

Klimaänderung II

6. Kurzlebige Klimaantriebe

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Oberpfaffenhofen

Vorlesung SS 2022

LMU München



Knowledge for Tomorrow

Klausur

- **Theresienstr. 39, B 040**
- **20.07.2023, 14:15**
- **Dauer: 2 h**



Technical information

- <http://www.pa.op.dlr.de/~RobertSausen/vorlesung/index.html>
 - Most recent update on the lecture
 - Slides of the lecture (with some delay)

 - See also LSF <https://lsf.verwaltung.uni-muenchen.de/>

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- Further information:
 - www.ipcc.ch
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Contents of IPCC AR 6 2021

Working Group I: the Physical Science Basis

| Chapters | |
|--|----------|
| Chapter 1: Framing, context, methods | DOWNLOAD |
| Chapter 2: Changing state of the climate system | DOWNLOAD |
| Chapter 3: Human influence on the climate system | DOWNLOAD |
| Chapter 4: Future global climate: scenario-based projections and near-term information | DOWNLOAD |
| Chapter 5: Global carbon and other biogeochemical cycles and feedbacks | DOWNLOAD |
| Chapter 6: Short-lived climate forcers | DOWNLOAD |
| Chapter 7: The Earth's energy budget, climate feedbacks, and climate sensitivity | DOWNLOAD |
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| Chapter 9: Ocean, cryosphere, and sea level change | DOWNLOAD |
| Chapter 10: Linking global to regional climate change | DOWNLOAD |
| Chapter 11: Weather and climate extreme events in a changing climate | DOWNLOAD |
| Chapter 12: Climate change information for regional impact and for risk assessment | DOWNLOAD |
| Atlas | DOWNLOAD |
| Supplementary Material | ▼ |
| Annexes | ▼ |

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Chapter 6: Short-lived Climate Forcers

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Chapter 6: Short-lived Climate Forcers

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Chapter 6: Short-lived Climate Forcers

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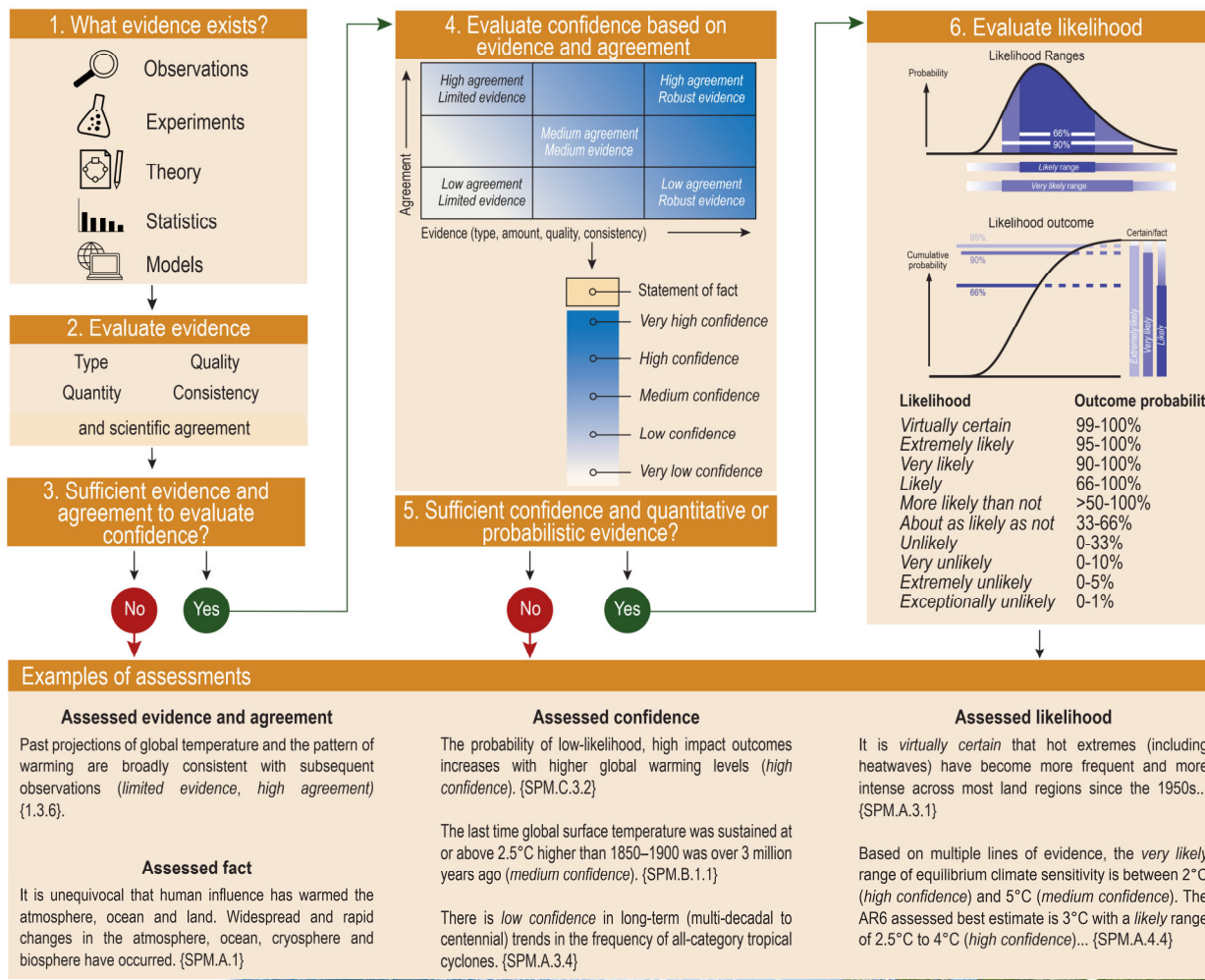
Chapter Scientist:

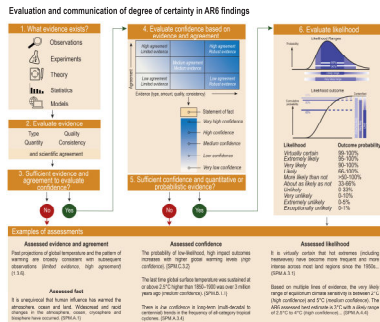
Chaincy Kuo (United States of America)





Evaluation and communication of degree of certainty in AR6 findings

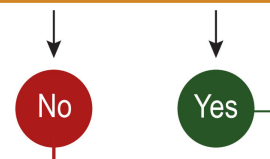




4. Evaluate confidence based on evidence and agreement

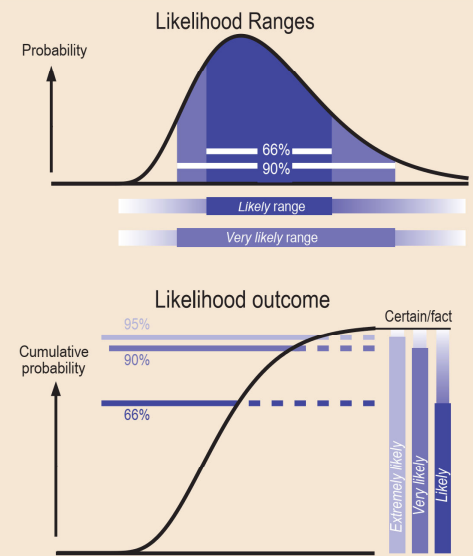
| | | | |
|-------------|---|-------------------------------------|-----------------------------------|
| Agreement ↑ | High agreement Limited evidence | | High agreement Robust evidence |
| | | Medium agreement Medium evidence | |
| | Low agreement Limited evidence | | Low agreement Robust evidence |
| | Evidence (type, amount, quality, consistency) → | | |

5. Sufficient confidence and quantitative or probabilistic evidence?



- Statement of fact
- Very high confidence
- High confidence
- Medium confidence
- Low confidence
- Very low confidence

6. Evaluate likelihood



| Likelihood | Outcome probability |
|------------------------|---------------------|
| Virtually certain | 99-100% |
| Extremely likely | 95-100% |
| Very likely | 90-100% |
| Likely | 66-100% |
| More likely than not | >50-100% |
| About as likely as not | 33-66% |
| Unlikely | 0-33% |
| Very unlikely | 0-10% |
| Extremely unlikely | 0-5% |
| Exceptionally unlikely | 0-1% |

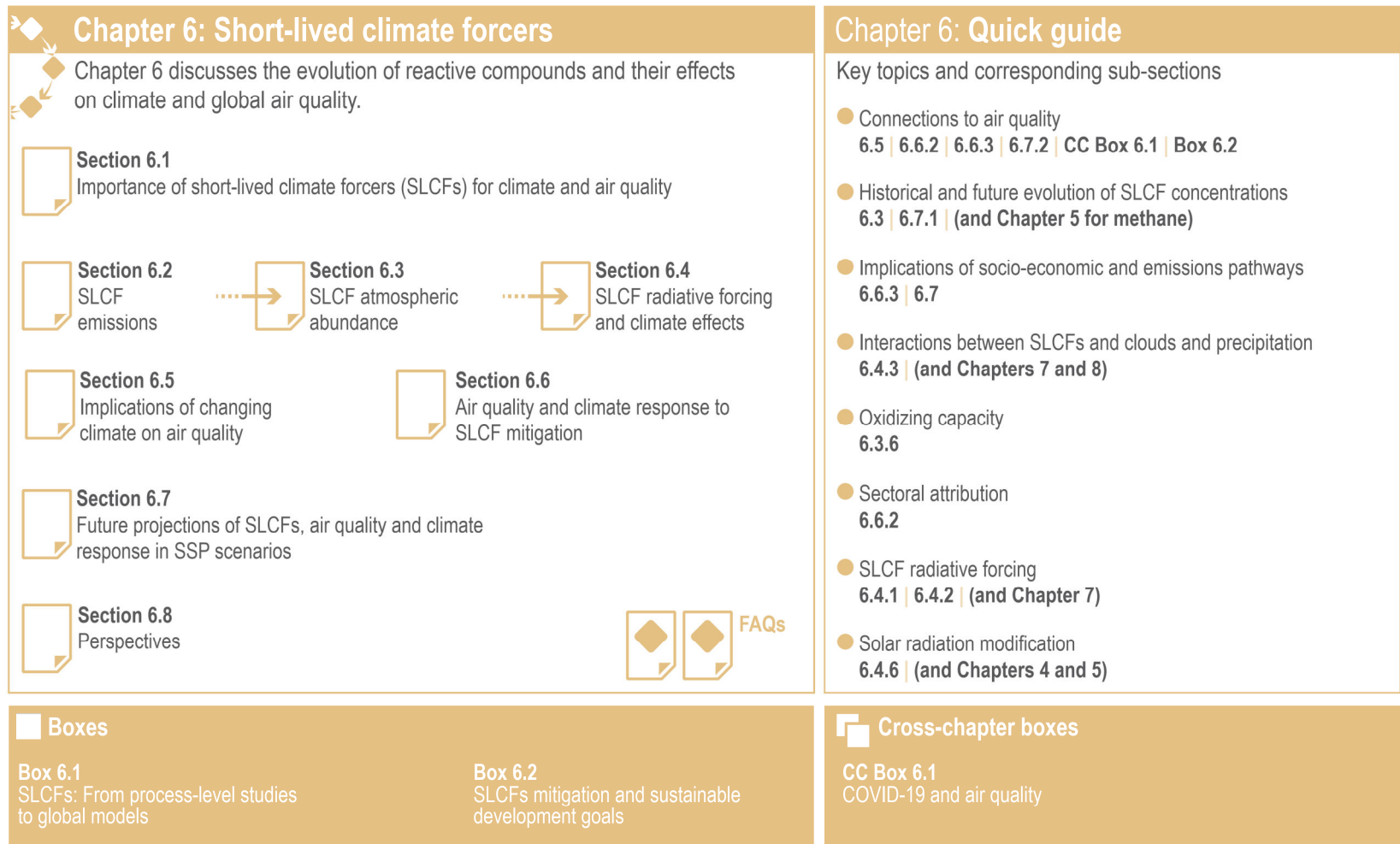


Figure 6.2 | Visual guide to Chapter 6. See Section 6.1.3 for additional description of the chapter.



Statements in the Executive Summary

Short-lived climate forcers (SLCFs) affect climate and are, in most cases, also air pollutants. They include aerosols (sulphate, nitrate, ammonium, carbonaceous aerosols, mineral dust and sea spray), which are also called particulate matter (PM), and chemically reactive gases (methane, ozone, some halogenated compounds, nitrogen oxides, carbon monoxide, non-methane volatile organic compounds, sulphur dioxide and ammonia).



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IPCC 2021, Chap. 6



Sources and processes leading to atmospheric short-lived climate forcer (SLCF) burden and their interactions with the climate system

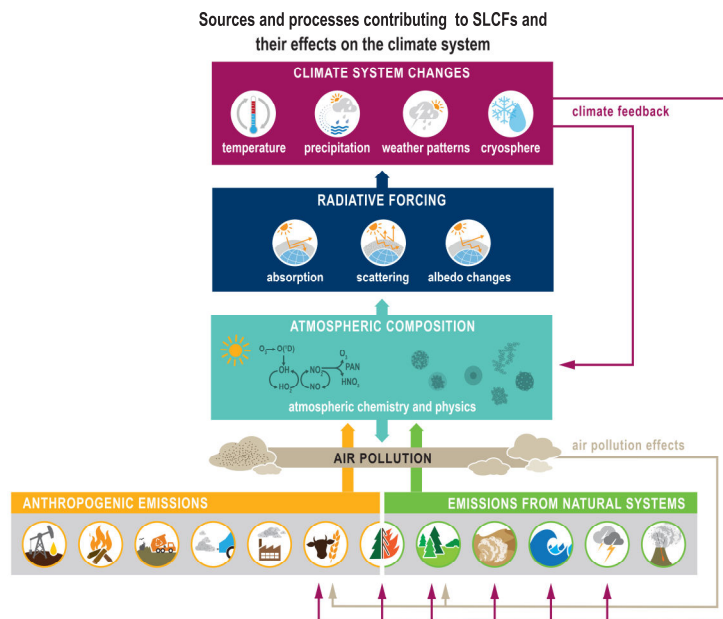
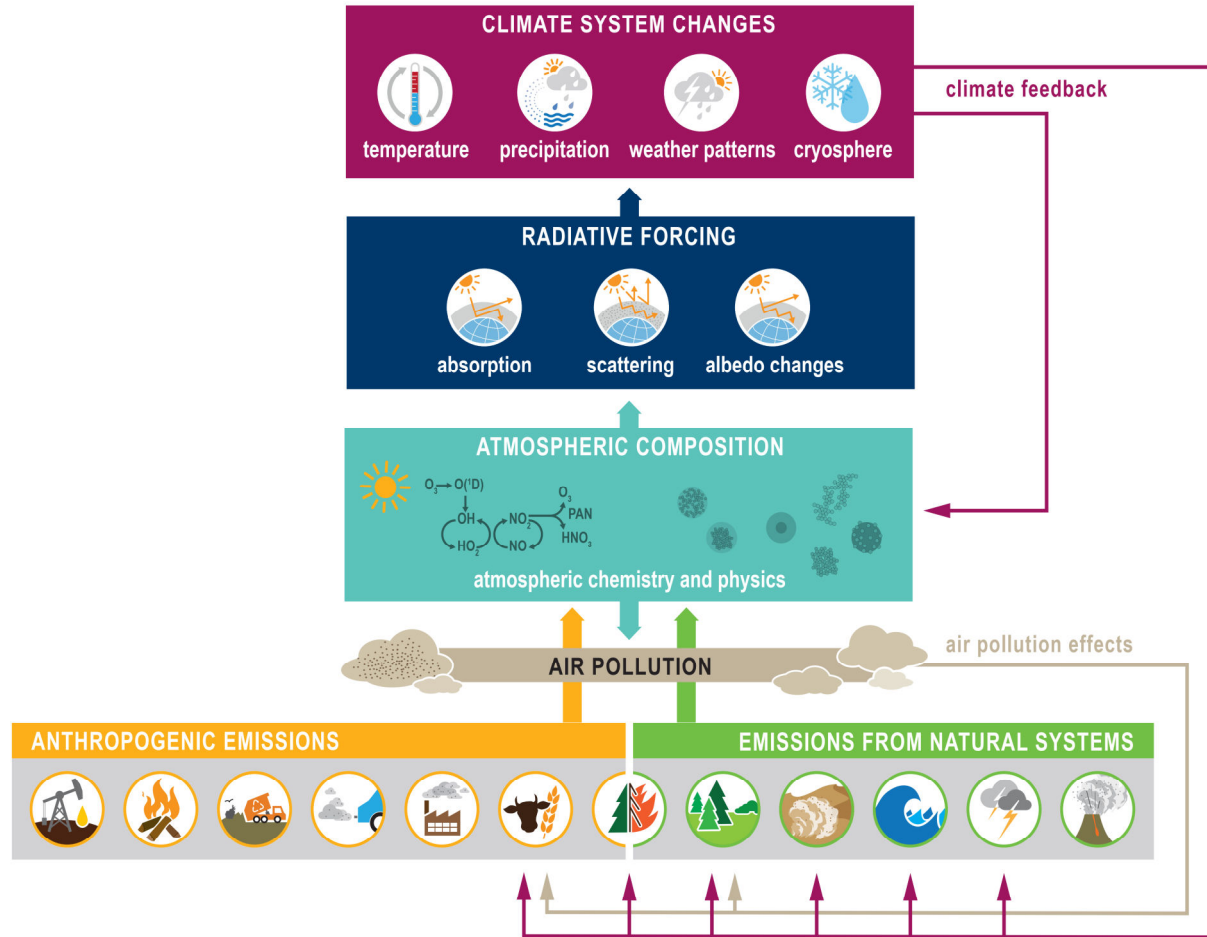


Figure 6.1 | Sources and processes leading to atmospheric short-lived climate forcer (SLCF) burden and their interactions with the climate system. Both direct and indirect SLCFs and the role of atmospheric processes for the lifetime of SLCFs are depicted. Anthropogenic emissions sectors illustrated are: fossil fuel exploration, distribution and use; biofuel production and use; waste; transport; industry; agricultural sources; and open biomass burning. Emissions from natural systems include those from open biomass burning, vegetation, soil, ocean, lightning and volcanoes. SLCFs interact with solar or terrestrial radiation, surface albedo, and cloud or precipitation systems. The radiative forcing due to individual SLCFs can be either positive or negative. Climate change induces changes in emissions from most natural systems as well as from some anthropogenic emissions sectors (e.g., agriculture) leading to a climate feedback (purple arrows). Climate change also influences atmospheric chemistry processes, such as chemical reaction rates or via circulation changes, thus affecting atmospheric composition leading to a climate feedback. Air pollutants influence emissions from terrestrial vegetation, including agriculture (grey arrow).



