

1 **Evaluating the Climate and Air Quality Impacts of Short-**
2 **Lived Pollutants**

4 **A. Stohl¹, B. Aamaas², M. Amann³, L. H. Baker⁴, N. Bellouin⁴, T. K. Berntsen², O.**
5 **Boucher⁵, R. Cherian⁶, W. Collins^{4, 7}, N. Daskalakis^{8, 9}, M. Dusinska¹, S. Eck-**
6 **hardt¹, J. S. Fuglestvedt², M. Harju¹, C. Heyes³, Ø. Hodnebrog², J. Hao¹⁰, U. Im^{8,}**
7 **11, M. Kanakidou^{8, 9}, Z. Klimont³, K. Kupiainen³, K. S. Law¹², M. T. Lund², R.**
8 **Maas¹³, C. R. MacIntosh⁴, G. Myhre², S. Myriokefalitakis^{8, 9}, D. Olivié¹⁴, J.**
9 **Quaas⁶, B. Quennerhen¹², J.-C. Raut¹², S. T. Rumbold⁷, B. H. Samset², M.**
10 **Schulz¹⁴, Ø. Seland¹⁴, K. P. Shine⁴, R. B. Skeie², S. Wang¹⁰, K. E. Yttri¹, T. Zhu¹⁵**

12 [1] {NILU - Norwegian Institute for Air Research, Kjeller, Norway}

13 [2] {Center for International Climate and Environmental Research – Oslo (CICERO), Oslo, Norway}

14 [3] {International Institute for Applied Systems Analysis (IIASA), Laxenburg, Austria}

15 [4] {Department of Meteorology, University of Reading, Reading, UK}

16 [5] {LATMOS, Université Pierre et Marie Curie (UPMC)/CNRS, Paris, France}

17 [6] {Institute for Meteorology, Universität Leipzig, Germany}

18 [7] {Met Office Hadley Centre, Exeter, UK}

19 [8] {Environmental Chemical Processes Laboratory, Department of Chemistry, University of
20 Crete, Heraklion, Crete, Greece}

21 [9] {FORTH, ICE-HT, Platani, Patras, Greece}

22 [10] {School of Environment, Tsinghua University, Beijing, China}

23 [11] {Now at: Aarhus University, Department of Environmental Science, Frederiksbergvej 399, 4000 Roskilde,
24 Denmark}

25 [12] {Sorbonne Universités, UPMC Univ. Paris 06 ; Université Versailles St-Quentin ; CNRS/INSU ; LAT-
26 MOS-IPSL, UMR 8190, Paris, France}

27 [13] {RIVM - National Institute for Public Health and the Environment, Bilthoven, Netherlands}

28 [14] {Norwegian Meteorological Institute, Oslo, Norway}

29 [15] {State Key Laboratory for Environmental Simulation and Pollution Control, College of Environmental
30 Sciences and Engineering, Peking University, Beijing, China}

1 **Correspondence to: A. Stohl (ast@nilu.no)**

2 **Abstract**

3 This paper presents a summary of the work done within the European Union's Seventh
4 Framework Programme project ECLIPSE (Evaluating the Climate and Air Quality Impacts of
5 Short-Lived Pollutants). ECLIPSE had a unique systematic concept for designing a realistic
6 and effective mitigation scenario for short-lived climate pollutants (SLCPs: methane, aerosols
7 and ozone, and their precursor species) and quantifying its climate and air quality impacts,
8 and this paper presents the results in the context of this overarching strategy. The first step in
9 ECLIPSE was to create a new emission inventory based on current legislation (CLE) for the
10 recent past and until 2050. Substantial progress compared to previous work was made by in-
11 cluding previously unaccounted types of sources such as flaring of gas associated with oil
12 production, and wick lamps. These emission data were used for present-day reference simula-
13 tions with four advanced Earth system models (ESMs) and six chemistry transport models
14 (CTMs). The model simulations were compared with a variety of ground-based and satellite
15 observational data sets from Asia, Europe and the Arctic. It was found that the models still
16 underestimate the measured seasonality of aerosols in the Arctic but to a lesser extent than in
17 previous studies. Problems likely related to the emissions were identified for Northern Russia
18 and India, in particular. To estimate the climate impacts of SLCPs, ECLIPSE followed two
19 paths of research: The first path calculated radiative forcing (RF) values for a large matrix of
20 SLCP species emissions, for different seasons and regions independently. Based on these RF
21 calculations, the Global Temperature change Potential metric for a time horizon of 20 years
22 (GTP₂₀) was calculated for each SLCP emission type. This climate metric was then used in an
23 integrated assessment model to identify all emission mitigation measures with a beneficial air
24 quality and short-term (20-year) climate impact. These measures together defined a SLCP
25 mitigation (MIT) scenario. Compared to CLE, the MIT scenario would reduce global methane
26 (CH₄) and black carbon emissions by about 50% and 80%, respectively. For CH₄, measures
27 on shale gas production, waste management and coal mines were most important. For non-
28 CH₄ SLCPs, elimination of high emitting vehicles and wick lamps, as well as reducing emis-
29 sions from gas flaring, coal and biomass stoves, agricultural waste, solvents and diesel en-
30 gines were most important. These measures lead to large reductions in calculated surface con-
31 centrations of ozone and particulate matter. We estimate that in the EU the loss of statistical
32 life expectancy due to air pollution was 7.5 months in 2010, which will be reduced to 5.2

1 months by 2030 in the CLE scenario. The MIT scenario would reduce this value by another
2 0.9 months to 4.3 months. Substantially larger reductions due to the mitigation are found for
3 China (1.8 months) and India (11-12 months). The climate metrics cannot fully quantify the
4 climate response. Therefore, a second research path was taken. Transient climate ensemble
5 simulations with these ESMs were run for the CLE and MIT scenarios, to determine the cli-
6 mate impacts of the mitigation. In these simulations, the CLE scenario resulted in a surface
7 temperature increase of 0.70 ± 0.14 K between the years 2006 and 2050. For the decade 2041-
8 2050, the warming was reduced by 0.22 ± 0.07 K in the MIT scenario, and this result was in
9 almost exact agreement with the response calculated based on the emission metrics (reduced
10 warming of 0.22 ± 0.09 K). The metrics calculations suggest that non-CH₄ SLCPs contribute
11 ~22% to this response and CH₄ 78%. This could not be fully confirmed by the transient simu-
12 lations, which attributed about 90% of the temperature response to CH₄ reductions. Attribu-
13 tion of the observed temperature response to non-CH₄ SLCP emission reductions and black
14 carbon (BC) specifically is hampered in the transient simulations by small forcing and co-
15 emitted species of the emission basket chosen. Nevertheless, an important conclusion is that
16 our mitigation basket as a whole would lead to clear benefits for both air quality and climate.
17 The climate response from BC reductions in our study is smaller than reported previously,
18 largely possibly because our study is one of the first to use fully coupled climate models,
19 where unforced variability and sea-ice responses cause relatively strong temperature fluctua-
20 tions which may counteract (and, thus, mask) the impacts of small emission reductions. The
21 temperature responses to the mitigation were generally stronger over the continents than over
22 the oceans, and with a warming reduction of 0.44 K (0.39-0.49) largest over the Arctic. Our
23 calculations suggest particularly beneficial climate responses in Southern Europe, where the
24 surface warming was reduced by about 0.3 K and precipitation rates were increased by about
25 15 (6-21) mm/yr (more than 4% of total precipitation) from spring to autumn. Thus, the miti-
26 gation could help to alleviate expected future drought and water shortages in the Medi-
27 terranean area. We also report other important results of the ECLIPSE project.

28

29 **1 Introduction**

30 The United Nations Framework Convention on Climate Change (UNFCCC) requires climate
31 policies to 'be cost-effective so as to ensure global benefits at the lowest possible cost' and

1 that ‘policies and measures should … be comprehensive … [and] … cover all relevant
2 sources, sinks and reservoirs’. This was made operational by the Kyoto Protocol, which sets
3 limits on emissions of six different greenhouse gases (GHGs), or groups of GHGs – carbon
4 dioxide (CO₂), methane (CH₄), nitrous oxide (N₂O), perfluorocarbons (PFCs), hydrofluoro-
5 carbons (HFCs) and sulfur hexafluoride (SF₆). Collectively these are often known as “the
6 Kyoto gases” or the “Kyoto basket” ¹. CO₂ is the most important anthropogenic driver of
7 global warming, with additional significant contributions from CH₄ and N₂O. However, other
8 anthropogenic emissions capable of causing climate change are not covered by the Kyoto Pro-
9 tocol. Some are covered by other protocols, e.g. emissions of chlorofluorocarbons (CFCs) and
10 hydrochlorofluorocarbons (HCFCs) are regulated by the Montreal Protocol, because of their
11 role in stratospheric ozone (O₃) depletion. But there are others, notably several short-lived
12 components that give strong contributions to climate change that are excluded from existing
13 climate agreements.

14 In the present study we investigate climate and air quality impacts of the emissions of CH₄,
15 which has a lifetime of about 9±1 years (Prather et al., 2012) and a number of much shorter-
16 lived components (atmospheric lifetimes of months or less) which directly or indirectly (via
17 formation of other short-lived species) influence the climate (Myhre et al., 2013a):

- 18 • Methane is a greenhouse gas with a radiative efficiency (in W m⁻² ppbv⁻¹) roughly 26
19 times stronger greater than that of CO₂ on a per molecule basis at current concentra-
20 tions. It is relatively well-mixed in the atmosphere and has both natural and anthro-
21 genic sources. It is also a precursor of O₃ and stratospheric water vapour.
- 22 • Black carbon (BC, also commonly known as soot), a product of incomplete combus-
23 tion of fossil fuels and biomass, affects climate via several mechanisms (Bond et al.,
24 2013). It causes warming through absorption of sunlight and by reducing surface albe-
25 do when deposited on snow. BC also affects clouds, with a consequent (but highly un-
26 certain) impact on their distribution and radiative properties (Boucher et al., 2013).

¹ Note that, formally, only species given values of Global Warming Potentials (GWP) in IPCC’s Second Assessment Report were controlled during the first commitment period (2008-2012) of the Kyoto Protocol. The second commitment period (2013-2020), via the Doha Amendment, also includes NF₃ in the list of greenhouse gases, and uses GWP values from the IPCC’s Fourth Assessment Report. The Doha Amendment is currently not in force, as it awaits ratification by a sufficient number of parties.

- Tropospheric O₃ is a greenhouse gas produced by chemical reactions from the emissions of the precursors CH₄, carbon monoxide (CO), non-CH₄ volatile organic compounds (NMVOCs) and nitrogen oxides (NO_x). Emissions of these same precursors also impact on hydroxyl radical (OH) concentrations with further impacts especially on CH₄.
- Several components have cooling effects on climate, mainly sulphate aerosol formed from sulphur dioxide (SO₂) and ammonia (NH₃), nitrate aerosol formed from NO_x and NH₃, and organic aerosol (OA) which can be directly emitted or formed from gas-to-particle conversion of NMVOCs. They cause a direct cooling by scattering solar radiation and alter the radiative properties of clouds, very likely leading to further cooling.

We refer to these substances as short-lived climate pollutants (SLCPs) as they also have detrimental impacts on air quality, directly or via formation of secondary pollutants (Kirtman et al., 2013). Notice that we include the precursors of O₃ and secondary aerosols in our definition of SLCPs. We also include CH₄ in our study even though it is included in the Kyoto Protocol, because of its relatively short lifetime compared to that of CO₂ and its importance for air quality via the formation of O₃. We do not include HFCs in our definition of SLCPs, as they have no significant impact on air quality and can be regulated from a climate policy perspective alone. For SLCPs, on the other hand, cost-effective environmental policy measures should be designed such that they optimize both the climate and air quality responses (Schmale et al., 2014). In some instances, control of the emissions of a species is expected to reduce future warming and improve air quality at the same time – a “win-win” situation (Anenberg et al., 2012); in others, the control of emissions may be conflicting, in the sense that it could increase warming while improving air quality (or vice versa) – in this case, emission control involves a “trade-off” between the impacts.

The net climate impact since pre-industrial times of all short-lived components other than CH₄ together is very likely to be cooling due primarily to sulphate aerosols (Myhre et al., 2013a). Whilst SLCP reductions are clearly beneficial for air quality, elimination of all current non-CH₄ SLCP emissions would thus very likely lead to extra warming. Nevertheless, targeted emission reductions of selected SLCPs which cause warming (either directly or via formation of secondary species) have the potential to reduce global warming on a short time-scale, as well as improving air quality. They may also reduce the rate of warming (Myhre et

1 al., 2011; Shindell et al., 2012) that is important, for example, for the adaptation of ecosys-
2 tems to climate change (as recognized by UNFCCC Art. 2) and is expected to accelerate in
3 the near future (Smith et al., 2015). Reducing these selected SLCP emissions might be effec-
4 tive to help avoid (or at least delay) certain undesired impacts of climate change (e.g., rapid
5 sea ice loss in the Arctic; Quinn et al., 2008). At least, optimized SLCP emission reductions
6 could help to reduce the undesired extra climate warming caused by air quality policy
7 measures that often do not consider climate impacts.

8 There are many studies that explore possibilities and effects of reductions of short-lived com-
9 ponents (e.g., Brasseur and Roeckner, 2005; Rypdal et al., 2009a; Kopp and Mauzerall, 2010;
10 Penner et al., 2010; Unger et al., 2010; Shindell et al., 2012; Bond et al., 2013; Bowerman et
11 al., 2013; Rogelj et al., 2014). Given the interest from policymakers in the abatement of
12 SLCPs, an urgent challenge is to determine the exact climate impacts of the different species
13 involved (e.g., Penner et al., 2010). BC has received particular attention as a component for
14 which a specific emission reduction might have an immediate climate benefit (e.g., Bond and
15 Sun, 2005; Boucher and Reddy, 2008; Grieshop et al., 2009; Rypdal et al., 2009b; Berntsen et
16 al., 2010; Bond et al., 2013).

17 For designing a successful SLCP emission abatement strategy, the key CH₄ sources are rela-
18 tively straightforward to deal with because their emission profile is dominated by CH₄ (e.g.,
19 venting of natural gas, rice paddies, etc.). Combustion sources, however, emit a mix of many
20 different SLCPs (e.g., BC, OA, NO_x, SO₂) as well as CO₂. This makes it difficult to reduce
21 the emissions of warming agents (e.g., BC) alone, as their control often also leads to removal
22 of co-emitted cooling agents (e.g., OA, SO₂). To achieve a climate benefit, abatement strate-
23 gies will be most effective if they target sources with a high fraction of warming species in
24 their emissions (e.g., diesel vehicles) (Unger et al., 2010).

25 **1.1 Climate effects of SLCPs**

26 There are several distinct issues that have to be addressed in considering the impact of any
27 proposed SLCP abatement strategy. First, there are large uncertainties in estimates of the cli-
28 mate effects of SLCPs (see e.g. Myhre et al., 2013a) and thus also in the effects of emission
29 reductions. These apply particularly to the impact of aerosols on cloud properties (e.g. Quaas
30 et al., 2009; Boucher et al. 2013), but there are also difficulties in evaluating direct radiative
31 effects of aerosols.

1 Second, the climate impact of short-lived components, even when averaged globally, can de-
2 pend strongly on location and time (e.g., summer vs. winter) of emissions (Fuglestvedt et al.,
3 1999; Wild et al., 2001; Berntsen et al., 2005, 2006; Koch et al., 2007; Naik et al., 2005; Red-
4 dy and Boucher, 2007; Shindell and Faluvegi, 2009). For well-mixed gases (e.g. Kyoto gases),
5 a single globally-valid value of Global Warming Potential (GWP – see Section 1.2 for
6 more details) can be calculated for a chosen time horizon, and then used to give the so-called
7 “CO₂-equivalent” emissions of a gas. By contrast, for the non-CH₄ SLCs, the GWP depends
8 significantly on when and where the emission occurs. Not only does this complicate the cal-
9 culation of GWPs, it also introduces an additional dimension into the framing of climate poli-
10 cy. For instance, the importance of location for BC emissions has received much attention in
11 this context (Ramanathan and Carmichael, 2008; Shindell and Faluvegi, 2009).

12 Third, inhomogeneity in the climate response to RF is important for SLCs. The geographical
13 pattern of RF due to the non-CH₄ SLCs is generally concentrated close to the source of
14 emission, and hence is quite distinct from the global-scale forcing due to the Kyoto gases. The
15 extent to which these heterogeneous forcing patterns will trigger different climate responses
16 compared to well-mixed gases is an unresolved scientific issue, even though the climate re-
17 sponse generally occurs on larger spatial scales (but mainly in the hemisphere where the forc-
18 ing takes place; Joshi et al. 2003; Shindell et al., 2010) than the forcing itself. One example of
19 the issue of inhomogeneity of response concerns the effects of absorption of solar radiation by
20 BC in the Arctic atmosphere. Flanner (2013) has shown that in the Arctic BC located at low
21 altitudes causes a strong local surface warming, but BC located at higher altitudes causes a
22 surface cooling, which is due to the reduced solar radiation reaching the surface. Another im-
23 portant example is emissions of NO_x as these lead to a shorter-lived (and hence more local-
24 ised) positive RF due to increases in O₃ and a longer-lived (and hence more global) negative
25 RF due to the increased rate of destruction of CH₄. This means that metrics based on global-
26 mean quantities may be poorly representative of the local impacts of an emission as the re-
27 sponse depends on both region and timescale (Shine et al., 2005; Lund et al., 2012).

28 Fourth, SLCs may have other effects on climate that go beyond global-mean temperature
29 (Andrews et al., 2010; Kvalevåg et al., 2013) such as through changes in the hydrological
30 cycle (Gedney et al., 2014) and in the atmospheric circulation. For example, in South East
31 Europe there are indications that changes in the radiation budget through direct and indirect
32 effects of aerosols have caused circulation, precipitation and evaporation changes (Lelieveld

1 et al., 2002; Tragou and Lascaratos, 2003). Thus, even a cooling component may cause un-
2 wanted climate impacts ([Shindell, 2015](#)).

3 Finally, there are important interdependencies between SLCPs and long-term climate change.
4 The climate (and air quality) impacts of SLCPs depend on the atmosphere into which they are
5 emitted – future changes in temperature, humidity, cloud amount, surface albedo, circulation
6 and atmospheric composition are likely to change these impacts (Isaksen et al., 2009). Acting
7 in the other direction, changes in SLCP emissions can impact vegetation via changes in air
8 quality (Sitch et al., 2007; Collins et al., 2010), nutrient deposition (Mahowald, 2011; Wang
9 et al., 2015) or photosynthetic active radiation (Mercado et al., 2009), thereby altering the
10 terrestrial carbon budget and hence future CO₂ concentrations and thus giving the SLCPs a
11 much longer-term impact.

12 Taking the above points into account, the short lifetimes and regional dependence of the cli-
13 mate impact of SLCP emissions make these species fundamentally different to the long-lived
14 GHGs regulated under the Kyoto Protocol and these impacts and metric values are much
15 more uncertain (Myhre et al., 2013a). Furthermore, cooling aerosols may have partly compen-
16 sated the warming due to well-mixed greenhouse gases in the past, and this masking effect
17 must be considered when determining the sensitivity of the climate system directly from ob-
18 servations (Knutti and Hegerl, 2008; Skeie et al., 2014). This also reduces our ability to calcu-
19 late future global warming (e.g. Andreae et al., 2005; Meinshausen et al., 2009; Penner et al.,
20 2010). Thus, there is an urgent need to understand and quantify the role that these components
21 may play in international efforts to reduce global warming (Jackson, 2009; Berntsen et al.,
22 2010; Arneth et al., 2009; Rypdal et al., 2009b; Molina et al., 2009; Unger et al., 2010).

23 **1.2 Climate metrics to characterise the effect of SLCPs**

24 The Kyoto Protocol to the UNFCCC is a multi-gas climate treaty that required a method to
25 place emissions of different gases on a common scale. It adopted the GWP with a hundred-
26 year time horizon, GWP₁₀₀, from the IPCC (Intergovernmental Panel on Climate Change)
27 Second Assessment Report as a metric in order to derive so-called CO₂-equivalents for non-
28 CO₂ gas emissions. The GWP has since then been widely used in implementing the Kyoto
29 Protocol, and for other purposes. However, it was not designed with a particular climate poli-
30 cy in mind, and as a result, GWP may not be the best choice for all particular policy objec-

1 tives (e.g., Tanaka et al., 2009; Fuglestvedt et al., 2010; Myhre et al. 2013[a](#); Pierrehumbert,
2 2014).

3 The GWP gives the RF due to a pulse emission of a gas or aerosol, integrated over some time
4 horizon, relative to that of CO₂. The choice of time-horizon has a significant impact on the
5 metric value of an emission (e.g. Skodvin and Fuglestvedt, 1997; Shine, 2009; Fuglestvedt et
6 al., 2010; Aamaas et al., 2013) and is a value-laden choice. The time-integrated nature of the
7 GWP means that it retains the memory of short-lived emissions even at long-time horizons,
8 when their forcing and most of the response have subsided.

9 Several alternatives to the GWP have been proposed and of these, the Global Temperature
10 change Potential (GTP) (Shine et al., 2005, 2007; Fuglestvedt et al., 2010) has attracted most
11 attention (e.g., Reisinger et al., 2010; Boucher and Reddy, 2008; Gillett and Matthews, 2010;
12 Collins et al., 2013). The GTP gives the global-mean surface temperature change some time
13 after a pulse emission, relative to that of CO₂. In contrast to the GWP, it uses temperature as
14 the indicator and is an “end point”, rather than an “integrative”, metric. Therefore, it does not
15 retain the memory of short-lived emissions in the same way as the GWP. Difficulties with the
16 GTP include its dependence on the climate sensitivity and on the method of incorporating the
17 ocean’s thermal response (Shine et al., 2007; Fuglestvedt et al., 2010; Olivié and Peters,
18 2013).

