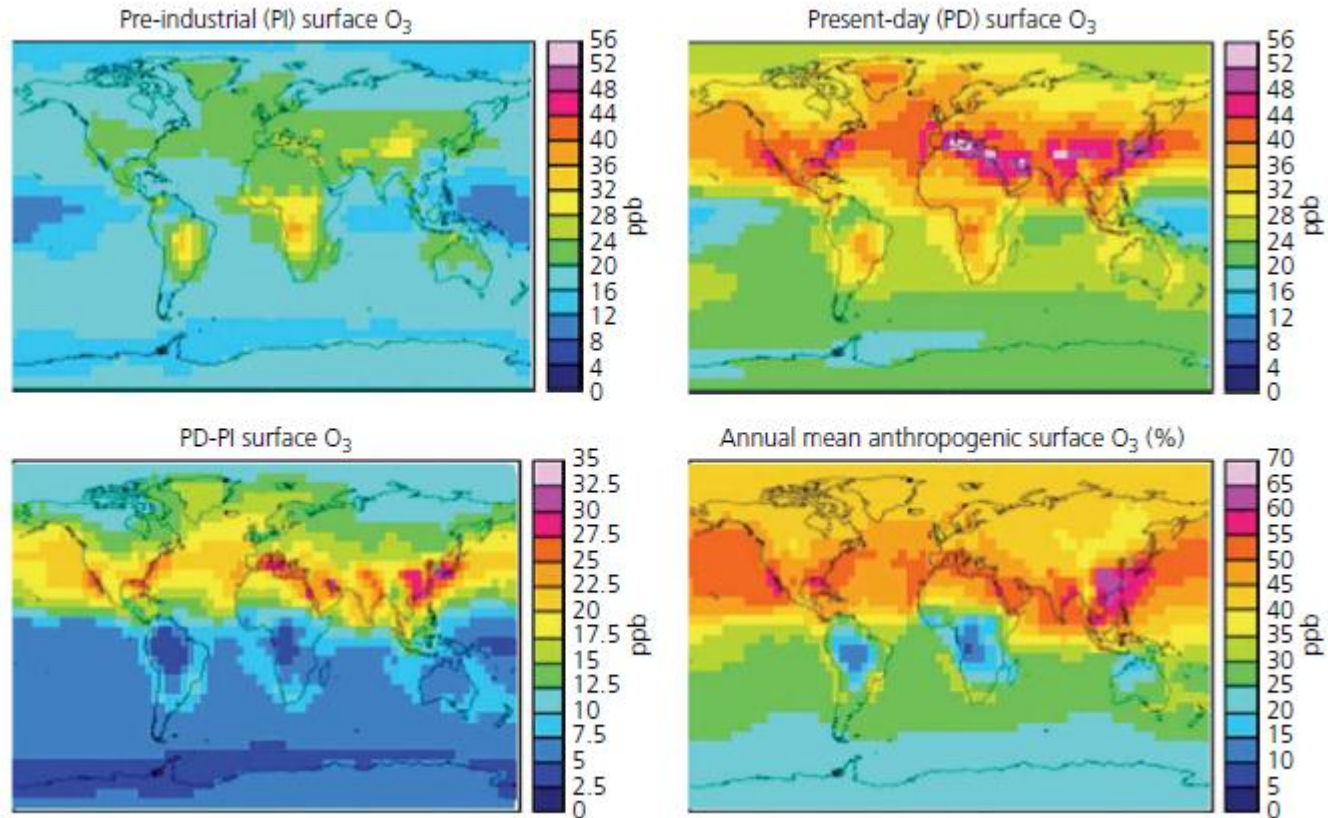
From W. CHAMEIDES and D.DAVIS, Chem. Eng. News 1982

Key chemical reactions involving the hydroxyl radical (OH)

More than 90 % of atmospheric methane (CH₄) is lost through the reaction with OH:

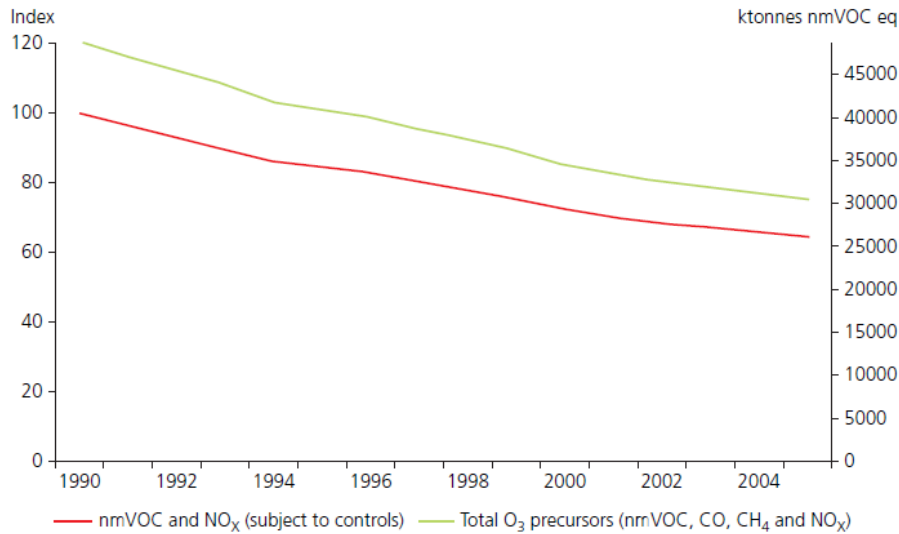


Surface ozone distribution



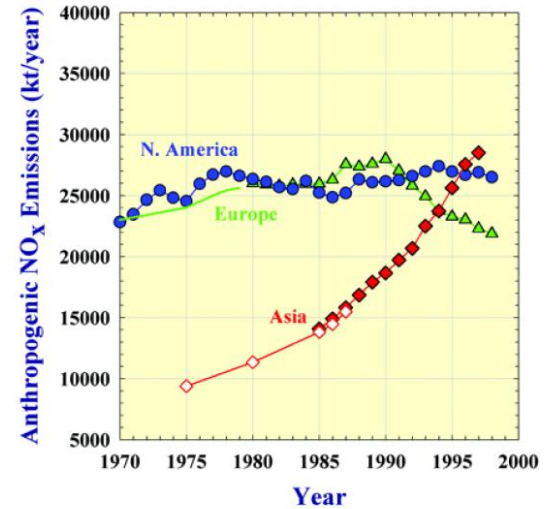
Modelled global changes in surface O₃ concentrations between pre-industrial times and present day. Multi-model mean surface layer annual mean O₃ (ppb) is presented. PI: pre-industrial, PD: present day, PD-PI: modelled increase in O₃, and in % of annual mean O₃ attributable to anthropogenic sources.

Trend and regional differences in pollution emissions



Emissions of O₃ precursors in Europe (EEA-32 member countries) from 1990–2005

Source: EEA (2007)



Changes in anthropogenic NO_x emissions in North America (United States and Canada), Europe (including Russia and the middle East), and Asia (East, Southeast, and South Asia).

Source: Akimoto, 2003

Tropospheric ozone column distribution

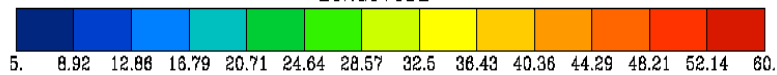
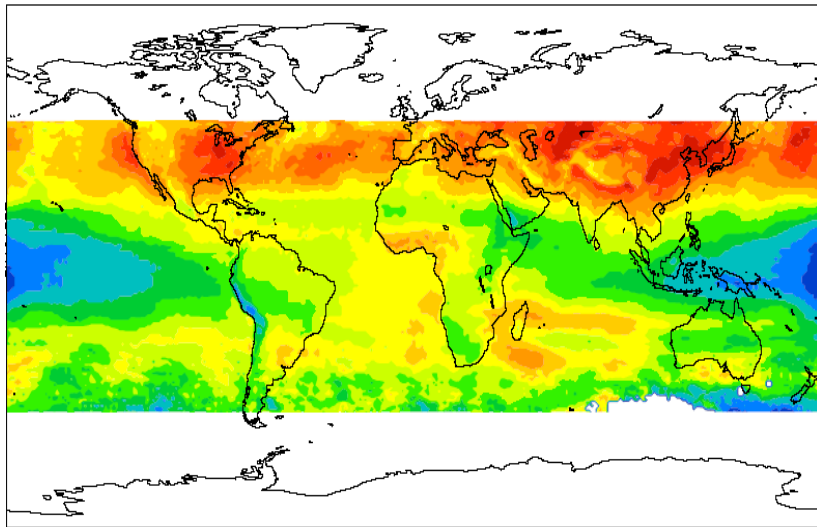
Question: How well can models represent values and patterns?

Tropospheric ozone residual

July 2000

From Fishman (NASA, Langley)

Dobson units

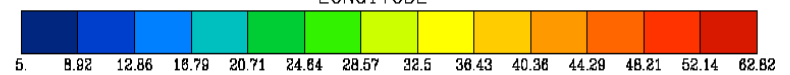
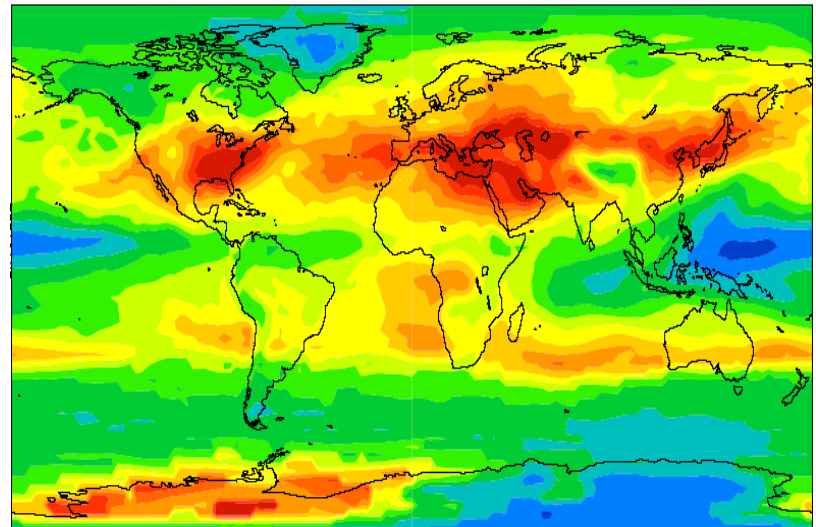