Sources and processes contributing to SLCFs and their effects on the climate system

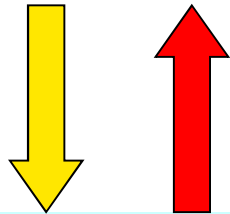


Was ist der "Strahlungsantrieb"? (vereinfacht)

What is "radiative forcing"? (simplified)

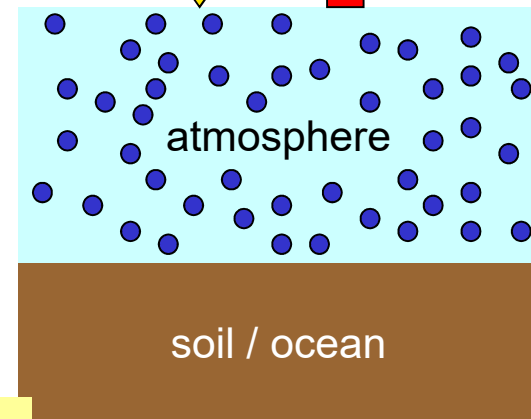
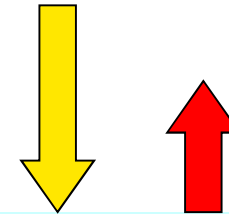
Gleichgewicht

$$RF = 0$$



Gestörte Situation

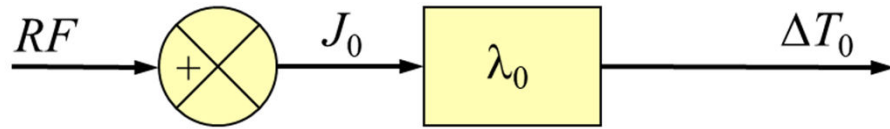
$$RF > 0 \rightarrow \Delta T \uparrow$$



$$\Delta T_{\text{surf}} = \lambda \cdot RF$$



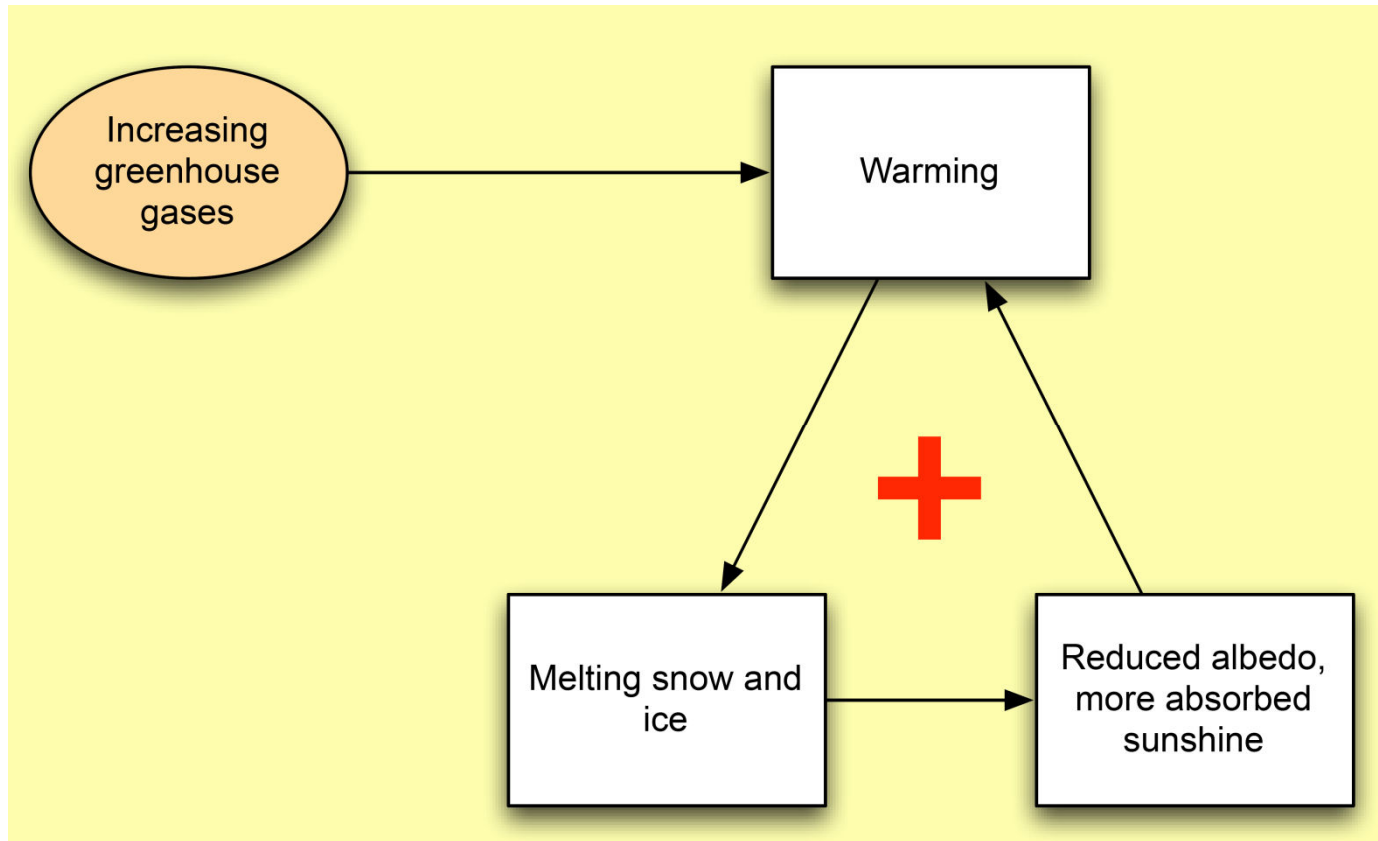
Forcing, Response and Feedback



$$\Delta T_0 = \lambda_0 \cdot J_0 = \lambda_0 \cdot RF$$



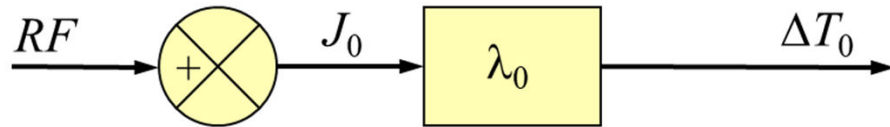
Example: Albedo feedback



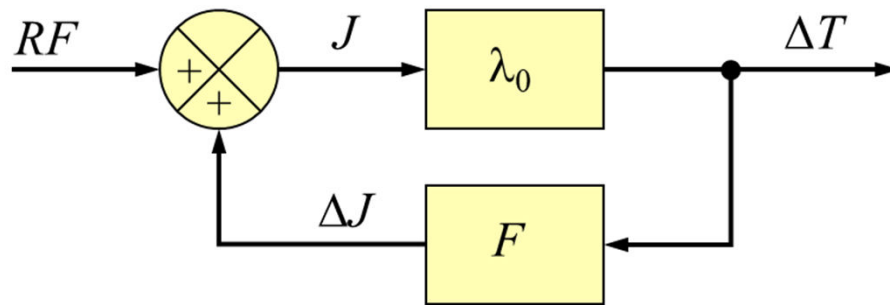
Randel, CSU, 2007



Forcing, Response and Feedback



$$\Delta T_0 = \lambda_0 \cdot J_0 = \lambda_0 \cdot RF$$



$$J = RF + \Delta J = RF + F \cdot \Delta T$$

$$\Delta T = \lambda_0 \cdot J = \lambda_0 \cdot (RF + F \cdot \Delta T) = \lambda \cdot RF$$

climate sensitivity parameter: λ

ratio of responses: $r = \Delta T / \Delta T_0 = \lambda_0 / \lambda = 1 / (1 - f)$

feedback factor: $f = F \cdot \lambda_0$

feedback: F



Schematic comparing RF calculation methodologies

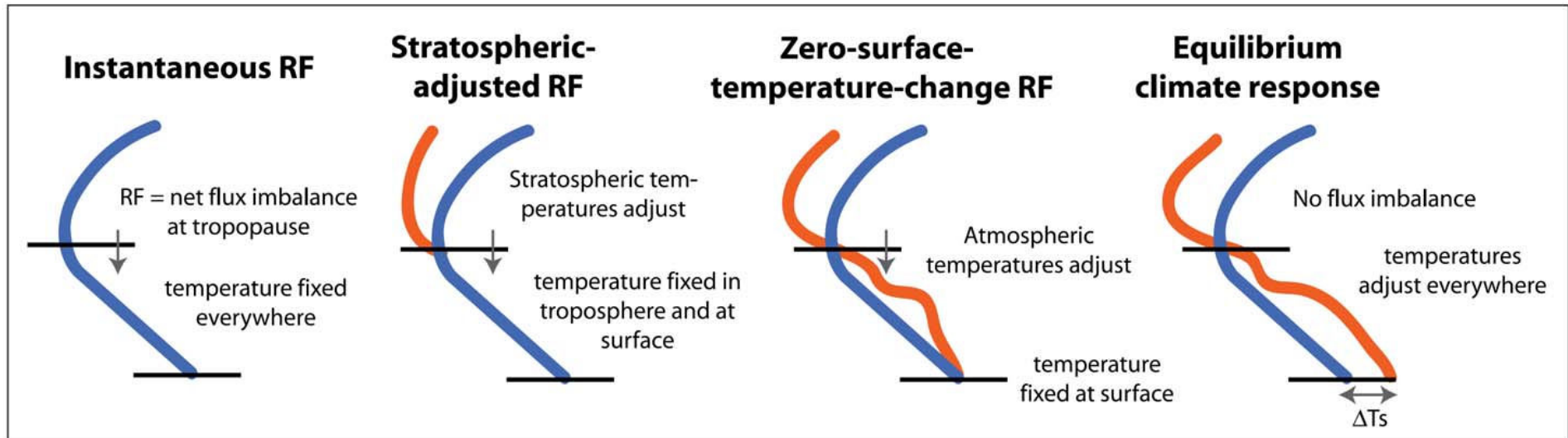


Figure 2.2. Schematic comparing RF calculation methodologies. Radiative forcing, defined as the net flux imbalance at the tropopause, is shown by an arrow. The horizontal lines represent the surface (lower line) and tropopause (upper line). The unperturbed temperature profile is shown as the blue line and the perturbed temperature profile as the red line. From left to right: Instantaneous RF: atmospheric temperatures are fixed everywhere; stratospheric-adjusted RF: allows stratospheric temperatures to adjust; zero-surface-temperature-change RF: allows atmospheric temperatures to adjust everywhere with surface temperatures fixed; and equilibrium climate response: allows the atmospheric and surface temperatures to adjust to reach equilibrium (no tropopause flux imbalance), giving a surface temperature change (ΔT_s).



Radiative forcing (RF) and effective radiative forcing (ERF) estimates

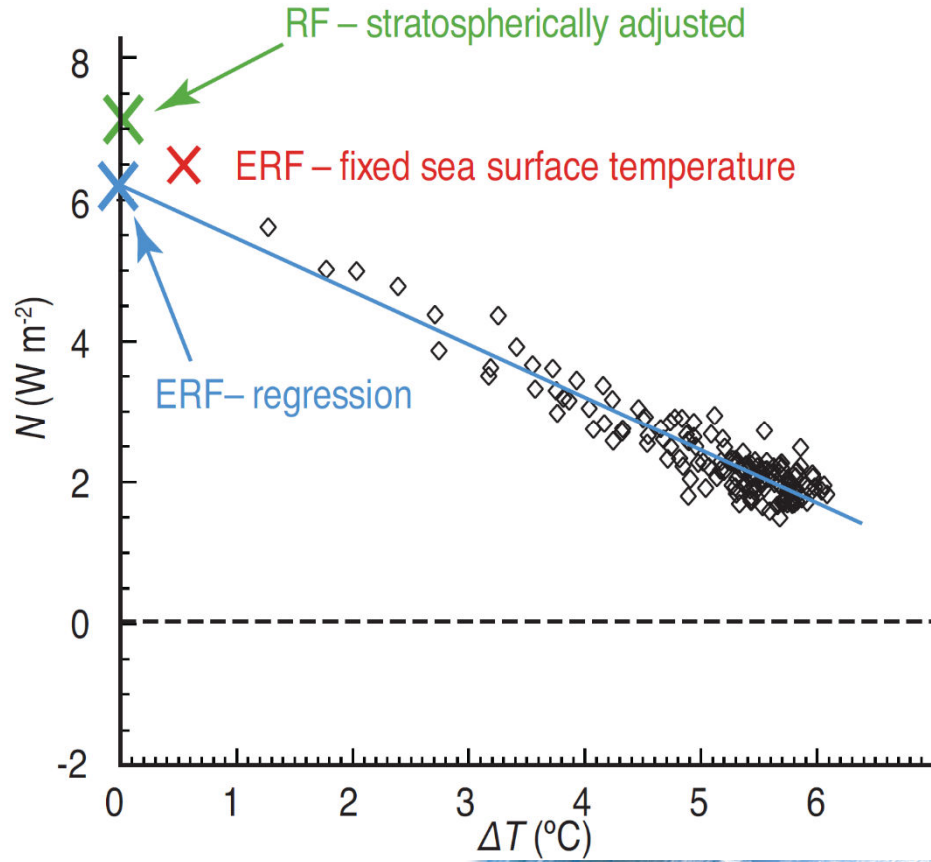


Figure 7.2 | Radiative forcing (RF) and effective radiative forcing (ERF) estimates derived by two methods, for the example of $4 \times \text{CO}_2$ experiments in one climate model. N is the net energy imbalance at the top of the atmosphere and ΔT the global mean surface temperature change. The fixed sea surface temperature ERF estimate is from an atmosphere-land model averaged over 30 years. The regression estimate is from 150 years of a coupled model simulation after an instantaneous quadrupling of CO_2 , with the N from individual years in this regression shown as black diamonds. The stratospherically adjusted RF is the tropopause energy imbalance from otherwise identical radiation calculations at $1 \times$ and $4 \times \text{CO}_2$ concentrations. (Figure follows Andrews et al., 2012.) See also Figure 8.1.

Statements in the Executive Summary

Recent Evolution in Short-lived Climate Forcer (SLCF) (1)

Over the last decade (2010–2019), strong shifts in the geographical distribution of emissions have led to changes in atmospheric abundances of highly variable SLCFs (high confidence). Evidence from satellite and surface observations shows strong regional variations in trends of ozone (O₃), aerosols and their precursors (*high confidence*).



