



UNIVERSITY OF LEEDS

This is a repository copy of *Declining uncertainty in transient climate response as CO2 forcing dominates future climate change*.

White Rose Research Online URL for this paper:  
<http://eprints.whiterose.ac.uk/85097/>

Version: Accepted Version

---

**Article:**

Myhre, G, Boucher, O, Bréon, F-M et al. (2 more authors) (2015) Declining uncertainty in transient climate response as CO2 forcing dominates future climate change. *Nature Geoscience*, 8 (3). 181 - 185. ISSN 1752-0894

<https://doi.org/10.1038/ngeo2371>

---

**Reuse**

Unless indicated otherwise, fulltext items are protected by copyright with all rights reserved. The copyright exception in section 29 of the Copyright, Designs and Patents Act 1988 allows the making of a single copy solely for the purpose of non-commercial research or private study within the limits of fair dealing. The publisher or other rights-holder may allow further reproduction and re-use of this version - refer to the White Rose Research Online record for this item. Where records identify the publisher as the copyright holder, users can verify any specific terms of use on the publisher's website.

**Takedown**

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing [eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk) including the URL of the record and the reason for the withdrawal request.



[eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk)  
<https://eprints.whiterose.ac.uk/>

# Declining uncertainty in transient climate response as CO<sub>2</sub> dominates future climate change

Gunnar Myhre<sup>1</sup>, Olivier Boucher<sup>2</sup>, François-Marie Bréon<sup>3</sup>, Piers Forster<sup>4</sup>, and Drew Shindell<sup>5</sup>

<sup>1</sup>Center for International Climate and Environmental Research – Oslo (CICERO), Oslo, Norway

<sup>2</sup>Laboratoire de Météorologie Dynamique, Institut Pierre Simon Laplace, Centre National de la Recherche Scientifique, Université Pierre et Marie Curie, 75252 Paris Cedex 05, France

<sup>3</sup>Laboratoire des Sciences du Climat et de l'Environnement, IPSL, laboratoire CEA-CNRS-UVSQ, 91191 Gif sur Yvette, France

<sup>4</sup>School of Earth and Environment, University of Leeds, UK

<sup>5</sup>Nicholas School of the Environment, Duke University, Durham, NC 27708, USA

*Carbon dioxide (CO<sub>2</sub>) has been the largest contributor to radiative forcing and surface temperature change over the industrial era but other anthropogenic drivers have had a significant role<sup>1,2</sup>. The large uncertainty in the total forcing makes it difficult to derive climate sensitivity from historical observations<sup>3-7</sup>. Based on data from Intergovernmental Panel of Climate Change (IPCC) reports, we show that the evolution of increased anthropogenic forcing and its reduced relative uncertainty between the Fourth and Fifth Assessment Reports<sup>1,8</sup> can be expected to continue into the future, driven by the greater ease of reducing air pollution than CO<sub>2</sub> emissions, long lifetime of CO<sub>2</sub>, and hence a stronger dominance of CO<sub>2</sub> forcing. Here we present, using a statistical model, that the relative uncertainty in anthropogenic forcing of more than 40% quoted in the latest IPCC report for 2011 will be reduced by almost half by 2030, even without further improvement in scientific understanding. Absolute forcing uncertainty will also decline for the first time assuming projected decreases in aerosols occur. Other factors being equal, this stronger constraint on forcing will bring a significant reduction in the uncertainty of observation-*

27 *based estimates of transient climate response, with a 50% reduction in its uncertainty range*  
28 *expected by 2030.*