CTM2 Tropospheric column

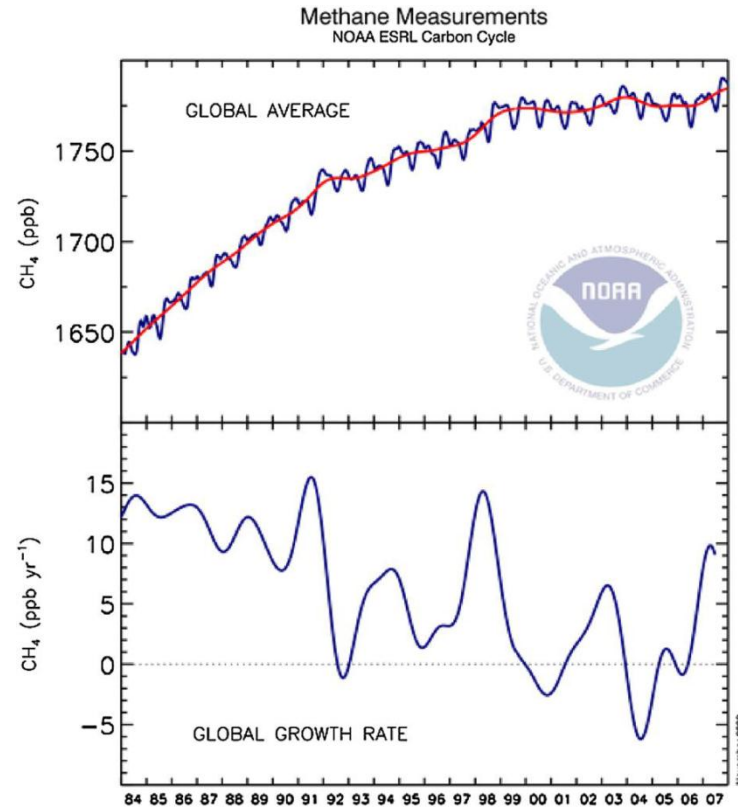
Jul 2000

From Dalsøren et al., 2006

Dobson units

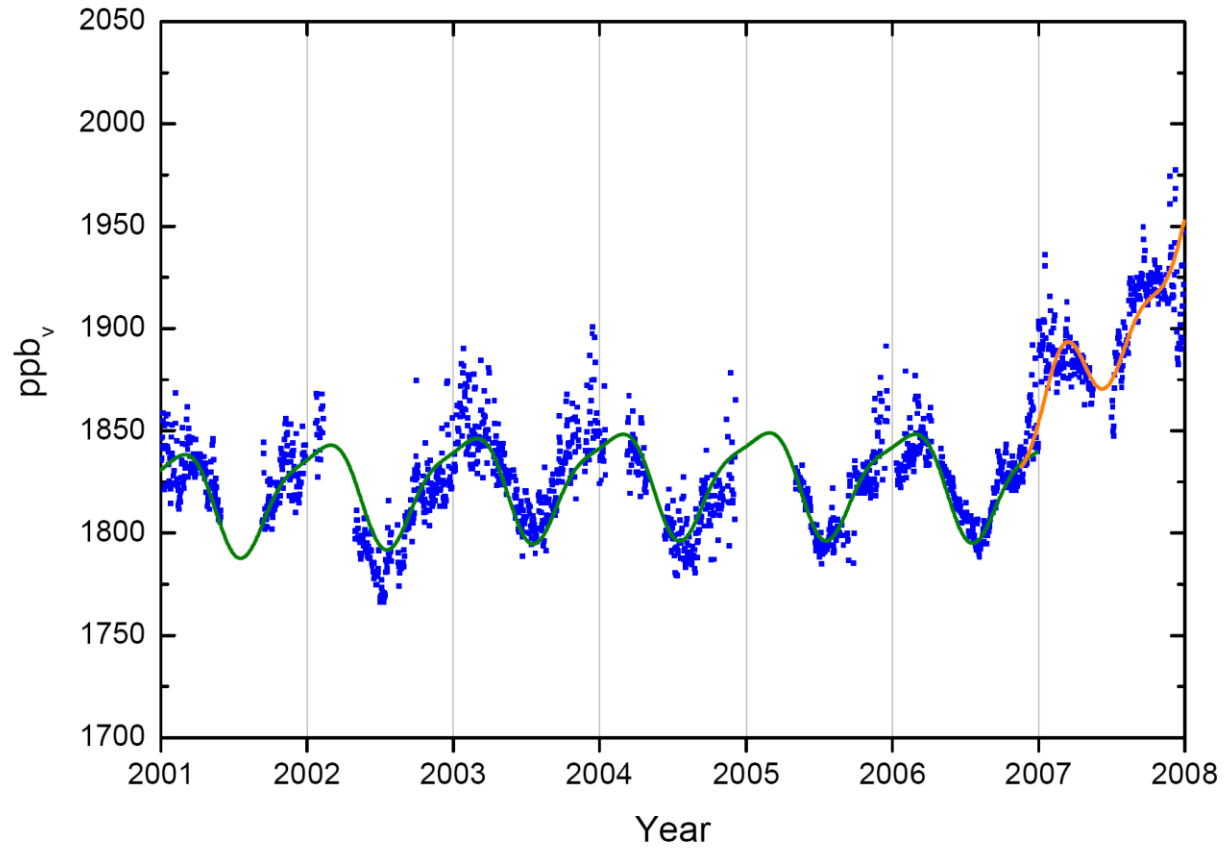


Global surface methane distribution



20 years of methane growth

Observations of methane for the period 2001-2007 at Zeppelin observatory, 79 deg. N



Significant methane increase in the Arctic after 2006

Emission from international shipping and aviation

- Land-based emissions have been reduced in many countries
- Emissions from international aviation and shipping are rapidly increasing
- Aviation contributes ~ 2% of global NO_x emission
- International shipping contributes ~15%, of global NO_x emission

Large increases are expected

- Global passenger air travel, growth by 5% per year between 2000 and 2020
- European NO_x emissions from shipping (32% of land-based NO_x emissions) will exceed these sources by 2020