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Recent Evolution in Short-lived Climate Forcer (SLCF) (1)

Over the last decade (2010–2019), strong shifts in the geographical distribution of emissions have led to changes in atmospheric abundances of highly variable SLCFs (high confidence). Evidence from satellite and surface observations shows strong regional variations in trends of ozone (O₃), aerosols and their precursors (*high confidence*). In particular, tropospheric columns of nitrogen dioxide (NO₂) and sulphur dioxide (SO₂) continued to decline over North America and Europe (*high confidence*), and to increase over Southern Asia (*medium confidence*), but have declined over Eastern Asia (*high confidence*). Global carbon monoxide (CO) abundance has continued to decline (*high confidence*). The concentrations of hydrofluorocarbons (HFCs) are increasing (*high confidence*). Global carbonaceous aerosol budgets and trends remain poorly characterized due to limited observations, but sites representative of background conditions have reported multi-year declines in black carbon (BC) over several regions of the Northern Hemisphere. {6.2, 6.3, 2.2.4, 2.2.5, 2.2.6}



Relative regional and sectoral contributions to the present day (year 2014) anthropogenic emissions of short-lived climate forcers (SLCFs)

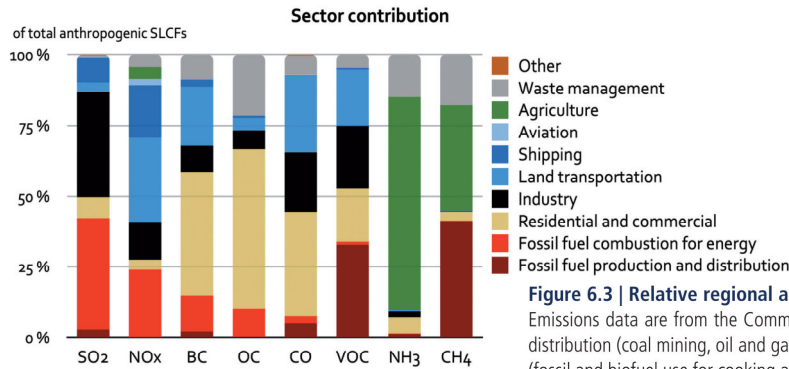
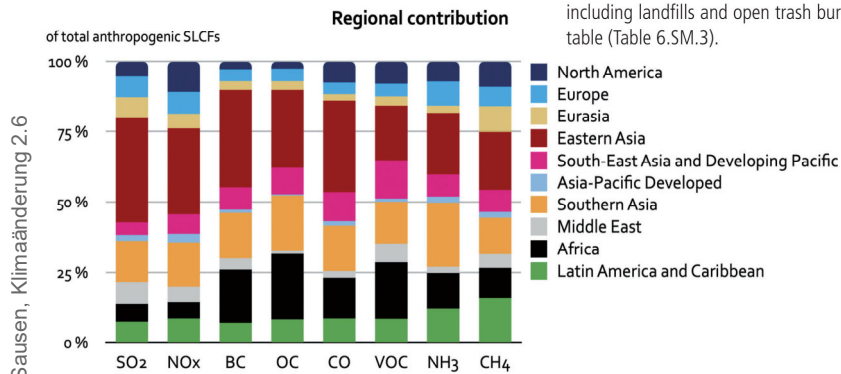
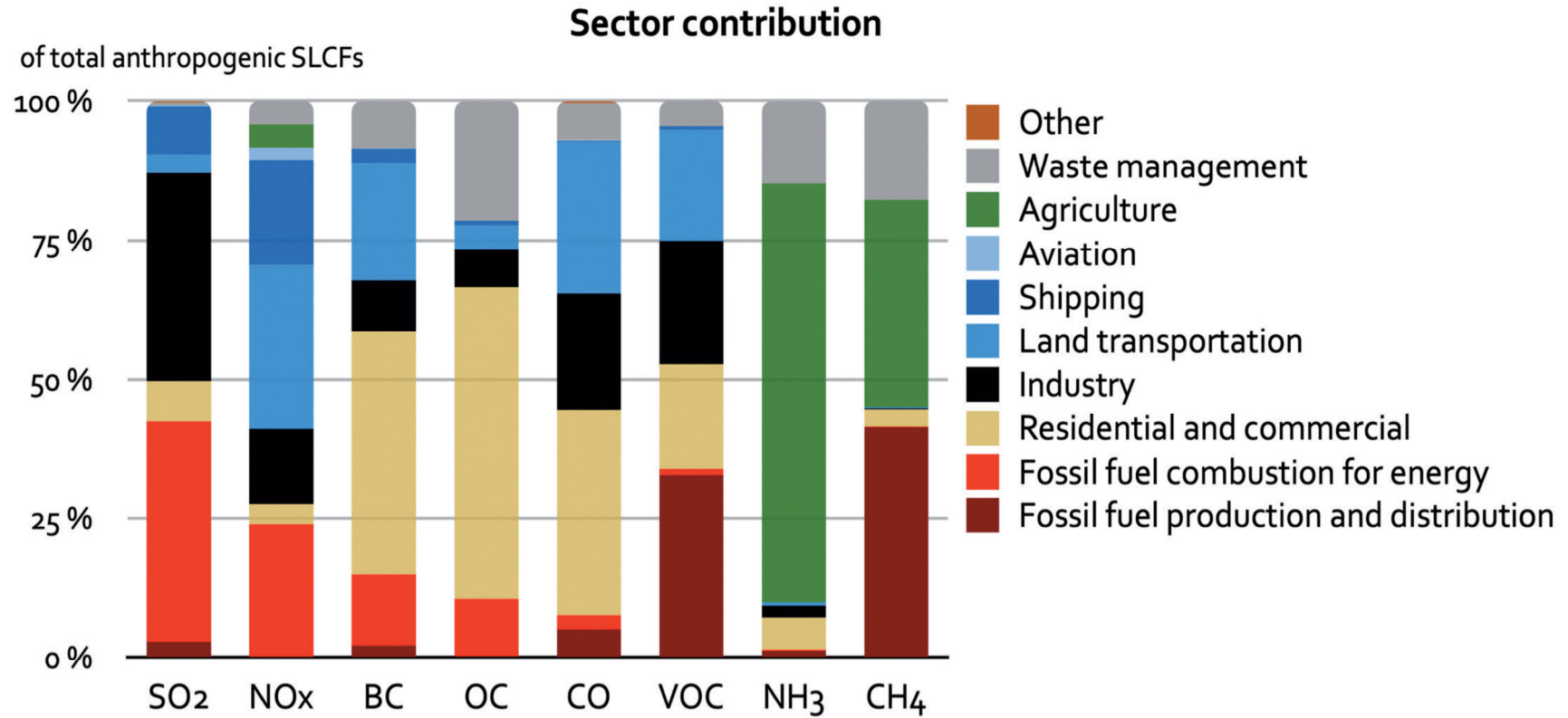


Figure 6.3 | Relative regional and sectoral contributions to the present day (year 2014) anthropogenic emissions of short-lived climate forcers (SLCFs). Emissions data are from the Community Emissions Data System (CEDS; Hoesly et al., 2018). Emissions are aggregated into the following sectors: fossil fuel production and distribution (coal mining, oil and gas production, upstream gas flaring, gas distribution networks), fossil fuel combustion for energy (power plants), residential and commercial (fossil and biofuel use for cooking and heating), industry (combustion and production processes, solvent-use losses from production and end use), transport (road and off-road vehicles), shipping (including international shipping), aviation (including international aviation), agriculture (livestock and crop production), waste management (solid waste, including landfills and open trash burning, residential and industrial waste water), and other. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).



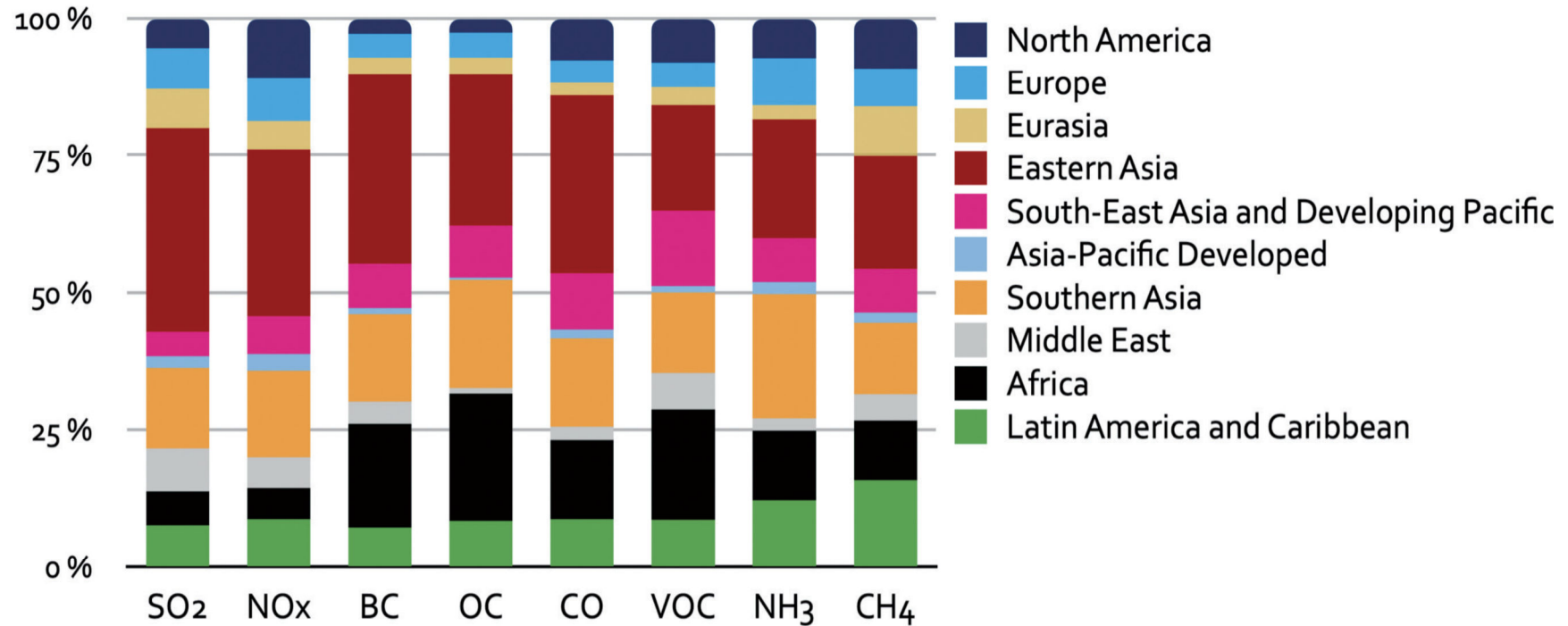
Relative regional and sectoral contributions to the present day (year 2014) anthropogenic emissions of short-lived climate forcers (SLCFs)



Relative regional and sectoral contributions to the present day (year 2014) anthropogenic emissions of short-lived climate forcers (SLCFs)

Regional contribution

of total anthropogenic SLCFs



Time evolution of global annual mean tropospheric ozone burden (in Tg) from 1850 to 2100

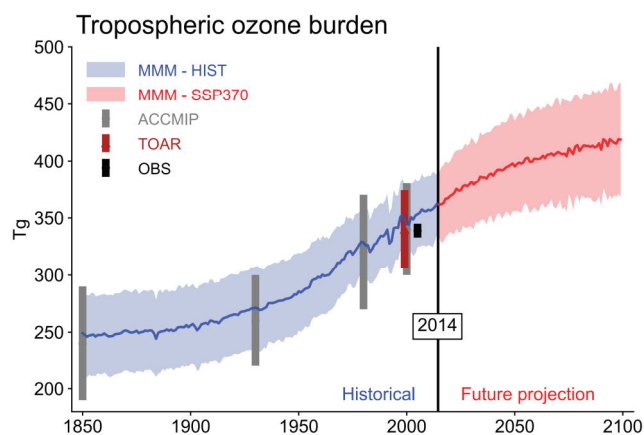
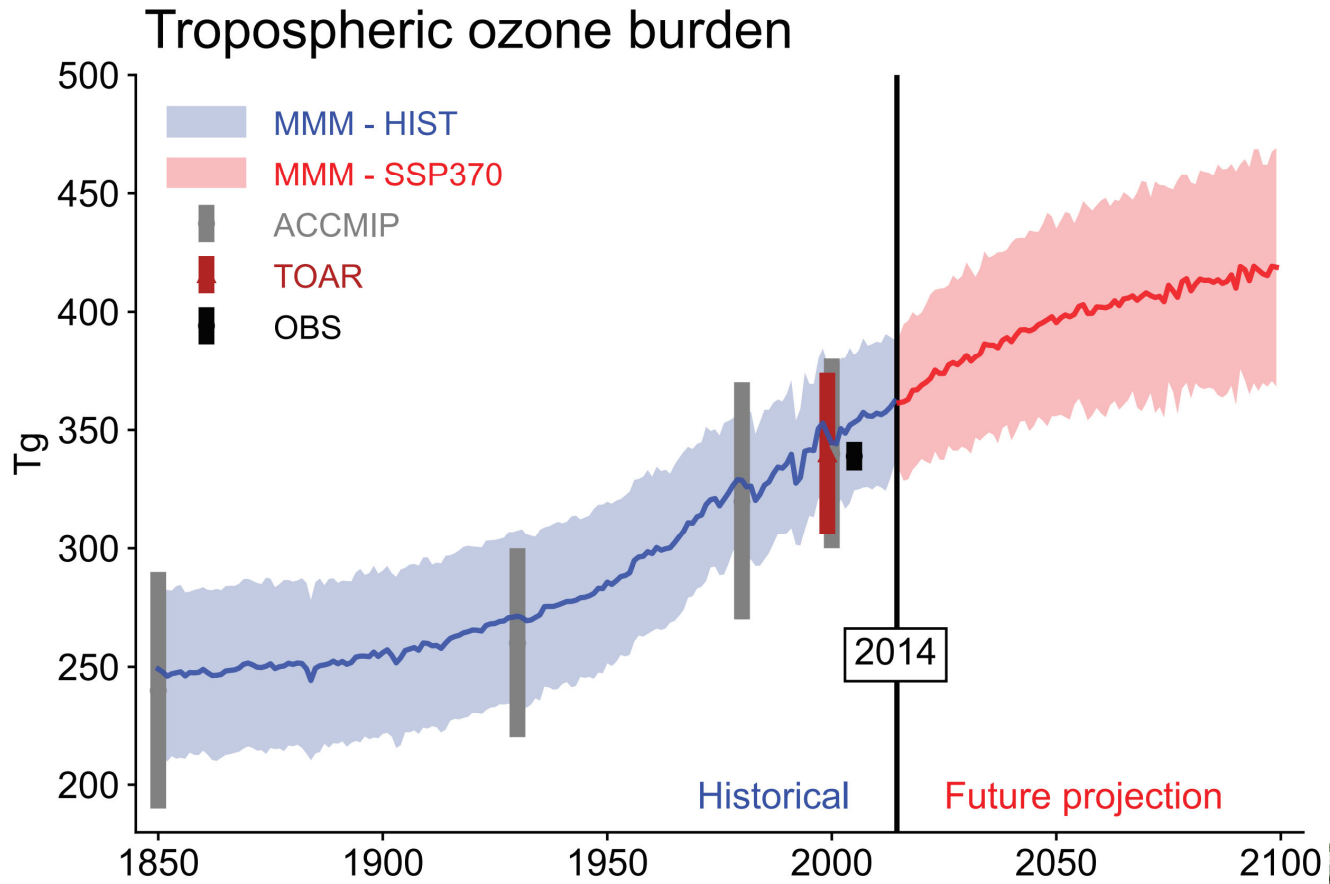


Figure 6.4 | Time evolution of global annual mean tropospheric ozone burden (in Tg) from 1850 to 2100. Multi-model means for CMIP6 historical experiment (1850–2014) from UKESM1-LL-0, CESM2-WACCM, MRI-ESM2-0, GISS-E2.1-G and GFDL-ESM4 and for ScenarioMIP SSP3-7.0 experiment (2015–2100) are represented with their inter-model standard deviation (± 1 standard deviation, shaded areas). Observation-based global tropospheric ozone burden estimate (from Table 6.3) is for 2010–2014. Tropospheric Ozone Assessment Report (TOAR) multi-model mean value (from Table 6.3) is for 2000 with a ± 1 standard deviation error-bar. Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP) multi-model means are for 1850, 1930, 1980 and 2000 time slices with ± 1 standard deviation error-bars. The troposphere is masked by the tropopause pressure calculated in each model using the WMO thermal tropopause definition. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).