19 The GTP may be more appropriate to target-based climate policies (UNEP/WMO, 2011)
20 where the aim is to keep temperature change below some given limit, such as the 2°C limit in
21 the UNFCCC’s Copenhagen Accord. The choice of time horizon is then no longer so arbit-
22 rary, but is linked to the time at which, for example, 2°C is likely to be reached. This use of
23 the GTP (Shine et al., 2007, Berntsen et al., 2010; Tanaka et al., 2013) mimics the behaviour
24 of more complex (but less transparent) metrics based on integrated assessment models
25 (Manne and Richels, 2001).

26 In its 5th Assessment Report, the IPCC assessed scientific aspects of climate metrics and their
27 applicability in policy making. It was emphasized that the most appropriate metric and time
28 horizon will depend on which aspects of climate change are considered most important to a
29 particular application. The assessment also pointed out that there are limitations and incon-
30 sistencies related to the treatment of indirect effects and feedbacks (e.g., climate-carbon cycle

1 feedbacks) in climate metrics. In this study, we have adopted GTP₂₀, the GTP over a 20-year
2 time horizon, as our key metric, after careful consideration of alternatives (see section 3.4).

3 **1.3 Air quality impacts of SLCPs**

4 The impact of SLCPs on air quality occurs at both the local and regional scale. While local
5 emissions contribute to episodes of high pollution levels which can cause acute health effects,
6 the long-range transport of air pollutants or their precursors even over intercontinental dis-
7 tances (e.g., Stohl and Trickl, 1999; Dentener et al., 2010) can increase the background con-
8 centrations upon which pollution episodes are superimposed. This is also important because
9 there is increasing evidence of harmful effects of long-term exposure to particulate matter
10 (PM), O₃, deposited acidifying compounds and nitrogen to human health and vegetation
11 (Anenberg et al., 2012). Thus, the impact of SLCPs on air quality is complex and requires
12 quantification on local to global scales. At international level, these aspects, including emis-
13 sion regulation, are covered by the UNECE Convention on Long-Range Transboundary Air
14 Pollution (CLRTAP) and its protocols including the Gothenburg Protocol and its amend-
15 ments.

16 The International Agency for Research on Cancer classified outdoor air pollution as *carcino-*
17 *genic to humans* with *sufficient evidence* that it causes lung cancer. A positive association
18 with an increased risk of bladder cancer was also demonstrated. It has been estimated that air
19 pollution caused 223,000 deaths from lung cancer worldwide in 2010 ([Unger Anenberg et al.,](#)
20 [2010](#)[2012; Lim et al., 2012](#)). Air quality guidelines for various substances published by dif-
21 ferent agencies are listed in Table 1.

22 Ozone and PM are the most problematic air pollutants with regard to effects on human health
23 (EEA, 2013). Ozone can, through impairment of lung function, lead to premature deaths and
24 increased hospitalization (West et al., 2006). PM was classified as *carcinogenic to humans*
25 (IARC, 2015, Grosse, 2013). It is estimated, for instance, that an increase of 10 µg/m³ in the
26 concentrations of PM₁₀ (PM with diameter smaller than 10 µm) will increase cardiopulmo-
27 nary mortality by 9% (Pope et al., 1995). Different aerosol types are considered when as-
28 ssuming climate impacts, whereas air quality legislation is based on the concept of total mass
29 concentrations of particulate matter – either as PM_{2.5} or PM₁₀. It is, however, likely that hu-
30 man health impacts also depend on PM composition. For instance, according to the World
31 Health Organization (WHO), epidemiological evidence indicates an association of daily varia-

1 tion in BC concentrations with short and long term adverse health effects such as all-cause
2 and cardiovascular mortality, and cardiopulmonary hospital admissions. Additionally, BC was
3 classified as *possibly carcinogenic to humans* (Group 2B) (WHO, 2012). However, concen-
4 tration-response functions for individual PM components still need to be established. Thus,
5 neither BC nor ultrafine particles are currently covered specifically by EU guidelines (WHO,
6 2013).

7 **2 Scope and overall concept**

8 The purpose of this paper is to present a summary of the work done within the European Un-
9 ion's Seventh Framework Programme project ECLIPSE (Evaluating the Climate and Air
10 Quality Impacts of Short-Lived Pollutants). ECLIPSE had a unique systematic concept for
11 designing a realistic and effective SLCP mitigation scenario and quantifying its climate and
12 air quality impacts, which is schematically shown in Fig. 1. Other papers describe particular
13 aspects of the ECLIPSE work in more detail, while we here present key ECLIPSE results in
14 the context of this overarching strategy and overall conclusions of the project.

15 The first step in ECLIPSE was to create a new set of global baseline emissions for the recent
16 past and future (see Section 3.1, top of Fig. 1). These emission data were used for present day
17 reference simulations with Earth system models (ESMs) and chemistry transport models
18 (CTMs). The model simulations were compared extensively with a variety of global ground-
19 based and satellite observational data sets, in particular in three target areas (China, Europe
20 and the Arctic, see Section 3.2) to evaluate their capabilities to simulate SLCP concentrations.

21 To study the climate impacts of SLCPs, ECLIPSE followed two paths of research. The first
22 path (the outer part of the spiral in Fig. 1) calculated RF values for a large matrix of SLCP
23 species emissions, for different seasons and regions independently by changing the emissions
24 of one species from one region and one season at a time (see Section 3.3). Based on these RF
25 calculations, suitable metrics were chosen to allow the estimation of the climate impact of
26 particular SLCP emissions over different time horizons (see Section 3.4).

27 These metrics were then used to generate an SLCP mitigation scenario to minimize climate
28 impacts that could be contrasted with the current legislation scenario (see the top of Fig. 1).
29 For this, the region-, season- and species-specific matrix of climate impact (as defined by the
30 chosen metric) was used as an input to an integrated assessment model. All region-, season-
31 and sector-specific emission mitigation measures with a beneficial air quality impact were

1 then evaluated according to their expected climate benefit. Notice here that emission measures
2 typically affect several SLCP species. For every mitigation measure, the emission reduction
3 of every SLCP species was therefore weighted with the chosen climate metric and summed
4 over all emitted SLCP species. Finally, all measures with beneficial air quality and climate
5 impacts were collected in a basket defining the SLCP mitigation scenario (see Section 3.5).

6 The metrics, however, cannot fully quantify the climate response, due to the underlying sim-
7 plifying assumptions, including linearity, the need to specify particular time horizons and,
8 most importantly, the focus on one single aspect of climate change (global mean temperature
9 for the chosen GTP metric). Therefore, a second research path (the inner part of the spiral in
10 Fig. 1) was taken to determine the climate response for a set of emission reductions for indi-
11 vidual SLCP species, using a small ensemble of four advanced Earth system models (ESMs).
12 Furthermore, transient climate ensemble simulations with these ESMs were run for the base-
13 line and emission mitigation scenarios, to calculate the transient climate and air quality im-
14 pacts of the mitigation scenario (see Section 3.6). A comparison between the climate impacts
15 expected from the metrics and those calculated with the transient simulations (left part of the
16 spiral in Fig. 1) closed the loop between the first and the second research path and allowed the
17 evaluation of the consistency of both approaches (see section 3.7).

18 In ECLIPSE, we used multi-model ensemble results wherever possible, as these are more ro-
19 bust than results from an individual model. However, certain calculations could only be per-
20 formed by a single model and are, thus, presented as such.

21

22 **3 Results**

23 3.1 The ECLIPSE emissions

24 The ECLIPSE emission data set was created with the GAINS (Greenhouse gas – Air pollution
25 Interactions and Synergies; <http://gains.iiasa.ac.at>) model (Amann et al., 2011), which pro-
26 vides emissions of long-lived greenhouse gases and shorter-lived species in a consistent
27 framework. The GAINS model holds essential information about key sources of emissions,
28 environmental policies, and mitigation opportunities for about 160 country-regions. The mod-
29 el relies on exogenous projections of energy use, industrial production, and agricultural activi-
30 ty (ECLIPSE scenarios draw on IEA, 2012, for energy and Alexandros and Bruinsma, 2012,

1 for agriculture) for which it distinguishes all key emission sources and control measures.
2 More than 2000 technologies to control air pollutant emissions and at least 500 options to
3 control GHG emissions are included.

4 Improvements in the emission model were made especially for China (Zhao et al., 2013;
5 Wang et al., 2014), where large changes have occurred recently, but also for Europe where
6 results of the consultation process during the development of scenarios for the review of the
7 EU National Emission Ceilings Directive (Amann and Wagner, 2014) were used. Further-
8 more, several sources like brick making, oil and gas production, non-ferrous metals and inter-
9 national shipping were reviewed and updated. Finally, a number of previously unaccounted
10 sources were added or specifically distinguished in the model, e.g., wick lamps, diesel genera-
11 tors, and high-emitting vehicles. The global SO₂ inventory used for IPCC's 5th Assessment
12 Report (Klimont et al., 2013) was also developed during ECLIPSE.

13 All emission data were gridded consistently to a resolution of 0.5°x0.5° longitude-latitude.
14 The spatial proxies used in GAINS for gridding are consistent with those applied within the
15 IPCC's Representative Concentration Pathways (RCPs) projections as described in Lamarque
16 et al. (2010) and as further developed within the Global Energy Assessment project (GEA,
17 2012). They were, however, modified to accommodate more recent year-specific information
18 where available, e.g., on population distribution, open biomass burning, location of oil and
19 gas production, and livestock-specific spatial production patterns (Klimont et al., 2013,
20 2015b). Emissions were also temporally allocated: monthly distribution was provided for all
21 sources and for the residential heating emissions were based on ambient air temperature (see
22 Stohl et al., 2013).

23 For the first time in a global emission inventory, emissions from flaring of associated gas in
24 oil production were considered directly, including spatial distribution. For BC, these emis-
25 sions constitute only about 3% of the global total. However, owing to emissions in Russia,
26 they constitute about one third of all BC emissions north of 60°N and two thirds of all emis-
27 sions north of 66°N. Stohl et al. (2013) found that the gas flaring emissions contribute 42% of
28 all BC found in the Arctic near the surface, and this has improved the performance of the
29 ECLIPSE models in the Arctic.

30 Figure 2 shows global anthropogenic ECLIPSE emissions for three developed scenarios
31 (Klimont et al., 2015a, b):

- Current legislation (CLE) includes current and planned environmental laws, considering known delays and failures up to now but assuming full enforcement in the future.
- No further control (NFC) uses the same assumptions as CLE until 2015 but no further legislation is introduced subsequently, even if currently committed. This leads to higher emissions than in CLE for most pollutants.
- The ECLIPSE SLCP mitigation scenario (MIT) includes all measures with beneficial air quality and climate impact (according to the climate metric; see sections 3.4 and 3.5).

Different versions of the ECLIPSE inventory (available on request from <http://eclipse.nilu.no>); also available from: http://www.iiasa.ac.at/web/home/research/researchPrograms/Global_emissions.html have been developed and were available at different times for different tasks (Klimont et al., 2015a, b). We describe here the version 5, which was used for the transient climate model simulations (section 3.6). For model evaluation (section 3.2) and climate perturbation simulations in section 3.6, versions 4 and 4a were used which, for the CLE scenario were very similar to version 5 (Klimont et al., 2015a, b).

During the past few decades, there was a strong growth in CO₂ emissions, but the SLCP emissions have followed a different trajectory, at least at the global level. For example, the SO₂ emissions have been decreasing since 1990, with a temporary increase between 2000 and 2005 (Klimont et al., 2013), owing to strong policies and drastic reductions in Europe and North America. The strong development in Asia was offset at the global level by these reductions but in the future, emissions of SO₂ grow again in the CLE scenario, primarily due to a strong increase in India (Klimont et al., 2013). In fact, also some other SLCPs (e.g., NO_x) show signs of a rebound about the years 2020-2025, when most of the existing policies will have been fully introduced (Klimont et al., 2015a, b). This is driven by increasing fossil fuel use and thus coupled to increasing CO₂ emissions. In the case of BC, GAINS does not predict further growth in emissions, mostly because current policies to reduce coal use in China for cooking and heating seem to be effective and because of the introduced diesel legislation.

The NFC scenario has higher SLCP emissions than the CLE scenario, showing the importance of actual introduction of already planned policies. However, the NFC scenario still

1 might be optimistic as it actually does not assume any failure or further delays in enforcement
2 of pre-2015 laws. The MIT scenario, which shows deep cuts in the emissions of some species,
3 is the result of a climate-optimized SLCP reduction scenario and is described in section 3.5.

4 Figure 2 indicates a large spread in possible future emission pathways, which for the air pollu-
5 tants is larger than anticipated in the RCP scenarios, shown by the grey shading. RCP scenar-
6 ios focused on building future emission scenarios with different radiative forcing (RF) from
7 long-lived GHGs while for air pollutants all assumed a very similar path, strongly linked with
8 the economic growth (Amann et al, 2013). Consequently all air pollutant emissions decline
9 strongly towards 2050 in all RCP scenarios. This is not the case for the ECLIPSE emissions,
10 and the spread is larger than the RCP spread despite the fact that all scenarios follow the same
11 energy use projection.

12 Emissions from international shipping differ between the ECLIPSE emission versions 4 and
13 5. Version 4a still drew on the work done for the RCP scenarios, while for the version 5 da-
14 taset, the historical emissions rely on the results of Endresen et al. (2007), with activity data
15 projected with growth rates from IEA (2012). This allowed to model region-specific regula-
16 tion, i.e., specifically in the Emission Control Areas (ECA), and long term targets to reduce
17 the sulphur content of fuels. For aviation, the emissions originate from Lee et al. (2009) and
18 are consistent with the RCP scenarios.

19 Non-agricultural, open biomass burning emissions are not calculated in the GAINS model
20 and, for the model simulations, were therefore taken from the Global Fire Emission Database
21 (GFED), version 3.1 (van der Werf et al., 2010) for the years 2008 and 2009 and held con-
22 stant in simulations of future scenarios. Biogenic emissions originate from the MEGAN data-
23 base (Guenther et al., 2012; <http://lar.wsu.edu/megan/>).

24 **3.2 Model evaluation**

25 Using the ECLIPSE version 4a CLE emissions, simulations were carried out with a range of
26 models. In addition to the four ESMs used in ECLIPSE (HadGEM3, ECHAM6-HAM2,
27 NorESM1-M and CESM1/CAM5.2; see Baker et al., 2015a for descriptions of these models),
28 three CTMs and a Lagrangian particle dispersion model were used (see Table 2). All models
29 were run for core periods in 2008 and 2009, when several aircraft campaigns took place in
30 China and the Arctic, but most models simulated the full 2008-2009 period. Some models

were also run for longer periods and were evaluated together with other models. For instance, in a comparison against aircraft measurements, Samset et al. (2014) found that the models systematically overpredict BC concentrations in the remote troposphere, especially at higher altitudes. They concluded that the BC lifetime in the models is too long. A follow-up study suggested that the best match to aircraft observations could be achieved with strongly increased BC emissions and decreased lifetimes (Hodnebrog et al., 2014). Daskalakis et al (2015) derived changes in the local lifetime of BC up to 150% associated with the use of different amounts and spatial distribution of fire emissions in the same chemistry transport model, demonstrating the dependence of BC lifetime on its emissions. Tsigaridis et al. (2014) found systematic underprediction of OA near the surface as well as a large model divergence in the middle and high troposphere. They attributed these discrepancies to missing or underestimated OA sources, the removal parameterisations as well as uncertainties in the temperature-dependent partitioning of secondary OA in the models. As a consequence of these studies, ECLIPSE models were improved in terms of emissions ([Klimont et al., 2015a, 2015b](#)), secondary OA formation ([Tsigaridis et al., 2014](#)) and removal parameterisations ([Samset et al., 2014; Hodnebrog et al., 2014](#)).

~~The improved ECLIPSE models were evaluated against global data sets such as aerosol optical depth (AOD), fine-mode AOD and absorption AOD derived from data of the Aeronet sun photometer network, as well as against various measurements of aerosol and gas-phase species (Schulz et al., 2015). Here, we focus on a more detailed regional model evaluation of the performance of the improved ECLIPSE models was made for Eastern Asia, Europe and the Arctic using satellite, airborne and ground-based measurements of pollutant gases (CO, NO₂, O₃ and SO₂) and aerosols ([Eckhardt et al., 2015](#); [Quennehen et al., 2015](#)). For Eastern Asia in August-September 2008 (Fig. 3, left two columns), data were averaged over three urban and five rural sites. The models have difficulties reproducing the urban concentrations, due to their coarse resolution. However, surprisingly most models overestimated the urban SO₂ mixing ratios. This could be related to power plant emissions that are actually occurring outside urban boundaries, being placed into the coarse urban model grid cells. For urban NO₂, models deviate less systematically from observations, with both overestimates and underestimates, thus and the model mean captures the observations. For rural NO₂, also the individual models deviate less from the measured concentrations, indicating that the individual model biases for urban NO₂ are very likely mainly due to the limited model resolution and not to~~

1 biases in emissions and/or chemical processes. The measured concentrations of NO_2 and O_3 at
2 the rural sites are matched relatively well (agreement within the range of the temporal distri-
3 bution at individual sites) but SO_2 is generally overestimated there as well. The most severe
4 problem at rural sites, however, is a systematic underestimation of CO mixing ratios, which
5 was attributed to underestimated CO lifetimes in the models (Quennehen et al., 2015).

6 A similar comparison was made for Europe with background measurements taken from sta-
7 tions of the European Monitoring and Evaluation Programme (EMEP) (Fig. 3, right two col-
8 umns for winter and summer; see also Schulz et al., 2015). Overall, over Europe the
9 ECLIPSE model mean captures the mean observations with the exception during summer for
10 CO that is underestimated (as in Asia). Summertime O_3 is overestimated by many models at
11 rural locations over Europe and Asia suggesting too much photochemical production down-
12 wind of emission regions.

13 Satellite-derived aerosol optical depth (AOD) measurements were reproduced quite well by
14 the models over China and Europe (Fig. 4). Evaluation of individual aerosol components over
15 Asia (Quennehen et al., 2015) shows an overestimation of the ECLIPSE model-mean surface
16 BC in urban China in summer 2008, which is probably due to the short-term mitigation
17 measures taken during the Olympic Games. Over Europe, ECLIPSE models satisfactorily
18 simulate surface BC observations both in winter and summer (Fig. 4). However, problems
19 were identified over India: Gadhavi et al. (2015) found that BC concentrations are strongly
20 underestimated in Southern India even when aerosol removal processes in one model were
21 completely switched off in the region. Furthermore, observed AOD values in Northern India
22 are larger than simulated by all but two of the ECLIPSE models (Fig. 4). This suggests that
23 the emissions of BC and precursors of other aerosols are underestimated for India in the
24 ECLIPSE emission data set. This could be related to the rapid recent growth of emissions in
25 India (Klimont et al., 2013), which may be underestimated in the inventories, as well as with
26 problems capturing the true spatial distribution of emissions in India.

27 The Arctic was shown previously to be a particularly challenging region for aerosol model
28 simulations (e.g., Shindell et al., 2008). Aerosol loadings in the Arctic are generally much
29 lower than in populated regions and the Arctic encompasses only a small fraction of the Earth.
30 Therefore, impacts of even large relative errors in the modelled aerosol concentrations in the
31 Arctic on global radiative forcing and global climate response are relatively small. Neverthe-

1 less, identification of model biases in this remote region is important as it can lead to im-
2 proved process understanding, especially of the aerosol removal mechanisms. An evaluation
3 of the ECLIPSE models over the Arctic was coordinated with the Arctic Monitoring and As-
4 sessment Programme (AMAP, 2015). Comparisons were made for BC and sulphate for six
5 ground stations and during six aircraft campaigns (Eckhardt et al., 2015). As an example, a
6 comparison of the BC concentrations simulated by the ECLIPSE models with measured
7 equivalent BC is shown in Fig. 5 for the stations Zeppelin on Svalbard, Pallas in Finland and
8 Tiksi in Siberia. For Zeppelin, most models clearly underestimate the observed concentrations
9 during winter and spring, whereas for Pallas which is closer to source regions, the models
10 tend to overestimate. In general, the model performance (also at other Arctic sites, not shown)
11 is better than in previous comparisons (e.g., Shindell et al., 2008). However, very large model
12 underestimates were found for Tiksi, from where measurement data have only recently be-
13 come available. Another ECLIPSE study showed that also the snow BC concentrations are
14 generally underestimated by models in Northern Russia but overestimated elsewhere in the
15 Arctic (Jiao et al., 2014). It is therefore likely that the model underestimates are caused by too
16 low BC emissions in Russia in the ECLIPSE CLE data set. Yttri et al. (2014) attribute this at
17 least partly to an underestimation of residential wood burning, based on levoglucosan meas-
18 urements made at Zeppelin. Eckhardt et al. (2015) suggest that also SO₂ emissions in North-
19 ern Russia are underestimated. ECLIPSE models participating in the AMAP (2015) assess-
20 ment also showed a systematic underestimation in CO concentrations in the Arctic and lack of
21 model skill in simulating reactive nitrogen species important for O₃ production.

22 An important finding of the model-measurement comparisons is that overall the ESMs show a
23 similar performance as the CTMs. This is encouraging for the further use of the ESMs for
24 determining the climate impacts (section 3.6). The comparisons led to some further improve-
25 ments of the ECLIPSE emissions for version 5, prior to their use for transient climate model
26 simulations. For instance, wick lamps were identified as an important emission source in In-
27 dia, the inclusion of which improved the agreement with the observations in a model sensi-
28 tivity study (Gadhavi et al., 2015). Other enhancements (e.g., re-gridding of non-ferrous smel-
29 ter emissions to improve SO₂ emissions in Russia as suggested by Eckhardt et al. (2015))
30 came too late for the climate impact studies and were only made in version 5a.