Ozone and methane perturbations by ship and aircraft

NO_x emission from aircraft → increased O₃ :

- Occurs at cruise altitudes (in the UTLS region, 10 to 12 km) where ozone is an efficient climate gas
- At UTLS heights ozone has a long lifetime and is transported to Arctic latitudes
- Background NO_x is important

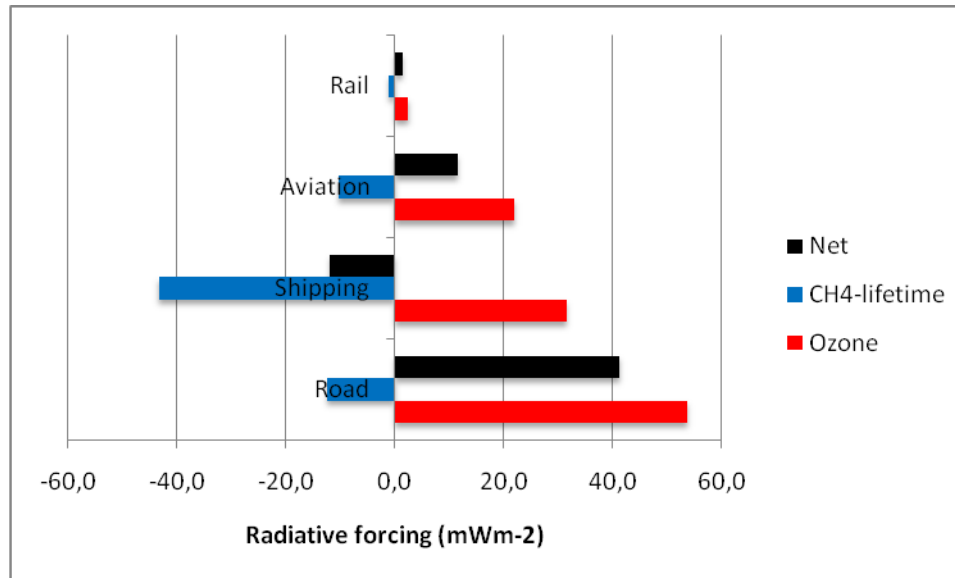
NO_x emission from ship → increased O₃ and reduced CH₄:

- Occurs in remote pristine regions with low background NO_x
- Emission index for aircraft: $NO_2/CO > 1$
- Emission index for ship: $NO_2/CO \gg 1$
- Surface emissions: $NO_2/CO \ll 1$

In particular ship emission have significant impact on OH (increase)

Impact from the transport sector

Estimate of radiative forcing for the different transport sectors



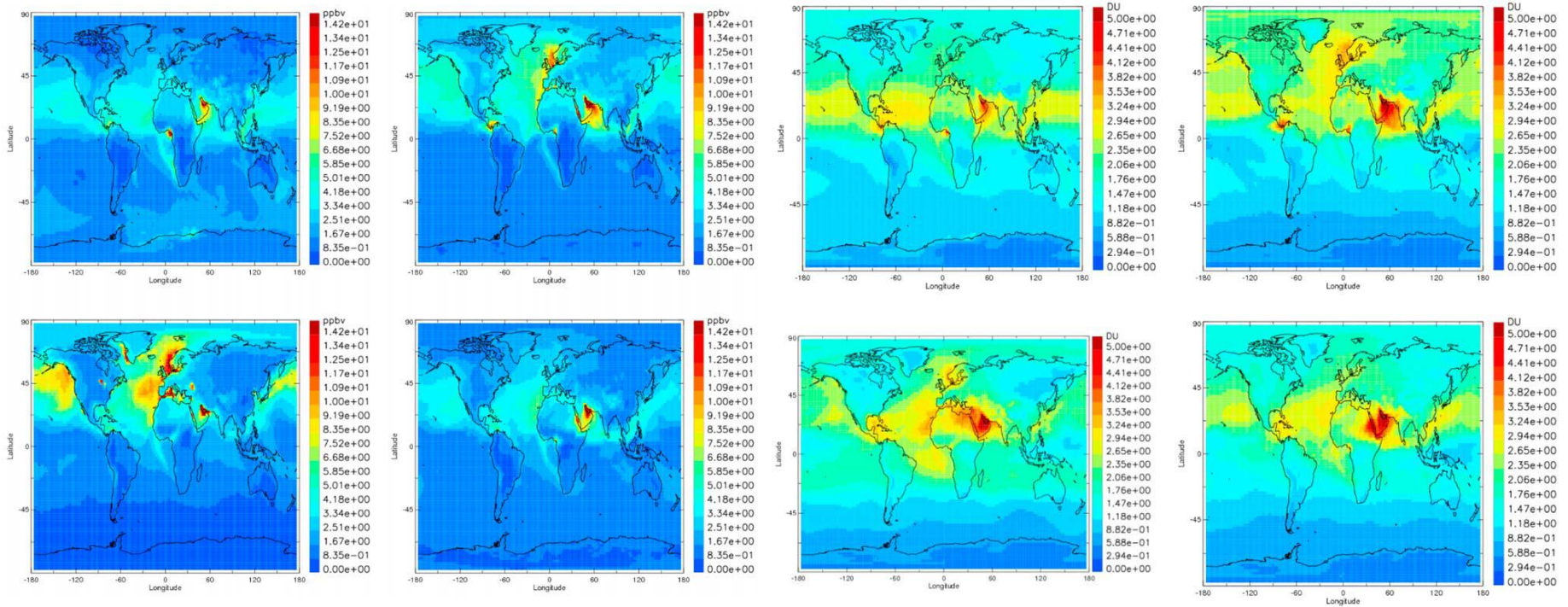
Fuglestvedt et al., 2008

Radiative forcing from CH₄ (changes in lifetime) and from O₃, and the corresponding net radiative forcing for four transport sectors from pre-industrial time to present due to NO_x emissions.