29

30 Equilibrium climate sensitivity (ECS) and transient climate response (TCR) are two key  
31 measures that are used to evaluate how much the world might warm. TCR, which corresponds  
32 to the warming at the time of a doubling CO<sub>2</sub> in a 1%- per-year CO<sub>2</sub> increase scenario, is  
33 more policy-relevant than ECS to gauge the strength of climate change over coming decades.  
34 Although there is high confidence in the human contribution to climate change, current IPCC  
35 estimates of TCR show a large uncertainty range of 1.0 to 2.5 °C (5-95% confidence interval)  
36 for a doubling of CO<sub>2</sub> (ref. <sup>2</sup>), which translates into an equivalent range of 0.27 to 0.68  
37 °C (W m<sup>-2</sup>)<sup>-1</sup> for the normalized definition of TCR that we adopt in this paper. Different  
38 methods and data sets have been used to derive estimates of TCR. Observation-based studies  
39 analyze the historical temperature record combined with information on the radiative forcing  
40 (RF)<sup>5</sup>. The high uncertainty in historical RF is the main contributor to the uncertainty in the  
41 estimate of TCR and ECS through such methods<sup>4,6</sup>. Recent studies have shown that  
42 uncertainties in climate sensitivity will be reduced in the future based on longer available time  
43 series of surface temperature<sup>9</sup>. Here, we show that narrowing the uncertainty in RF can have a  
44 larger effect on the diagnosed TCR uncertainty.

45

46 Recently the IPCC 5<sup>th</sup> Assessment Report assessed historical RF and its uncertainty<sup>1</sup>. In this  
47 paper we evaluate how uncertainty estimates have evolved between IPCC reports and  
48 estimate how we expect uncertainty estimates in RF to change in the future. We then evaluate  
49 the consequences of these trends on future uncertainty in diagnosing TCR from the available  
50 temperature observations record.

51 Climate change can be driven by a wide range of emitted compounds as well as by physical  
52 and chemical processes<sup>1,8</sup>. The increase in well-mixed greenhouse gas abundances leads to a  
53 documented RF with small relative uncertainty ( $\approx 10\%$ ) with all uncertainties presented here  
54 covering 5-95% ranges and all relative uncertainties given as half the 5-95% relative ranges  
55 unless otherwise stated. However, several of these greenhouse gases affect atmospheric  
56 chemistry leading to indirect effects that add to the RF uncertainty<sup>10-12</sup>. The positive RF from  
57 greenhouse gas increases since pre-industrial time has partly been counteracted by an overall  
58 negative RF by anthropogenic aerosols<sup>13-16</sup>; however the scattering and absorbing effects of  
59 atmospheric aerosols, including the component due to aerosol-cloud interactions, have  
60 uncertainties<sup>17</sup> that are much larger ( $\sim 100\%$  relative uncertainty) than those associated with  
61 CO<sub>2</sub> (see Supplementary Figure 1).

62 In the two most recent IPCC assessment reports (AR4 and AR5) best estimates for  
63 anthropogenic RF, together with their uncertainties, have been provided<sup>1,8</sup>. Similar estimates  
64 have been provided for earlier IPCC assessments<sup>18,19</sup>. The changes in RF estimates and their  
65 uncertainties between the reports are combinations of evolution in our scientific  
66 understanding and temporal change of the forcing agents between the RF evaluation years.  
67 Forcing estimates in the IPCC assessment are based on observations and modelling, and  
68 estimates constrained from observed climate change<sup>20,21</sup> are ignored.

69 Relative to pre-industrial (1750) the total anthropogenic RF in AR5 (for year 2011) is larger  
70 than in AR4 (for year 2005) and TAR (for year 1998) and Figure 1 shows that further  
71 increases are expected under two extreme Representative Concentration Pathways (RCPs) for  
72 2030 (see Methods). This increased total anthropogenic forcing from TAR and AR4 to AR5 is  
73 due to increases in greenhouse gases as well as increased scientific evidence for a less  
74 negative aerosol forcing<sup>1</sup>. In AR5 more aerosol processes are included in the forcing estimates