31 Another aspect of model evaluation is to determine the capability of models to reproduce past
32 trends, and this was tested over Europe. Strong reductions of aerosol emissions occurred over

1 Europe, since the 1980s due to air quality legislation in Western Europe, and since the early
2 1990s due to economic restructuring in Eastern Europe. This emission reduction is manifest
3 e.g. in strongly increasing trends in surface solar radiation (“solar brightening”) and visibility
4 (Stjern et al., 2011), but also in a stronger warming trend compared to the earlier period in
5 which aerosol emissions increased (Cherian et al., 2014). The “historical” simulations con-
6 tributed to the 5th Coupled Model Intercomparison Project (CMIP5, Taylor et al. 2012) using
7 previous versions of the ECLIPSE ESMs were assessed for continental Europe, and compared
8 to observations from the Global Energy Balance Archive (Gilgen et al., 1998) and the [Climate](#)
9 [Climatic](#) Research Unit (CRU) of the University of East Anglia (CRU-TS-3.10, Mitchell and
10 Jones, 2005). The 1960-1980 period shows a strong “solar dimming” (reduction in surface
11 solar radiation) and small warming, since the greenhouse-gas-induced warming is offset by
12 the aerosol forcing. The period 1990-2005, in turn, shows the solar brightening, and a much
13 stronger warming. All three tested models are able to reproduce this strong increase in warm-
14 ing trend to within their uncertainties (Fig. 6), suggesting that the climate response to aerosol
15 changes is captured [despite the masking influence of natural climate variability on these](#)
16 [trends. However, the absolute amplitude of the trends is not equally well captured by all mod-](#)
17 [els, indicating that the skill of the ECLIPSE \(and other\) ESMs to simulate temperature trends](#)
18 [responding to changing aerosol emissions is limited. This is due to both limitations in the](#)
19 [models themselves, the emission input, as well as the influence of natural climate variability.](#)

20 **3.3 Radiative forcing**

21 To provide input to the metrics needed for designing a mitigation scenario, dedicated model
22 simulations by three ESMs (ECHAM6-HAM2, HadGEM3, NorESM) and a CTM
23 (OsloCTM2) were used to establish a matrix of specific RF (Bellouin et al., 2015) by season
24 and region of emission. Specific RF (SRF) is defined as the RF per unit change in emission
25 rate once the constituents have reached equilibrium and is given in $\text{mW m}^{-2} (\text{Tg yr}^{-1})^{-1}$. To
26 estimate SRF, the emissions of eight short-lived species (BC, OA, SO_2 , NH_3 , NO_x , CO, CH_4
27 and NMVOCs) were reduced by 20% compared to their ECLIPSE baseline. These species
28 cause RF themselves and/or lead to the perturbation of radiative forcers (e.g., O_3). The re-
29 gional reductions were made for Europe and China, as well as for the global shipping sector
30 and for a rest-of-the-world region. To account for seasonal differences in SRF, separate reduc-
31 tions were applied for May-October and November-April. Henceforth we will refer to these as

1 Northern Hemisphere (NH) “summer” and “winter”. Notice that in our case the sign of SRF is
2 opposite to that of RF, because the imposed emission changes are negative. A reduction of a
3 warming species gives negative RF values but positive SRF values. It is important to note that
4 SRF excludes rapid adjustments in the atmosphere, with the exception that BC semi-direct
5 effects were calculated explicitly, and stratospheric temperature adjustments were included
6 for O₃ and CH₄.

7 Models generally agreed on the sign of RF and the ranking of the efficiency of the different
8 emitted species, but disagreed quantitatively (see Bellouin et al., 2015 for details). The best
9 estimate of a species’ RF was considered to be the average of all models, with the model
10 spread indicating its uncertainty. However, not all models have calculated RF for all species
11 or have accounted for all processes. For instance, all models were able to quantify the aerosol
12 direct effect but only three quantified the 1st indirect effect. For BC aerosols only one model
13 quantified the snow albedo effect and the semi-direct effect explicitly. Therefore, mean RF
14 values were determined by averaging across all available models for each process separately.
15 In most cases, all four models were available for this, but for some processes fewer models
16 had to be used.

17 Figure 7 shows the resulting SRF for reductions in the emissions of SO₂, NO_x, CH₄ and BC
18 and the processes contributing to the total forcing, for Europe, China, and on global average.
19 The globally-averaged SRF was obtained by adding RF for Europe, China, and Rest of the
20 World, then normalising to global emission change. The SRF values are largest for BC but
21 note that global emissions of BC are smaller than for the other species. In addition, the semi-
22 direct effect of BC potentially offsets a considerable fraction of the aerosol direct RF and RF
23 due to deposition on snow. Quantifying the semi-direct effect has large uncertainties, howev-
24 er, because internal variability of the climate system masks tropospheric adjustments to BC
25 perturbations. This means that the sign of total SRF exerted by decreases in BC emissions
26 may be negative if a weak BC direct effect is more than compensated by a strong semi-direct
27 effect. Nevertheless, the ECLIPSE BC SRF best estimate of about 50 mW m⁻² (Tg[C] yr⁻¹)⁻¹
28 when semi-direct effects are included is not an outlier compared to previous estimates, which
29 range from 24 to 108 mW m⁻² (Tg [C] yr⁻¹)⁻¹ according to Table 23 of Bond et al. (2013).
30 Moreover, ECLIPSE simulations indicate that the magnitude of the semi-direct effect is
31 smaller than the direct aerosol effect (Hodnebrog et al., 2014; Samset and Myhre, 2015), in
32 agreement with most, but not all, previous studies (again, see Table 23 of Bond et al. (2013)).

1 Reductions in the emissions of light scattering aerosols such as sulphate stemming from its
2 precursor SO_2 induce a negative SRF. The RF values of aerosols are generally larger for
3 summer emissions than for winter emissions because of the stronger insolation. However,
4 there are exceptions to this. For instance, the BC deposition on snow is more effective for
5 winter emissions because of the larger snow extent in winter and spring and partial preserva-
6 tion of deposited BC into spring. Aerosol SRF is also larger in magnitude for Europe than for
7 China, most likely because of different cloud regimes which are differently affected by semi-
8 direct and indirect aerosol effects.

9 For NO_x , SRF is uncertain because decreases in NO_x emissions perturb tropospheric chemis-
10 try in two opposite ways, working on different timescales: First, they reduce tropospheric O_3
11 concentrations, thus exerting a positive SRF. Second, they increase CH_4 concentrations, thus
12 exerting a negative SRF, with an additional CH_4 -induced change in O_3 . ECLIPSE accounts
13 for those two pathways, and quantifies a third, whereby reductions in NO_x emissions suppress
14 nitrate aerosol formation and its associated RF. ECLIPSE is therefore able to state-~~confirm~~
15 with confidence the earlier quantification (Myhre et al., 2013a) that NO_x exerts a negative
16 SRF, because the O_3 response is not sufficient to offset the combined CH_4 and nitrate re-
17 sponse. For CH_4 , ECLIPSE finds a relatively large range of SRF estimates from the models,
18 reflecting the differences in methane lifetime and methane's effects on ozone and aerosols.
19 This range does not reflect uncertainties in the absorbing properties of CH_4 . Rather, the range
20 reflects the different CH_4 lifetimes in the models—and the changes in these lifetimes—and
21 hence the sensitivities of the burdens to emission changes.

22 The BC radiative forcing is very uncertain and needs some discussion. When scaled to 100%
23 BC reductions and reported annually, ECLIPSE BC total RF and its range are 0.2814 (0.020-
24 0.4621) W m^{-2} . Neglecting semi-direct effects, those numbers become 0.4120 (0.0711-0.248)
25 W m^{-2} . These values are consistent with the stronger end of the multi-model ACCMIP (At-
26 mospheric Chemistry and Climate Model Intercomparison Project) estimate of $0.24 \pm$
27 0.1 W m^{-2} (Shindell et al., 2013) and the AeroCom estimate of $0.18 \pm 0.07 \text{ W m}^{-2}$ (Myhre et
28 al., 2013b). So in the context of global modelling, ECLIPSE BC forcing estimates are repre-
29 sentative of both magnitude and range. In contrast, ECLIPSE estimates are at the middle or
30 weak end of the relatively large values estimates that were reported in several recent assess-
31 ments: 0.45 (0.30-0.60) W m^{-2} in UNEP/WMO (2011), 0.71 (0.08-1.27) W m^{-2} in Bond et al.
32 (2013), and 0.40 (0.08-0.8) W m^{-2} in Boucher et al. (2013). The estimates in UNEP/WMO

(2011) are skewed towards higher values by accounting for semi-empirical estimates, which include a brown-carbon contribution, the possibility that BC semi-direct forcing is the same sign as direct forcing (even though semi-direct forcing opposes the sign of direct forcing in most models) and assuming a high efficacy of BC deposition forcing. The large observationally-constrained estimates of BC forcing by Bond et al. (2013), which influenced the IPCC estimate (Boucher et al., 2013) are due to strong scaling using Aeronet absorbing AOD measurements. However, these measurements are highly uncertain and probably biased and the resulting high BC RF values have recently been called into question by several studies (Wang et al., 2014; Samset et al., 2014; Wang et al., 2015). There are several reasons for this. Firstly, the scaled models have substantially higher BC abundances globally than supported by BC measurements. Wang et al. (2014) and Samset et al. (2014) argue that BC residence time has to be relatively short to fit observed remote BC concentrations. Furthermore, Wang et al. (2015) showed that the fairly low resolution of global models induces an artificial negative bias when comparing to AERONET stations in Asia. Consequently, in ECLIPSE we have not scaled simulated RF values for BC but have used native model results.

3.4 Climate metrics

ECLIPSE explored various options for climate metrics. In addition to selecting the most useful metric for designing the mitigation scenario, the project also made conceptual developments. Collins et al. (2013) developed further the application of Regional Temperature change Potential (RTP) presented by Shindell et al. (2012) by accounting for the location of emissions, thereby opening for regionality for both *drivers* and *responses*. Collins et al. (2013) also expanded the GTP concept by tentatively including the climate-carbon feedback for the non-CO₂ gases. So far, this feedback has only been included for the reference gas CO₂ which means that GTPs (and GWPs) tend to underestimate the relative effect of non-CO₂ components (Myhre et al., 2013a). Furthermore, Shine et al. (2015) present a new metric named the Global Precipitation change Potential (GPP), which is designed to gauge the effect of emissions on the global water cycle. Of particular relevance for SLCPs is their demonstration of a strong near-term effect of CH₄ on precipitation change and the role of sustained emissions of BC and sulphate in suppressing precipitation.

1 Aamaas et al. (2013) investigated several different metrics and showed that emissions of CO₂
2 are important regardless of what metric and time horizon is used, but that the importance of
3 SLCP varies greatly depending on the metric choices made. MacIntosh et al. (2015) consid-
4 ered the errors made when calculating RF and climate metrics from multi-model ensembles in
5 different ways. They showed that the mean metric values are relatively robust but the estima-
6 tion of uncertainties is very dependent on the methodology adopted. Finally, Lund et al.
7 (2014a) applied climate metrics to quantify the climate impacts of BC and co-emitted SLCPs
8 from on-road diesel vehicles, and Lund et al. (2014b) considered the special case of a fuel
9 switch from diesel to biodiesel.

10 As explained in section 3.5, climate metrics were used in ECLIPSE to identify specific sets of
11 air pollution reduction measures that result in net positive climate effects (i.e., reduced warm-
12 ing), considering the impacts on all co-controlled substances. Based on the RF results shown
13 in section 3.3, Aamaas et al. (2015) calculated regional and seasonal GTP and GWP metrics
14 for the SLCP emissions, for various time horizons, and explored their properties. Of all the
15 explored metrics, the pollution control analysis was carried out for GWP₁₀₀ and GTP₂₀, as
16 these two metrics showed large differences in their quantifications. It was found, however,
17 that the emerging basket of emission control measures was very similar for both, although the
18 ranking of the potential climate impacts of individual measures was sometimes different, with
19 larger effect of CH₄-related measures for the GWP₁₀₀ metric (due to the higher value this met-
20 ric with longer time horizon assigns to CH₄).

21 In the following, we concentrate our analysis on the GTP₂₀ metric, which is shown in Fig. 8
22 for a pulse emission relative to an equal mass emission of CO₂ for a selected number of spe-
23 cies. The metric builds on and reflects important aspects of the RF forcing values shown in
24 Fig. 7. For example, for most species GTP₂₀ values for summer are larger than values for win-
25 ter (see, e.g., results for SO₂), and values for Europe are larger than values for Asia. While
26 SO₂ only has negative values (i.e., reduction of SO₂ always leads to warming), the opposite is
27 true for CH₄. Fig. 8 also shows the contribution from different processes to the total GTP₂₀
28 metric, and for BC and NOx they are often of different sign and their magnitude strongly de-
29 pends on the emission region and season. While all of this was accounted for in GAINS (see
30 section 3.5), it is clear that this makes the choice of mitigation options more uncertain, and
31 this is further complicated by the fact that BC and NOx emissions are always associated with

1 co-emissions of other SLCPs (e.g., OA). The GTP₂₀ metric was determined also for all other
2 SLCPs (not shown).

3 For the implementation of mitigation measures, a pulse emission metric is not very realistic,
4 as measures are normally introduced gradually, become more effective with time, and are
5 usually maintained indefinitely. Therefore, we also considered versions of the GTP and GWP
6 metrics for sustained emission measures and a linear ramp-up of emission measures over a
7 15-year time period (Aamaas et al., 2015; see also Boucher and Reddy, 2008). We chose the
8 ramp-up version of the GTP₂₀ metric for designing our mitigation scenario with the GAINS
9 model (see section 3.5), as it most realistically reflects the implementation of mitigation
10 measures. Notice that the ramp-up version of the GTP₂₀ metric can be derived directly from
11 its pulse emission version, by linear combination of emission pulses (see Aamaas et al., 2015,
12 for details). Fig. 8 shows the pulse emission version because pulse emissions are building
13 blocks for various versions of the metrics, including the ramp-up version.

14 To go beyond global mean temperature, it is also possible to calculate regional surface tem-
15 perature changes using the regional temperature potentials (RTP) concept of Shindell and
16 Faluvegi (2009). This concept maps RF in one given latitude band to the temperature re-
17 sponse in several other latitude bands and was adopted by us (Aamaas et al., paper in prepara-
18 tion). For the mapping, we used the pre-calculated RTP-coefficients of Shindell and Faluvegi
19 (2009). Even though the coefficients are likely model-dependent, we had to use these values
20 because they are not available from any other model (and specifically not from the ECLIPSE
21 models). The coefficients also seem fairly robust in comparison with the response to historical
22 aerosol forcing in several other models (Shindell, 2012). We made two additions, however,
23 for BC in the Arctic and BC on snow using the method of Lund et al. (2014a) with results
24 from Samset et al. (2013) and Flanner (2013). Using the RTP concept and RF calculations
25 from section 3.3 (Bellouin et al., 2015), we calculated absolute regional temperature change
26 potentials (ARTP) for the set of components, regions and seasons for which ECLIPSE has
27 determined RF values (Fig. 7). Following Collins et al. (2013), the ARTPs were used to esti-
28 mate transient surface warming using an impulse response function for temperature response
29 from Boucher and Reddy (2008) (as was also used for the GTP calculations). Details of the
30 method and ARTP values will be given in Aamaas et al. (paper in preparation) and results of
31 the ARTP calculations will be used in section 3.7 for comparisons with transient ESM runs.

3.5 Emission mitigation and air quality impacts

For designing a climate-optimized SLCP emission mitigation scenario, the CO₂-equivalent SLCP emissions were minimized using the GAINS model. For this, the numerical values of the GTP₂₀ metric for the final year 2035 for each species, region and season were implemented into GAINS. The ramp-up version of this metric assumes a linear implementation of mitigation measures between the years 2015 and 2030 and thereafter a full implementation. Mitigation measures typically affect several species at the same time. For instance, controlling BC emissions leads to a “co-control” of OA and other species. The GTP₂₀ metric values are different for all of these species and can be of different sign, and it is important to determine the net impact of a mitigation measure across all affected species. In practice, the species-specific metric values were weighted with the emission factors to obtain the net metric value for each of the ~2000 mitigation measures in every one of the 168 regions considered in GAINS, and for summer and winter periods separately. These measures were subsequently ranked according to their CO₂-equivalent total net climate impact as measured by the GTP₂₀ metric, and all measures with a beneficial climate and air quality impact were included in the mitigation (MIT) scenario basket (Klimont et al., 2015b); this approach is consistent with that taken by UNEP/WMO (2011) and Shindell et al. (2012).

The mitigation basket contains three groups of measures: 1) Measures that affect emissions of CH₄ that can typically be centrally implemented (e.g., by large energy companies, municipalities, etc.) and which also impact background O₃; 2) Technical measures that reduce the emissions of BC, mainly for small stationary and mobile sources; 3) Non-technical measures to eliminate BC emissions, e.g., through economic and technical assistance to the poorest population. While about 220 GAINS model measures were collected in the mitigation basket, they were merged into representative measure groups. For example, high emitters are calculated for each of the GAINS transport subsectors and fuels while here removing high emitters is represented by one ‘measure’. Similarly for cooking stoves, GAINS estimates mitigation potential for various types of fuels but all of these are included further into one category ‘clean cooking stoves’. Also for CH₄, losses from gas distribution are calculated for several GAINS end use sectors while here the mitigation potential is reported under one measure where leaks from low pressure pipelines are reduced. Finally, for NMVOCs all of the solvent related options in GAINS (over 50) are categorized as one measure reducing losses from solvent use activities. Considering the above, about 50 ‘measures’ represent the about 220 GAINS op-

1 tions that were included in the mitigation basket. The 17 most effective measures contribute
2 80% of the total climate benefit, according to the GTP₂₀ metric, with CH₄ measures contrib-
3 uting about 47% and BC-focused measures contributing 33%. These measures are listed in
4 Table 3. It is interesting to notice that the top CH₄ and BC-focused measures both concern the
5 oil and gas industry and specifically the venting or flaring of associated gas.

6 As can be seen in Fig. 2, the mitigation has only minor effects on CO₂ emissions, but reduces
7 most SLCPs strongly compared to the CLE scenario. By 2030, CH₄ emissions are reduced by
8 about 50% and BC emissions by nearly 80%. OA is co-controlled with BC, causing a nearly
9 70% (not shown) reduction of its emissions, as for some sectors BC outweighs the cooling
10 effects of OA. While NO_x emission reductions are in most cases also not preferred by the
11 GTP₂₀ metric (see Fig. 8), reductions stem from the co-control when higher Euro standards
12 are introduced; they reduce significantly several pollutants such as BC, CO, NMVOC and
13 also NO_x (Klimont et al., 2015a, b). By contrast, SO₂ emissions are nearly the same in both
14 the CLE and MIT scenarios, as for the key sectors emitting SO₂, the warming by SO₂ reduc-
15 tions (see Fig. 8) cannot be outweighed by co-control of species whose reduction would lead
16 to cooling. Thus, SO₂ reductions are largely avoided.

17 The global CO₂-equivalent emissions (calculated using the GTP₂₀ metric values) are shown as
18 a function of time in Fig. 9. Values are shown for CLE and MIT and are split into contribu-
19 tions from CH₄ and other SLCP emissions. For comparison, CO₂ emissions are also shown.
20 On the short time scale of the GTP₂₀ metric, CH₄ and CO₂ emissions are nearly equally im-
21 portant in the CLE scenario. The CO₂-equivalent emissions of CH₄ are, however, reduced by
22 50% in MIT. The CO₂-equivalent emissions for the other SLCPs are negative in both scenari-
23 os, indicating that in total they have a cooling impact. As the mitigation reduces preferentially
24 warming components, this cooling becomes even stronger in the MIT case. The total CO₂-
25 equivalent emissions (including CO₂ emissions) are reduced substantially in the MIT scenario
26 (blue shaded area in Fig. 9 shows the reduction), for example by about 70% in the year 2030.
27 About 56% of this reduction is due to CH₄ measures and ~44% is due to other measures. It is
28 important to notice that the effect of the SLCP mitigation is relatively large for the rather
29 short time horizon of the GTP₂₀ metric; it would be smaller for a longer time horizon. Similar-
30 ly, the relative importance of CH₄ compared to other SLCP emissions would be increased for
31 a longer time horizon.

1 Surface concentrations of SLCPs resulting from the CLE and MIT scenarios were determined
2 from the various model simulations. Fig. 10 shows maps of the model-mean relative differ-
3 ences for O₃ and PM_{2.5} for the last decade (2041-2050) of the transient simulations and Ta-
4 ble 4 reports differences for several regions shown with boxes in Fig. 10. Concentrations of
5 O₃ (Fig. 10, upper panel) are reduced globally, with reductions of more than 12% in most of
6 the Northern Hemisphere and the strongest reductions of up to about 20% occurring in East
7 Asia. For instance, in Eastern China (see Table 4), O₃ is reduced by 19.3% (16.0-24.4%). BC
8 and OA concentrations (not shown) were also globally reduced, with BC reductions reaching
9 more than 80%. For sulphate (not shown), the relative changes are much smaller than for BC
10 and OA and both increases and decreases occur - a consequence of the relatively small global
11 SO₂ emission reductions (see Fig. 2). Changes in PM_{2.5} concentrations (Fig. 10, lower panel)
12 are smaller because of large contributions from natural sources (e.g., sea salt, dust, wildfire
13 emissions). PM_{2.5} concentrations in the SLCP source regions were reduced by typically 10-
14 20% and up to nearly 50% in smaller regions. Reductions are strongest in Asia, for instance
15 19.8% (17.9-22.5%) in India (Table 4).

16 In summary, air pollutant concentrations in the MIT scenario are dramatically reduced com-
17 pared to the CLE scenario, especially in the polluted (and heavily populated) source regions.
18 This indicates the beneficial effect of the SLCP mitigation on air quality. Nevertheless, a
19 word of caution is needed for O₃. The O₃ concentrations increase strongly (typically between
20 5 and 20%, depending on region and season) between now and 2050 in the CLE scenario,
21 because of increasing CH₄ and NO_x emissions. Therefore, global mean O₃ concentrations
22 even in the MIT scenario do not decrease substantially with time. However, strong relative O₃
23 reductions by the mitigation are simulated in the SLCP source regions (Table 4), which more
24 than outweigh the overall concentration increase in the CLE scenario. Therefore, population
25 exposure to O₃ decreases with time in the MIT scenario.