Large differences between the sectors:

- Road transport large positive contribution
- Ship emission negative contribution
- Aircraft have noticeable positive and negative contributions

Impact of ship emission on tropospheric ozone

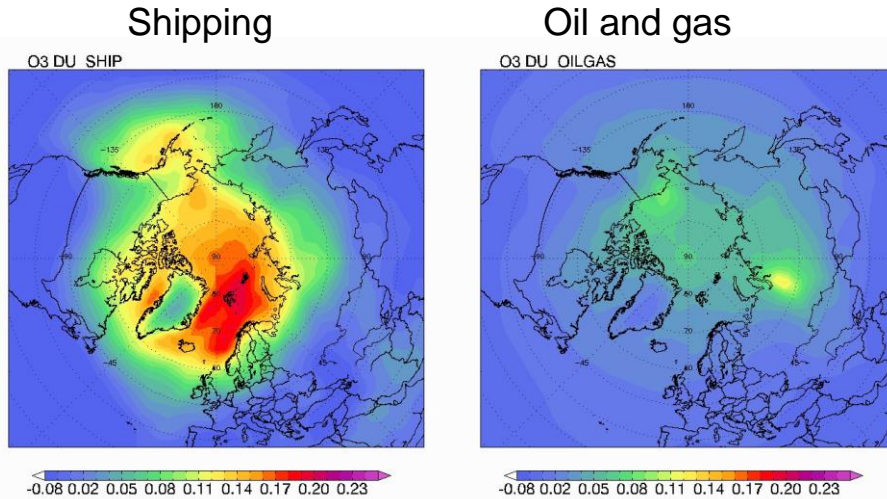


Surface (left) and column (right) ozone change due to year 2000 ship emissions for the months (top left) January, (top right) April, (bottom left) July and (bottom right) October.

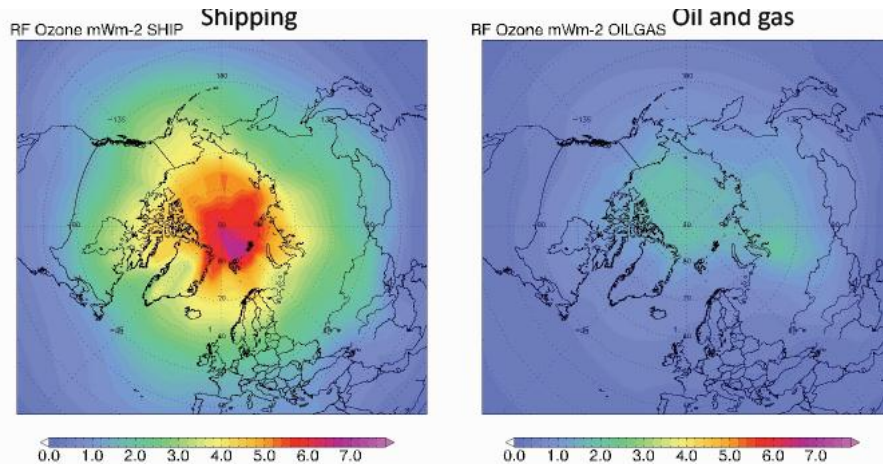
Max tropospheric column change from ship: 10 to 20 %.

Activity in the Arctic:

Current changes in tropospheric ozone columns and in methane

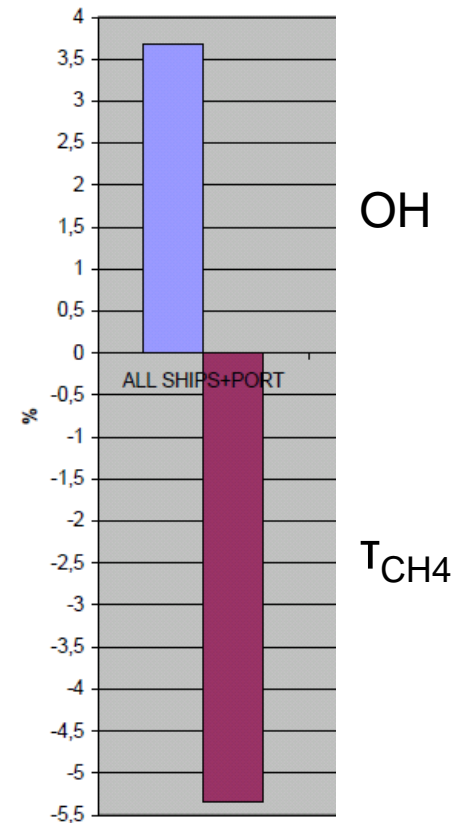


Max ozone column change from shipping ~ 1% of trop. column



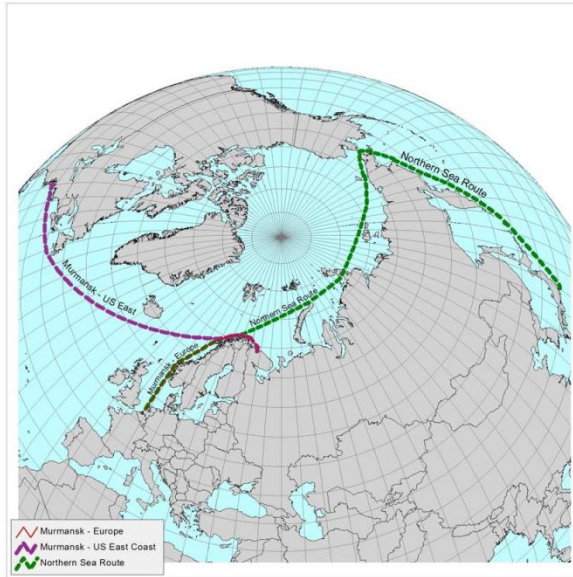
Radiative Forcing from ozone increases in mW/m²