75 (allowing for rapid adjustments in the atmosphere) relative to previous IPCC assessments,  
76 which resulted in a less negative RF estimate (-0.9 (-1.9 to -0.1) W m<sup>-2</sup> in AR5 versus -1.2 (-  
77 2.4 to -0.6) W m<sup>-2</sup>). Importantly, the relative uncertainty is reduced from TAR and AR4 to  
78 AR5 as shown in the right panel of Figure 1. Projections using the RCPs indicate that this  
79 reduction will continue and, by 2030, the relative uncertainty in the total anthropogenic RF  
80 will be approximately halved relative to the latest IPCC assessments assuming no change in  
81 the scientific understanding of the forcing mechanisms (based on RCP2.6 and RCP8.5).  
82 Despite improvements in understanding, individual RF uncertainties changed relatively little  
83 between the assessment reports; yet the relative uncertainty in total RF has narrowed and can  
84 be expected to exhibit an even stronger decrease by 2030.

85 The main cause of reduction in the relative uncertainty of the total RF is due to the increasing  
86 share of the CO<sub>2</sub> contribution to the total as shown in Figure 2. In the last decades of the 20<sup>th</sup>  
87 century, non-CO<sub>2</sub> greenhouse gases made substantial contributions to the total, with rapid RF  
88 increases, while aerosols offset part of the greenhouse gas RF. The first decade in this century  
89 and projections for the next few decades show limited RF changes for non-CO<sub>2</sub> greenhouse  
90 gases and a decrease in the offsetting negative aerosol forcing combined with an enhancement  
91 in the CO<sub>2</sub> RF. This dramatic change in the relative RF contributions is due to fairly stable or  
92 declining recent and projected emissions of short-lived aerosols and aerosol precursors and  
93 most non-CO<sub>2</sub> GHGs, in contrast with continuous increases in CO<sub>2</sub> emissions<sup>22</sup> coupled with  
94 its long lifetime. The forcing due to aerosols including their influence on clouds is better  
95 understood and quantified than in AR4, but uncertainties remain large<sup>17</sup>. Over the last two  
96 decades, there has been a large change in the distribution of aerosols, linked to reduced  
97 anthropogenic emissions in Europe and North America and increased emissions over South

98 and East Asia. These opposite trends over the last decades are expected to more-or-less  
99 balance each other in terms of global mean RF<sup>1,23</sup>.

100 Figure 3 illustrates the time evolution of forcing used in this paper and its standard deviation.  
101 A maximum in the standard deviation (and hence absolute uncertainty) was reached around  
102 2011 and is expected to decline further despite the increase in forcing. This leads to continued  
103 reduction in the relative uncertainty which, based on AR5 estimates, has been declining since  
104 about 1970. Figure 3 clearly shows how the reduction in relative uncertainty is caused by the  
105 increasing dominance of CO<sub>2</sub> in the total RF. Overall, the combination of enhanced CO<sub>2</sub>  
106 forcing and weak magnitude of the non-CO<sub>2</sub> and aerosols forcing both contributed to the  
107 recent reduction in uncertainty in anthropogenic forcing and likewise for the trend in the  
108 coming decades. Whereas the change in the relative uncertainty in the anthropogenic forcing  
109 from AR4 to AR5 is a combination of larger CO<sub>2</sub> domination and improved quantification of  
110 the aerosol forcing, estimated further change by 2030 is solely due to a change in atmospheric  
111 abundances and no assumed change in scientific understanding about the individual drivers of  
112 climate change.

113 In the following we take the trends in anthropogenic RF estimates described above and  
114 examine their implications for estimating TCR from the historical record. TCR, when derived  
115 from historical observations or simulations relates the temperature change ( $\Delta T$ ) and the RF at  
116 a given time as follows:

$$117 \quad \text{TCR} = \Delta T / \text{RF} \quad (1)$$

118 The method used to estimate TCR here is similar to that used in recent studies<sup>6</sup>. Note that in  
119 the above equation TCR is expressed per unit of RF rather than for a doubling of CO<sub>2</sub>  
120 abundance. It assumes quasi linear changes in  $\Delta T$  and RF over a chosen time period. Often