Time evolution of global annual mean tropospheric ozone burden (in Tg) from 1850 to 2100



Decadal tropospheric ozone trends since 1994

Decadal ozone trends since 1994

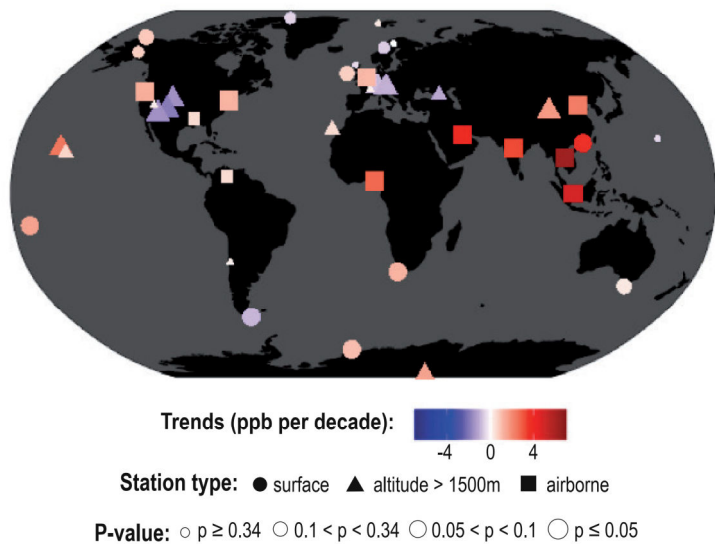
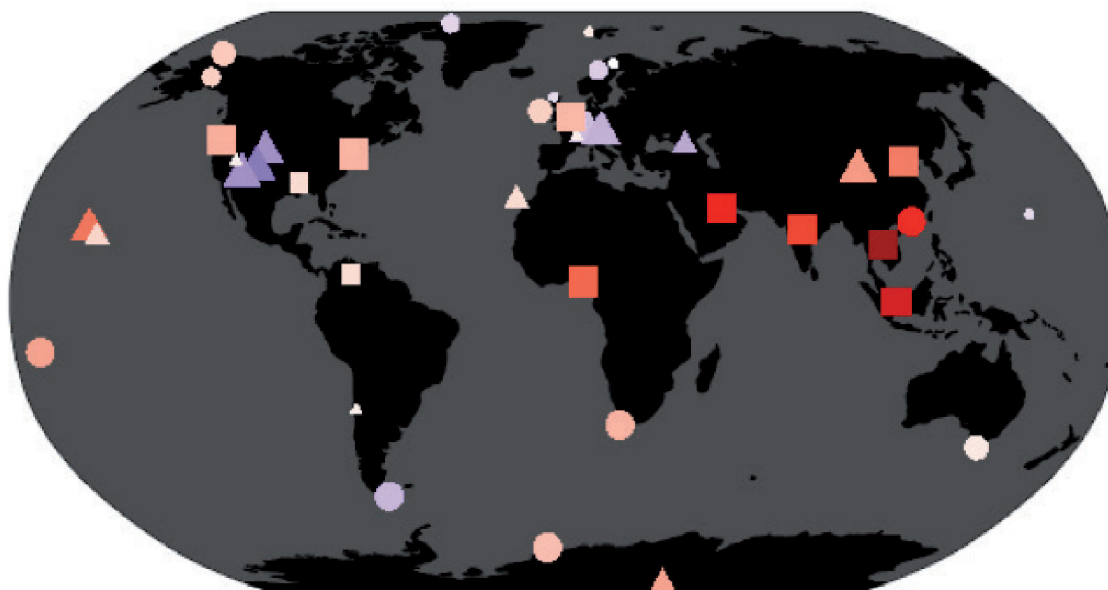


Figure 6.5 | Decadal tropospheric ozone trends since 1994. Trends are shown at 28 remote and regionally representative surface sites (Cooper et al., 2020) and in 11 regions of the lower free troposphere (650 hPa, about 3.5 km) as measured by In-Service Aircraft for a Global Observing System (IAGOS) above Europe, north-eastern USA, south-eastern USA, western North America, north-east China, South East Asia, southern India, the Persian Gulf, Malaysia/Indonesia, the Gulf of Guinea and northern South America (Gaudel et al., 2020). High-elevation surface sites are >1500 m above sea level. All trends end with the most recently available year but begin in 1995 or 1994. The sites and datasets are the same as those used in Figure 2.8, further details on data sources and processing are available in the Chapter 2 data table (Table 2.SM.1).



Decadal tropospheric ozone trends since 1994

Decadal ozone trends since 1994



Trends (ppb per decade):  -4 0 4

Station type: ● surface ▲ altitude > 1500m ■ airborne

P-value: ○ $p \geq 0.34$ ○ $0.1 < p < 0.34$ ○ $0.05 < p < 0.1$ ○ $p \leq 0.05$



Long-term climatological mean (a) and time evolution (b) of tropospheric nitrogen dioxide (NO₂) vertical column density

Long term average and trends in tropospheric NO₂ column over 1996-2017

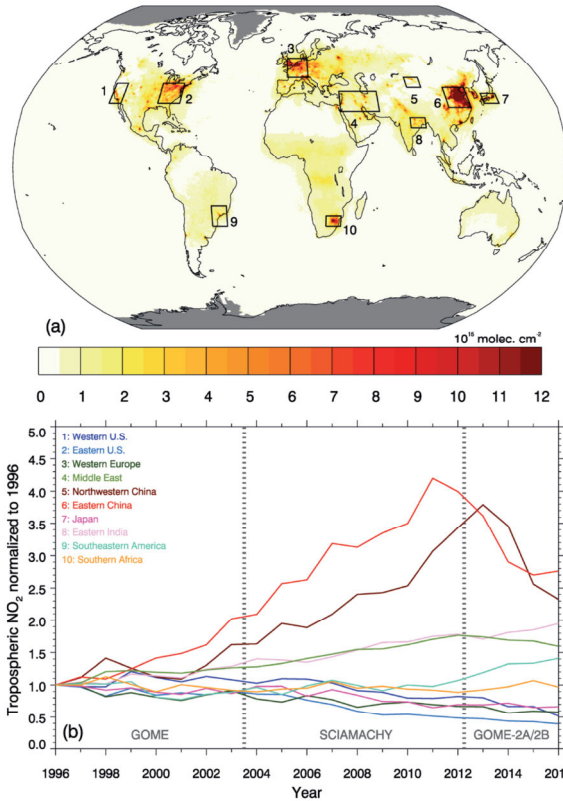
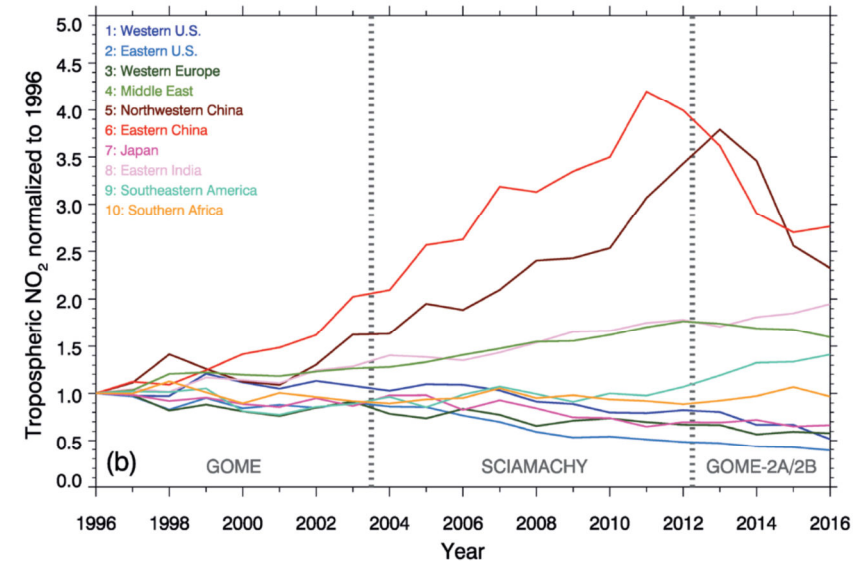
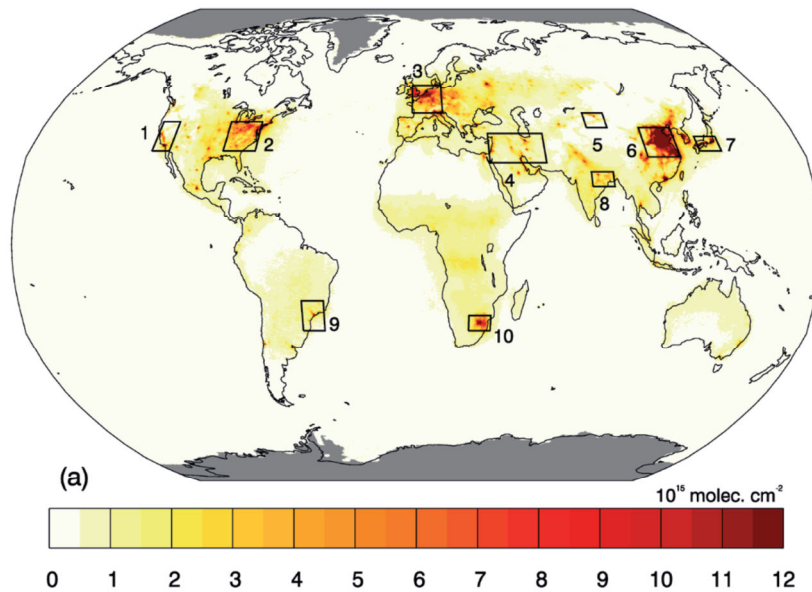


Figure 6.6 | Long-term climatological mean (a) and time evolution (b) of tropospheric nitrogen dioxide (NO₂) vertical column density. Data are from the merged GOME/SCIAMACHY/GOME-2 (TM4NO2A version 2.3) dataset for the period 1996–2016 (Georgoulias et al., 2019). Time evolution of NO₂ column shown in panel (b) is normalized to the fitted 1996 levels for the 10 regions shown as boxes in panel (a). Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).



Long-term climatological mean (a) and time evolution (b) of tropospheric nitrogen dioxide (NO₂) vertical column density

Long term average and trends in tropospheric NO₂ column over 1996-2017



Distribution of PM_{2.5} composition mass concentration (in $\mu\text{g m}^{-3}$) for the major PM_{2.5} aerosol components

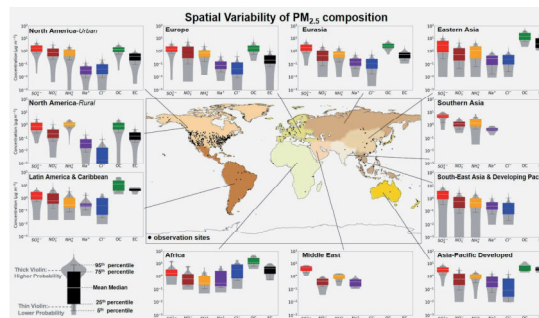
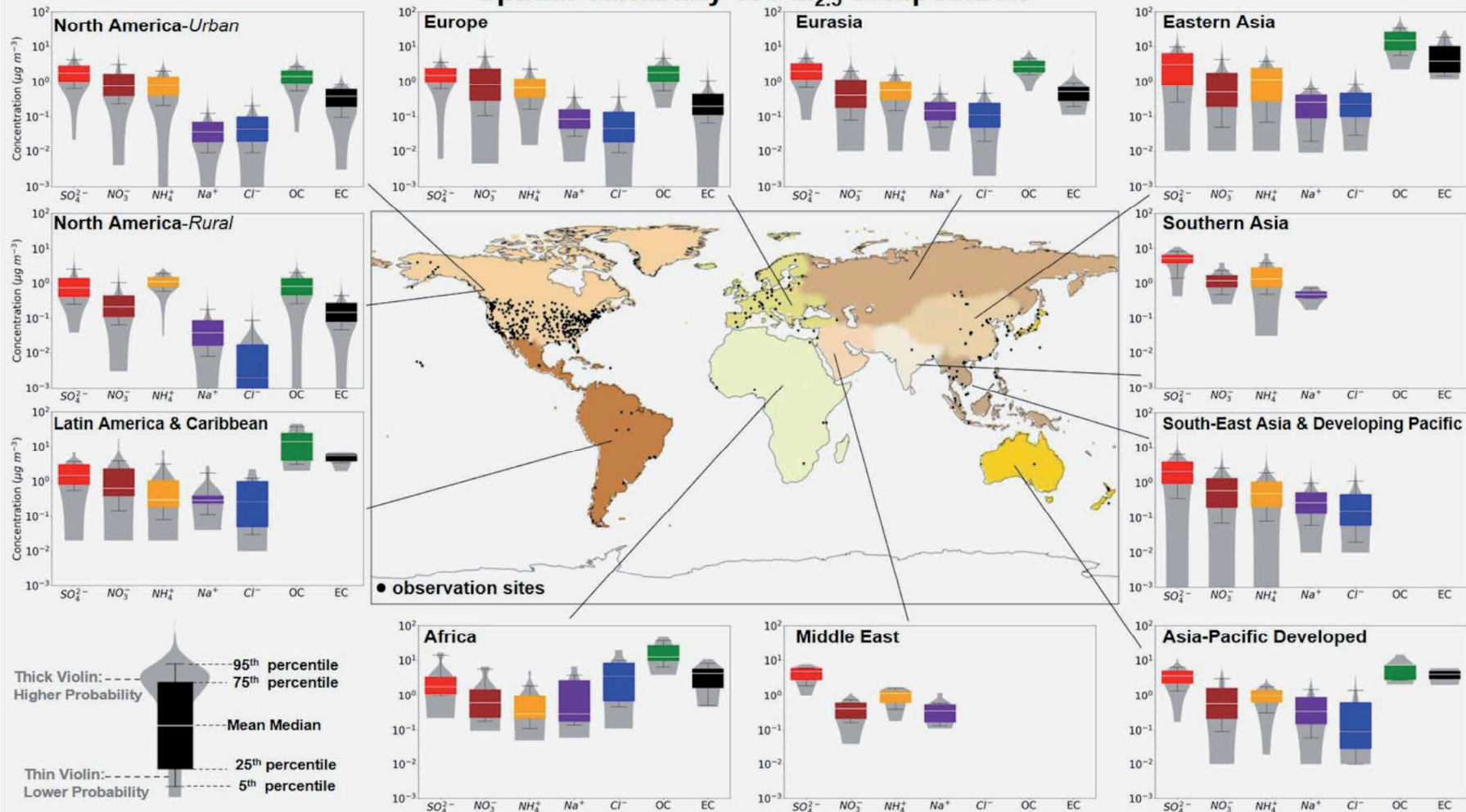


Figure 6.7 | Distribution of PM_{2.5} composition mass concentration (in $\mu\text{g m}^{-3}$) for the major PM_{2.5} aerosol components. Those aerosol components are sulphate, nitrate, ammonium, sodium, chloride, organic carbon and elemental carbon. The central world map depicts the intermediate-level regional breakdown of observations (10 regions) following the IPCC Sixth Assessment Report Working Group III (AR6 WGIII). Monthly averaged PM_{2.5} aerosol component measurements are from: **(i)** the Environmental Protection Agency (EPA) network which include 211 monitor sites primarily in urban areas of North America during 2000–2018 (Solomon et al., 2014), **(ii)** the Interagency Monitoring of Protected Visual Environments (IMPROVE) network during 2000–2018 over 198 monitoring sites representative of the regional haze conditions over North America, **(iii)** the European Monitoring and Evaluation Programme (EMEP) network over 70 monitoring in Europe and (eastern) Eurasia during 2000–2018, **(iv)** the Acid Deposition Monitoring Network in Eastern Asia (EANET) network with 39 (18 remote, 10 rural, 11 urban) sites in Eurasia, Eastern Asia, South East Asia and Developing Pacific, and Asia-Pacific Developed during 2001–2017, **(v)** the global Surface Particulate Matter Network (SPARTAN) during 2013–2019 with sites primarily in highly populated regions around the world (i.e., North America, Latin America and Caribbean, Africa, Middle East, Southern Asia, Eastern Asia, South East Asia and Developing Pacific; Snider et al., 2015, 2016), and **(vii)** individual observational field campaign averages over Latin America and Caribbean, Africa, Europe, Eastern Asia, and Asia-Pacific Developed (Celis et al., 2004; Feng et al., 2006; Bourotte et al., 2007; Fuzzi et al., 2007; Mariani and de Mello, 2007; Molina et al., 2007, 2010; Favez et al., 2008; Mkoma, 2008; Aggarwal and Kawamura, 2009; Mkoma et al., 2009; de Souza et al., 2010; Li et al., 2010; Martin et al., 2010; Radhi et al., 2010; Weinstein et al., 2010; Batmunkh et al., 2011; Gioda et al., 2011; Pathak et al., 2011; F. Zhang et al., 2012; Cho and Park, 2013; Zhao et al., 2013; Wang et al., 2019; Kuzu et al., 2020). Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).