Ship emission impact on methane lifetime

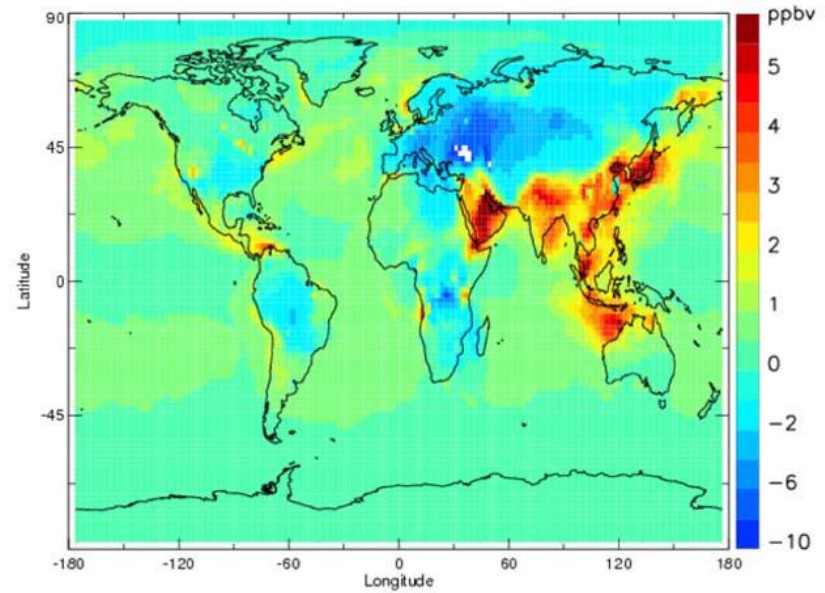


Changes in global average OH and CH₄ lifetime from ship emissions (2000)

Impact of future sea transport



Northern Sea Route, expected oil trades and main ports for the tanker transport in 2015.



Ozone surface change from July 2000 to 2015 from a sensitivity study including both changes in ship emissions and land based anthropogenic sources.

Future chemical impact

- Issues:
 - Climate change
 - Air pollution
 - Interaction air pollution/climate
- Key compounds:
 - O₃, OH (CH₄)
- Processes related to:
 - Emission and emission scenarios (NO_x, CO)
 - Regional and sectorwise processes
 - Modelling capability
 - Climate change (Atmospheric processes, emission, deposition)