121 TCR is assumed to be similar for all climate forcing mechanisms, although this may not  
122 be the case<sup>24</sup> (see further discussion below and in the Supplementary Material). It may also  
123 depend on the rate of change of forcing<sup>25</sup>, which introduces a further small uncertainty term  
124 not accounted for here. The relative uncertainty in TCR ( $d \text{ TCR} / \text{TCR}$ ), where  $d$  refers to half  
125 the uncertainty of the 5-95% uncertainty range, is shown in Figure 4a for RF relative  
126 uncertainties for AR4, AR5 and two RCPs for 2030 as a function of temperature change using  
127 the Monte Carlo simulations described and discussed in the Methods and Supplementary  
128 Material. Figure 4a shows that the relative uncertainty in TCR for the two RCPs is about half  
129 that found for AR5 data. The uncertainties related to temperature decrease as the temperature  
130 change increases, as can be seen for the two RCPs. However, the contribution from  
131 temperature uncertainties is less than 10% of the change in the relative uncertainty in TCR  
132 between AR5 and the RCPs for 2030, emphasizing that changes in the RF uncertainties are  
133 the dominant cause of the differences between AR5 and the RCPs for 2030.

134 The difference in RF uncertainty between AR4, AR5 and the two RCPs for 2030 translates  
135 into a large difference in the 5-95% uncertainty range of the TCR for AR5 present-day  
136 temperature changes and best estimate RF as shown in the inset in Figure 4. The only  
137 difference in these ranges in TCR is caused by the declining uncertainty in RF. The better  
138 quantification of RF has the largest impact on the upper range of the derived TCR in absolute  
139 terms. Upper ranges of TCR are associated with low values of RF for which the lower bound  
140 of the aerosol RF is particularly relevant. A relatively symmetric distribution of RF leads to  
141 more asymmetric shape of the distribution of TCR<sup>7</sup>. The two RCPs uncertainty estimates in  
142 2030 provide rather similar uncertainty ranges for TCR. The AR5 likely range of TCR can be  
143 reduced by about 50% based on climate data from two additional decades solely due to  
144 expected RF trends without further improvements in understanding (subject to continued

145 availability of global surface temperature observations), see inset in Figure 4b. The absolute  
146 change in TCR is more dependent on temperature changes than the relative change in TCR.  
147 Figure 4b shows that the absolute range in TCR will be at least 25% lower than the AR5  
148 range over the RCP8.5 temperature ranges for 2030. For small temperature changes, the  
149 absolute uncertainty in TCR will see a greater reduction than the relative uncertainty in TCR  
150 between AR5 and the RCP for 2030 (up to 56%).

151 Including an enhanced response to forcing in the Northern Hemisphere extratropics<sup>24</sup> would  
152 increase the uncertainty in present-day TCR calculations, but would lead to an even greater  
153 narrowing of the TCR uncertainty moving to 2030 RCP conditions (see Supplementary Figure  
154 2). The combination of air quality policies, the Montreal Protocol, trends in emission of  
155 climate related compounds, and most importantly the differentiated lifetime of the compounds  
156 suggests that the current evolution is likely to continue over the next few decades. Our  
157 findings illustrate that the stronger domination of CO<sub>2</sub> RF over the other forcing terms leads  
158 to a better quantification of TCR.

159 A better quantification of TCR will have a pronounced impact on the probability distribution  
160 of estimates of the amount of permissible CO<sub>2</sub> emissions for a given temperature target, e.g.  
161 the 2 °C target agreed to under the UNFCCC. Currently these emissions are highly uncertain<sup>26</sup>,  
162 but the expected CO<sub>2</sub> domination will bring about (by itself) a better quantification of TCR  
163 and future projections of climate change.