26 For Europe and Asia, the GAINS model also contains source-receptor relationships which
27 allow the estimation of the impacts of emissions on human health. GAINS quantifies the im-
28 pacts of changes in SLCP emissions on the long-term population exposure to PM_{2.5} in Europe,
29 China and India and estimates the resulting premature mortality, in terms of reduced statistical
30 life expectancy and cases of premature deaths (Amann et al. 2011). Calculations follow the
31 recommendations of the findings of the WHO review on health impacts of air pollution and
32 recent analyses conducted for the Global Burden of Disease project (Lim et al. 2012), relying

1 on the results of the American Cancer Society cohort study (Pope et al., 2002) and its re-
2 analysis (Pope et al., 2009). It uses cohort- and country-specific mortality data extracted from
3 life table statistics to calculate for each cohort the baseline survival function over time. No-
4 Notice, however, that, in contrast to the changes of pollutant concentrations presented in Fig. 10,
5 the quantification of human health impacts is based on results from a single model.

6 Using GAINS, we estimate that in the EU the loss of statistical life expectancy will be re-
7duced from 7.5 months in 2010 to 5.2 months in 2030 in the CLE scenario. The ECLIPSE
8 mitigation measures (MIT) would reduce statistical life shortening by another 0.9 months
9 (Fig. 11, upper panel), resulting in 4.3 months of reduction in life expectancy. This value is
10 only slightly above the target of 4.1 months that has been set by the European Commission in
11 its 2013 Clean Air Policy proposal (EC 2013). Population in non-EU countries would gain
12 approximately one month life expectancy from the implementation of the ECLIPSE measures
13 in 2030 (Fig. 11, upper panel).

14 In China and India, the potential health gains from the implementation of the ECLIPSE
15 measures are significantly larger, however, starting from a substantially higher level of life
16 shortening due to PM_{2.5} (Fig. 11, lower panel). In China, the ECLIPSE measures would in the
17 year 2030 extend the life expectancy of the population by approximately 1.8 months and re-
18 duce the premature deaths attributable to PM_{2.5} by 150,000 – 200,000 cases per year.

19 In India, rapid increase in energy consumption, together with lacking regulations on emission
20 controls for important sources (e.g., power generation) and poor enforcement of existing laws
21 (e.g., for vehicle pollution controls) will lead to a steep increase in PM_{2.5} levels. If no satura-
22 tion of health impacts is assumed for such high levels (there are no cohort studies available
23 for such high concentrations), with conservative assumptions GAINS estimates approximately
24 850,000 cases annually of premature deaths from air pollution in 2010. For 2030, PM_{2.5} expo-
25 sure would increase by more than 50%, and at the same time the population would increase
26 and age. Combined, these factors would let the premature deaths from air pollution grow by
27 approximately 125% to 1.9 million cases in 2030, with another doubling to 3.7 million cases
28 in 2050. Against this background, the ECLIPSE measures would avoid more than 400,000
29 cases of premature deaths in 2030 and almost 700,000 in 2050. Using the loss in statistical life
30 expectancy as an alternative metric, the ECLIPSE measures would gain 11-12 months to the
31 Indian population (Fig. 11, lower panel).

1 **3.6 Climate impacts**

2 The climate impacts of SLCPs were determined with four ESMs (HadGEM3, NorESM,
3 ECHAM6-HAM2/MPIOM, and CESM-CAM4) in two different experiments. In the first ex-
4 periment, still part of the outer loop of the spiral in Fig. 1, all land-based anthropogenic emis-
5 sions of each of SO₂, OA and BC were removed one at a time and the models were run to
6 *equilibrium*; in the other experiment, the mitigation scenario described in section 3.5 was fol-
7 lowed in a series of *transient* ensemble model runs, constituting the inner spiral loop in Fig. 1.

8 Note that all ESMs implicitly include the semi-direct effect, while for the radiative forcing
9 calculations this was calculated explicitly by only one model.

10 For the first experiment, ~~described in Baker et al. (2015a)~~, the four ESMs, with full ocean
11 coupling were run for a control simulation and a perturbation run for 50 years, after a spin-up
12 period to equilibrium. The control simulation used ECLIPSE V4a emissions for the year 2008
13 (except for CESM-CAM4, which used year 2000 emissions). For the perturbation, 100% of
14 the land-based emissions of the three individual species were removed in turn to achieve dis-
15 cernible climate responses. While only three ESMs ran the experiments for SO₂ and OA, all
16 four ESMs ran the BC experiment and three of these used two or three ensemble members
17 each. Only NorESM included the effect of albedo reduction by BC deposited on snow and
18 ice. This experiment is described in detail in Baker et al. (2015a), where more detailed de-
19 scriptions of the models used can also be found. Here, we provide a synthesis of the results.

20 When removing SO₂ emissions, all three models show an increase of global mean surface
21 temperature (Fig. 12a) by 0.69 K (0.40-0.84 K) on average. Here, the first value is the multi-
22 model mean, whereas the values in brackets give the full range of results obtained with the
23 individual models. We will keep this notation throughout the rest of the paper, unless other-
24 wise noted. The zonal mean temperature change is positive at all latitudes and increases with
25 latitude in the Northern Hemisphere, reaching 2.46 K (1.38-3.31 K) at the North Pole (Fig.
26 13a). It is also positive for most regions of the Earth and is more positive over the continents
27 than over the oceans (Baker et al., 2015a). The models also agree that removing SO₂ emis-
28 sions results in an increase in global mean precipitation (Fig. 12b), which is in line with the
29 expected impact from a global temperature increase. The precipitation increases are particu-
30 larly strong over India and China because of a northward shift of the Intertropical Conver-

1 gence Zone (ITCZ, see Fig. 13b), which is in accordance with previous studies (e.g. Broccoli
2 et al., 2006).

3 The response to removing anthropogenic BC emissions, -0.05 K (-0.15 to $+0.08$ K), is much
4 smaller than the SO_2 response and the models do not all agree on the sign of the global mean
5 response. The multi-model mean is slightly negative (Fig. 12a) but within ± 0.5 K everywhere
6 on the globe (not shown) and the zonal mean temperature response differs from model to
7 model (Fig. 13c). The NorESM model shows the strongest cooling in the Arctic, likely be-
8 cause it is the only model accounting for snow albedo changes. Precipitation changes from
9 removing BC emissions are also small, but consistently positive in all models (Fig. 12b) de-
10 spite the cooling in most models. This is consistent with calculations of the relationship be-
11 tween atmospheric RF and precipitation change in Andrews et al. (2010) and Kvalevåg et al.
12 (2013) (see also Shine et al., 2015). The multi-model temperature response to removing OA
13 emissions is similar to that for removing SO_2 , but much weaker overall (Fig. 12a and
14 Fig. 13e).

15 In summary, the emission perturbation studies show that elimination of anthropogenic SO_2
16 emissions leads to robust warming, elimination of OA also leads to – albeit much weaker –
17 warming, whereas elimination of BC leads to a small temperature response with substantial
18 differences between the models but slight cooling in the multi-model mean. This could be due
19 to the different sizes of the indirect and semi-direct effects of BC in different models, shown
20 in ECLIPSE (Hodnebrog et al., 2014), and possibly to unforced responses of climate system
21 components, especially sea ice, that happen to counteract the small temperature response.

22 The second ESM experiment simulated the *transient* responses to the climate-optimized
23 SLCP mitigation of section 3.5 (Baker et al., 2015b). For this, the four ESMs ran three en-
24 semble members each for both the CLE and MIT scenario until the year 2050. The CLE sce-
25 nario resulted in an increase of global mean surface temperature of 0.70 ± 0.14 K (the value
26 following the “ \pm ” sign gives the standard deviation obtained from all ensemble members)
27 between the years 2006 and 2050 in the multi-model ensemble mean. A large part of this in-
28 crease is a response to increasing CO_2 concentrations, and consequently the MIT scenario also
29 showed an (albeit smaller) temperature increase. As we here are mainly interested in the re-
30 sponse to the SLCP mitigation, we only consider the difference between the MIT and the CLE
31 scenario in the following. Time series of the global mean temperature difference between the

1 two scenarios are shown in Fig. 14. There is a considerable spread between individual ensemble
2 members even from the same model. This reflects simulated natural climate variability
3 superimposed on the response to the SLCP mitigation, which makes diagnosis of the latter
4 difficult. Systematically negative global mean temperature responses emerge only when the
5 mitigation measures are fully implemented. The multi-model mean global temperature (MIT-
6 CLE) difference is -0.22 ± 0.07 K for the final 10 years, confirming that the mitigation could
7 be successful in reducing the warming of the CLE scenario.

8 The relative cooling is particularly strong over the continents and weakest over the Northern
9 North Atlantic (not shown), similar to the response patterns seen in the perturbation simulations.
10 Figure 15 shows mean temperature responses for the last decade of the simulation, for
11 various regions. The strongest relative cooling between MIT and CLE of about 0.44 K (0.39-
12 0.49 K) are found for the Arctic, with peak values of about 0.62 K (0.37-0.84 K) occurring in
13 autumn (winter values are similar). Over Europe, differences are more consistent between
14 models and more strongly negative in the southern parts than in the northern parts (see also
15 Fig. 16). This can be explained by the larger natural climate variability in Northern Europe.
16 Mainly small and inconsistent results are found over India, due to model differences in the
17 shift of the ITCZ, whereas changes over China and North America are consistently negative
18 from (almost) all models and for all seasons.

19 The precipitation responses are less robust which, in the tropics, is due to model differences in
20 the migration of the ITCZ. Nevertheless, there are regions with consistent responses of all
21 models. Of particular interest is the precipitation increase over Southern Europe (Fig. 16).
22 Seen against the background of expected warming and drying in the Mediterranean area due
23 to CO₂-driven climate change, the precipitation increase due to the SLCP mitigation would be
24 beneficial, especially given the fact that it is strongest from spring to autumn (Fig. 17), with
25 15 (6-21) mm/yr increase (corresponding to more than 4% (2-6%) of total precipitation) and
26 combined with a temperature reduction that is also largest during that period (Fig. 15). Thus,
27 our mitigation approach would help to alleviate drought and water shortages in the Mediterranean
28 area in summer, as they are expected for the future (Orlowsky and Seneviratne, 2012).

29 In the aerosol perturbation experiments, we have seen that SO₂ emission reductions lead to
30 strong warming and OA reductions to weaker warming, whereas BC emission reductions lead
31 to much weaker and model-dependent cooling. While the GTP₂₀ metric attributes ~44% of the

1 reduction in CO₂-equivalent emissions to BC-related measures, it was interesting to clarify
2 whether this is reflected in the transient climate model simulations, and how this fraction
3 changes with time. For this, two additional experiments with four ensemble members each
4 were run with one ESM (CESM-CAM4) with a slab ocean representation (for greater compu-
5 tational efficiency): one with all emission reductions (MIT), and one with only CH₄ emission
6 reductions (MIT-CH₄-only). Ozone reductions resulting from CH₄ emission measures are in-
7 cluded in the MIT-CH₄-only simulations, whereas O₃ changes from non-CH₄ measures are
8 omitted. The two experiments yielded a relatively similar temporal characteristic of global
9 mean temperature difference (see Fig. 18), where the MIT scenario led to 0.45±0.04 K less
10 warming than CLE, and the MIT-CH₄-only scenario led to 0.41±0.04 K less warming than
11 CLE for the final decade of the simulation (2041-2050). The difference between the two sce-
12 narios is not significant, owing to the small number of ensemble members. Thus, the simula-
13 tions indicate a dominant (~90%) role of CH₄ emission reductions and a small (~10%) but
14 perhaps non-negligible role of non-CH₄ mitigation measures for reducing the warming by
15 2041-2050.

16 Notice that the warming reduction for the MIT scenario with the slab-ocean version of the
17 CESM-CAM4 model is twice as strong as with the more realistic full-ocean version shown in
18 Fig. 14. However, given the results with the slab-ocean version, a dominant role of CH₄ in the
19 mitigation is likely also for the full-ocean simulations. In fact, this adds to the explanation
20 why the temperature response in Fig. 14 emerges from natural variability only about 10 years
21 (i.e., approximately the lifetime of CH₄) after the start of the mitigation measures.

22 We have seen in section 3.5 that the chosen set of mitigation measures also leads to consider-
23 able reductions in PM and O₃ and thus improves air quality. If implemented as such, the miti-
24 gation package as a whole would therefore have beneficial impacts on both air quality and
25 climate. However, the experiments conducted here also suggest that the co-benefits for cli-
26 mate and air quality result mainly from CH₄ mitigation, which improves air quality (via clear
27 reductions of the background surface O₃ concentrations) and reduces warming considerably.
28 The co-benefits of the non-CH₄ SLCP mitigation measures, on the other hand, are quite lim-
29 ited. These measures improve air quality strongly (via reductions of PM concentrations and
30 O₃) but reduce warming only slightly. This does of course not mean that all of the individual
31 measures have small co-benefits. For instance, it is likely that mitigating sources with the
32 highest BC/OA ratio would lead to larger co-benefits. Partly, the small co-benefits for the

1 non-CH₄ SLCP measures are a result of the design of our MIT scenario, which was based on
2 the GTP₂₀ metric and this seems to suggest a stronger warming reduction due to non-CH₄
3 SLCP measures than seen in the transient ESM simulation results. The consistency between
4 the two approaches will be discussed in the next section, but it is clear that if the metric was
5 too “optimistic” with respect to the achievable warming reductions by non-CH₄ SLCP mitigation
6 measures, some individual measures will have been included in the MIT basket, which in
7 the transient simulations might have actually led to warming enhancement instead of warming
8 reduction.

9 **3.7 Closing the loop: Climate impacts from metric calculations and transient**
10 **model simulations, and applications of the metrics**

11 The purpose of this section is to compare the climate impacts of the SLCP mitigation as estimated
12 with the metrics and as obtained from the transient simulations, as well as to show applications
13 of the metrics. A perfect agreement between the two methods cannot be expected, as the metrics assume linearity of the climate response to all individual forcing contributions
14 and are also valid for a specific time horizon. Furthermore, after using the RF values calculated
15 in ECLIPSE, the methodology adopted to calculate the GTP values uses a representation of
16 both the size and time dependence of the response derived from one particular (pre-ECLIPSE)
17 ESM calculation. Since the different ESMs used in ECLIPSE have a diverse range of climate
18 sensitivities and different representations of uptake of heat by the oceans, this is a further important
19 reason why exact agreement should not be expected. The metrics are also calculated
20 for a specific climate and are less accurate for periods with a changed climate. One example is
21 the BC snow albedo effect, which is gradually reduced over time because the snow and ice
22 extent decreases as the climate warms. On the other hand, diagnosis of temperature responses
23 from the transient climate simulations is also associated with large error bars because of natural
24 climate variability. Nevertheless, a comparison of the two methods is an important consistency
25 check but, to our knowledge, has never been done before for a SLCP mitigation scenario and using an ensemble of ESMs.

26 The ARTP metric described in section 3.5, applied over a series of timescales, allows calculation
27 of the regional (in broad latitude bands) temperature response from the reduction of individual
28 species in the mitigation basket as a function of time. Based on these regional respons-

1 es a total global mean response is derived which can be compared directly with the global
2 mean temperature difference between the MIT and CLE transient simulations in the ESMs.

3 The ARTPs depend on the global climate sensitivity assumed for the impulse response func-
4 tion used (Shindell et al., 2012). To provide a consistent comparison with the ESM transient
5 simulations, the ARTPs have been scaled by the individual climate sensitivities for each
6 ESM. Furthermore, not all the ESMs include all components and forcing mechanisms in their
7 transient simulations, e.g., only NorESM includes the forcing due to BC deposited on snow
8 and ice. To make a consistent comparison, ARTP contributions were summed only over the
9 components and processes included in each ESM. Figure 19 shows the global mean tempera-
10 ture responses using the ARTP method, which can be compared directly with the temperature
11 changes obtained by the transient ESM simulations shown in Fig. 14. The solid lines in Fig.
12 19 show the total global temperature response of the mitigation, while the dashed lines show
13 the contribution from CH₄ reductions only. For the last decade of the simulation (2041-2050)
14 both the ESMs and ARTPs give a mean global response of -0.22 K. The ranges of the esti-
15 mates based on the four models are also very similar (-0.15 to -0.29 K for the ESMs, and -
16 0.13 to -0.33 K for the ARTPs). Note, however, that the factors contributing to these ranges
17 are not the same in these two estimates. For the ESMs, the differences in radiative forcing, the
18 models' climate sensitivities and internal variability determine the range, while for the ARTP-
19 based estimate the differences in which processes causing RF are included in each model and
20 the model's climate sensitivities are accounted for. The ranking of the responses between the
21 individual models are, however, not identical. Nevertheless, based on these results we con-
22 clude that for the global mean response to SLCP mitigation the ARTP-based method simu-
23 lates well the full ESM simulations to estimate the impact of SLCP mitigation.

24 The ARTP-based method suggests a larger contribution of non-CH₄ SLCPs to the temperature
25 response for the 2041-2050 decade (~22%, see dashed lines in Fig. 19) than the transient sim-
26 ulations using the slab-ocean version of the CESM-CAM4 model discussed in the previous
27 section (~10%). One reason for this disagreement is that many of the BC-related measures
28 included in the mitigation basket were relatively OC-rich and this makes their net temperature
29 response extremely uncertain, especially when aerosol indirect effects and the semi-direct
30 effect are considered. An even larger contribution (44%) of non-CH₄ SLCPs to the CO₂-
31 equivalent emission reductions of the MIT scenario was obtained with the GTP₂₀ metric di-
32 rectly (see section 3.5). However, this value is valid only for the temperature response in the

1 year 2035, for which the relative contribution of the non-CH₄ SLCPs (which are shorter-lived
2 than CH₄) is larger than for later years. As the diagnostic uncertainties for a single year are
3 very large for the transient ESM simulations, we abstain from comparisons. Direct compari-
4 sons are anyway not meaningful, since the metrics include all recognized RF mechanisms,
5 whereas most ESMs lack some of these (e.g., snow albedo changes by BC).

6 To further investigate the ability of a simplified metrics method to represent the regional re-
7 sponse simulated by the ESMs, we use the ARTPs to estimate the response in four broad lati-
8 tude bands and these results can be compared to the corresponding results from the ESMs.
9 Figure 20 shows the 20-year mean results (2031-2050) for both methods. The general pattern
10 of the responses with largest impact for the main source region at Northern Hemisphere Mid-
11 Latitudes (NHML) and in the Arctic is well captured by the ARTP method. However, as ex-
12 pected the agreement between the estimates is not as good as for the global mean (correlation
13 coefficient of 0.68 for the 16 data points). The ARTP-based estimates are close to the ESM
14 means for the NHML and the Tropics, while for the Arctic and for the SH the ARTP method
15 underestimates the response simulated by the ESMs. The reason for the more pronounced
16 Arctic amplification in our ESMs than for the ARTPs is probably a less pronounced Arctic
17 amplification in the model of Shindell and Faluvegi (2009), from where the RTP-coefficients
18 were taken. For future development and use of the ARTP method, RTP-coefficients are need-
19 ed also from other ESMs.

20 The motivation for establishing these ARTPs is that after quality-control they provide policy-
21 makers with a relatively simple tool to quantify how sectorial emissions (i.e., by region, sector
22 and component) contribute to temperature change over time in broad latitude bands. Figure 21
23 illustrates this potential for the MIT scenario. It shows the responses (mean of the ARTP-
24 based estimates for the four ESMs) for mitigation taking place in Europe, China and globally.
25 The larger relative impact on the Arctic by mitigation of European emissions compared to
26 mitigation in China is clearly seen, while the impact for NHML is about three times larger in
27 absolute terms for mitigation of SLCPs in China. These results can then easily be further ana-
28 lysed by separating the impact by emitted species (Fig. 21, panel d), or by separating the im-
29 pact of a single emission sector. Figure 22, for instance, shows the estimated impact on Arctic
30 temperatures by global mitigation of SLCPs from the residential heating and cooking sector,
31 broken down to contributions from individual SLCP species. Notice that it would be prohibi-

tively expensive to run an ensemble of ESM simulations that is large enough to detect the small temperature response resulting from such minute emission changes.

Conclusions

ECLIPSE has come to a number of important scientific conclusions, which are also of high relevance for climate and air quality policy:

- ECLIPSE has created a new inventory for anthropogenic SLCP emissions, including scenarios for the future. An important finding is the large range of possible future developments of anthropogenic SLCP emissions, which even for a single future energy pathway substantially exceeds the range of SLCP emissions given in IPCC's RCPs. The large range results from the uncertainties of future air quality policies, as well as from the expected level of implementation and enforcement of existing policies (Klimont et al., 2015a,b).
- Detailed comparisons between measured and modelled distributions of aerosol, O₃ and other SLCP gases have shown that for many substances the models are in good agreement with available background observations. The model performance of the ESMs is similar to that of CTMs. For BC, in particular, the agreement between models and measurements has improved for the Arctic (Eckhardt et al., 2015), which is partly the result of accounting for emissions from gas flaring and emission seasonality (Stohl et al., 2013). Outside the Arctic, a reduction of the BC lifetime led to improvements (Samset et al., 2014). Nevertheless, our comparisons suggest underestimates of BC and aerosol precursor emissions in high-latitude Russia and in India. Furthermore, it was found that SO₂ concentrations are overestimated and CO concentrations are underestimated by the models (Quennehen et al., 2015) at the surface in Asia and Europe during summer and autumn. The CO underestimate is likely associated with a too short CO lifetime in the models. Ozone, on the other hand, is generally overestimated at rural locations. Such discrepancies may affect model responses to emission perturbations and thus radiative forcing.
- Earth System Models can reproduce the accelerated upward trend of surface temperature over Europe that was observed when aerosol precursor emissions were reduced in

1 the 1990s (leading to solar brightening), after a period of emission growth in the
2 1960s-1980s (leading to solar dimming) (Cherian et al., 2014).