Spatial Variability of PM_{2.5} composition



Statements in the Executive Summary

Recent Evolution in Short-lived Climate Forcer (SLCF) (2)

There is no significant trend in the global mean tropospheric concentration of hydroxyl (OH) radical – the main sink for many SLCFs, including methane (CH₄) – from 1850 up to around 1980 (*low confidence*) but OH has remained stable or exhibited a positive trend since the 1980s (*medium confidence*). Global OH cannot be measured directly and is inferred from Earth system and chemistry–climate models (ESMs, CCMs) constrained by emissions and from observationally constrained inversion methods. There is conflicting information from these methods for the 1980–2014 period. ESMs and CCMs concur on a positive trend since 1980 (about a 9% increase over 1980–2014) and there is *medium confidence* that this trend is mainly driven by increases in global anthropogenic (human-caused) nitrogen oxide (NO_x) emissions and decreases in anthropogenic CO emissions. The observation-constrained methods suggest either positive trends or the absence of trends based on *limited evidence* and *medium agreement*. Future changes in global OH, in response to SLCF emissions and climate change, will depend on the interplay between multiple offsetting drivers of OH. {6.3.6 and Cross-Chapter Box 5.1}



Time evolution of global annual mean tropospheric hydroxyl (OH) over the historical period, expressed as a percentage anomaly relative to the mean over 1998–2007

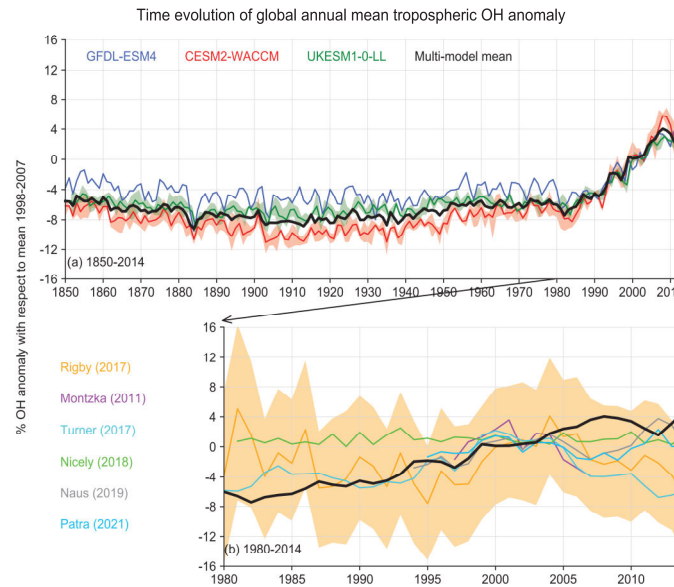
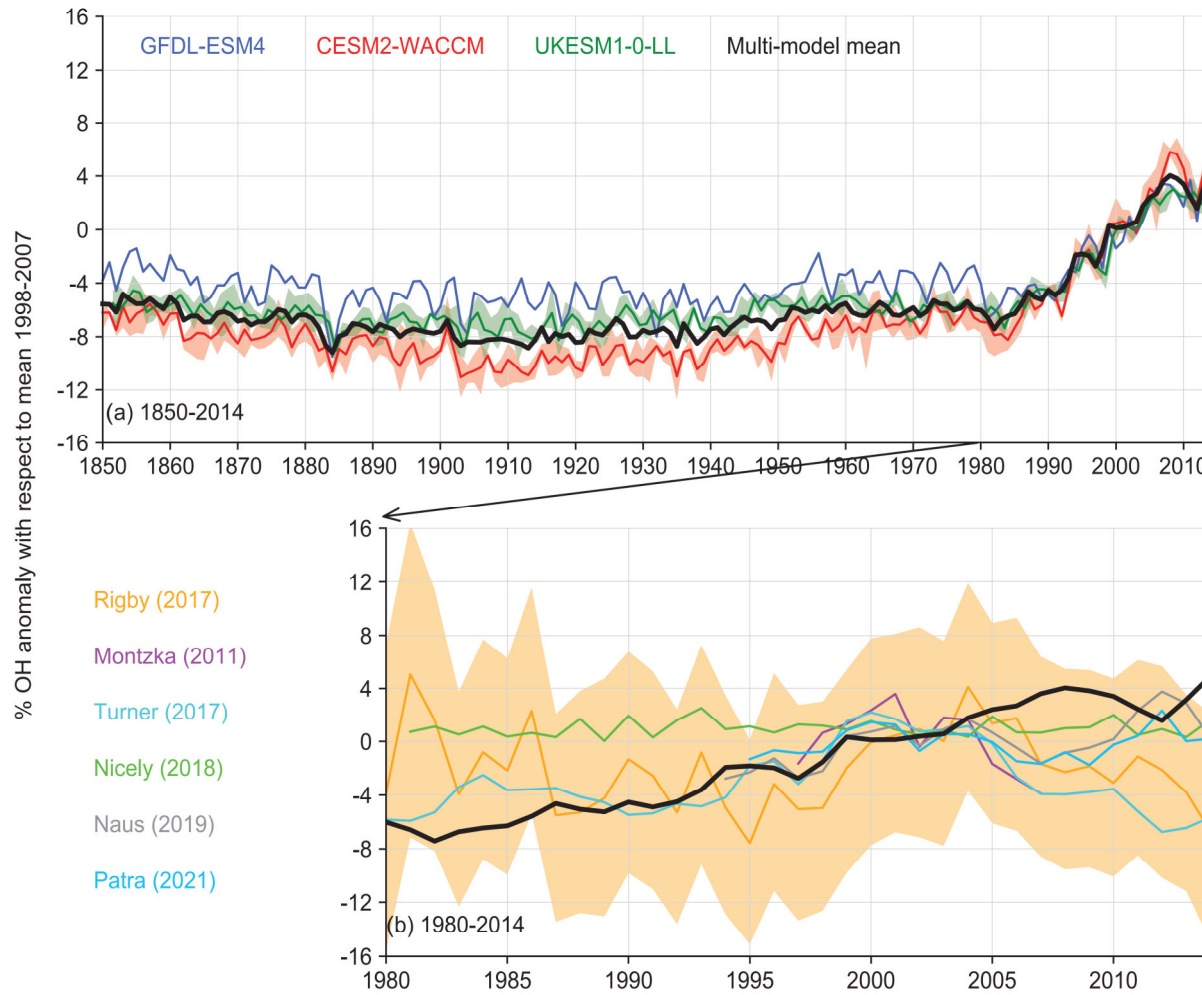


Figure 6.9 | Time evolution of global annual mean tropospheric hydroxyl (OH) over the historical period, expressed as a percentage anomaly relative to the mean over 1998–2007. (a) Results from three CMIP6 models, including UKESM1-0LL (green), GFDL-ESM4 (blue), and CESM2-WACCM (red), are shown; the shaded light green and light red bands show mean over multiple ensemble members for UKESM1-0LL (3) and CESM2-WACCM (3) models, respectively with the multi-model mean anomalies shown in thick black line. (b) Multi-model mean OH anomalies for the 1980–2014 period compared with those derived from observational-based inversions from Montzka et al., (2011); Rigby et al., (2017); Turner et al., (2017); Nicely et al., (2018); Naus et al., (2019); Patra et al., (2021) in the zoomed box. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).





Time evolution of global annual mean tropospheric OH anomaly



Statements in the Executive Summary

Effect of SLCFs on Climate and Biogeochemical Cycles (1)

Over the historical period, changes in aerosols and their effective radiative forcing (ERF) have primarily contributed to a surface cooling, partly masking the greenhouse gas-driven warming (*high confidence*). Radiative forcings induced by aerosol changes lead to both local and remote temperature responses (*high confidence*). The temperature response preserves the south to north gradient of the aerosol ERF – hemispherical asymmetry – but is more uniform with latitude and is strongly amplified towards the Arctic (*medium confidence*). {6.4.1, 6.4.3}

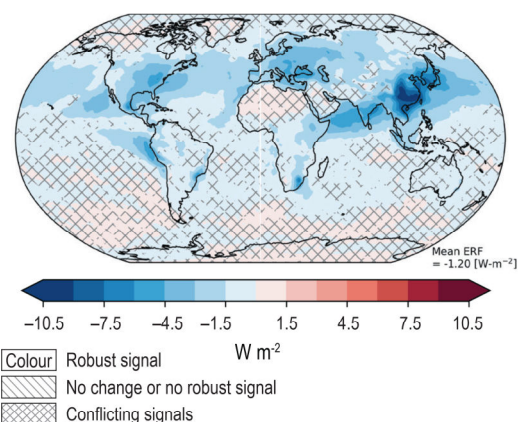
Since the mid-1970s, trends in aerosols and their precursor emissions have led to a shift from an increase to a decrease of the magnitude of the negative globally averaged net aerosol ERF (*high confidence*). However, the timing of this shift varies by continental-scale region and has not occurred for some finer regional scales. The spatial and temporal distribution of the net aerosol ERF from 1850 to 2014 is highly heterogeneous, with stronger magnitudes in the Northern Hemisphere (*high confidence*). {6.4.1}

IPCC 2021, Chap. 6



Multi-model mean effective radiative forcings (ERFs) over the recent-past (1995-2014) induced by aerosol changes since 1850

(a) Net effective radiative forcing due to aerosols



(b) Mean regional effective radiative forcing due to aerosols

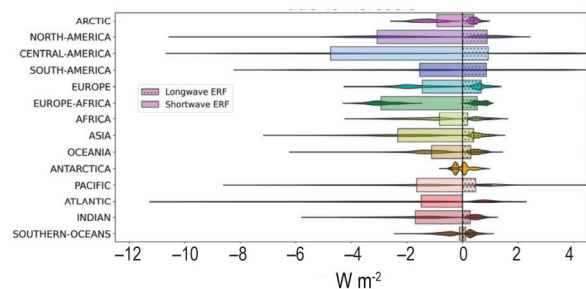
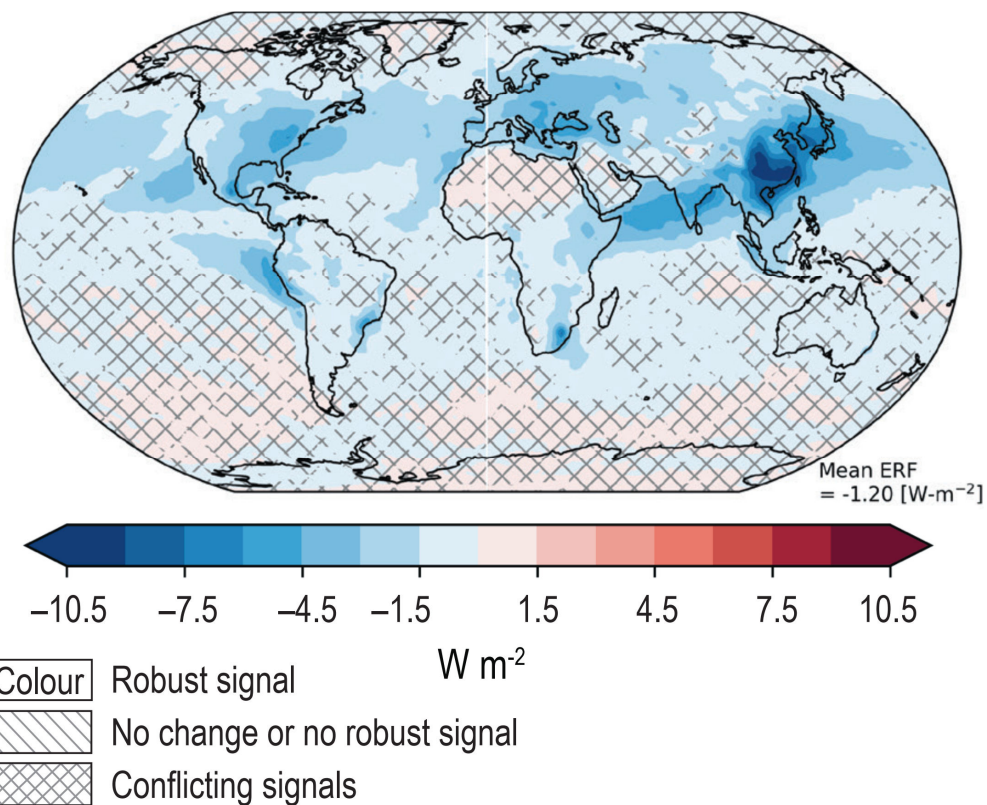


Figure 6.10 | Multi-model mean effective radiative forcings (ERFs) over the recent-past (1995-2014) induced by aerosol changes since 1850. Panel (a) shows the spatial distribution of the net ERF with area-weighted global mean ERF shown at the lower right corner. Uncertainty is represented using the advanced approach: no overlay indicates regions with robust signal, where $\geq 66\%$ of models show change greater than variability threshold and $\geq 80\%$ of all models agree on sign of change; diagonal lines indicate regions with no change or no robust signal, where $< 66\%$ of models show a change greater than the variability threshold; crossed lines indicate regions with conflicting signal, where $\geq 66\%$ of models show change greater than variability threshold and $< 80\%$ of all models agree on sign of change. For more information on the advanced approach, please refer to the Cross-Chapter Box Atlas.1. Panel (b) shows the mean shortwave and longwave ERF for each of the 14 regions defined in the Atlas. Violins in panel (b) show the distribution of values over regions where ERFs are significant. ERFs are derived from the difference between top of the atmosphere (TOA) radiative fluxes for Aerosol Chemistry Model Intercomparison Project (AerChemMIP) experiments *histSST* and *histSST-piAer* (Collins et al., 2017) averaged over 1995–2014 (Box 1.4, Chapter 1). The results come from seven Earth system models: MIROC6, MPI-I-ESM-1-2-HAM, MRI-ESM2-0, GFDL-ESM4, GISS-E2-1-G, NorESM2-LM and UKESM-0-LL. These data can be seen in the Interactive Atlas. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).

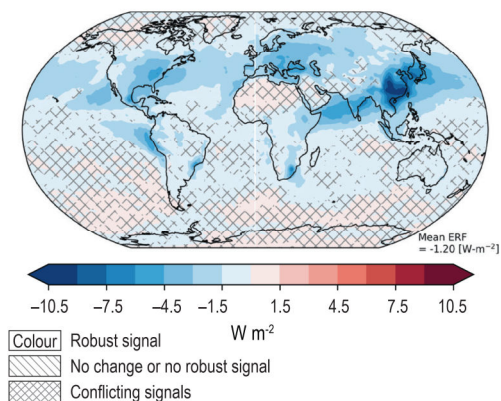
Multi-model mean effective radiative forcings (ERFs) over the recent-past (1995-2014) induced by aerosol changes since 1850

(a) Net effective radiative forcing due to aerosols

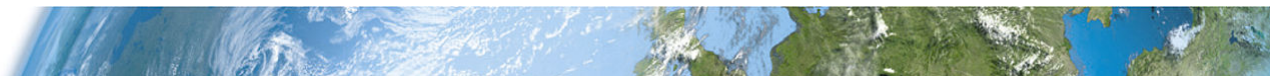
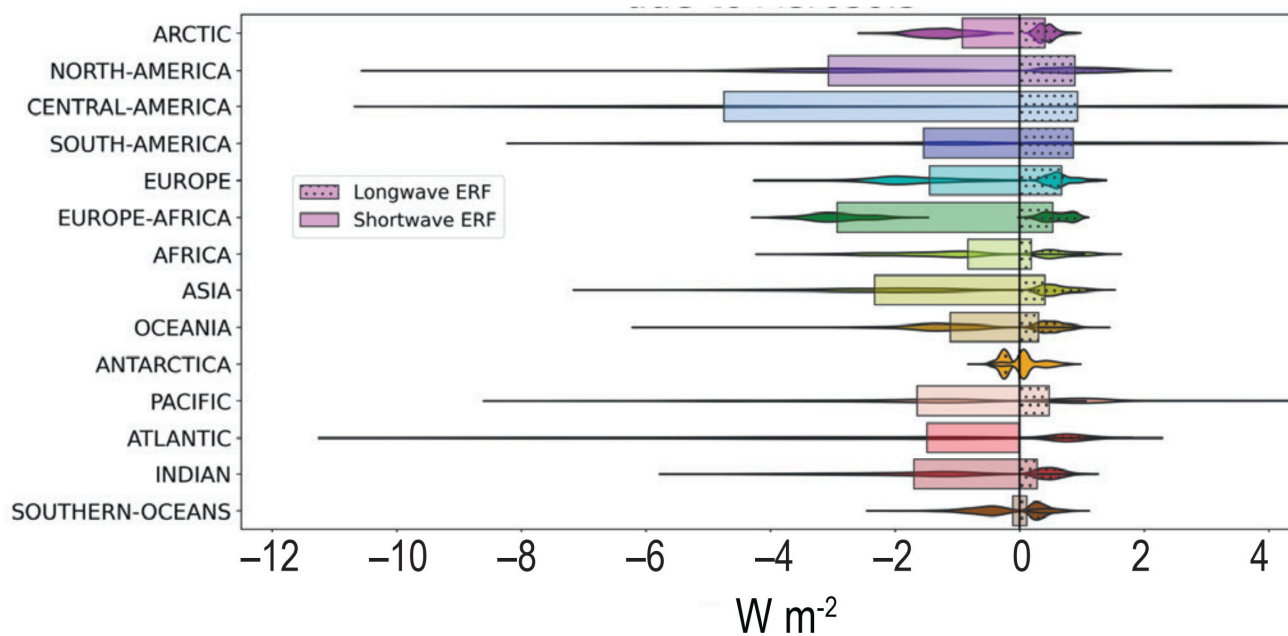


Multi-model mean effective radiative forcings (ERFs) over the recent-past (1995-2014) induced by aerosol changes since 1850

(a) Net effective radiative forcing due to aerosols



(b) Mean regional effective radiative forcing due to aerosols



Time evolution of 20-year multi-model mean averages of the annual area-weighted mean regional net effective radiative forcings (ERFs) due to aerosols for each of the 14 major regions in the Atlas, and global mean, using the models and model experiments as in Figure 6.10