164

165

## 166 **Methods**

167 All forcing values and their uncertainties used for figures and analysis are given in the IPCC  
168 AR5 in chapter text, supplementary or annex<sup>1,27</sup>, except for one case as described below for

169 RCP2.6. The time evolution of historical and future RF is also from IPCC AR5. Projections  
170 for 2030 are based on the two most extreme RCPs, namely RCP2.6 and RCP8.5<sup>28</sup>. These two  
171 RCP represent lower and upper projections over the next decades, respectively in terms of  
172 CO<sub>2</sub> emissions and to some extent other climate relevant species. The development over the  
173 last decade is closest to RCP8.5 in terms of CO<sub>2</sub> emissions. Other emission scenarios based  
174 on realistic development until 2030 have little impact on our findings. AR5 forcing estimates  
175 for aerosols and contrail induced cirrus include rapid adjustments and thus use the effective  
176 radiative forcing concept<sup>1,17</sup>, whereas in AR4 and previous IPCC reports rapid adjustments  
177 were not quantified. This makes some of the forcing estimates not entirely comparable, but  
178 allowing for the difference in treatment of rapid adjustment is the most consistent method for  
179 the aerosols between the IPCC reports. Forcing estimates for the two RCPs and AR5 are  
180 derived consistently with the same forcing concept and relative uncertainty for the individual  
181 drivers. The combined forcing from ozone and stratospheric water vapour in Figure 1 has a  
182 small change in the relative uncertainty between AR5 and the RCPs caused solely by  
183 abundance changes. The best estimate of the total anthropogenic RF for the various IPCC  
184 reports and the two RCPs is calculated based on the sum of the best estimate of each  
185 component. The range of the total anthropogenic RF is derived from the square root of the  
186 sum of the square of the upper and lower range deviation from the best estimate for the  
187 individual component. This allows for a consistent treatment of the best estimate and range,  
188 but may differ slightly from the report values in previous IPCC reports. The best estimate and  
189 uncertainty ranges for the two RCPs for 2030 are derived consistently with AR5 estimates,  
190 where the only change in estimate arise from atmospheric compositional change. Aerosols RF  
191 for RCP2.6 is not provided in the IPCC AR5 annex<sup>27</sup> and is derived based on the difference in  
192 aerosol forcing from 2010 to 2030 as derived from one model (OsloCTM2)<sup>29</sup> and thus also

193 made consistent with the AR5 estimate. The main source of the time evolution of historical  
194 and future forcing of aerosols and ozone for IPCC AR5 was a multi-model study<sup>30</sup>.  
195 The time series of uncertainty used in Figure 3 are derived from a Monte-Carlo method, based  
196 on converting IPCC AR5 uncertainty ranges in RF for 2011 into fractional error PDFs. We  
197 then sample these to generate plausible RF time series.  
198 For the calculations of changes in uncertainties in TCR probability distribution functions  
199 (PDFs) of TCR from PDFs of temperature change and RF is derived using a Monte Carlo  
200 random sampling approach. The values adopted to derive the PDFs are given in  
201 Supplementary Table 1. Supplementary Figure 3 shows PDFs of TCR derived in this way  
202 from PDFs of RF and temperature change. The 5-95% interval is derived from the PDFs of  
203 TCR.  
204 Relative uncertainties are given as half the 5-95% relative ranges. In Figure 4 the full 5-95%  
205 relative range is added to the relative uncertainty. In Figure 1 the relative uncertainties are  
206 calculated as half the 5-95% confidence range, divided by the best estimate.

207 **Figure 1:** Anthropogenic forcing for four phases of IPCC reports and two RCPs. Aerosols,  
208 ozone and stratospheric water vapour, well-mixed greenhouse gases (WMGHG), land use  
209 change and total forcing are given for SAR (1750-1993), TAR (1750-1998), AR4 (1750-  
210 2005) and AR5 (1750-2011)) and two RCPs for 2030. . All the forcing values are based on  
211 best estimates reported in the IPCC reports, but with a consistent approach to calculate the  
212 total forcing which may differ slightly from reported values (see Methods). In SAR land use  
213 change was not estimated and thus not included in the total. Further, the RF of a given CO<sub>2</sub>  
214 concentration was estimated to be 15% higher in SAR compared to the recent IPCC reports,  
215 adding 0.24 Wm<sup>-2</sup> to the total RF quoted in SAR. Estimate for AR5 and the two RCPs for  
216 2030 includes rapid adjustments in the RF, whereas these have not been quantified earlier in  
217 SAR, TAR, and AR4. The probability density function for SAR and TAR are based on  
218 Boucher and Haywood<sup>19</sup> and their simulation C1.5. The relative uncertainties are shown in the  
219 right panel. All uncertainty ranges correspond to 5-95% confidence intervals with relative  
220 uncertainties given as half the 5-95% relative range.