- 3 • ECLIPSE performed detailed multi-model calculations of RF for all considered SLCP
4 species, as a function of emission region and season. It is found that the absolute val-
5 ues of specific RF for aerosols are generally larger in summer than in winter (Bellouin
6 et al., 2015). It is found that the semi-direct effect on clouds, although highly uncer-
7 tain, can potentially offset a considerable fraction of the direct positive RF of BC.
8 This, together with reduced BC lifetimes, causes the net RF for BC calculated in
9 ECLIPSE to be only weakly positive, which is different from most previous studies.
- 10 • NO_x emissions affect the concentrations of O_3 , CH_4 and nitrate aerosols. The first ef-
11 fect leads to positive RF, while the latter two cause negative RF. We have quantified
12 all these effects and can state with confidence that the current net RF of global histori-
13 cal NO_x emissions is negative. The forward looking metrics GWP and GTP for NO_x
14 are negative as well, except for short time horizons.
- 15 • ECLIPSE had a focus on calculation and testing of emission metrics, which led to a
16 better understanding of existing metrics (Aamaas et al., 2013), further development of
17 the applications of the RTP concept (Collins et al., 2013) and introduction of new met-
18 ric concepts such as the Global Precipitation change Potential (GPP) by Shine et al.
19 (2015). After careful consideration of the alternatives, we chose a 15-year ramp-up
20 version (i.e., assuming a linear implementation of measures) of the GTP_{20} metric for
21 designing a SLCP mitigation scenario.
- 22 • The GTP_{20} metric was implemented into the GAINS model to identify mitigation
23 measures (MIT) that have beneficial impacts on both air quality and climate. We find
24 that the 17 most important mitigation measures would contribute more than 80% of
25 the climate benefits according to the GTP_{20} metric. The top measures both for CH_4
26 and BC mitigation are to prevent the venting (for CH_4) and flaring (for BC) of gas as-
27 sociated with the oil production. For CH_4 , measures on shale gas production, waste
28 management and coal mines were also important. For non- CH_4 SLCPs, elimination of
29 high emitting vehicles and wick lamps, as well as reducing emissions from coal and
30 biomass stoves, agricultural waste, solvents and diesel engines were also important.
- 31 • Full implementation of these measures (the MIT scenario) would reduce global an-
32 thropogenic emissions of CH_4 and BC by 50% and 80%, respectively. As a result of

1 co-control with BC, emissions of organic aerosols would also be reduced by 70%,
2 whereas emissions of CO₂ and SO₂ would hardly be changed. Based on the GTP₂₀
3 metric, the CO₂-equivalent emissions (including CO₂ emissions) would be decreased
4 by about 70% in the year 2030, with about 56% of the decrease caused by CH₄
5 measures and 44% caused by non-CH₄ SLCP measures.

- 6 • The mitigation scenario would reduce surface concentrations of O₃ and PM_{2.5} globally
7 compared to the CLE scenario, with BC reductions of more than 80% in some areas.
8 We estimate that in the EU the loss of statistical life expectancy due to air pollution
9 will be reduced from 7.5 months in 2010 to 5.2 months in 2030 in the CLE scenario.
10 The MIT measures would gain another 0.9 months. Substantially larger health im-
11 provements from SLCP measures are estimated for China (1.8 months) and India (11-
12 months).
- 13 • Climate impacts of SLCP emissions were simulated with four ESMs with full ocean
14 coupling. Equilibrium simulations that removed all land-based anthropogenic emis-
15 sions of SO₂, BC and OA in turn showed robust global-mean increase in surface tem-
16 peratures of 0.69 K (0.40-0.84 K) for SO₂ removal and smaller warming for OA re-
17 moval (Baker et al., 2015a). The global mean temperature response to BC removal
18 was slightly negative: -0.05 K (-0.15 to 0.08 K). The relatively small global response
19 to BC emission reductions was attributed to strong, while uncertain, indirect and semi-
20 direct effects, which partly offset the direct aerosol radiative effect.
- 21 • Climate impacts of the MIT scenario were investigated with ESM ensemble transient
22 simulations of both the CLE and MIT scenario (Baker et al., 2015b). Multi-model en-
23 semble mean global mean surface temperature in the CLE scenario increased by
24 0.70±0.14 K between the years 2006 and 2050. The ensemble mean global mean sur-
25 face warming for the last decade of the simulation (2041-2050) was, however,
26 0.22±0.07 K weaker for the MIT scenario, demonstrating the effect of the SLCP miti-
27 gation. The response was strongest in the Arctic, with warming reduced by about
28 0.44 K (0.39-0.49 K).
- 29 • In addition to global annual mean temperature change, there are other climate parame-
30 ters that are of relevance for policy decisions (e.g., changes in precipitation, regional
31 temperatures, etc.). The SCLP reductions in the MIT scenario led to particularly bene-
32 ficial climate responses in Southern Europe, where the surface warming was reduced

1 by about 0.3 K from spring to autumn and precipitation rates were increased by about
2 15 (6-21) mm/yr (15 mm/yr corresponding to more than 4% of total precipitation),
3 compared to the CLE scenario. Thus, the mitigation could help to alleviate expected
4 future drought and water shortages in the Mediterranean region.

- 5 • Additional ESM transient simulations, which only included the CH₄ emission reductions,
6 led to a global warming reduction that amounted to about 90% of the reduction
7 produced by the simulations using the full set of measures for the final decade of the
8 simulations (2041-2050). This suggests that, for longer time horizons, the net climate
9 benefits from our chosen non-CH₄ SLCP mitigation measures in terms of global annual
10 mean temperature change are limited, probably due to small forcing and co-emitted
11 cooling species. Nevertheless, if implemented as such, the mitigation package as a
12 whole would have beneficial impacts on both air quality and climate, and for the latter,
13 also in other climate variables than global annual mean temperature change such as
14 regional changes in temperature and precipitation.
- 15 • For the first time, ECLIPSE compared the temperature response to an SLCP mitigation
16 scenario as it is given by climate metrics (using the ARTP method) and as it is
17 simulated with transient ESM simulations. This is crucial for the application of metrics,
18 which – because of their simplicity and flexibility – are very relevant in a policy
19 context where they can substitute full ESM simulations which are expensive and im-
20 practical for small perturbations. Both approaches give a global mean reduced warm-
21 ing of the surface temperatures by 0.22 K (and similar uncertainty ranges) for the pe-
22 riod 2041-2050. Also the large-scale pattern of the response (with strongest warming
23 reductions in the Arctic) is reproduced similarly by both methods, even though the
24 agreement is less good than for the global mean.
- 25 • The metrics-based approach and the transient model simulations agree less well on the
26 relative contribution of CH₄ and non-CH₄ SLCP mitigation measures to the reduced
27 warming. While the metrics-based approach suggests that the non-CH₄ measures ac-
28 count for 22% of the global-mean temperature response for 2041-2050, the transient
29 simulations result in a contribution from non-CH₄ measures of only about 10%. One
30 reason for this disagreement is that many of the BC-related measures included in the
31 mitigation basket were relatively OC-rich and this makes their net temperature re-
32 sponse more uncertain, especially when aerosol indirect effects are considered. Fur-

thermore, small cooling influences are easily masked by unforced variability in fully-coupled climate simulations.

- The major share of the cooling effect in our SLCP mitigation scenario is contributed by CH₄ reductions, with 20-30% of the difference in near-term global mean climate warming from the reduction of non-CH₄ SLCPs (multi-model range 0.01-0.06 K for time periods from 2021-2040, according to metrics-based estimates). Thus, to maximize climate co-benefits of non-CH₄ SLCPs, sources with the highest BC/OA emission ratios should be addressed with priority. At the same time, air pollution policies should consider mitigation of CH₄, with clear co-benefits for climate warming and air quality via reduced surface O₃ concentrations.
- The ECLIPSE mitigation scenario has been developed to be representative of a mitigation strategy that considers both climate and air quality, assuring reduced climate forcing without detrimental impact on air quality; as a matter of fact, strong air quality co-benefits were identified. Real-world scenarios are likely to favour particular policy objectives and will also consider the costs for the mitigation measures. More work is therefore needed to explore a larger range of scenarios. By demonstrating the efficiency and capacity of the metrics-based approach to quantify temperature changes and its consistency with transient climate model simulations, ECLIPSE has opened the way to explore a large number of such scenarios. This would be an impossible task if transient climate ensemble model simulations were needed for each.
- The number of models contributing to the ECLIPSE project was relatively small. While the models were shown to be largely representative of results obtained from larger model ensembles, this makes quantification of mean values and especially uncertainties (e.g., of RF or temperature response) dependent on the particular properties of the ECLIPSE models. For a more comprehensive quantification of uncertainties, it is therefore recommended to repeat the modelling exercises presented in this paper with a larger international model ensemble.

Acknowledgements

The research leading to these results has received funding from the European Union Seventh Framework Programme (FP7/2007-2013) under grant agreement no 282688 – ECLIPSE. **We**

1 thank D. Shindell and an anonymous referee for their comments, which helped to improve
2 this paper.

3

4 **References (refs from the proposal are missing here)**

5 Aamaas, B., Peters, G. P., Fuglestvedt, J. S.: Simple emission metrics for climate impacts,
6 Earth Syst. Dyn., 4, 145-170, 2013.

7 Aamaas B., T.K. Berntsen, J. S. Fuglestvedt, K. P. Shine, N. Bellouin: Multimodel emission
8 metrics for regional emissions of short lived climate forcers, Atmos. Chem. Phys., ~~in prepara-~~
9 ~~tionsubmitted~~, 2015.

10 Alexandratos, N., and Bruinsma, J.: World agriculture towards 2030/2050, the 2012 revision
11 (No. 12-03), ESA Working Paper. World Food and Agricultural Organization, Rome, Italy,
12 2012.

13 Amann, M., Bertok, I., Borken-Kleefeld, J., Cofala, J., Heyes, C., Höglund-Isaksson, L.,
14 Klimont, Z., Nguyen, B., Posch, M., Rafaj, P., Sandler, R., Schöpp, W., Wagner, F., and Win-
15 iwarter, W.: Cost-effective control of air quality and greenhouse gases in Europe: Modeling
16 and policy applications, Env. Mod. & Software, 26, 1489-1501,
17 <http://dx.doi.org/10.1016/j.envsoft.2011.07.012>, 2011.

18 Amann, M., Klimont, Z., and Wagner, F.: Regional and global emissions of air pollutants:
19 Recent trends and future scenarios, Ann. Rev. Environ. Res., 38, 31-55, 2013.

20 Amann, M., and Wagner, F.: A Flexibility Mechanism for Complying with National Emission
21 Ceilings for Air Pollutants, TSAP Report #15, Version 1.0, International Institute for Applied
22 Systems Analysis IIASA, Laxenburg, Austria, 2014.

23 AMAP: AMAP Assessment 2015: Black Carbon and Ozone as Arctic Climate Forcers. Arctic
24 Monitoring and Assessment Programme (AMAP), Oslo, Norway.(in preparation), 2015.

25 Andreae, M. O., Jones, C. D., and Cox, P. M.: Strong present-day aerosol cooling implies a
26 hot future, Nature, 435, 1187-1190, 2005.

27 Andrews, T., Forster, P. M., Boucher, O., Bellouin, N., and Jones, A.: Precipitation, radiative
28 forcing and global temperature change, Geophys. Res. Lett., 37, L14701,
29 doi:10.1029/2010GL043991, 2010.

1 Anenberg, S. C., Schwartz, J., Shindell, D., Amann, M., Faluvegi, G., Klimont, Z., Janssens-
2 Maenhout, G., Pozzoli, L., van Dingenen, R., Vignati, E., Emberson, L., Muller, N. Z., West,
3 J. J., Williams, M., Demkine, V., Hicks, W. K., Kuylensierna, J., Raes, F., and Ramanathan,
4 V.: Global air quality and health co-benefits of mitigating near-term climate change through
5 methane and black carbon emission controls, Env. Health Perspectives, 120, 831-839,
6 <http://dx.doi.org/10.1289/ehp.1104301>, 2012.

7 Arneth, A., Unger, N., Kulmala, M., and Andreae, M.O.: Clean the air, heat the planet? Sci-
8 ence, 326, 672-673, doi: 10.1126/science.1181568, 2009.

9 Baker, L. H., Collins, W. J., Olivié, D. J. L., Cherian, R., Hodnebrog, Ø., Myhre, G., and
10 Quaas, J.: Climate responses to anthropogenic emissions of short-lived climate pollutants,
11 Atmos. Chem. Phys., 15, 8201-8216, doi:10.5194/acp-15-8201-2015, 2015a.

12 Baker, L. H., Collins, W. J., Olivié, D. J. L., Cherian, R., Hodnebrog, Ø., Myhre, G., and
13 Quaas: Climate responses to anthropogenic emissions of short lived climate pollutants, At-
14 mos. Chem. Phys. Discuss., 15, 3823-3862, doi:10.5194/acpd-15-3823-2015, 2015a.

15 Baker, L. H., et al.: Transient simulations of the ECLIPSE mitigation scenario, in preparation,
16 2015b.

17 Bellouin, N., L. Baker, Ø. Hodnebrog, D. Oliviè, R. Cherian, C. BarberMacIntosh, B. Samset,
18 A. Esteve, B. Aamaas, J. Quaas, and G. Myhre: Regional and seasonal radiative forcing by
19 perturbations to aerosol and ozone precursor emissions, Atmos. Chem. Phys., in preparation,
20 2015.

21 Bentsen, M., Bethke, I., Debernard, J. B., Iversen, T., Kirkevag, A., Seland, O., Drange, H.,
22 Roelandt, C., Seierstad, I. A., Hoose, C., and Kristjansson, J. E.: The Norwegian Earth Sys-
23 tem Model, NorESM1-M - Part 1: Description and basic evaluation of the physical climate,
24 Geosci. Mod. Dev., 6, 687-720, doi:10.5194/gmd-6-687-2013, 2013.

25 Berntsen, T. K., Fuglestvedt, J. S., Joshi, M. M., Shine, K. P., Stuber, N., Ponater, M., Sausen,
26 R., Hauglustaine, D. A., and Li, L.: Climate response to regional emissions of ozone precur-
27 sors: sensitivities and warming potentials, Tellus, 57B, 283–304, 2005.

28 Berntsen, T., Fuglestvedt, J., Myhre, G., Stordal, F., and Berglen, T. F.: Abatement of green-
29 house gases: does location matter? Clim. Change, 74, 377–411, 2006.

1 Berntsen, T. K., Tanaka, K., and Fuglestvedt, J. S.: Does black carbon abatement hinder CO₂
2 abatement? A letter, *Climatic Change*, 103, 627–633, doi:10.1007/s10584-010-9941-3, 2010.

3 Bond, T. C., Doherty, S. J., Fahey, D. W., Forster, P. M., Berntsen, T., DeAngelo, B. J., Flan-
4 ner, M. G., Ghan, S., Kärcher, B., Koch, D., Kinne, S., Kondo, Y., Quinn, P. K., Sarofim, M.
5 C., Schultz, M. G., Schulz, M., Venkataraman, C., Zhang, H., Zhang, S., Bellouin, N., Gut-
6 tikunda, S. K., Hopke, P. K., Jacobson, M. Z., Kaiser, J.W., Klimont, Z., Lohmann, U.,
7 Schwarz, J. P., Shindell, D., Storelvmo, T., Warren, S. G., and Zender, C. S.: Bounding the
8 role of black carbon in the climate system: A scientific assessment, *J. Geophys. Res.*, 118,
9 5380–5552, doi:10.1002/jgrd.50171, 2013.

10 Bond, T. C., and Sun, H. L.: Can reducing black carbon emissions counteract global warm-
11 ing? *Environ. Sci. Tech.*, 39, 5921–5926, 2005.

12 Boucher, O., and Reddy, M. S.: Climate trade-off between black carbon and carbon dioxide
13 emissions, *Energy Policy*, 36, 193–200, 2008. See corrigendum on
14 http://www.lmd.jussieu.fr/~obolmd/PDF/Corrigendum_Boucher_Reddy_Energy_Policy_2008.pdf

16 Boucher, O., Randall, D., Artaxo, P., Bretherton, C., Feingold, G., Forster, P., Kerminen, V.-
17 M., Kondo, Y., Liao, H., Lohmann, U., Rasch, P., Satheesh, S. K., Sherwood, S., Stevens, B.,
18 and Zhang, X. Y.: Clouds and Aerosols. In: *Climate Change 2013: The Physical Science Ba-*
19 *sis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental*
20 *Panel on Climate Change* [Stocker, T. F., Qin, D., Plattner, G.-K., Tignor, M., Allen, S. K.,
21 Boschung, J., Nauels, A., Xia, Y., Bex, V., and Midgley, P. M. (eds.)]. Cambridge University
22 Press, Cambridge, United Kingdom and New York, NY, USA, pg. 571-657, 2013.

23 Bowerman, N. H. A., Frame, D. J., Huntingford, C., Lowe, J. A., Smith, S.M., and Allen, M.
24 R.: The role of short-lived climate pollutants in meeting temperature goals. *Nature Clim.*
25 *Change*, 3, 1021-1024, doi:10.1038/nclimate2034, 2013.

26 Brasseur, G. P., and Roeckner, E.: Impact of improved air quality on the future evolution of
27 climate, *Geophys. Res. Lett.*, 32, L23704, doi:10.1029/2005GL023902, 2005.

28 Broccoli, A. J., Dahl, K. A., and Stouffer, R. J.: Response of the ITCZ to Northern Hemi-
29 sphere cooling, *Geophys. Res. Lett.*, 33, L01702, doi:10.1029/2005GL024546, 2006.

1 Cherian, R., Quaas, J., Salzmann, M., and Wild, M.: Pollution trends over Europe constrain
2 global aerosol forcing as simulated by climate models, *Geophys. Res. Lett.*, 41, 2176-2181,
3 doi:10.1002/2013GL058715, 2014.

4 Collins, W. J., Sitch, S., and Boucher, O.: How vegetation impacts affect climate metrics for
5 ozone precursors. *J. Geophys. Res.*, 115, D23308, doi:10.1029/2010JD014187, 2010.

6 Collins, W. J., Fry, M. M., Yu, H., Fuglestvedt, J. S., Shindell, D. T., and West, J. J.: Global
7 and regional temperature-change potentials for near-term climate forcers, *Atmos. Chem.*
8 *Phys.*, 13, 2471-2485, doi:10.5194/acp-13-2471-2013, 2013.

9 Daskalakis, N., Myriokefalitakis, S., and Kanakidou, M.: Sensitivity of tropospheric loads and
10 lifetimes of short lived pollutants to fire emissions, *Atmos. Chem. Phys.*, 15, 3543-3563,
11 doi:10.5194/acp-15-3543-2015, 2015.

12 Dentener, F., Keating, T., and Akimoto, H. (editors): *Hemispheric Transport of Air Pollution*
13 2010 - Part A: Ozone and Particulate Matter, *Air Pollution Studies* No. 17, United Nations,
14 New York and Geneva, 2010.

15 EC: *Proposal for a Directive of the European Parliament and of the Council on the reduction of*
16 *national emissions of certain atmospheric pollutants and amending Directive 2003/35/EC*. European
17 Commission (EC), Brussels, Belgium, 2013.

18 ~~Eckhardt, S., Quennchen, B., Olivié, D. J. L., Berntsen, T. K., Cherian, R., Christensen, J. H.,~~
19 ~~Collins, W., Crepinsek, S., Daskalakis, N., Flanner, M., Herber, A., Heyes, C., Hod-~~
20 ~~nebrog, Ø., Huang, L., Kanakidou, M., Klimont, Z., Langner, J., Law, K. S., Massling, A.,~~
21 ~~Myriokefalitakis, S., Nielsen, I. E., Nøjgaard, J. K., Quaas, J., Quinn, P. K., Raut, J. C., Rum-~~
22 ~~bøld, S. T., Schulz, M., Skeie, R. B., Skov, H., Lund, M. T., Uttal, T., von Salzen, K.,~~
23 ~~Mahmood, R., and Stohl, A.: Current model capabilities for simulating black carbon and sul-~~
24 ~~fate concentrations in the Arctic atmosphere: a multi model evaluation using a comprehensive~~
25 ~~measurement data set, *Atmos. Chem. Phys. Discuss.*, 15, 10425-10477, doi:10.5194/acpd-15-~~
26 ~~10425-2015, 2015.~~

27 Eckhardt, S., Quennchen, B., Olivié, D. J. L., Berntsen, T. K., Cherian, R., Christensen, J. H.,
28 Collins, W., Crepinsek, S., Daskalakis, N., Flanner, M., Herber, A., Heyes, C., Hod-
29 nebrog, Ø., Huang, L., Kanakidou, M., Klimont, Z., Langner, J., Law, K. S., Lund, M. T.,
30 Mahmood, R., Massling, A., Myriokefalitakis, S., Nielsen, I. E., Nøjgaard, J. K., Quaas, J.,

1 [Quinn, P. K., Raut, J.-C., Rumbold, S. T., Schulz, M., Sharma, S., Skeie, R. B., Skov, H., Ut-](#)
2 [tal, T., von Salzen, K., and Stohl, A.: Current model capabilities for simulating black carbon](#)
3 [and sulfate concentrations in the Arctic atmosphere: a multi-model evaluation using a com-](#)
4 [prehensive measurement data set, Atmos. Chem. Phys., 15, 9413-9433, doi:10.5194/acp-15-](#)
5 [9413-2015, 2015.](#)

6 EEA, 2013: Air quality in Europe — 2013 report, EEA report 9/2013, ISBN 978-92-9213-
7 406-8, <http://www.eea.europa.eu/publications/air-quality-in-europe-2013>, accessed
8 28.04.2014.

9 Endresen, Ø., Sørgård, E., Behrens, H. L., Brett, P. O., and Isaksen, I. S. A.: A historical re-
10 construction of ships' fuel consumption and emissions, *J. Geophys. Res.*, 112, D12301,
11 doi:10.1029/2006JD007630, 2007.

12 Flanner, M. G.: Arctic climate sensitivity to local black carbon, *J. Geophys. Res. Atmos.*, 118,
13 1840–1851, doi:10.1002/jgrd.50176, 2013.