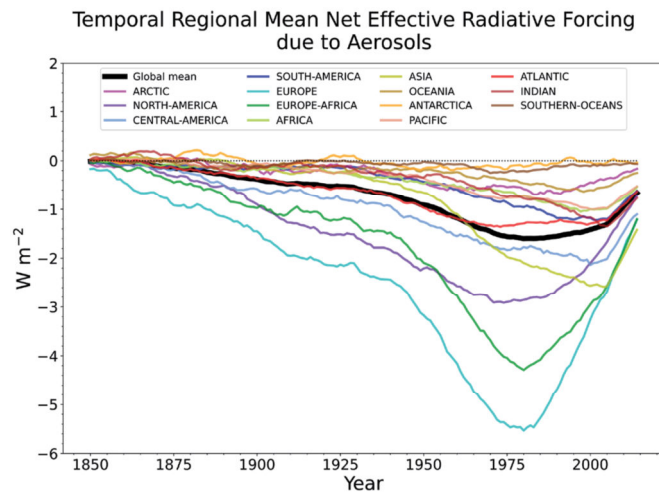
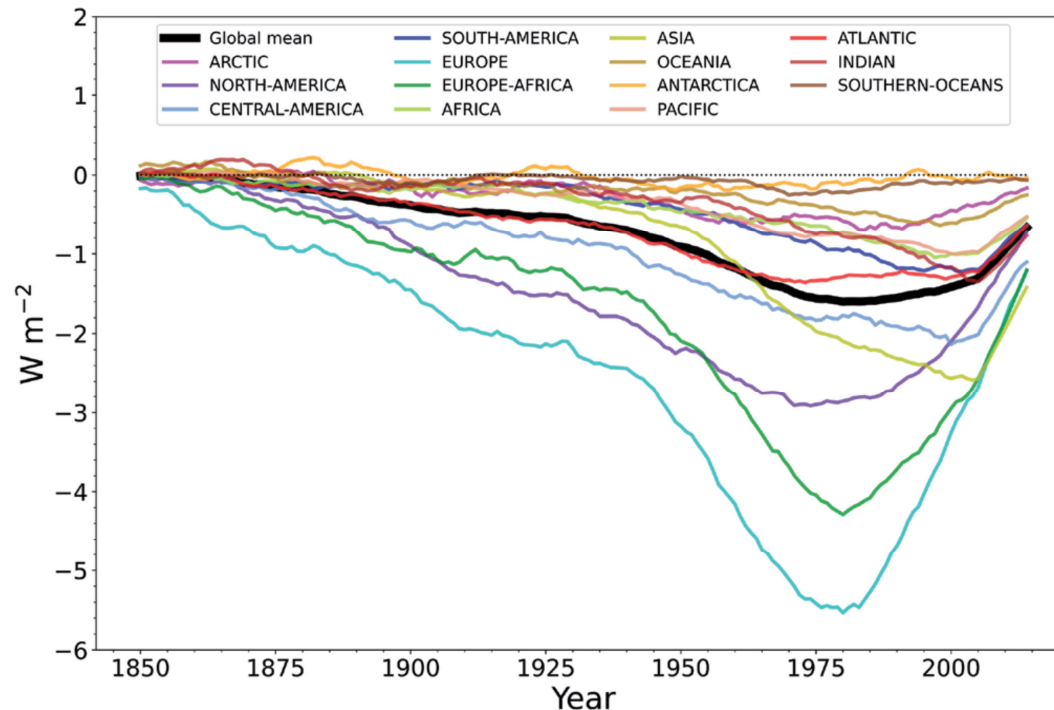


Figure 6.11 | Time evolution of 20-year multi-model mean averages of the annual area-weighted mean regional net effective radiative forcings (ERFs) due to aerosols for each of the 14 major regions in the Atlas, and global mean, using the models and model experiments as in Figure 6.10. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).



Time evolution of 20-year multi-model mean averages of the annual area-weighted mean regional net effective radiative forcings (ERFs) due to aerosols for each of the 14 major regions in the Atlas, and global mean, using the models and model experiments as in Figure 6.10

Temporal Regional Mean Net Effective Radiative Forcing due to Aerosols



Contribution to effective radiative forcing (ERF) (a) and global mean surface air temperature (GSAT) change (b) from component emissions between 1750 to 2019 based on CMIP6 models (Thornhill et al., 2021b)

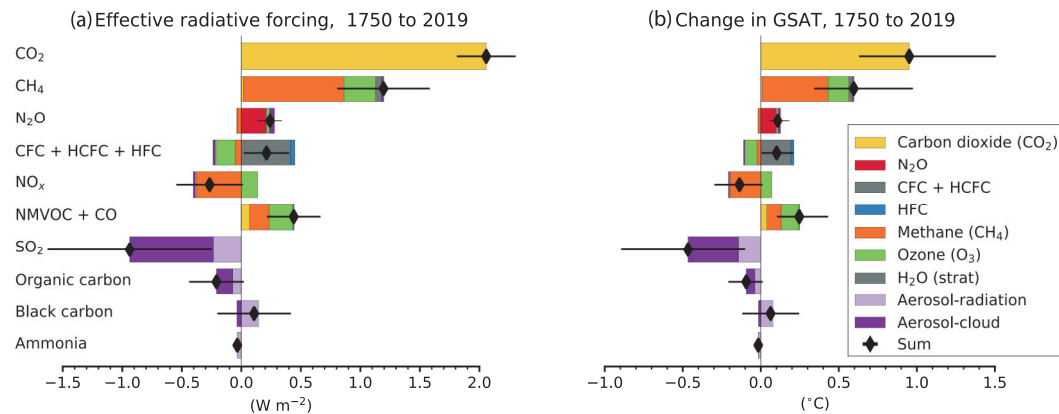
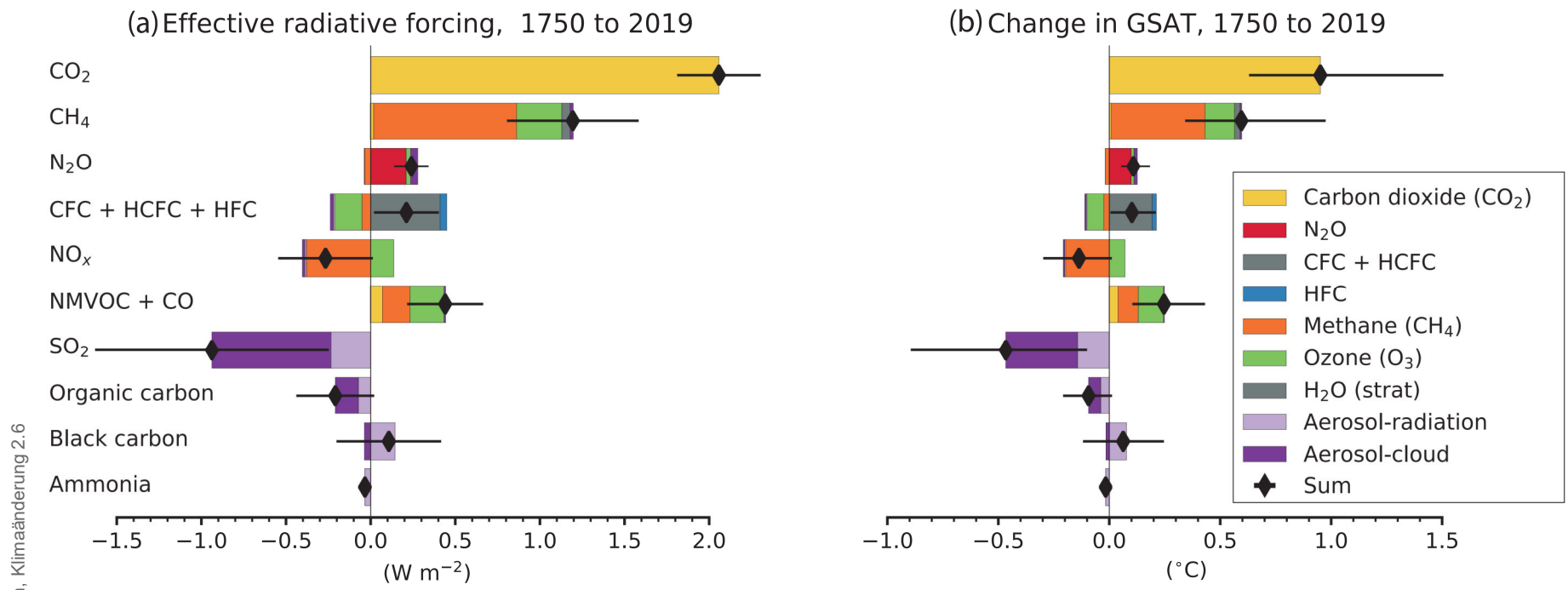


Figure 6.12 | Contribution to effective radiative forcing (ERF) (a) and global mean surface air temperature (GSAT) change (b) from component emissions between 1750 to 2019 based on CMIP6 models (Thornhill et al., 2021b). ERFs for the direct effect of well-mixed greenhouse gases (WMGHGs) are from the analytical formulae in section 7.3.2, H₂O (strat) is from Table 7.8. ERFs for other components are multi-model means from Thornhill et al. (2021b) and are based on ESM simulations in which emissions of one species at a time are increased from 1850 to 2014 levels. The derived emissions-based ERFs are rescaled to match the concentration-based ERFs in Figure 7.6. Error bars are 5–95% and for the ERF account for uncertainty in radiative efficiencies and multi-model error in the means. ERFs due to aerosol–radiation (ERF_{ari}) and cloud effects are calculated from separate radiation calls for clear-sky and aerosol-free conditions (Ghan, 2013; Thornhill et al., 2021b). ‘Cloud’ includes cloud adjustments (semi-direct effect) and ERF from indirect aerosol-cloud to $-0.22 W m^{-2}$ for ERF_{ari} and $-0.84 W m^{-2}$ interactions (ERF_{aci}). The aerosol components (SO₂, organic carbon and black carbon) are scaled to sum to $-0.22 W m^{-2}$ for ERF_{ari} and $-0.84 W m^{-2}$ for ‘cloud’ (Section 7.3.3). For GSAT estimates, time series (1750–2019) for the ERFs have been estimated by scaling with concentrations for WMGHGs and with historical emissions for SLCFs. The time variation of ERF_{aci} for aerosols is from Chapter 7. The global mean temperature response is calculated from the ERF time series using an impulse response function (Cross-Chapter Box 7.1) with a climate feedback parameter of $-1.31 W m^{-2} ^{\circ}C^{-1}$. Contributions to ERF and GSAT change from contrails and light-absorbing particles on snow and ice are not represented, but their estimates can be seen on Figure 7.6 and 7.7, respectively. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).

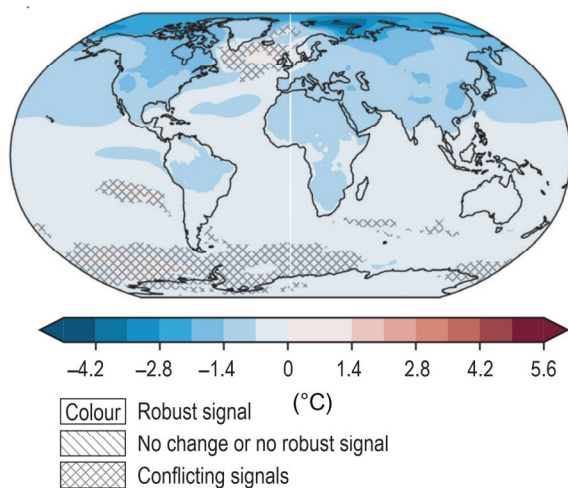


Contribution to effective radiative forcing (ERF) (a) and global mean surface air temperature (GSAT) change (b) from component emissions between 1750 to 2019 based on CMIP6 models (Thornhill et al., 2021b)



Multi-model mean surface air temperature response over the recent past (1995–2014) induced by aerosol changes since 1850

(a) Surface air temperature response due to aerosols



(b) Zonal mean change in surface air temperature due to aerosols

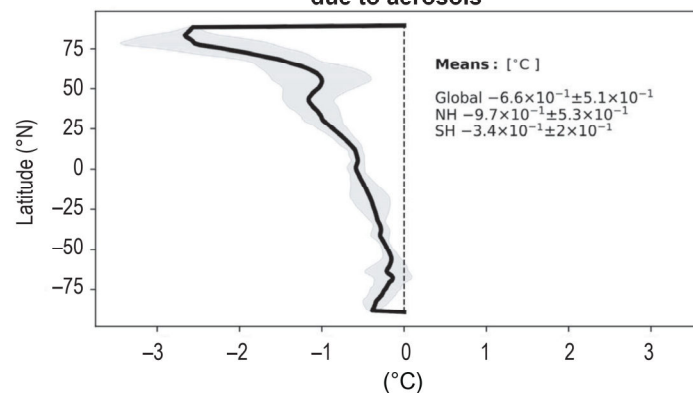
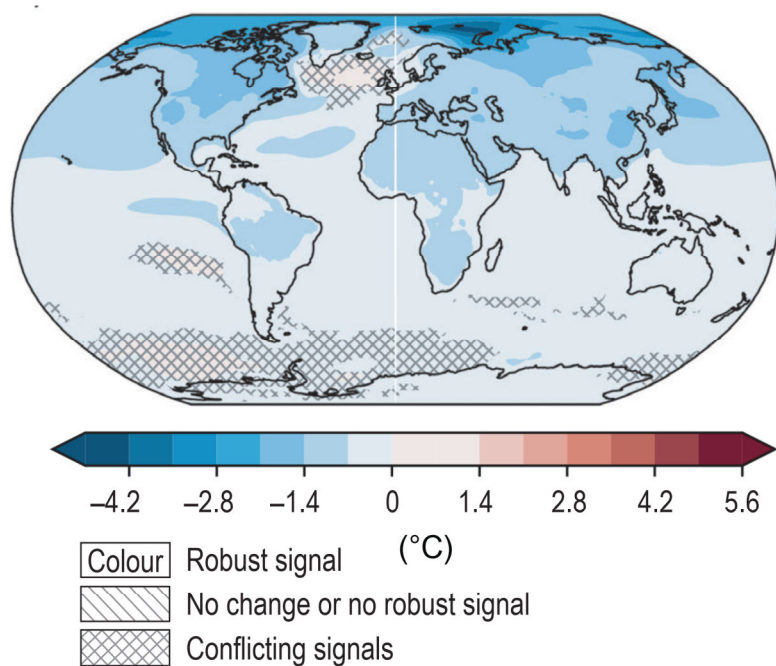


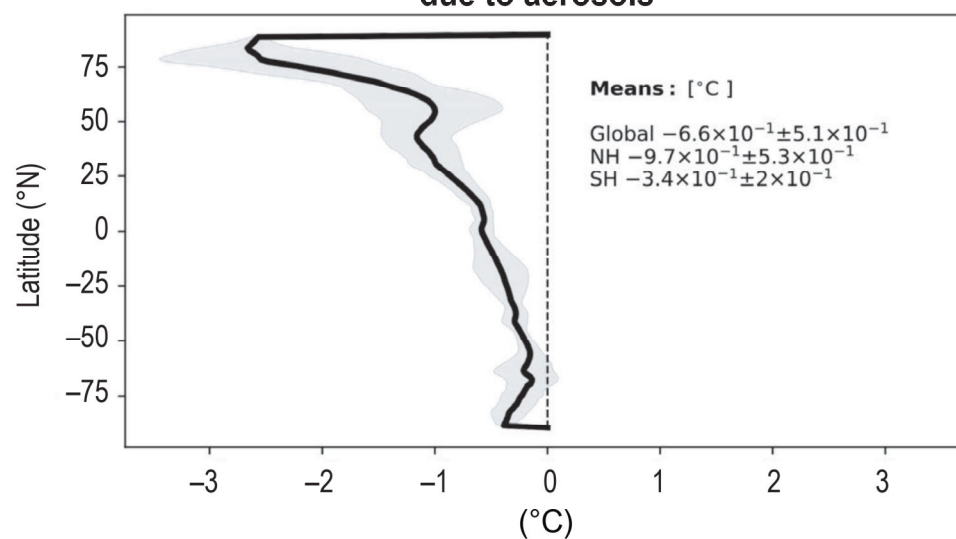
Figure 6.13 | Multi-model mean surface air temperature response over the recent past (1995–2014) induced by aerosol changes since 1850. Calculation is based on the difference between CMIP6 ‘historical’ and AerChemMIP ‘hist-piAer’ experiments averaged over 1995–2014, where (a) is the spatial pattern of the annual mean surface air temperature response, and (b) is the mean zonally averaged response. Model means are derived from the years 1995–2014. Uncertainty is represented using the advanced approach: No overlay indicates regions with robust signal, where $\geq 66\%$ of models show change greater than variability threshold and $\geq 80\%$ of all models agree on sign of change; diagonal lines indicate regions with no change or no robust signal, where $< 66\%$ of models show a change greater than the variability threshold; crossed lines indicate regions with conflicting signal, where $\geq 66\%$ of models show change greater than variability threshold and $< 80\%$ of all models agree on sign of change. For more information on the advanced approach, please refer to the Cross-Chapter Box Atlas.1. AerChemMIP models MIROC6, MRI-ESM2-0, NorESM2-LM, GFDL-ESM4, GISS-E2-1-G and UKESM1-0-LL are used in the analysis. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).

Multi-model mean surface air temperature response over the recent past (1995–2014) induced by aerosol changes since 1850

(a) Surface air temperature response due to aerosols



(b) Zonal mean change in surface air temperature due to aerosols



Statements in the Executive Summary

Effect of SLCFs on Climate and Biogeochemical Cycles (2)

For forcings with short lifetimes (e.g., months) and not considering chemical adjustments, the response in surface temperature occurs strongly as soon as a sustained change in emissions is implemented, and that response continues to grow for a few years, primarily due to thermal inertia in the climate system (*high confidence*). Near its maximum, the response slows down but will then take centuries to reach equilibrium (*high confidence*). For SLCFs with longer lifetimes (e.g., a decade), a delay equivalent to their lifetimes is appended to the delay due to thermal inertia (*high confidence*). {6.6.1}



Global mean surface air temperature (GSAT) response to an abrupt reduction in emissions (at time t=0) of idealized climate forcing agents with different lifetimes.