Impact on chemically active climate gases in a warmer climate

Permafrost thawing with enhanced emission of CH₄

- Long term impact on global climate

Ozone increase during frequent high pressure systems

- Regional pollution

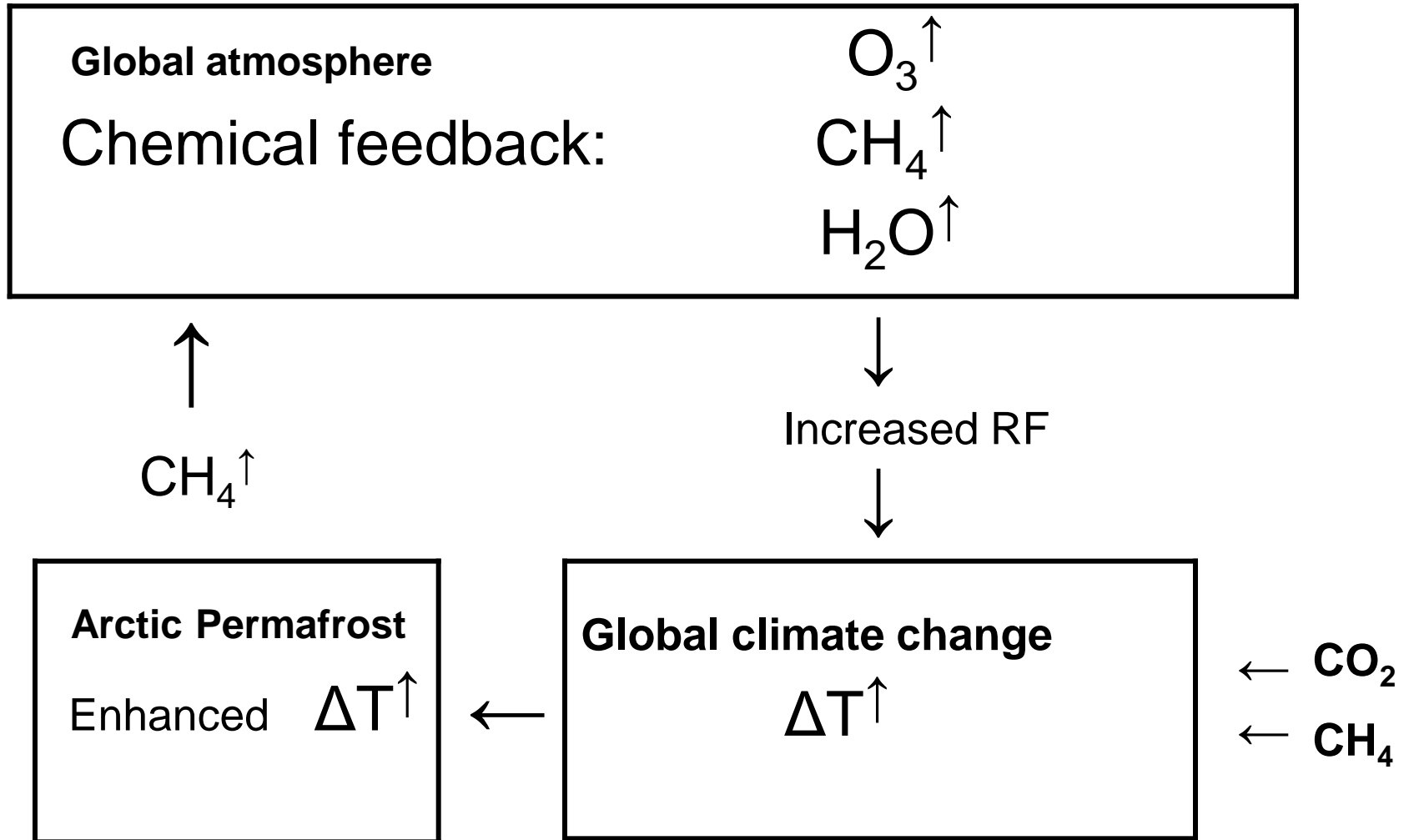
Warmer climate

- Enhanced permafrost thawing → Release of greenhouse compounds (CH₄, CO₂)
- More severe pollution episodes (ozone, particles)
- Changes in emission of biogenic compounds (isoprene, nitrogen oxides)
- Changes in atmospheric oxidation (increased OH → reduced CH₄)

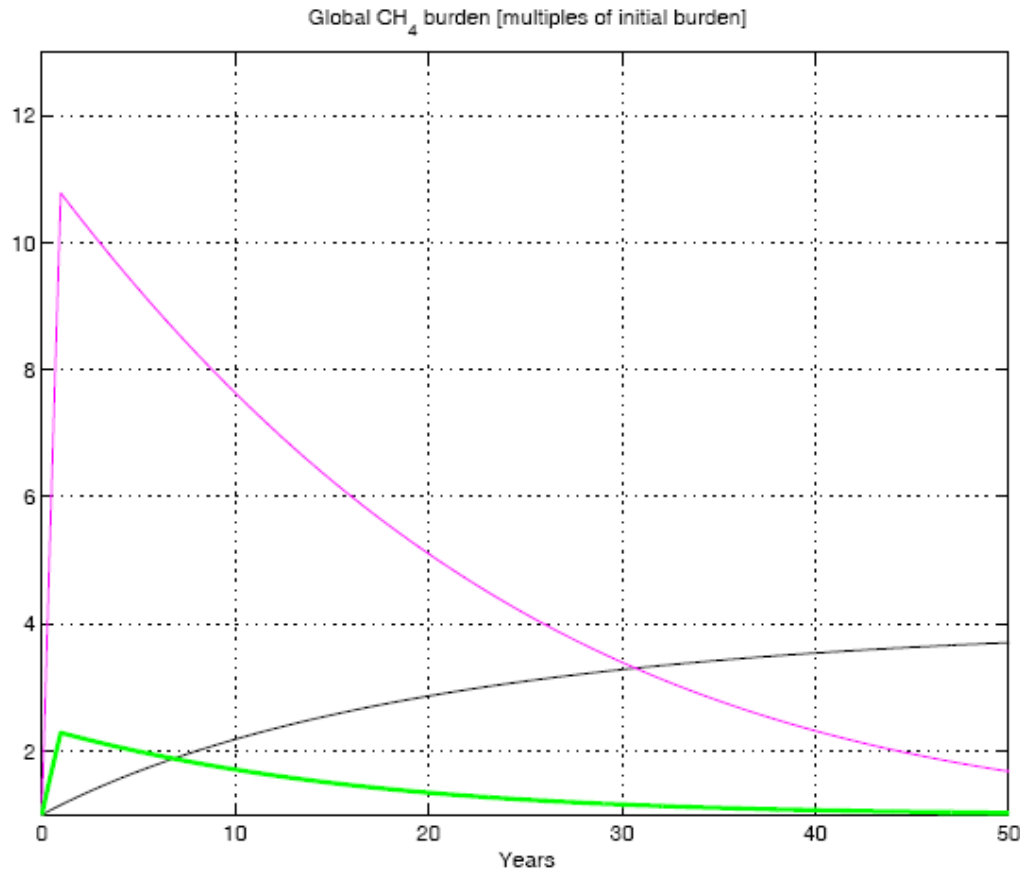
Atmospheric chemical changes non linear

- Not a 1 to 1 relation between release and atmospheric changes (CH₄ a factor 1.5)
- Formation of secondary compounds (O₃, sulfat) depends on precursors