14 Fuglestvedt, J. S., Berntsen, T. K., Isaksen, I. S. A., Mao, H. T., Liang, X. Z., and Wang, W.
15 C.: Climatic forcing of nitrogen oxides through changes in tropospheric ozone and methane;
16 global 3D model studies, *Atmos. Environ.*, 33, 961–977, 1999.

17 Fuglestvedt, J. S., Shine, K. P., Berntsen, T., Cook, J., Lee, D. S., Stenke, A., Skeie, R. B.,
18 Velders, G. J. M., and Waitz, I. A.: Transport impacts on atmosphere and climate: Metrics,
19 *Atmos. Environ.*, 44, 4648-4677, 10.1016/j.atmosenv.2009.04.044, 2010.

20 Gadhavi, H. S., Renuka, K., Ravi Kiran, V., Jayaraman, A., Stohl, A., Klimont, Z., and
21 Beig, G.: Evaluation of black carbon emission inventories using a Lagrangian dispersion
22 model – a case study over Southern India, *Atmos. Chem. Phys.* 15, 1447–1461,
23 doi:10.5194/acp-15-1447-2015, 2015.

24 GEA: Global Energy Assessment: Toward a Sustainable Future, Cambridge University Press,
25 UK, 2012.

26 Gedney, N., Huntingford, C., Weedon, G. P., Bellouin, N., Boucher, O., and Cox, P. M.: De-
27tection of solar dimming and brightening effects on Northern Hemisphere river flow, *Nature*
28 *Geosci.*, 7, 796–800, doi:10.1038/ngeo2263, 2014.

29 Gent, P. R., Danabasoglu, G., Donner, L. J., Holland, M. M., Hunke, E. C., Jayne, S. R., Law-
30 rence, D. M., Neale, R. B., Rasch, P. J., Vertenstein, M., Worley, P. H., Yang, Z.-L., and

1 Zhang, M.: The Community Climate System Model version 4, *J. Clim.*, 24, 4973-4991,
2 doi:10.1175/2011jcli4083.1, 2011.

3 Gilgen, H., Wild, M., and Ohmura, A.: Means and trends of shortwave irradiance at the sur-
4 face estimated from global energy balance archive data, *J. Clim.*, 11, 2042–2061, 1998.

5 Gillett, N. P. and Matthews, H. D.: Accounting for carbon cycle feedbacks in a comparison of
6 the global warming effects of greenhouse gases, *Environ. Res. Lett.*, 5, 034011,
7 doi:10.1088/1748-9326/5/3/034011, 2010.

8 Grell, G. A., Peckham, S. E., Schmitz, R., McKeen, S. A., Frost, G., Skamarock, W. C., and
9 Eder, B.: Fully coupled online chemistry within the WRF model, *Atmos. Environ.*, 39, 6957-
10 6975, 2005.

11 Grieshop, A. P., Reynolds, C. C. O., Kandlikar, M., and Dowlatbadi, H.: A black-carbon
12 mitigation wedge, *Nature Geosci.*, 2, 533-534, 2009.

13 Grosse, L. Y., Lauby-Secretan, B., El Ghissassi, F., Bouvard, V., Benbrahim-Tallaa, L.,
14 Guha, N., Baan, R., Mattock, H., and Straif, K.: The carcinogenicity of outdoor air pollution,
15 *Lancet*, 14, 1262–1263, 2013.

16 Guenther, A. B., Jiang, X., Heald, C. L., Sakulyanontvittaya, T., Duhl, T., Emmons, L. K.,
17 and Wang, X.: The Model of Emissions of Gases and Aerosols from Nature version 2.1
18 (MEGAN2.1): an extended and updated framework for modeling biogenic emissions, *Geosci.*
19 *Model Dev.*, 5, 1471-1492, doi:10.5194/gmd-5-1471-2012, 2012.

20 Hewitt, H. T., Copsey, D., Culverwell, I. D., Harris, C. M., Hill, R. S. R., Keen, A. B.,
21 McLaren, A. J., and Hunke, E. C.: Design and implementation of the infrastructure of HadG-
22 EM3: the next-generation Met Office climate modelling system, *Geosci. Mod. Dev.*, 4, 223-
23 253, doi:10.5194/gmd-4-223-2011, 2011.

24 Hodnebrog, Ø., Myhre, G., and Samset, B.: How shorter black carbon lifetime alters its cli-
25 mate effect, *Nature Communications*, 5, 5065, doi:10.1038/ncomms6065, 2014.

26 IARC: IARC monographs on the evaluation of carcinogenic risks to humans. Volume 109.
27 Outdoor air pollution. Lyon: International Agency for Research on Cancer (in press).

28 IEA (International Energy Agency): Energy Technology Perspectives. 2012 - Pathways to a
29 Clean Energy System. OECD/IEA, Paris, 2012.

1 Im, U., Christodoulaki, S., Violaki, K., Zampas, P., Kocak, M., Daskalakis, N., Mihalopoulos,
2 N., and Kanakidou, M.: Atmospheric deposition of nitrogen and sulfur over southern Europe
3 with focus on the Mediterranean and the Black Sea, *Atmos. Environ.*, 81, 660-670,
4 doi:10.1016/j.atmosenv.2013.09.048, 2013.

5 Isaksen, I. S. A., Granier, C., Myhre, G., Berntsen, T. K., Dalsøren, S. B., Gauss, M.,
6 Klimont, Z., Benestad, R., Bousquet, P., Collins, W., Cox, T., Eyring, V., Fowler, D., Fuzzi,
7 S., J'ockel, P., Laj, P., Lohmann, U., Maione, M., Monks, P., Prevot, A. S. H., Raes, F., Rich-
8 ter, A., Rognerud, B., Schulz, M., Shindell, D., Stevenson, D. S., Storelvmo, T., Wang, W.-
9 C., van Weele, M., Wild, M., and Wuebbles, D.: Atmospheric composition change: Climate-
10 chemistry interactions, *Atmos. Environ.*, 43, 5138-5192, 2009.

11 Jackson, S. C.: Parallel pursuit of near-term and long-term climate mitigation, *Science*, 326,
12 526-527, 2009.

13 Jiao, C., Flanner, M. G., Balkanski, Y., Bauer, S. E., Bellouin, N., Berntsen, T. K., Bian, H.,
14 Carslaw, K. S., Chin, M., De Luca, N., Diehl, T., Ghan, S. J., Iversen, T., Kirkevåg, A.,
15 Koch, D., Liu, X., Mann, G. W., Penner, J. E., Pitari, G., Schulz, M., Seland, Ø., Skeie, R. B.,
16 Steenrod, S. D., Stier, P., Takemura, T., Tsigaridis, K., van Noije, T., Yun, Y., and Zhang, K.:
17 An AeroCom assessment of black carbon in Arctic snow and sea ice, *Atmos. Chem. Phys.*,
18 14, 2399-2417, doi:10.5194/acp-14-2399-2014, 2014.

19 Joshi, M., Shine, K., Ponater, M., Stuber, N., Sausen, R., and Li, L.: A comparison of climate
20 response to different radiative forcings in three general circulation models: towards an im-
21 proved metric of climate change, *Climate Dynam.*, 20, 843-854, 2003.

22 Kanakidou, M., Duce, R. A., Prospero, J. M., Baker, A. R., Benitez-Nelson, C., Dentener, F.
23 J., Hunter, K. A., Liss, P. S., Mahowald, N., Okin, G. S., Sarin, M., Tsigaridis, K., Uematsu,
24 M., Zamora, L. M., and Zhu, T.: Atmospheric fluxes of organic N and P to the global ocean,
25 *Glob. Biogeochem. Cyc.*, 26, doi:10.1029/2011gb004277, 2012.

26 Kawamoto, T., Pham, T. T., Matsuda, T., Oyama, T., Tanaka, M., Yu, H.S., Uchiyama, I.:
27 Historical review on development of environmental quality standards and guideline values for
28 air pollutants in Japan, *Int. J. Hyg. Environ. Health*, 214, 296-304,
29 doi:10.1016/j.ijheh.2011.05.007, 2011.

1 Kirkevåg, A., Iversen, T., Seland, Ø., Hoose, C., Kristjánsson, J. E., Struthers, H., Ek-
2 man, A. M. L., Ghan, S., Griesfeller, J., Nilsson, E. D., and Schulz, M.: Aerosol–climate in-
3 teractions in the Norwegian Earth System Model – NorESM1-M, *Geosci. Model Dev.*, 6, 207-
4 244, doi:10.5194/gmd-6-207-2013, 2013.

5 Kirtman, B., Power, S. B., Adedoyin, J. A., Boer, G. J., Bojariu, R., Camilloni, I., Doblas-
6 Reyes, F. J., Fiore, A. M., Kimoto, M., Meehl, G. A., Prather, M., Sarr, A., Schär, C., Sutton,
7 R., van Oldenborgh, G. J., Vecchi, G., and Wang, H. J.: Near-term Climate Change: Projec-
8 tions and Predictability. In: *Climate Change 2013: The Physical Science Basis. Contribution*
9 *of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Cli-
mate Change* [Stocker, T. F., Qin, D., Plattner, G.-K., Tignor, M., Allen, S. K., Boschung, J.,
11 Nauels, A., Xia, Y., Bex, V., and Midgley, P. M. (eds.)]. Cambridge University Press, Cam-
12 bridge, United Kingdom and New York, NY, USA, pg. 953-1028, 2013.

13 Klimont, Z., Smith, S. J., and Cofala, J.: The last decade of global anthropogenic sulfur diox-
14 ide: 2000-2011 emissions, *Environ. Res. Lett.*, 8, 014003, doi:10.1088/1748-
15 9326/8/1/014003, 2013.

16 Klimont, Z., Kupiainen, K., Heyes, Ch., Purohit, P., Cofala, J., Rafaj, P., Borken-Kleefeld, J.,
17 Schoepp, W.: Global anthropogenic emissions of particulate matter, in preparation, 2015a.

18 Klimont, Z., Hoglund, L., Heyes, Ch., Rafaj, P., Schoepp, W., Cofala, J., Borken-Kleefeld, J.,
19 Purohit, P., Kupiainen, K., Winiwarter, W., Amann, M., Zhao, B., Wang, S.X., Bertok, I., and
20 Sander, R.: Global scenarios of air pollutants and methane: 1990-2050, in preparation, 2015b.

21 Knutti, R., and Hegerl, G. C.: The equilibrium sensitivity of the Earth’s temperature to radia-
22 tion changes, *Nature Geosci.*, 1, 735-753, doi:10.1038/ngeo337, 2008.

23 Koch, D., Bond, T. C., Streets, D., Unger, N., and van der Werf, G. R.: Global impacts of
24 aerosols from particular source regions and sectors, *J. Geophys. Res.*, 112, D02205,
25 10.1029/2005JD007024, 2007.

26 Kopp R. E., and Mauzerall, D. L.: Assessing the climatic benefits of black carbon mitigation,
27 *Proc. Natl. Acad. Sci. USA*, 107, 11703-11708, 2010.

28 Kvalevåg, M. M., Samset, B., and Myhre, G.: Hydrological sensitivity to greenhouse gases
29 and aerosols in a global climate model, *Geophys. Res. Lett.*, 40, 1432-1438, 2013.

1 Lamarque, J. F., Bond, T. C., Eyring, V., Granier, C., Heil, A., Klimont, Z., Lee, D., Liousse,
2 C., Mieville, A., Owen, B., Schultz, M. G., Shindell, D., Smith, S. J., Stehfest, E., Van
3 Aardenne, J., Cooper, O. R., Kainuma, M., Mahowald, N., McConnell, J. R., Naik, V., Riahi,
4 K., and van Vuuren, D. P.: Historical (1850-2000) gridded anthropogenic and biomass burn-
5 ing emissions of reactive gases and aerosols: methodology and application, *Atmos. Chem.*
6 *Phys.*, 10, 7017-7039, 10.5194/acp-10-7017-2010, 2010.

7 Lee, D. S., Fahey, D. W., Forster, P. M., Newton, P. J., Wit, R. C. N., Lim, L. L., Owen, B.,
8 and Sausen, R.: Aviation and global climate change in the 21st century, *Atmos. Environ.*, 43,
9 3520–3537, 2009.

10 Lelieveld, J., Berresheim, H., Borrmann, S., Crutzen, P. J., Dentener, F. J., Fischer, H.,
11 Feichter, J., Flatau, P. J., Heland, J., Holzinger, R., Kormann, R., Lawrence, M. G., Levin, Z.,
12 Markowicz, K. M., Mihalopoulos, N., Minikin, A., Ramanathan, V., De Reus, M., Roelofs, G.
13 J., Scheeren, H. A., Sciare, J., Schlager, H., Schultz, M., Siegmund, P., Steil, B., Stephanou,
14 E. G., Stier, P., Traub, M., Warneke, C., Williams, J., and Ziereis, H.: Global air pollution
15 crossroads over the Mediterranean, *Science*, 298, 794-799, 2002.

16 Lim, S. S., Vos, T., Flaxman, A. D., Danaei, G., Shibuya, K., Adair-Rohani H, Amann M,
17 Anderson HR, Andrews KG, Aryee M, Atkinson C, Bacchus LJ, Bahalim AN, Balakrishnan
18 K, Balmes J, Barker-Collo S, Baxter A, Bell ML, Blore JD, Blyth F, Bonner C, Borges G,
19 Bourne R, Boussinesq M, Brauer M, Brooks P, Bruce NG, Brunekreef B, Bryan-Hancock C,
20 Bucello C, Buchbinder R, Bull F, Burnett RT, Byers TE, Calabria B, Carapetis J, Carnahan E,
21 Chafe Z, Charlson F, Chen H, Chen JS, Cheng AT, Child JC, Cohen A, Colson KE, Cowie
22 BC, Darby S, Darling S, Davis A, Degenhardt L, Dentener F, Des Jarlais DC, Devries K,
23 Dherani M, Ding EL, Dorsey ER, Driscoll T, Edmond K, Ali SE, Engell RE, Erwin PJ, Fa-
24 himi S, Falder G, Farzadfar F, Ferrari A, Finucane MM, Flaxman S, Fowkes FG, Freedman
25 G, Freeman MK, Gakidou E, Ghosh S, Giovannucci E, Gmel G, Graham K, Grainger R,
26 Grant B, Gunnell D, Gutierrez HR, Hall W, Hoek HW, Hogan A, Hosgood HD 3rd, Hoy D,
27 Hu H, Hubbell BJ, Hutchings SJ, Ibeaneusi SE, Jacklyn GL, Jasrasaria R, Jonas JB, Kan H,
28 Kanis JA, Kassebaum N, Kawakami N, Khang YH, Khatibzadeh S, Khoo JP, Kok C, Laden
29 F, Laloo R, Lan Q, Lathlean T, Leasher JL, Leigh J, Li Y, Lin JK, Lipshultz SE, London S,
30 Lozano R, Lu Y, Mak J, Malekzadeh R, Mallinger L, Marcenes W, March L, Marks R, Mar-
31 Martin R, McGale P, McGrath J, Mehta S, Mensah GA, Merriman TR, Micha R, Michaud C,

1 Mishra V, Mohd Hanafiah K, Mokdad AA, Morawska L, Mozaffarian D, Murphy T, Naghavi
2 M, Neal B, Nelson PK, Nolla JM, Norman R, Olives C, Omer SB, Orchard J, Osborne R, Ost-
3 ro B, Page A, Pandey KD, Parry CD, Passmore E, Patra J, Pearce N, Pelizzari PM, Petzold M,
4 Phillips MR, Pope D, Pope CA 3rd, Powles J, Rao M, Razavi H, Rehfuss EA, Rehm JT, Ritz
5 B, Rivara FP, Roberts T, Robinson C, Rodriguez-Portales JA, Romieu I, Room R, Rosenfeld
6 LC, Roy A, Rushton L, Salomon JA, Sampson U, Sanchez-Riera L, Sanman E, Sapkota A,
7 Seedat S, Shi P, Shield K, Shivakoti R, Singh GM, Sleet DA, Smith E, Smith KR, Stapelberg
8 NJ, Steenland K, Stöckl H, Stovner LJ, Straif K, Straney L, Thurston GD, Tran JH, Van Din-
9 genen R, van Donkelaar A, Veerman JL, Vijayakumar L, Weintraub R, Weissman MM, Whi-
10 te RA, Whiteford H, Wiersma ST, Wilkinson JD, Williams HC, Williams W, Wilson N,
11 Woolf AD, Yip P, Zielinski JM, Lopez AD, Murray CJ, Ezzati M, AlMazroa MA, Memish
12 ZA: A comparative risk assessment of burden of disease and injury attributable to 67 risk fac-
13 tors and risk factor clusters in 21 regions, 1990–2010: a systematic analysis for the Global
14 Burden of Disease Study 2010, *The Lancet*, 380, 2224–2260, doi:10.1016/S0140-
15 6736(12)61766-8, 2012.

16 Lund, M., Berntsen, T., Fuglestvedt, J., Ponater, M., and Shine, K: How much information is
17 lost by using global-mean climate metrics? an example using the transport sector, *Clim.*
18 *Change*, 113, 949–963, 2012.

19 Lund, T. M., Berntsen, T. K., Heyes, C., Klimont, Z., and Samset, B. H.: Global and regional
20 climate impacts of black carbon and co-emitted species from the on-road diesel sector, *At-
21 mos. Env.*, 98, 50-58, 2014a.

22 Lund, T. M., Berntsen, T. K., and Fuglestvedt, J. S.: Climate impacts of short-lived climate
23 forcers versus CO₂ from biodiesel: a case of the EU on-road sector, *Environ. Sci. Technol.*,
24 48, 14445–14454, doi:10.1021/es505308g, 2014b.

25 MacIntosh, C. R., Shine, K. P., and Collins, W. J.: Radiative forcing and
26 climate metrics for ozone precursor emissions: the impact of multi-model
27 averaging, *Atmos. Chem. Phys.*, 15, 3957-3969, doi:10.5194/acp-15-3957-2015,
28 2015.

1 Mahowald, N.: Aerosol indirect effect on biogeochemical cycles and climate, *Science*, 334,
2 794-796, doi:10.1126/science.1207374, 2011.

3 Mann, G. W., Carslaw, K. S., Spracklen, D. V., Ridley, D. A., Manktelow, P. T., Chipper-
4 field, M. P., Pickering, S. J., and Johnson, C. E.: Description and evaluation of GLOMAP-
5 mode: a modal global aerosol microphysics model for the UKCA composition-climate model,
6 *Geosci. Model Dev.*, 3, 519-551, doi:10.5194/gmd-3-519-2010, 2010.

7 Manne, A.S., and Richels, R.G.: An alternative approach to establishing trade-offs among
8 greenhouse gases, *Nature*, 410, 675–677, 2001.

9 Meinshausen, M., Meinshausen, N., Hare, W., Raper, S. C. B., Frieler, K., Knutti, R., Frame,
10 D. J., and Allen, M. R.: Greenhouse-gas emission targets for limiting global warming to 2
11 degrees C, *Nature*, 458, 1158-1162, 2009.

12 Mercado, L. M., Bellouin, N., Sitch, S., Boucher, O., Huntingford, C., Wild, O., and Cox, P.
13 M.: Impact of changes in diffuse radiation on the global land carbon sink, *Nature*, 458, 1014-
14 1017, doi:10.1038/nature07949, 2009.

15 Mitchell, T. D., and Jones, P. D.: An improved method of constructing a database of monthly
16 climate observations and associated high-resolution grids, *Int. J. Climatol.*, 25, 693–712,
17 doi:10.1002/joc.1181, 2005.

18 Molina, M., Zaelke, D., Sarma, K. M., Andersen, S. O., Ramanathan, V., and Kaniaru, D.:
19 Reducing abrupt climate change risk using the Montreal Protocol and other regulatory actions
20 to complement cuts in CO₂ emissions, *Proc. Natl Acad. Sci. USA*, 106, 20616–20621, 2009.

21 Myhre, G., Berglen, T. F., Johnsrud, M., Hoyle, C. R., Berntsen, T. K., Christopher, S. A.,
22 Fahey, D. W., Isaksen, I. S. A., Jones, T. A., Kahn, R. A., Loeb, N., Quinn, P., Remer, L.,
23 Schwarz, J. P., and Yttri, K. E.: Modelled radiative forcing of the direct aerosol effect with
24 multi-observation evaluation, *Atmos. Chem. Phys.*, 9, 1365-1392, doi:10.5194/acp-9-1365-
25 2009, 2009.

26 Myhre, G., Fuglestvedt, J. S., Berntsen, T. K., and Lund, M. T.: Mitigation of short-lived
27 heating components may lead to unwanted long-term consequences, *Atmos. Environ.*, 45,
28 6103-6106, doi:10.1016/j.atmosenv.2011.08.009, 2011.

29 Myhre, G., Shindell, D., Breon, F.-M., Collins, W., Fuglestvedt, J., Huang, J., Koch, D., La-
30 marque, J.-F., Lee, D., Mendoza, B., Nakajima, T., Robock, A., Stephens, G., Takemura, T.,

1 and Zhang, H.: Anthropogenic and Natural Radiative Forcing. In: Climate Change 2013: The
2 Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of
3 the Intergovernmental Panel on Climate Change [Stocker, T. F., Qin, D., Plattner, G.-K.,
4 Tignor, M., Allen, S. K., Boschung, J., Nauels, A., Xia, Y., Bex, V., and Midgley, P. M.
5 (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA,
6 pg. 659-740, 2013a.

7 Myhre, G., Samset, B. H., Schulz, M., Balkanski, Y., Bauer, S., Berntsen, T. K., Bian, H.,
8 Bellouin, N., Chin, M., Diehl, T., Easter, R. C., Feichter, J., Ghan, S. J., Hauglustaine, D.,
9 Iversen, T., Kinne, S., Kirkevåg, A., Lamarque, J.-F., Lin, G., Liu, X., Lund, M. T., Luo, G.,
10 Ma, X., van Noije, T., Penner, J. E., Rasch, P. J., Ruiz, A., Seland, Ø., Skeie, R. B., Stier, P.,
11 Takemura, T., Tsigaridis, K., Wang, P., Wang, Z., Xu, L., Yu, H., Yu, F., Yoon, J.-H.,
12 Zhang, K., Zhang, H., and Zhou, C.: Radiative forcing of the direct aerosol effect from Aero-
13 Com Phase II simulations, Atmos. Chem. Phys., 13, 1853-1877, doi:10.5194/acp-13-1853-
14 2013, 2013b.