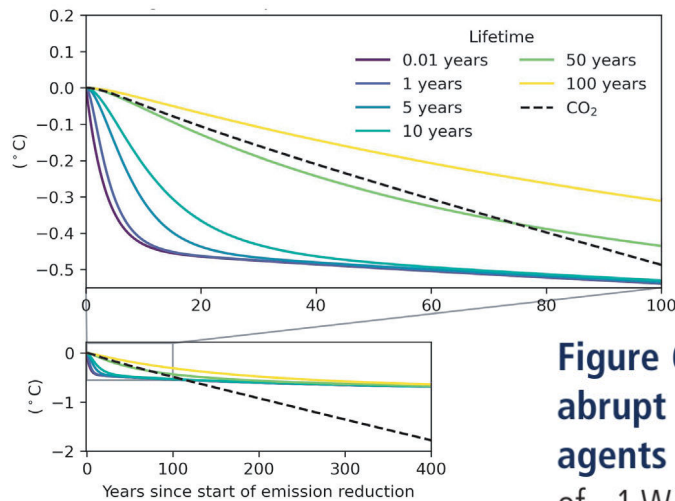
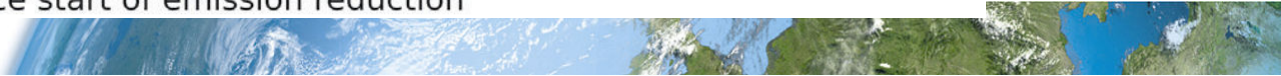
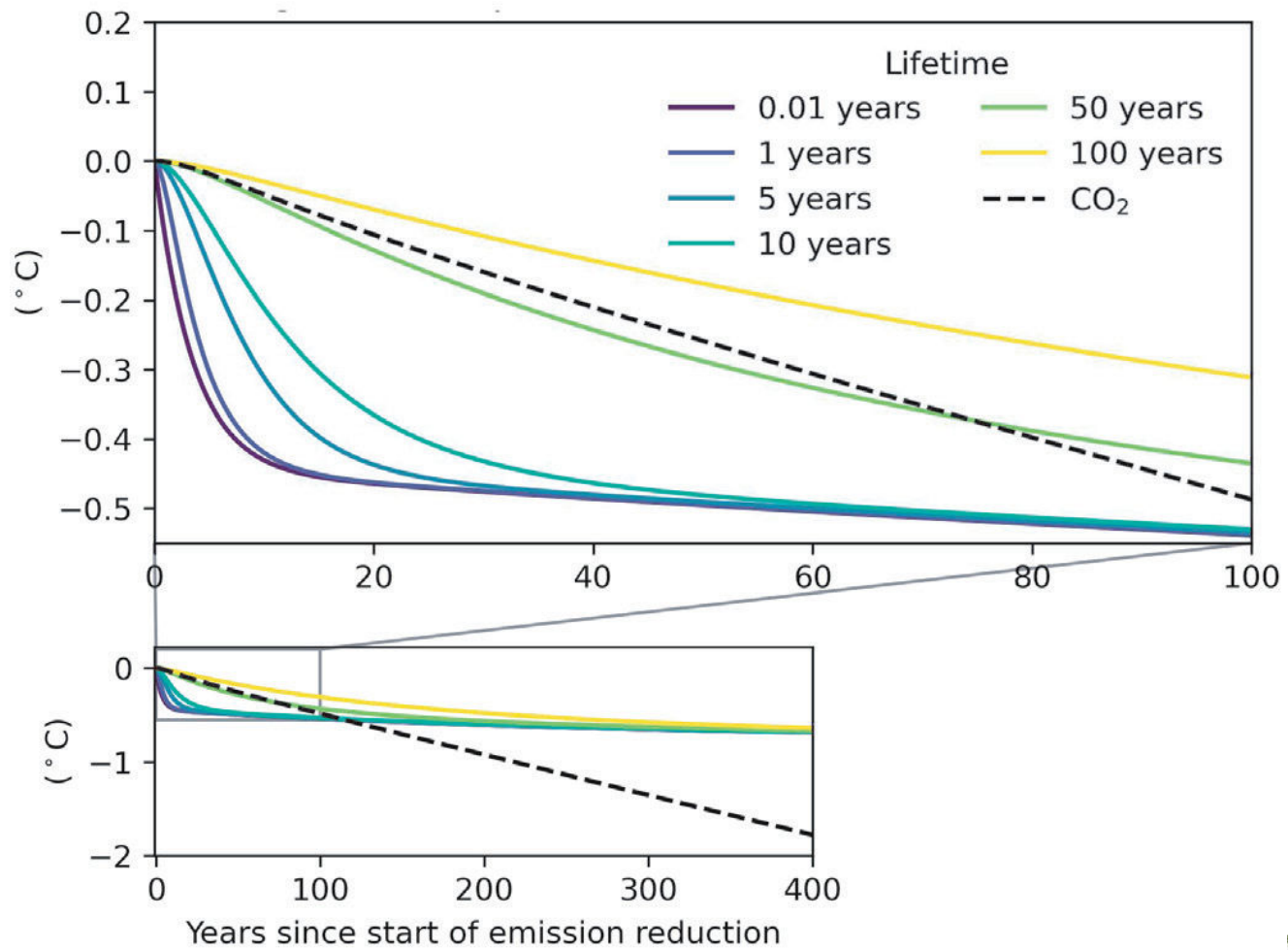


Figure 6.15 | Global mean surface air temperature (GSAT) response to an abrupt reduction in emissions (at time t=0) of idealized climate forcing agents with different lifetimes. All emissions are cut to give a radiative forcing of -1 W m^{-2} at a steady state (except for CO_2). In other words, if the yearly emissions are E_0 before the reduction, they will have a fixed lower value $E_{\text{year}>0} = (E_0 - \Delta E)$ for all succeeding years. For comparison, the GSAT response to a sustained reduction in CO_2 emissions resulting in an RF of -1 W m^{-2} in year 100 is included (dashed line). The temperature response is calculated using an impulse response function (Cross-Chapter Box 7.1) with a climate feedback parameter of $-1.31 \text{ W m}^{-2} \text{ }^\circ\text{C}^{-1}$. Further details on data sources and processing are available in the chapter data table (Table 6. SM.3).





Global mean temperature response 10 and 100 years following one year of present-day (year 2014) emissions

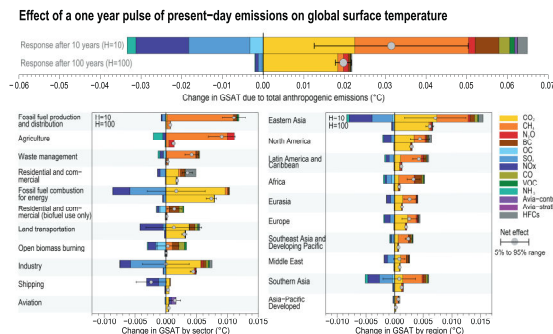
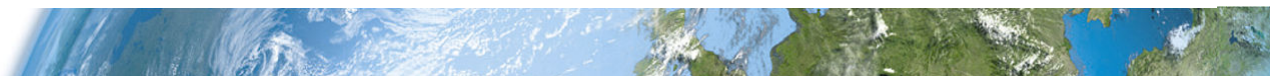
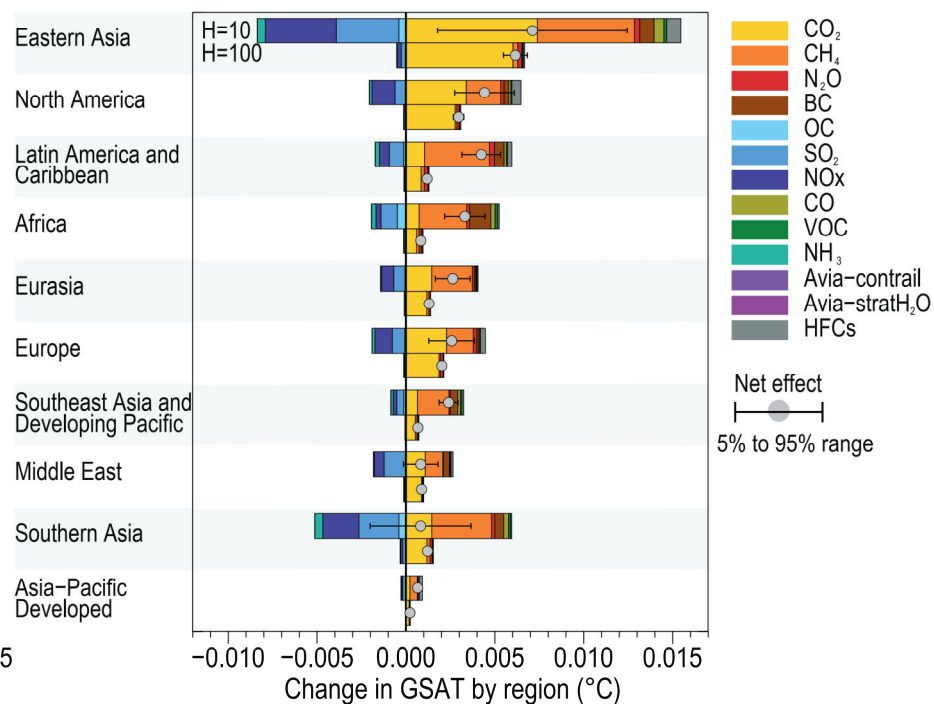
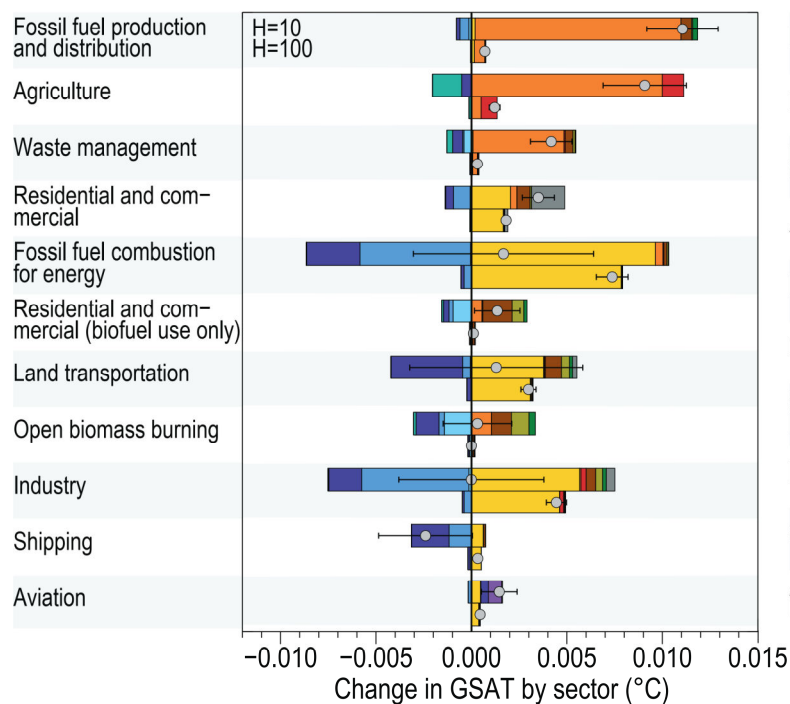
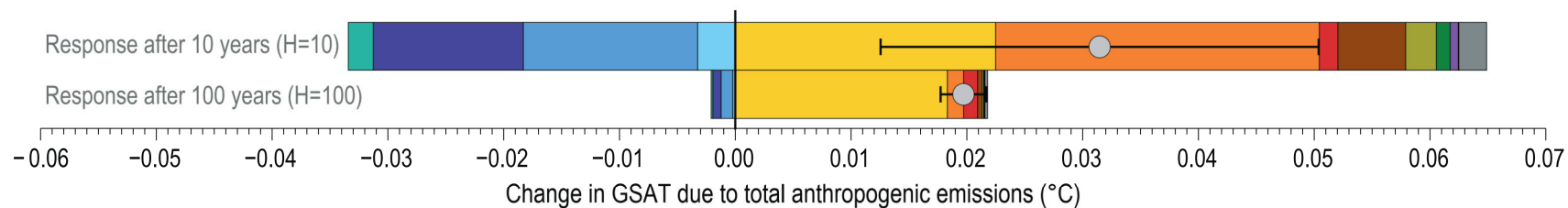


Figure 6.16 | Global mean temperature response 10 and 100 years following one year of present-day (year 2014) emissions. The temperature response is broken down by individual species and shown for total anthropogenic emissions (top), sectoral emissions (left) and regional emissions (right). Sectors and regions are sorted by (high-to-low) net temperature effect on the 10-year time scale. Error bars in the top panel show uncertainty (5–95% interval) in net temperature effect due to uncertainty in radiative forcing *only* (calculated using a Monte Carlo approach and best estimate uncertainties from the literature – see Lund et al. (2020) for details). CO₂ emissions are excluded from open biomass burning and residential biofuel use due to their unavailability in the Community Emissions Data System (CEDS) and uncertainties around non-sustainable emission fraction. Emissions for 2014 originate from the CEDS (Hoesly et al., 2018), except for HFCs which are from Purohit et al. (2020), open biomass burning from van Marle et al. (2017), and aviation H₂O which is from Lee et al. (2021). The split of fossil fuel production and distribution and combustion for energy and residential and commercial fuel use into fossil fuel and biofuel components is obtained from the GAINS model (ECLIPSE version 6b dataset). Open biomass burning emissions are not included for the regions. Emissions are aggregated into fossil fuel production and distribution (coal mining, oil and gas production, upstream gas flaring and gas distribution networks), agriculture (livestock and crop production), fossil fuel combustion for energy (power plants), industry (combustion and production processes, solvent-use losses from production and end use), residential and commercial (fossil fuel use for cooking and heating as well is HFCs leakage from A/C and refrigeration), waste management (solid waste, including landfills and open trash burning, residential and industrial waste water), transport (road and off-road vehicles, and HFC leakage from A/C and refrigeration equipment), residential and commercial (biofuels use for cooking and heating), open biomass burning (forest, grassland, savanna fires and agricultural waste burning), shipping (including international shipping), and aviation (including international aviation). Further details on data sources and processing are available in the chapter data table (Table 6. SM.3).



Effect of a one year pulse of present-day emissions on global surface temperature



Statements in the Executive Summary

Effect of SLCFs on Climate and Biogeochemical Cycles (3)

Over the 1750–2019 period, changes in SLCF emissions, especially of methane, NO_x and SO₂, have substantial effects on effective radiative forcing (ERF) (*high confidence*). The net global emissions-based ERF of NO_x is negative and that of non-methane volatile organic compounds (NMVOCs) is positive, in agreement with the AR5 Assessment (*high confidence*). For methane, the emissions-based ERF is twice as high as the abundance-based ERF (*high confidence*) attributed to chemical adjustment mainly via ozone production. SO₂ emissions changes make the dominant contribution to the ERF from aerosol–cloud interactions (*high confidence*). Over the 1750–2019 period, the contributions from the emitted compounds to changes in global surface air temperature (GSAT) broadly match their contributions to the ERF (*high confidence*). Since a peak in emissions-induced SO₂ ERF has already occurred recently and since there is a delay in the full GSAT response, changes in SO₂ emissions have a slightly larger contribution to GSAT change than CO₂ emissions, relative to their respective contributions to ERF. {6.4.2, 6.6.1 and 7.3.5}



Statements in the Executive Summary

Effect of SLCFs on Climate and Biogeochemical Cycles (4)

Reactive nitrogen, ozone and aerosols affect terrestrial vegetation and the carbon cycle through deposition and effects on large-scale radiation (*high confidence*). However, the magnitude of these effects on the land carbon sink, ecosystem productivity and hence their indirect CO₂ forcing remain uncertain due to the difficulty in disentangling the complex interactions between the individual effects. As such, these effects are assessed to be of second order in comparison to the direct CO₂ forcing (*high confidence*), but effects of ozone on terrestrial vegetation could add a substantial (positive) forcing compared with the direct ozone forcing (*low confidence*). {6.4.4}



Statements in the Executive Summary

Effect of SLCFs on Climate and Biogeochemical Cycles (5)

Climate feedbacks induced from changes in emissions, abundances or lifetimes of SLCFs mediated by natural processes or atmospheric chemistry are assessed to have an overall cooling effect (*low confidence*), that is, a total negative feedback parameter of -0.20 [-0.41 to $+0.01$] $\text{W m}^{-2} \text{ } ^\circ\text{C}^{-1}$. These non-CO₂ biogeochemical feedbacks are estimated from ESMs, which have advanced since AR5 to include a consistent representation of biogeochemical cycles and atmospheric chemistry. However, process-level understanding of many chemical and biogeochemical feedbacks involving SLCFs, particularly natural emissions, is still emerging, resulting in low confidence in the magnitude and sign of most SLCF climate feedback parameters. {6.2.2, 6.4.5}



Statements in the Executive Summary

Future Projections for Air Quality Considering Shared Socio-economic Pathways (SSPs) (1)