221

222

223 **Figure 2:** Decadal RF change between 1970 and 2010 and for 2020 to 2030 for two RCPs.  
224 The forcing is given for 1970 to 1980, 1980 to 1990, 1990 to 2000, and 2000 to 2010 and for  
225 2020 to 2030 based on IPCC AR5 forcing values (see Methods). RF for ozone includes  
226 changes in the troposphere as well as in the stratosphere. Other WMGHG includes CH<sub>4</sub>, N<sub>2</sub>O  
227 and halocarbons. All process associated with aerosol-radiation and aerosol-cloud interactions  
228 taken into account in the IPCC assessments are included for aerosol, except black carbon on  
229 snow. Consistent treatment is applied for RCPs for 2030 and AR5. Forcing mechanisms other  
230 than those shown in the figure are small (see Supplementary Information).

231

232

233 **Figure 3:** Time evolution in RF and standard deviation in RF. RF for total anthropogenic,  
234 CO<sub>2</sub>, the combined non-CO<sub>2</sub> greenhouse gases (GHG) such as CH<sub>4</sub>, N<sub>2</sub>O, halocarbons, ozone,  
235 and stratospheric water vapour, the others such as land use changes, black carbon on snow  
236 and ice, and contrails, and finally aerosols over the period 1850 and 2030 (a); the time  
237 evolution of the standard deviation of RF (b) and the ratio of the standard deviation of RF to  
238 the total RF (c). All the time evolutions of forcing are taken from IPCC AR5 (see Methods).  
239 RCP8.5 and RCP2.6 are shown with solid and dashed lines, respectively.

240

241

242

243

244 **Figure 4:** The relative a) and absolute b) uncertainty in TCR for the indicated conditions as a  
245 function of temperature change. The results are based on Monte Carlo simulations of the PDF  
246 of TCR as a function of temperature change and relative uncertainty in RF for AR4, AR5, and  
247 RCP2.6 and RCP8.5 for 2030 (see methods for source of RF values). The uncertainties in RF  
248 for the two RCPs are based on the scientific knowledge in AR5 and projected abundance  
249 changes. Observed temperature changes and their uncertainties from AR4 and AR5 are  
250 adopted in the calculations of the relative uncertainty in TCR, whereas for RCP2.6 and  
251 RCP8.5 results are shown for CMIP5 simulated temperature changes. The absolute  
252 uncertainty in temperature change for 2030 is assumed to be same as in AR5. The relative  
253 uncertainties and best estimates of TCR are shown as horizontal lines with ranges shown for  
254 lower and upper bound of the relative uncertainty in TCR and TCR (lines for AR4 and AR5  
255 whereas bands for the two RCPs for 2030). The inset shows the TCR and the uncertainty  
256 range from Equation 1 for temperature changes and RF at the time of AR5 but with different  
257 relative uncertainty in RF from AR4, AR5, and RCP2.6 and RCP8.5 for 2030. The diamond  
258 symbol shows the TCR with the best estimate RF and is thus constant to illustrate solely the  
259 difference in relative uncertainty in RF.