15 Naik, V., Mauzerall, D., Horowitz, L., Schwarzkopf, M. D., Ramaswamy, V., and Oppenheimer, M.: Net radiative forcing due to changes in regional emissions of tropospheric ozone
16 precursors, J. Geophys. Res., 110, D24306, doi:10.1029/2005JD005908, 2005.

18
19 Olivié, D. J. L., and Peters, G. P.: Variation in emission metrics due to variation in CO₂ and
20 temperature impulse response functions, Earth Syst. Dynam., 4, 267-286, 10.5194/esd-4-267-
21 2013, 2013.

22 Orlowsky, B., and Seneviratne, S. I.: Global changes in extreme events: regional and seasonal
23 dimension, Climatic Change, 110, 669-696, 10.1007/s10584-011-0122-9, 2012.

24 Penner, J. E., Prather, M. J., Isaksen, I. S. A., Fuglestvedt, J. S., Klimont, Z., and Stevenson,
25 D. S.: Short-lived uncertainty? Nature Geosci., 3, 587 – 588, doi:10.1038/ngeo932, 2010.

26 Pierrehumbert, R. T.: Short-Lived Climate Pollution, Ann. Rev. Earth Planetary Sci., 42, 341-
27 379, 10.1146/annurev-earth-060313-054843, 2014.

28 Pope, C. A., III, Thun, M. J., Namboodiri, M. M., Dockery, D. W., Evans, J. S., and Speizer,
29 F. E.: Particulate air pollution as a predictor of mortality in a prospective study of U.S. adults,
30 Am. J. Respir. Crit. Care Med., 151, 669–674, 1995.

1 Pope, C. A., III, Burnett, R.T., Thun, M.J., Calle, E. E., Krewski, D., Ito, K., Thurston, G. D.:
2 Lung cancer, cardiopulmonary mortality, and long-term exposure to fine particulate air pollu-
3 tion, *JAMA*, 287, 1132-1141, 2002.

4 Pope, C. A., III, Ezzati, M., Dockery, D. W.: Fine-particulate air pollution and life expectancy
5 in the United States, *New Engl. J. Med.*, 360, 376-386, doi:10.1056/NEJMsa0805646, 2009.

6 Prather, M. J., Holmes, C. D., and Hsu, J.: Reactive greenhouse gas scenarios: Systematic
7 exploration of uncertainties and the role of atmospheric chemistry, *Geophys. Res. Lett.*, 39,
8 L09803, doi:10.1029/2012GL051440, 2012.

9 Quaas, J., Ming, Y., Menon, S., Takemura, T., Wang, M., Penner, J. E., Gettelman, A.,
10 Lohmann, U., Bellouin, N., Boucher, O., Sayer, A. M., Thomas, G. E., McComiskey, A.,
11 Feingold, G., Hoose, C., Kristjánsson, J. E., Liu, X., Balkanski, Y., Donner, L. J.,
12 Ginoux, P. A., Stier, P., Grandey, B., Feichter, J., Sednev, I., Bauer, S. E., Koch, D., Grain-
13 ger, R. G., Kirkevåg, A., Iversen, T., Seland, Ø., Easter, R., Ghan, S. J., Rasch, P. J., Morri-
14 son, H., Lamarque, J.-F., Iacono, M. J., Kinne, S., and Schulz, M.: Aerosol indirect effects –
15 general circulation model intercomparison and evaluation with satellite data, *Atmos. Chem.*
16 *Phys.*, 9, 8697-8717, doi:10.5194/acp-9-8697-2009, 2009.

17 Quennehen, B., Raut, J.-C., Law, K. S., Ancellet, G., Clerbaux, C., Kim, S.-W., Lund, M. T.,
18 Myhre, G., Olivié, D. J. L., Safieddine, S., Skeie, R. B., Thomas, J. L., Tsyro, S., Bazu-
19 reau, A., Bellouin, N., Daskalakis, N., Hu, M., Kanakidou, M., Klimont, Z., Kupiainen, K.,
20 Myrokefalitakis, S., Quaas, J., Rumbold, S. T., Schulz, M., Cherian, R., Shimizu, A.,
21 Wang, J., Yoon, S.-C., and Zhu, T.: Multi-model evaluation of short-lived pollutant distribu-
22 tions over East Asia during summer 2008, *Atmos. Chem. Phys. Discuss.*, 15, 11049-11109,
23 doi:10.5194/acpd-15-11049-2015, 2015.

24 Quinn, P. K., Bates, T. S., Baum, E., Doubleday, N., Fiore, A. M., Flanner, M., Fridlind, A.,
25 Garrett, T. J., Koch, D., Menon, S., Shindell, D., Stohl, A., and Warren, S. G.: Short-lived
26 pollutants in the Arctic: their climate impact and possible mitigation strategies, *Atmos. Chem.*
27 *Phys.*, 8, 1723-1735, doi:10.5194/acp-8-1723-2008, 2008.

28 Ramanathan, V., and Carmichael, G.: Global and regional climate changes due to black car-
29 bon, *Nature Geosci.*, 1, 221-222, 2008.

1 Reddy, M. S., and Boucher, O.: Climate impact of black carbon emitted from energy con-
2 sumption in the world's regions, *Geophys. Res. Lett.*, 34, L11802,
3 doi:10.1029/2006GL028904, 2007.

4 Reisinger, A., Maltehausen, M., Manning, M. and Bodeker, G.: Uncertainties of global warm-
5 ing metrics: CO₂ and CH₄, *Geophys. Res. Lett.*, 47, L14707, doi:10.1029/2010GL043803,
6 2010.

7 Rogelj, J., Schaeffer, M., Meinshausen, M., Shindell, D. T., Hare, W., Klimont, Z., Velders,
8 G. J. M., Amann, M., and Schellnhuber, H. J.: Disentangling the effects of CO₂ and short-
9 lived climate forcer mitigation, *Proc. Natl. Acad. Sci. USA*, 111, 16325-16330, doi:
10 10.1073/pnas.1415631111, 2014.

11 Rypdal, K., Rive, N., Berntsen, T., Fagerli, H., Klimont, Z., Mideksa, T. K., and Fuglestvedt,
12 J.: Climate and air quality-driven scenarios of ozone and aerosol precursor abatement, *Envi-
13 ron. Sci. Policy*, 12, 855-869, 2009a.

14 Rypdal, K., Rive, N., Berntsen, T., Klimont, Z., Mideksa, T.K., Myhre, G., and Bieltvedt
15 Skeie, R.: Costs and global impacts of black carbon abatement strategies. *Tellus Ser. B*, 61,
16 625-641, 2009b.

17 Samset, B. H., Myhre, G., Schulz, M., Balkanski, Y., Bauer, S., Berntsen, T. K., Bian, H.,
18 Bellouin, N., Diehl, T., Easter, R. C., Ghan, S. J., Iversen, T., Kinne, S., Kirkevag, A., La-
19 marque, J. F., Lin, G., Liu, X., Penner, J. E., Seland, O., Skeie, R. B., Stier, P., Takemura, T.,
20 Tsigaridis, K. and Zhang, K.: Black carbon vertical profiles strongly affect its radiative forc-
21 ing uncertainty, *Atmos. Chem. Phys.*, 13, 2423-2434, 2013.

22 Samset, B. H., Myhre, G., Herber, A., Kondo, Y., Li, S.-M., Moteki, N., Koike, M., Oshi-
23 ma, N., Schwarz, J. P., Balkanski, Y., Bauer, S. E., Bellouin, N., Berntsen, T. K., Bian, H.,
24 Chin, M., Diehl, T., Easter, R. C., Ghan, S. J., Iversen, T., Kirkevåg, A., Lamarque, J.-F.,
25 Lin, G., Liu, X., Penner, J. E., Schulz, M., Seland, Ø, Skeie, R. B., Stier, P., Takemura, T.,
26 Tsigaridis, K., and Zhang, K.: Modelled black carbon radiative forcing and atmospheric life-
27 time in AeroCom Phase II constrained by aircraft observations, *Atmos. Chem. Phys.*, 14,
28 12465-12477, 2014.

1 Samset, B. H., and Myhre, G.: Climate response to externally mixed black carbon as a func-
2 tion of altitude, *Journal of Geophysical Research-Atmospheres*, 120, 2913-2927,
3 doi:10.1002/2014jd022849, 2015.

4 Schmale, J., Shindell, D., von Schneidemesser, E., Chabay, I., and Lawrence, M.: Air pollu-
5 tion: Clean up our skies, *Nature*, 515, 335-337, 2014.

6 Schulz, M., Olivié, D., Tsyro, S., Kanakidou, M., Myriokefalitakis, S., Daskalakis, N., Im, U.,
7 Fanourgakis, G., Hodnebrog, Ø., Skeie, R., Lund, M., Myhre, G., Bellouin, N., Rumbold, S.,
8 Collins, B., Cherian, R., and Quaas, J.: ECLIPSE Deliverable 2.1: Report on model accuracy,
9 available from <http://eclipse.nilu.no/language/en-GB/ProjectOverview/Deliverables.aspx>.

10 Shindell, D. T.: The social cost of atmospheric release, *Climatic Change*, 130, 313–326, doi:
11 [10.1007/s10584-015-1343-0](https://doi.org/10.1007/s10584-015-1343-0), 2015.

12 Shindell, D. T.: Evaluation of the absolute regional temperature potential, *Atmos. Chem.*
13 *Phys.*, 12, 7955-7960, doi:10.5194/acp-12-7955-2012, 2012.

14 Shindell, D. T., Chin, M., Dentener, F., Doherty, R. M., Faluvegi, G., Fiore, A. M., Hess, P.,
15 Koch, D. M., MacKenzie, I. A., Sanderson, M. G., Schultz, M. G., Schulz, M., Stevenson, D.
16 S., Teich, H., Textor, C., Wild, O., Bergmann, D. J., Bey, I., Bian, H., Cuvelier, C., Duncan,
17 B. N., Folberth, G., Horowitz, L. W., Jonson, J., Kaminski, J. W., Marmer, E., Park, R., Prin-
18 gle, K. J., Schroeder, S., Szopa, S., Takemura, T., Zeng, G., Keating, T. J., and Zuber, A.: A
19 multi-model assessment of pollution transport to the Arctic, *Atmos. Chem. Phys.*, 8, 5353-
20 5372, 2008.

21 Shindell, D., and Faluvegi, G.: Climate response to regional radiative forcing during the twen-
22 tieth century, *Nature Geosci.*, 2, 294-300, doi:10.1038/ngeo473, 2009.

23 Shindell, D., Schulz, M., Ming, Y., Takemura, T., Faluvegi, G., and Ramaswamy, R.: Spatial
24 scales of climate response to inhomogeneous radiative forcing, *J. Geophys. Res.*, 115,
25 D19110, doi:10.1029/2010JD014108, 2010.

26 Shindell, D., Kuylenstierna, J. C. I., Vignati, E., van Dingenen, R., Amann, M., Klimont, Z.,
27 Anenberg, S. C., Muller, N., Janssens-Maenhout, G., Raes, F., Schwartz, J., Faluvegi, G.,
28 Pozzoli, L., Kupiainen, K., Höglund-Isaksson, L., Emberson, L., Streets, D., Ramanathan, V.,
29 Hicks, K., Kim Oanh, N. T., Milly, G., Williams, M., Demkine, V., and Fowler, D.: Simulta-

1 neously mitigating near-term climate change and improving human health and food security,
2 Science, 335, 183-189, doi:10.1126/science.1210026, 2012.

3 [Shindell, D. T., Lamarque, J.-F., Schulz, M., Flanner, M., Jiao, C., Chin, M., Young, P. J.,](#)
4 [Lee, Y. H., Rotstayn, L., Mahowald, N., Milly, G., Faluvegi, G., Balkanski, Y., Collins, W. J.,](#)
5 [Conley, A. J., Dalsoren, S., Easter, R., Ghan, S., Horowitz, L., Liu, X., Myhre, G., Nagashima, T., Naik, V., Rumbold, S. T., Skeie, R., Sudo, K., Szopa, S., Takemura, T., Voulgarakis, A., Yoon, J.-H., and Lo, F.: Radiative forcing in the ACCMIP historical and future climate simulations, Atmos. Chem. Phys., 13, 2939-2974, doi:10.5194/acp-13-2939-2013, 2013.](#)
6
7
8

9
10

11 Shine, K. P.: The global warming potential: the need for an interdisciplinary retrial. An editorial comment, Climatic Change, 96, 467-472, 2009.

13 Shine, K. P., Berntsen, T. K., Fuglestvedt, J. S., Bieltvedt Skeie, R. and Stuber, N.: Comparing the climate effect of emissions of short- and long-lived climate agents. Philos. Trans. R. Soc., A, 365, 1903–1914, 2007.

16 Shine, K. P., Berntsen, T. K., Fuglestvedt, J. S., and Sausen, R.: Scientific issues in the design 17 of metrics for inclusion of oxides of nitrogen in global climate agreements, Proc. Natl. Acad. 18 Sci. USA, 102, 15768–15773, 2005.

19 Shine, K. P., Allan, R. P., Collins, W. J., and Fuglestvedt, J. S.: Metrics for linking emissions 20 of gases and aerosols to global precipitation changes, Earth Syst. Dynam. Discuss., 6, 719- 21 760, doi:10.5194/esdd-6-719-2015, 2015. [Accepted for final publication.](#)

22 Simpson, D., Benedictow, A., Berge, H., Bergstrom, R., Emberson, L. D., Fagerli, H., 23 Flechard, C. R., Hayman, G. D., Gauss, M., Jonson, J. E., Jenkin, M. E., Nyiri, A., Richter, 24 C., Semeena, V. S., Tsyro, S., Tuovinen, J. P., Valdebenito, A., and Wind, P.: The EMEP 25 MSC-W chemical transport model - technical description, Atmospheric Chemistry and Physics, 12, 7825-7865, 10.5194/acp-12-7825-2012, 2012.

27 Sitch, S., Cox, P. M., Collins, W. J., and Huntingford, C.: Indirect radiative forcing of climate 28 change through ozone effects on the land-carbon sink, Nature, 448, 791-794, 2007.

1 Skeie, R. B., Berntsen, T. K., Myhre, G., Tanaka, K., Kvalevag, M. M., and Hoyle, C. R.:
2 Anthropogenic radiative forcing time series from pre-industrial times until 2010, *Atmos.*
3 *Chem. Phys.*, 11, 11827-11857, doi:10.5194/acp-11-11827-2011, 2011.

4 Skeie, R. B., Berntsen, T., Aldrin, M., Holden, M., and Myhre, G.: A lower and more con-
5 strained estimate of climate sensitivity using updated observations and detailed radiative forc-
6 ing time series, *Earth Syst. Dynam.*, 5, 139-175, doi:10.5194/esd-5-139-2014, 2014.

7 Skodvin, T., and Fuglestvedt, J. S.: A comprehensive approach to climate change: Political
8 and scientific considerations, *Ambio*, 26, 351-358, 1997.

9 Smith, S. J., Edmonds, J., Hartin, C. A., Mundra, A., and Calvin, K.: Near-term acceleration
10 in the rate of temperature change, *Nature Clim. Change*, 5, 333–336,
11 doi:10.1038/nclimate2552, 2015.

12 Stevens, B., Giorgetta, M., Esch, M., Mauritsen, T., Crueger, T., Rast, S., Salzmann, M.,
13 Schmidt, H., Bader, J., Block, K., Brokopf, R., Fast, I., Kinne, S., Kornblueh, L., Lohmann,
14 U., Pincus, R., Reichler, T., and Roeckner, E.: Atmospheric component of the MPI-M Earth
15 System Model: ECHAM6-HAM2, *Journal of Advances in Modeling Earth Systems*, 5, 146-
16 172, doi:10.1002/jame.20015, 2013.

17 Stjern, C. W., Stohl, A., and Kristjánsson, J. E.: Have aerosols affected trends of visibility and
18 precipitation in Europe? *J. Geophys. Res.*, 116, D02212, doi:10.1029/2010JD014603, 2011.

19 Stohl, A., and Trickl, T.: A textbook example of long-range transport: Simultaneous observa-
20 tion of ozone maxima of stratospheric and North American origin in the free troposphere over
21 Europe, *J. Geophys. Res.*, 104, 30445-30462, 1999.

22 Stohl, A., Hittenberger, M., and Wotawa, G.: Validation of the Lagrangian particle dispersion
23 model FLEXPART against large scale tracer experiments. *Atmos. Environ.* **32**, 4245-4264,
24 1998.

25 Stohl, A., Forster, C., Frank, A., Seibert, P., and Wotawa, G.: Technical note: The Lagrangian
26 particle dispersion model FLEXPART version 6.2, *Atmos. Chem. Phys.*, 5, 2461-2474, 2005.

27 Stohl, A., Klimont, Z., Eckhardt, S., Kupiainen, K., Shevchenko, V. P., Kopeikin, V. M., and
28 Novigatsky, A. N.: Black carbon in the Arctic: the underestimated role of gas flaring and resi-
29 dential combustion emissions, *Atmos. Chem. Phys.*, 13, 8833-8855, doi:10.5194/acp-13-
30 8833-2013, 2013.

1 Tanaka, K., O'Neill, B. C., Rokityanskiy, D., Obersteiner, M., and Tol, R. S. J.: Evaluating
2 Global Warming Potentials with historical temperature, *Climatic Change*, 96, 443-466, 2009.

3 Tanaka, K., Johansson, D. J. A., O'Neill, B. C., and Fuglestvedt, J. S.: Emission metrics under
4 the 2°C climate stabilization target, *Climatic Change*, 117, 933-941, doi:10.1007/s10584-013-
5 0693-8, 2013.

6 Taylor, K. E., Stouffer, R. J., and Meehl, G. A.: An overview of CMIP5 and the experiment
7 design, *Bull. Am. Meteorol. Soc.*, 93, 485–498, doi:10.1175/BAMS-D-11-00094.1, 2012.

8 Tragou, E., and Lascaratos, A.: Role of aerosols on the Mediterranean solar radiation, *J. Ge-
9 ophys. Res.*, 108, 3025, doi: 10.1029/2001JC001258, 2003.

10 Tsigaridis, K., Daskalakis, N., Kanakidou, M., Adams, P. J., Artaxo, P., Bahadur, R., Balkan-
11 ski, Y., Bauer, S. E., Bellouin, N., Benedetti, A., Bergman, T., Berntsen, T. K., Beukes, J. P.,
12 Bian, H., Carslaw, K. S., Chin, M., Curci, G., Diehl, T., Easter, R. C., Ghan, S. J.,
13 Gong, S. L., Hodzic, A., Hoyle, C. R., Iversen, T., Jathar, S., Jimenez, J. L., Kaiser, J. W.,
14 Kirkevåg, A., Koch, D., Kokkola, H., Lee, Y. H., Lin, G., Liu, X., Luo, G., Ma, X.,
15 Mann, G. W., Mihalopoulos, N., Morcrette, J.-J., Müller, J.-F., Myhre, G., Myrioke-
16 falitakis, S., Ng, N. L., O'Donnell, D., Penner, J. E., Pozzoli, L., Pringle, K. J., Russell, L. M.,
17 Schulz, M., Sciare, J., Seland, Ø., Shindell, D. T., Sillman, S., Skeie, R. B., Spracklen, D.,
18 Stavrakou, T., Steenrod, S. D., Takemura, T., Tiitta, P., Tilmes, S., Tost, H., van Noije, T.,
19 van Zyl, P. G., von Salzen, K., Yu, F., Wang, Z., Wang, Z., Zaveri, R. A., Zhang, H.,
20 Zhang, K., Zhang, Q., and Zhang, X.: The AeroCom evaluation and intercomparison of or-
21 ganic aerosol in global models, *Atmos. Chem. Phys.*, 14, 10845-10895, doi:10.5194/acp-14-
22 10845-2014, 2014.

23 UNEP/WMO: Integrated Assessment of Black Carbon and Tropospheric Ozone. United Na-
24 tions Environment Programme (UNEP), Nairobi, Kenya, 2011.

25 Unger, N., Bond, T. C., Wang, J. S., Koch, D. M., Menon, S., Shindell, D. T., and Bauer, S.:
26 Attribution of climate forcing to economic sectors, *Proc. Natl. Acad. Sci. USA*, 107, 3382–
27 3387, 2010.

28 van der Werf, G. R., Randerson, J. T., Giglio, L., Collatz, G. J., Mu, M., Kasibhatla, P. S.,
29 Morton, D. C., DeFries, R. S., Jin, Y., and van Leeuwen, T. T.: Global fire emissions and the

1 contribution of deforestation, savanna, forest, agricultural, and peat fires (1997–2009), Atmos.
2 Chem. Phys., 10, 11707-11735, doi:10.5194/acp-10-11707-2010, 2010.

3 Wang, S. X., Zhao, B., Cai, S. Y., Klimont, Z., Nielsen, C. P., Morikawa, T., Woo, J. H.,
4 Kim, Y., Fu, X., Xu, J. Y., Hao, J. M., and He, K. B.: Emission trends and mitigation options
5 for air pollutants in East Asia, Atmos. Chem. Phys., 14, 6571-6603, doi:10.5194/acp-14-
6 6571-2014, 2014.

7 Wang, R., Balkanski, Y., Boucher, O., Ciais, P., Penuelas, J., and Tao, S.: Significant contri-
8 bution of combustion-related emissions to the atmospheric phosphorus budget, Nature Ge-
9 osci., 8, 48-54, doi:10.1038/ngeo2324, 2015.

10 West, J. J., Fiore, A. M., Horowitz, L. W., Mauzerall, D. L.: Global health benefits of mitigat-
11 ing ozone pollution with methane emission controls, Proc. Natl. Acad. Sci. USA, 103, 3988-
12 3993, 2006.