Future air quality (in term of surface ozone and PM concentrations) on global to local scales will be primarily driven by changes in precursor emissions as opposed to climate change (*high confidence*) and climate change is projected to have mixed effects. A warmer climate is expected to reduce surface ozone in regions remote from pollution sources (*high confidence*) but is expected to increase it by a few parts per billion over polluted regions, depending on ozone precursor levels (*medium to high confidence*). Future climate change is expected to have mixed effects, positive or negative, with an overall low effect, on global surface PM and more generally on the aerosol global burden (*medium confidence*), but stronger effects are not excluded in regions prone to specific meteorological conditions (*low confidence*). Overall, there is *low confidence* in the response of surface ozone and PM to future climate change due to the uncertainty in the response of the natural processes (e.g., stratosphere–troposphere exchange, natural precursor emissions, particularly including biogenic volatile organic compounds, wildfire-emitted precursors, land and marine aerosols, and lightning NO_x) to climate change. {6.3, 6.5}

IPCC 2021, Chap. 6



Statements in the Executive Summary

Future Projections for Air Quality Considering Shared Socio-economic Pathways (SSPs) (2)

The SSPs span a wider range of SLCF emissions than the Representative Concentration Pathways (RCPs), representing better the diversity of future options in air pollution management (*high confidence*). In the SSPs, the socio-economic assumptions and climate change mitigation levels primarily drive future emissions, but the SLCF emissions trajectories are also steered by varying levels of air pollution control originating from the SSP narratives, independently from climate change mitigation. Consequently, SSPs consider a large variety of regional ambitions and effectiveness in implementing air pollution legislation and result in wider range of future air pollution levels and SLCF-induced climate effects. {6.7.1}



Statements in the Executive Summary

Future Projections for Air Quality Considering Shared Socio-economic Pathways (SSPs) (3)

Air pollution projections range from strong reductions in global surface ozone and PM (e.g., SSP1-2.6, with strong mitigation of both air pollution and climate change) to no improvement and even degradation (e.g., SSP3-7.0 without climate change mitigation and with only weak air pollution control) (*high confidence*). Under the SSP3-7.0 scenario, PM levels are projected to increase until 2050 over large parts of Asia, and surface ozone pollution is projected to worsen over all continental areas through 2100 (*high confidence*). Without climate change mitigation but with stringent air pollution control (SSP5-8.5), PM levels decline through 2100, but high methane levels hamper the decline in global surface ozone at least until 2080 (*high confidence*). {6.7.1}



Statements in the Executive Summary

Future Projections of the Effect of SLCFs on GSAT in the Core SSPs (1)

In the next two decades, it is *very likely* that the SLCF emissions changes in the WGI core set of SSPs will cause a warming relative to 2019, whatever the SSPs, in addition to the warming from long-lived greenhouse gases. The net effect of SLCF and hydrofluorocarbon (HFC) changes on GSAT across the SSPs is a *likely* warming of 0.06°C – 0.35°C in 2040 relative to 2019. Warming over the next two decades is quite similar across the SSPs due to competing effects of warming (methane, ozone) and cooling (aerosols) SLCFs. For the scenarios with the most stringent climate and air pollution mitigations (SSP1-1.9 and SSP1-2.6), the likely near-term warming from the SLCFs is predominantly due to sulphate aerosol reduction, but this effect levels off after 2040. In the absence of climate change policies and with weak air pollution control (SSP3-7.0), the likely near-term warming due to changes in SLCFs is predominantly due to increases in methane, ozone and HFCs, with smaller contributions from changes in aerosols. SSP5-8.5 has the highest SLCF-induced warming rates due to warming from methane and ozone increases and reduced aerosols due to stronger air pollution control compared to the SSP3-7.0 scenario

IPCC 2021, Chap. 6



Statements in the Executive Summary

Future Projections of the Effect of SLCFs on GSAT in the Core SSPs (2)

At the end of the century, the large diversity of GSAT response to SLCF changes among the scenarios robustly covers the possible futures, as the scenarios are internally consistent and span a range from very high to very low emissions. In the scenarios without climate change mitigation (SSP3-7.0 and SSP5-8.5) the *likely* range of the estimated warming due to SLCFs in 2100 relative to 2019 is 0.4°C–0.9°C {6.7.3, 6.7.4}. In SSP3-7.0 there is a near-linear warming due to SLCFs of 0.08°C per decade, while for SSP5-8.5 there is a more rapid warming in the first half of the century. For the scenarios considering the most stringent climate and air pollution mitigations (SSP1-1.9 and SSP1-2.6), the reduced warming from reductions in methane, ozone and HFCs partly balances the warming from reduced aerosols, and the overall SLCF effect is a *likely* increase in GSAT of 0.0°C–0.3°C in 2100, relative to 2019. The SSP2-4.5 scenario (with moderate climate change and air pollution mitigations) results in a *likely* warming of 0.2°C–0.5°C in 2100 due to SLCFs, with the largest warming from reductions in aerosols. {6.7.3}

IPCC 2021, Chap. 6



Time evolution of the effects of changes in short-lived climate forcers (SLCFs) and hydrofluorocarbons (HFCs) on global surface air temperature (GSAT) across the WGI core set of Shared Socio-Economic Pathways (SSPs)

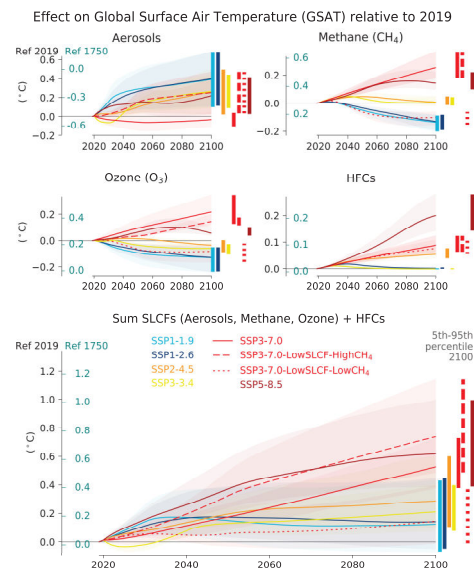
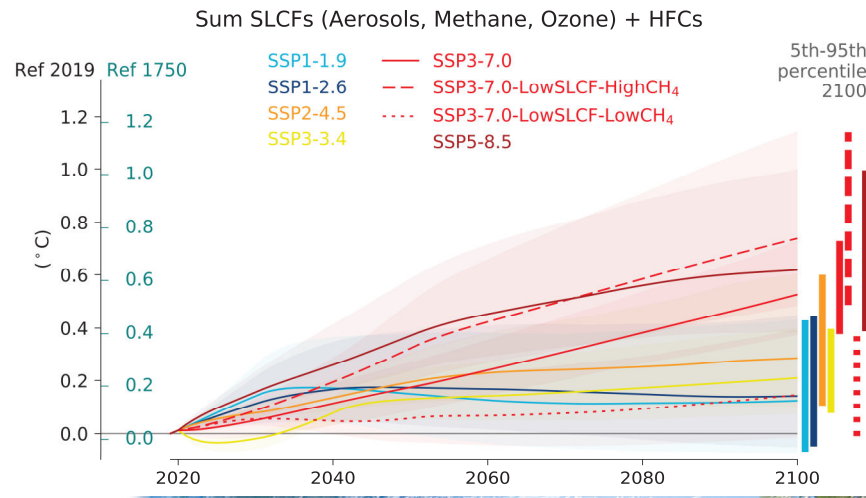
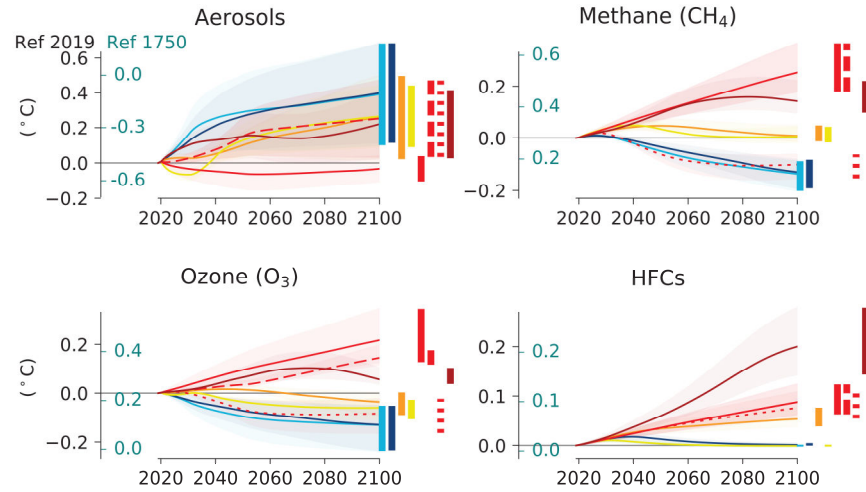


Figure 6.22 | Time evolution of the effects of changes in short-lived climate forcers (SLCFs) and hydrofluorocarbons (HFCs) on global surface air temperature (GSAT) across the WGI core set of Shared Socio-Economic Pathways (SSPs). Effects of net aerosols, methane, tropospheric ozone and hydrofluorocarbons (HFCs; with lifetimes <50years), and the sum of these, relative to the year 2019 and to the year 1750. The GSAT changes are based on the assessed historic and future evolution of effective radiative forcing (ERF; Section 7.3.5). The temperature responses to the ERFs are calculated with an impulse response function with an equilibrium climate sensitivity of 3.0°C for a doubling of atmospheric CO₂ (feedback parameter of $-1.31 \text{ W m}^{-2} \text{ °C}^{-1}$, see Cross-Chapter Box 7.1). The vertical bars to the right in each panel show the uncertainties (5–95% ranges) for the GSAT change between 2019 and 2100. Further details on data sources and processing are available in the chapter data table (Table 6.SM.3).





Effect on Global Surface Air Temperature (GSAT) relative to 2019



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (1)

Over time scales of 10 to 20 years, the global temperature response to a year's worth of current emissions of SLCFs is at least as large as that due to a year's worth of CO₂ emissions (*high confidence*). Sectors producing the largest SLCF-induced warming are those dominated by methane emissions: fossil fuel production and distribution, agriculture and waste management (*high confidence*). On these time scales, SLCFs with cooling effects can significantly mask the CO₂ warming in the case of fossil fuel combustion for energy and land transportation, or completely offset the CO₂ warming and lead to an overall net cooling in the case of industry and maritime shipping (prior to the implementation of the revised fuel-sulphur limit policy for shipping in 2020) (*medium confidence*). Ten years after a one-year pulse of present-day aviation emissions, SLCFs induce strong but short-lived warming contributions to the GSAT response (*medium confidence*), while CO₂ both gives a warming effect in the near term and dominates the long-term warming impact (*high-confidence*). {6.6.1, 6.6.2}



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (2)

The effects of SLCFs decay rapidly over the first few decades after pulse emission. Consequently, on time scales longer than about 30 years, the net long-term global temperature effects of sectors and regions are dominated by CO₂ (*high confidence*). The global mean temperature response following a climate change mitigation measure that affects emissions of both short- and long-lived climate forcers depends on their atmospheric decay times, how fast and for how long the emissions are reduced, and the inertia in the climate system. For SLCFs including methane, the rate of emissions drives the long-term global temperature effect, as opposed to CO₂ for which the long-term global temperature effect is controlled by the cumulative emissions. About 30 years or more after a one-year emission pulse occurs, the sectors contributing the most to global warming are industry, fossil fuel combustion for energy and land transportation, essentially through CO₂ (*high confidence*). Current emissions of SLCFs, CO₂ and N₂O from Eastern Asia and North America are the largest regional contributors to additional net future warming on both short (medium confidence) and long time scales (*high confidence*). {6.6.1, 6.6.2}

IPCC 2021, Chap. 6



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (3)

At present, emissions from the residential and commercial sectors (fossil and biofuel use for cooking and heating) and the energy sector (fossil fuel production, distribution and combustion) contribute the most to the world population's exposure to anthropogenic fine PM (*high confidence*), whereas emissions from the energy and land transportation sectors contribute the most to ozone exposure (*medium to high confidence*). The contribution of different sectors to PM varies across regions, with the residential sector being the most important in Southern Asia and Africa, agricultural emissions dominating in Europe and North America, and industry and energy production dominating in Central and Eastern Asia, Latin America and the Middle East. Energy and industry are important PM_{2.5} contributors in most regions, except Africa (*high confidence*). Sector contributions to surface ozone concentrations are similar for all regions. {6.6.2}



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (4)

Assuming implementation and efficient enforcement of both the Kigali Amendment to the Montreal Protocol on Ozone Depleting Substances and current national plans to limit emissions (as in SSP1-2.6), the effects of HFCs on GSAT, relative to 2019, would remain below $+0.02^{\circ}\text{C}$ from 2050 onwards versus about $+0.04^{\circ}\text{C}$ to $+0.08^{\circ}\text{C}$ in 2050 and $+0.1^{\circ}\text{C}$ to $+0.3^{\circ}\text{C}$ in 2100 considering only national HFC regulations decided prior to the Kigali Amendment (as in SSP5-8.5) (*medium confidence*). Further improvements in the efficiency of refrigeration and air-conditioning equipment during the transition to low-global-warming-potential refrigerants would bring additional greenhouse gas reductions (*medium confidence*) resulting in benefits for climate change mitigation and to a lesser extent for air quality due to reduced air pollutant emissions from power plants. {6.6.3, 6.7.3}



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (5a)

Future changes in SLCFs are expected to cause additional warming. This warming is stable after 2040 in scenarios leading to lower global air pollution as long as methane emissions are also mitigated, but the overall warming induced by SLCF changes is higher in scenarios in which air quality continues to deteriorate (induced by growing fossil fuel use and limited air pollution control) (*high confidence*). If a strong air pollution control resulting in reductions in anthropogenic aerosols and non-methane ozone precursors was considered in SSP3-7.0, it would lead to a likely additional near-term global warming of 0.08 [0.00 to 0.10] °C in 2040. An additional concomitant methane mitigation (consistent with SSP1's stringent climate change mitigation policy implemented in the SSP3 world) would not only alleviate this warming but would turn this into a cooling of 0.07°C with a likely range of [−0.02 to +0.14] °C (compared with SSP3-7.0 in 2040). ...



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (5b)

... Across the SSPs, the collective reduction of methane, ozone precursors and HFCs can make a difference of 0.2°C with a very likely range of [0.1 to 0.4] °C in 2040 and 0.8°C with a very likely range of [0.5 to 1.3] °C at the end of the 21st century (comparing SSP3-7.0 and SSP1-1.9), which is substantial in the context of the Paris Agreement. Sustained methane mitigation, wherever it occurs, stands out as an option that combines near- and long-term gains on surface temperature (high confidence) and leads to air-quality benefits by reducing surface ozone levels globally (high confidence). {6.6.3, 6.7.3, 4.4.4}



Statements in the Executive Summary

Potential Effects of SLCF Mitigation (6)

Rapid decarbonization strategies lead to air-quality improvements but are not sufficient to achieve, in the near term, air-quality guidelines set for fine PM by the World Health Organization (WHO), especially in parts of Asia and in some other highly polluted regions (*high confidence*). Additional methane and BC mitigation would contribute to offsetting the additional warming associated with SO₂ reductions that would accompany decarbonization (*high confidence*). Strong air pollution control as well as strong climate change mitigation, implemented separately, lead to large reductions in exposure to air pollution by the end of the century (*high confidence*). Implementation of air pollution controls, relying on the deployment of existing technologies, leads more rapidly to air quality benefits than climate change mitigation (*high confidence*), which requires systemic changes. However, in both cases, significant parts of the population are projected to remain exposed to air pollution exceeding the WHO guidelines (*high confidence*). Additional policies envisaged to attain Sustainable Development Goals (SDGs; e.g., access to clean energy, waste management) bring complementary SLCF reduction. Only strategies integrating climate, air quality, and development goals are found to effectively achieve multiple benefits. {6.6.3, 6.7.3, Box 6.2}

Statements in the Executive Summary

Implications of COVID-19 Restrictions for Emissions, Air Quality and Climate

Emissions reductions associated with COVID-19 containment led to a discernible temporary improvement of air quality in most regions, but changes to global and regional climate are undetectable above internal variability. Global anthropogenic NO_x emissions decreased by a maximum of about 35% in April 2020 (*medium confidence*). There is *high confidence* that, with the exception of surface ozone, these emissions reductions have contributed to improved air quality in most regions of the world. Global fossil CO₂ emissions decreased by 7% (with a range of 5.8–13.0%) in 2020 relative to 2019, largely due to reduced emissions from the transportation sector (medium confidence). Overall, the net ERF, relative to ongoing trends, from COVID-19 restrictions was likely small and positive for 2020 (less than 0.2 W m⁻²), thus temporarily adding to the total anthropogenic climate influence, with positive forcing from aerosol changes dominating over negative forcings from CO₂, NO_x and contrail cirrus changes. Consistent with this small net radiative forcing, and against a large component of internal variability, Earth system model simulations show no detectable effect on global or regional surface temperature or precipitation (high confidence). {Cross-Chapter Box 6.1}

IPCC 2021, Chap. 6