Impact on climate from permafrost thawing



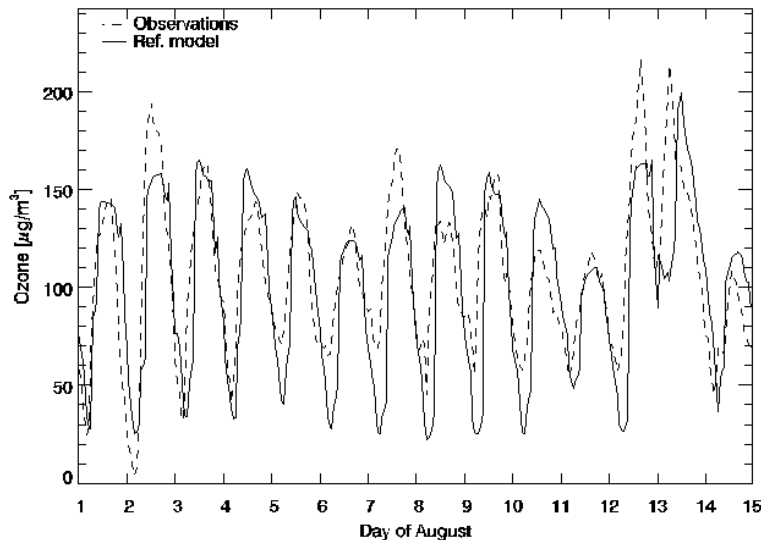
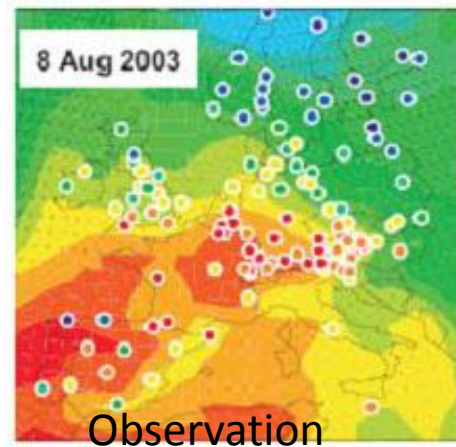
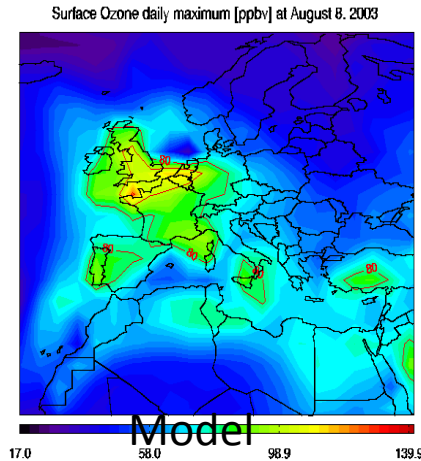
Methane emissions in the Arctic during permafrost thawing



Max RF for a pulse of
5 Gt: 1.1 W/m²

Global methane increase for pulses of 5 Gt (green), 50 Gt (lilac), and continuous emission of 0.8 Gt/year (black)

Possible effect of a warmer climate: Enhanced surface ozone during the August 2003 heat wave



Extremely high surface ozone and particle levels over Western Europe

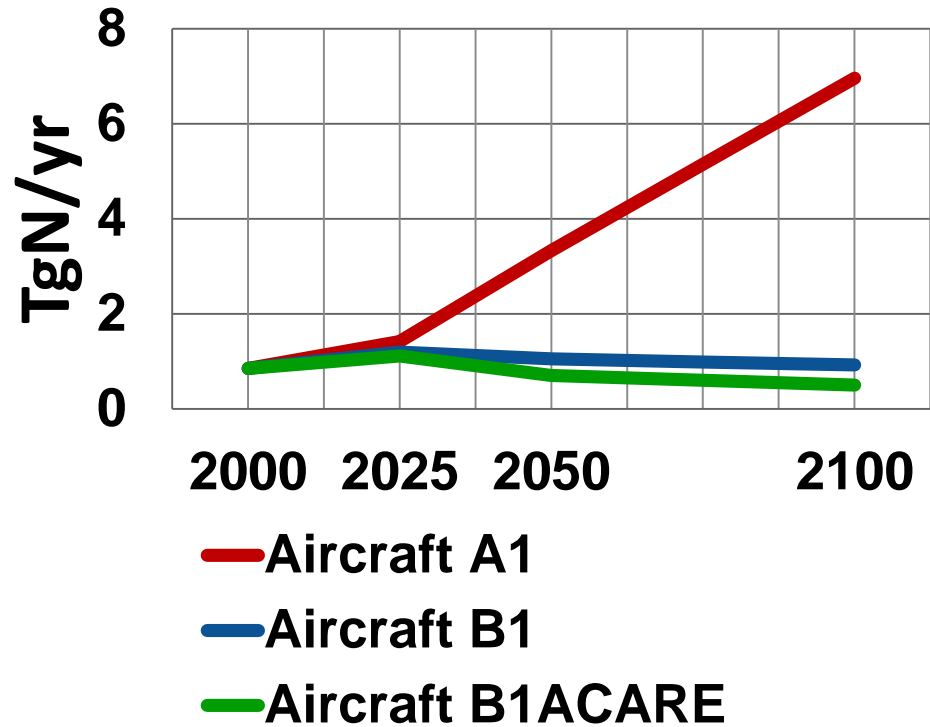
Enhanced mortality

More frequent during a future warmer climate?

Observed and modeled surface ozone for one European stations (in ppb)

Søvde (2008)

Large uncertainties with future scenarios: Aircraft NO_x emission scenarios ifrom the QUANTIFY project



Large differences in scenarios of future emissions

- Potential for significant emission reductions

Final comment on future studies of impact and adaptation of control measures

A key question for estimates of future impact is the adoption of realistic emission scenarios

- Even on a time scale of a few decades scenarios show a wide spread in the scenarios

Estimates of climate contribution require studies of the compounds with highly different lifetimes

- Appropriate metrics need to be adopted

Several key compounds involved in climate change are affected by atmospheric chemistry

- The role of climate-chemistry interaction need to be better understood

Adaptation of control measures to reduce the climate impact require a sectorwise approach

- Sectors emit different compounds with different climate impact, and emissions occur in different regions