13 WHO: WHO Air quality guidelines for particulate matter, ozone, nitrogen dioxide and sulfur
14 dioxide: Global update 2005 - Summary of risk assessment, 2006
15 (http://www.euro.who.int/_data/assets/pdf_file/0005/78638/E90038.pdf?ua=1, accessed
16 2014-04-22).

17 WHO: Health effects of black carbon, Joint WHO/UNECE Task Force on Health Aspects of
18 Air Pollutants under UNECE's Long-Range Transboundary Air Pollution Convention
19 (LRTAP), World Health Organization, Regional Office for Europe, Copenhagen, 2012
20 (http://www.unece.org/fileadmin/DAM/env/documents/2012/air/Health_Effects_of_Black_Ca
21 rbon_report.pdf), accessed 14 May 2013.

22 WHO: Review of evidence on health aspects of air pollution – REVIHAAP project: final
23 technical report, 2013. (<http://www.euro.who.int/en/health-topics/environment-and-health/air->
24 [publications/2013/review-of-evidence-on-health-aspects-of-air-pollution-revihaap-](http://www.euro.who.int/en/health-topics/environment-and-health/air-quality/publications/2013/review-of-evidence-on-health-aspects-of-air-pollution-revihaap-)
25 [project-final-technical-report](http://www.euro.who.int/en/health-topics/environment-and-health/air-quality/publications/2013/review-of-evidence-on-health-aspects-of-air-pollution-revihaap-project-final-technical-report), accessed 29-04-2014)

26 Wild, O., Prather, M. J., and Akimoto, H.: Indirect long-term global radiative cooling from
27 NO_x emissions, Geophys. Res. Lett., 28, 1719–1722, 2001.

28 Yttri, K. E., Myhre, C. L., Eckhardt, S., Fiebig, M., Dye, C., Hirdman, D., Stroem, J.,
29 Klimont, Z., and Stohl, A.: Quantifying black carbon from biomass burning by means of

1 levoglucosan - a one-year time series at the Arctic observatory Zeppelin, *Atmos. Chem.*
2 *Phys.*, 14, 6427-6442, 10.5194/acp-14-6427-2014, 2014.

3 Zaveri, R. A., Easter, R. C., Fast, J. D., and Peters, L. K.: Model for Simulating Aerosol Inter-
4 actions and Chemistry (MOSAIC), *J. Geophys. Res.*, 113, D1320429,
5 doi:10.1029/2007jd008782, 2008.

6 Zhang, K., O'Donnell, D., Kazil, J., Stier, P., Kinne, S., Lohmann, U., Ferrachat, S., Croft, B.,
7 Quaas, J., Wan, H., Rast, S., and Feichter, J.: The global aerosol-climate model ECHAM-
8 HAM, version 2: sensitivity to improvements in process representations, *Atmos. Chem. Phys.*,
9 12, 8911-8949, doi:10.5194/acp-12-8911-2012, 2012.

Zhao, B., Wang, S. X., Liu, H., Xu, J. Y., Fu, K., Klimont, Z., Hao, J. M., He, K. B., Co-
fala, J., and Amann, M.: NO_x emissions in China: historical trends and future perspectives,
Atmos. Chem. Phys., 13, 9869-9897, doi:10.5194/acp-13-9869-2013, 2013.

10

1 Table 1. Air quality standards for Europe (European Union reference values), WHO air quality
 2 guidelines (AQG), US-EPA National Ambient Air Quality Standards (NAAQS) and the
 3 Environmental Quality Standards (EQS) and guideline values for air pollutants in Japan. Val-
 4 ues in brackets give time period for which the guideline is defined.

Pollutants	EU reference levels ^a	WHO AQG ^b	USEPA NAAQS ^c	Japan EQS ^d
PM _{2.5}	20 µg/m ³ (year)	10 µg/m ³ (year)	12 µg/m ³ (year)	15 µg/m ³ (year)
PM ₁₀	40 µg/m ³ (day)	20 µg/m ³ (year)	150 µg/m ³ (day)	100 µg/m ³ (day, SPM ^e)
O ₃	120 µg/m ³ (8-hour)	100 µg/m ³ (8-hour)	0.075 ppm(8-hour)	118 µg/m ³ (1-hour ^f)
NO ₂	40 µg/m ³ (year)	40 µg/m ³ (year)	53 ppb (year)	75-113 µg/m ³ (1-hour)
SO ₂	125 µg/m ³ (day)	20 µg/m ³ (day)	75 ppb (1-hour)	105 µg/m ³ (1-day)
CO	10 mg/m ³ (8-hour)	10 mg/m ³ (8-hour)	9 ppm (8-hour)	10 ppm (1-hour)

5 ^aEEA 2013, Indicator CSI 004

6 ^bWHO Air Quality Guidelines (WHO 2006)

7 ^cUS-EPA National Ambient Air Quality Standards (<http://www.epa.gov/air/criteria.html#3>, accessed
 8 16/4/2014).

9 ^dEnvironmental Quality Standards (EQS) and guideline values for air pollutants in Japan (Kawamoto
 10 et al. 2011).

11 ^e100% efficiency cut-off at 10 µm while PM₁₀ is defined as 50% efficiency cut-off at 10 µm aerody-
 12 namic diameter (Kawamoto et al., 2011)

13 ^fPhotochemical oxidants (Ox) (Kawamoto et al., 2011)

1 Table 2. Overview of the ECLIPSE models and how they were set up for the years 2008-
 2 2009.

Model Name	Model Type ¹	Horizontal/vertical resolution	Meteorological fields	Periods simulated / output	References
				temporal resolution	
FLEXPART	LPDM	Meteorological input $1^\circ \times 1^\circ$, 92L	ECMWF analyses	operational 2008-2009 3h	Stohl et al. (1998, 2005)
OsloCTM2	CTM	$2.8^\circ \times 2.8^\circ$, 60L	ECMWF IFS forecasts	2008-2009 3h	Myhre et al. (2009), Skeie et al. (2011)
EMEP	CTM	$1^\circ \times 1^\circ$, 20L	ECMWF operational	2008-2009, 24 h	Simpson et al. (2012)
TM4-ECPL	CTM	$2^\circ \times 3^\circ$, 34L	ECMWF ERA-interim	2008-2009 24h	Kanakidou et al. (2012); Daskalakis et al. (2015)
WRF-CMAQ	CTM	50x50 km, 23L	NCEP	2008, 24 h	Im et al. (2013)
WRF-Chem	CTM	50 km x 50 km, 49L	Nudged to FNL	March-August 2008 3h	Grell et al. (2005), Zaveri et al. (2008)
NorESM	ESM	$1.9^\circ \times 2.5^\circ$, 26L	Internal, SST prescribed	observed 2008-2009 3h	Kirkevåg et al. (2013), Bentsen et al. (2013)
ECHAM6-HAM2	ESM	$1.8^\circ \times 1.8^\circ$, 31L	ECMWF reanalysis	March-August, 2008, 1h	Stevens et al. (2013), Zhang et al. (2012)
HadGEM3	ESM	$1.9^\circ \times 1.3^\circ$, 63L	ECMWF ERA-interim	March-June, 2008, January, 2009 November, 2009 2h	Hewitt et al. (2011), Mann et al. (2010)
CESM-CAM4	ESM	$1.9^\circ \times 2.5^\circ$, 26L	Internal	Was not evaluated for 2008-2009; only used for 2000-2050 simulations	Gent et al. (2011)

3 ¹Chemistry transport model (CTM), Lagrangian particle dispersion model (LPDM), Earth system model (ESM).
 4
 5

1 Table 3: Top-17 mitigation measures contributing together more than 80% of the climate ben-
 2 efit. Measures are ranked by importance starting from the top. Methane measures contribute
 3 about 47% of the benefits according to the GTP₂₀ metric, while 33% are attributed to BC fo-
 4 cused measures; 20% are contributed by measures not listed here.

Methane measures	Measures targeting BC reduction
Oil and gas industry: Recovery and use (rather than venting or flaring) of associated gas	Oil and gas industry: Improving efficiency and reducing gas flaring
Oil and gas industry (unconventional): Reducing emissions from unintended leaks during production (extraction) of shale gas	Transport: Eliminating high emitting vehicles (super-emitters)
Coal mining: Reducing (oxidizing) emissions released during hard coal mining (ventilation air CH ₄)	Residential-commercial: Clean biomass cooking stoves
Waste: Municipal waste – waste paper separation, collection, and recycling	Residential-commercial: Replacement of kerosene wick lamps with LED lamps
Waste: Municipal food waste separation, collection and treatment in anaerobic digestion (biogasification) plants	Transport: Widespread Euro VI emission standards (incl. particle filters) on diesel vehicles
Coal mining: Hard and brown coal -pre-mining emissions – Degasification	Industrial processes: Modernized (mechanized) coke ovens
Gas distribution: Replacement of grey cast iron gas distribution network	Agriculture: Effective ban of open field burning of agricultural residues
Waste: Industrial solid waste (food, wood, pulp and paper, textile) – recovery and incineration	
Waste: Wastewater treatment from paper and pulp, chemical, and food industries - anaerobic treatment in digester, reactor or deep lagoon with gas recovery, upgrading and use. For residential wastewater centralized collection with anaerobic secondary and/or tertiary treatment (incl. treatment with bacteria and/or flaring of residual CH ₄)	
Oil and gas industry (conventional): Reducing emissions from unintended leaks during production (extraction)	

1 Table 4: Relative differences (%) in O₃ and PM_{2.5} surface concentrations between simulations
 2 using the mitigation (MIT) and current legislation (CLE) scenarios (given as $100 \times (\text{MIT} - \text{CLE})/\text{CLE}$) for several regions and for the final decade of the simulations (2041-2050). The
 3 regions correspond to the boxes shown in Figure 10. Except for the Mediterranean, only the
 4 land-based grid cells inside the boxes were used. Mean values and full model ranges for the
 5 three models NorESM, HadGEM and OsloCTM2 are reported. To exclude effects of changes
 6 in sea salt and dust emissions caused by natural variability and emission responses to forced
 7 changes in meteorological conditions, sea salt and dust concentrations were kept at CLE lev-
 8 els for the purpose of these calculations.

Region	O ₃ mean (%)	O ₃ range (%)	PM _{2.5} mean (%)	PM _{2.5} range (%)
Northern Europe	-13.7	[-15.9, -11.1]	-12.5	[-17.6, -7.3]
Southern Europe	-14.9	[-17.0, -12.3]	-2.3	[-3.1, -1.5]
Mediterranean	-15.1	[-17.8, -12.6]	-1.6	[-2.3, -1.0]
Eastern China	-19.3	[-24.4, -16.0]	-16.3	[-23.2, -11.8]
Western China	-15.8	[-17.9, -12.8]	-2.2	[-4.3, -1.0]
India	-17.1	[-22.8, -12.7]	-19.8	[-22.5, -17.9]
Eastern United States	-13.6	[-15.7, -10.3]	-8.8	[-11.7, -3.6]
Western United States	-14.3	[-16.1, -11.5]	-4.5	[-8.7, -1.4]

10
 11

1 **Figure captions**

2 Figure 1: Schematic of the ECLIPSE overall methodology. Numbers in brackets correspond
3 to section numbers in this paper.

4

5 Figure 2. Global annual anthropogenic emissions of CO₂, CH₄ and key air pollutants (SO₂,
6 NO_x and BC) for the current legislation (CLE), no further controls (NFC) and ECLIPSE
7 SLCP mitigation scenario. Units are Gt for CO₂ and Mt for the SLCPs. Also shown for com-
8 parison is the range of the RCP emission scenarios (grey shading).

9

10 Figure 3: Box and whiskers plots showing the frequency distribution of measured and mod-
11 eled CO, NO₂, O₃ and SO₂ mixing ratios or concentrations representative for background
12 stations in urban and rural areas in East Asia during August and September 2008 (two left
13 panel columns) and for rural background stations in Europe for winter (December-February,
14 DJF) and summer (June-August, JJA) 2008 (two right panel columns). Circles and central
15 lines show the means and the medians, respectively; box edges represent the 25th and the 75th
16 percentiles. For East Asia, results are averaged over several sites: Beijing, Inchon and Seoul
17 for the urban areas, Gosan, Kunsan, Kangwha, Mokpo and Taean for rural areas. Results for
18 individual sites can be found in Quennehen et al. (2015). For Europe, daily mean observed
19 values are averaged over all stations of the EMEP network with available data. Model data are
20 treated like the observations and only the days with available observations are taken into ac-
21 count.

22

23 Figure 4: Comparison of satellite-derived (MODIS) and modelled aerosol optical depth
24 (AOD) at wavelengths of 550 nm over Eastern China and Northern India (for area definition,
25 see Quennehen et al., 2015) in August-September 2008, and Europe (14.5°W-34.5°E, 35.5°-
26 74.5°N) in winter (December-February, DJF) and summer (June-August, JJA) of 2008. Mean
27 values (circles), medians (central lines), 25th and 75th percentiles (boxes) and range of other
28 data excluding outliers (whiskers) are shown.

29

1 Figure 5. Monthly (month is displayed on the abscissa) median observed and modelled BC
2 concentrations for the stations Zeppelin on Svalbard (11.9°E, 78.9°N, top), Pallas in Finland
3 (24.12°E, 67.97°N, middle) and Tiksi in Siberia (128.9° E, 71.6°N, bottom), for late win-
4 ter/spring (left column) and summer/autumn (right column) for the years 2008-2009 (for
5 Tiksi, measured values were available only from July 2009 to June 2010). The red dashed
6 lines connect the observed median values, the light red shaded areas span from the 25th to the
7 75th percentile of the observations. Modelled median values are shown with lines of different
8 colour according to the legend. Notice that different concentration scales are used for individ-
9 ual panels and also for January-May (axis on left hand side) and June-December (axis on right
10 hand side) periods. Modified from Eckhardt et al. (2015).

11

12 Figure 6. Linear trends in (left) surface solar radiation and (right) near-surface temperature
13 increase per decade over continental Europe from the „historical“ simulations in the CMIP5
14 archive contributed by previous versions of three ECLIPSE ESMs. The period 1960–1980 is
15 shown in red, the period 1990–2005 in blue.

16

17 Figure 7: The ECLIPSE estimates of specific radiative forcing (SRF, mWm^{-2} per Tg yr^{-1} of
18 emission rate change) for reductions in the emissions of SO_2 , NO_x , CH_4 , and BC aerosols in
19 Europe, China and the global average, separately for NH summer (Sum., May-October) and
20 NH winter (Win., November-April). Shown are values averaged over all five models, with the
21 error bars indicating the full range of the model estimates. Colours indicate the contribution of
22 different forcing mechanisms. Notice that for CH_4 regionality was not accounted for because,
23 due to its longer lifetime, forcing does not depend on the emission region.

24

25 Figure 8: GTP₂₀ values for SLCPs, relative to an equal mass emission of CO_2 , for all regions
26 and seasons, decomposed by processes and based on the RF values shown in Fig. 7. The re-
27 gions included are Europe (EUR), China (CHN), global (GLB), and the shipping sector con-
28 sidered separately (SHP), all for both NH summer (s, May-October) and NH winter (w, No-
29 vember-April). Uncertainty bars reflect model spread. Only two models calculated effects of
30 emissions from shipping and there no uncertainty range is given.

31

1 Figure 9: CO₂-equivalent emissions (Gt) determined with the GTP₂₀ metric, as a function of
2 time, for global emissions of the current legislation (CLE) and SLCP mitigation (MIT) sce-
3 narios. Lines show the values for individual forcing components (black CO₂, red CH₄, blue
4 other SLCPs), while the shaded areas show the total CO₂ equivalent emissions from all
5 SLCPs and CO₂. The blue shading indicates the mitigation potential of MIT, compared to
6 CLE.

7

8 Figure 10: Relative difference maps (in %) for the surface concentrations of O₃ and PM_{2.5}, as
9 obtained from simulations based on the emission mitigation (MIT) scenario and the current
10 legislation (CLE) scenario, i.e. $(100 \times (\text{MIT}-\text{CLE})/\text{CLE})$. Shown are mean concentration dif-
11 ferences for the period 2041-2050 and averaged over model results from OsloCTM2,
12 NorESM and HadGEM. The black boxes define the regions used in Table 4.

13

14 Figure 11: Loss of statistical life expectancy (months) due to the exposure to PM_{2.5} in Europe,
15 for EU-28 and Non-EU countries (upper panel) and for China and India (lower panel). The
16 black bars give the values for the mitigation (MIT) scenario, whereas the blue increments
17 show the difference to the current legislation (CLE) scenario. For India and China, green bars
18 indicate the gains from the implementation of the CLE scenario.

19

20 Figure 12: Global mean annual average changes in (a) surface temperature and (b) precipita-
21 tion averaged over all land areas excluding ice sheets, from the four ESMs (one was run only
22 for BC), for a complete removal of all land-based emissions of a particular species, compared
23 to the ECLIPSE version 4 baseline emissions for the year 2008 (CESM-CAM4 used the year
24 2000). The plot shows results averaged over 50-year model simulations; for HadGEM and
25 NorESM (CESM-CAM4), two (three) ensemble members each were run for BC (named e1,
26 e2, e3 in the labels). Error bars are 95% confidence intervals in the mean, based on 50 annual
27 means from each model and thus only reflect the uncertainty of the mean caused by natural
28 variability within each model.

29

1 Figure 13: Annual mean changes in zonal-mean surface temperature (left panels a, c, e) and
2 precipitation (right panels b, d, f) for a complete removal of all land-based emissions of (a-b)
3 SO_2 , (c-d) BC, and (e-f) OA. For BC, some models ran two or three ensemble members (e1,
4 e2, e3). Notice the differences in scales between different panels.

5

6 Figure 14: Time evolution of differences in global mean surface temperature between transi-
7 ent simulations following the mitigation (MIT) and the current legislation (CLE) scenario, i.e.
8 (MIT-CLE), for the four ECLIPSE models. Negative values mean that temperatures are lower
9 in the MIT than in the CLE scenario. The ensemble means for each model are shown as thick
10 lines, whereas the individual ensemble members are shown with thin lines.

11

12 Figure 15: Seasonal and annual mean differences in surface temperatures (in K) in various
13 regions (a-f) and for the whole globe (g) between transient simulations of the mitigation
14 (MIT) and the current legislation (CLE) scenario, i.e. (MIT-CLE), averaged over the last 10
15 years of the simulations (2041-2050). Regions are defined as (a) 45°N - 65°N , 10°W - 65°E , (b)
16 30°N - 45°N , 10°W - 65°E , (c) 7°N - 35°N , 68°E - 90°E , (d) 24°N - 48°N , 80°E - 132°E , (e) 60°N -
17 90°N , (f) 30°N - 60°N , 120°W - 50°W . Results are shown for the four ECLIPSE models individ-
18 ually and for the multi-model mean. Negative values mean that temperatures are lower in the
19 MIT than in the CLE scenario. Error bars on the model mean values show the standard devia-
20 tions of the individual model results.

21

22 Figure 16: Annual average differences in (top) surface temperature and (bottom) precipitation
23 over Europe between the transient simulations based on the mitigation (MIT) and the current
24 legislation (CLE) scenario, averaged over the last 10 years of the simulations (2041-2050).
25 Stippling shows where all four models agree on the sign of the response. The thick horizontal
26 line distinguishes the southern and northern Europe boxes. In the top panel, negative values
27 mean that temperature in the MIT scenario is lower than in the CLE scenario; in the bottom
28 panel, positive values mean that there is more precipitation in the MIT than in the CLE sce-
29 nario.

30

1 Figure 17: Seasonal and annual mean differences in precipitation (in mm y^{-1}) over Northern
2 (a) and Southern (b) Europe between transient simulations of the mitigation (MIT) scenario
3 and the current legislation (CLE) scenario, averaged over the last 10 years of the simulations
4 (2041-2050). Regions are defined as for Figure 15 (a, b). Results are shown for the four
5 ECLIPSE models individually and for the multi-model mean. Positive values mean that MIT
6 simulations have more precipitation than CLE simulations. Error bars on the model mean val-
7 ues show the standard deviations of the individual model results.

8

9 Figure 18: Time evolution of differences in global mean surface temperature between transi-
10 ent simulations following the mitigation (MIT) and the current legislation (CLE) scenario
11 (i.e., MIT-CLE), as simulated by the CESM-CAM4 model with a slab-ocean representation.
12 One experiment (red lines) included all emission reductions of the MIT scenario, whereas
13 another experiment (blue lines) included only the CH_4 emission reductions of the MIT scenar-
14 io. Ensemble mean results are shown with thick lines, individual ensemble members with thin
15 lines.

16

17 Figure 19: Time evolution of difference in global mean surface temperature for the MIT and
18 CLE scenario calculated with the ARTP-based method. Solid lines are the total difference
19 (MIT-CLE), while the dashed lines give the responses to CH_4 mitigation only. The solid lines
20 can be compared directly with the results from the transient model simulations shown in Fig-
21 ure 14.

22

23 Figure 20: Temperature changes (MIT-CLE) in four latitude bands (Arctic, Northern Hemis-
24 phere Mid-Latitudes (NHML), Tropics and Southern Hemisphere) calculated with the ESMs
25 and ARTP-based method. Mean changes over the 2031-2050 period are shown for individual
26 models and the mean over the models. Notice the different temperature scales.

27

28 Figure 21: Annual mean surface temperature changes estimated by the ARTP method. Panels
29 a-c: Changes in four latitude bands due to ECLIPSE mitigation scenario (MIT-CLE) for miti-
30 gation in Europe (a), China (b) and globally (c). Panel d: Arctic temperature changes due to

1 mitigation of individual components from Europe. CE-Aero: Co-emitted aerosol (precursor)
2 species (OA, SO₂ and NH₃), OP: Ozone precursors (NO_x, CO and NMVOCs). Note the dif-
3 ferent scales on the vertical axes.

4

5 Figure 22: Total Arctic surface temperature change (K) by global mitigation of residential
6 burning (heating and cooking) in the ECLIPSE emission scenario (MIT-CLE), as obtained
7 with the ARTP method.