260

261

- 263 <sup>1</sup> Myhre, G. et al., *Anthropogenic and Natural Radiative Forcing*, in *Climate Change 2013: The*  
264 *Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the*  
265 *Intergovernmental Panel on Climate Change*, edited by T. F. Stocker et al. (Cambridge  
266 University Press, Cambridge, United Kingdom and New York, NY, USA, 2013), pp. 659-740.
- 267 <sup>2</sup> Stocker, T. F., D. Qin, G.-K. Plattner, M. Tignor, S. K. Allen, J. Boschung, A. Nauels, Y. Xia, V.  
268 Bex and P. M. Midgley ed., *IPCC, 2013: Summary for Policymakers*. (Cambridge University  
269 Press, Cambridge, United Kingdom and New York, NY, USA, 2013).
- 270 <sup>3</sup> Aldrin, M. et al., Bayesian estimation of climate sensitivity based on a simple climate model  
271 fitted to observations of hemispheric temperatures and global ocean heat content.  
272 *Environmetrics* **23**, 253-271 (2012).
- 273 <sup>4</sup> Andreae, M. O., Jones, C. D., and Cox, P. M., Strong present-day aerosol cooling implies a hot  
274 future. *Nature* **435**, 1187-1190 (2005).
- 275 <sup>5</sup> Knutti, R. and Hegerl, G. C., The equilibrium sensitivity of the Earth's temperature to  
276 radiation changes. *Nature Geoscience* **1**, 735-743 (2008).
- 277 <sup>6</sup> Otto, A. et al., Energy budget constraints on climate response. *Nature Geoscience* **6**, 415-416  
278 (2013).
- 279 <sup>7</sup> Roe, G. H. and Armour, K. C., How sensitive is climate sensitivity? *Geophysical Research*  
280 *Letters* **38**, L14708 (2011).
- 281 <sup>8</sup> Forster, P. et al., *Changes in Atmospheric Constituents and in Radiative Forcing*, in *Climate*  
282 *Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth*  
283 *Assessment Report of the Intergovernmental Panel on Climate Change* (Cambridge University  
284 Press, Cambridge, United Kingdom and New York, NY, USA, 2007).
- 285 <sup>9</sup> Urban, N. M. et al., Historical and future learning about climate sensitivity. *Geophysical*  
286 *Research Letters* **41**, 2543-2552 (2014).
- 287 <sup>10</sup> Isaksen, I. S. A. et al., Atmospheric composition change: Climate-Chemistry interactions.  
288 *Atmospheric Environment* **43**, 5138-5192 (2009).
- 289 <sup>11</sup> Raes, F., Liao, H., Chen, W. T., and Seinfeld, J. H., Atmospheric chemistry-climate feedbacks.  
290 *Journal of Geophysical Research-Atmospheres* **115** (2010).
- 291 <sup>12</sup> Shindell, D. T. et al., Improved Attribution of Climate Forcing to Emissions. *Science* **326**, 716-  
292 718 (2009).
- 293 <sup>13</sup> Crook, J. A. and Forster, P. M., A balance between radiative forcing and climate feedback in  
294 the modeled 20th century temperature response. *Journal of Geophysical Research-*  
295 *Atmospheres* **116**, D17108 (2011).
- 296 <sup>14</sup> Huber, M. and Knutti, R., Anthropogenic and natural warming inferred from changes in  
297 Earth's energy balance. *Nature Geoscience* **5**, 31-36 (2012).
- 298 <sup>15</sup> Hansen, J., Sato, M., Kharecha, P., and von Schuckmann, K., Earth's energy imbalance and  
299 implications. *Atmospheric Chemistry and Physics* **11**, 13421-13449 (2011).
- 300 <sup>16</sup> Ramanathan, V., Crutzen, P. J., Kiehl, J. T., and Rosenfeld, D., Atmosphere - Aerosols, climate,  
301 and the hydrological cycle. *Science* **294**, 2119-2124 (2001).
- 302 <sup>17</sup> Boucher, O. et al., *Clouds and Aerosols*, in *Climate Change 2013: The Physical Science Basis.*  
303 *Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental*  
304 *Panel on Climate Change*, edited by T. F. Stocker et al. (Cambridge University Press,  
305 Cambridge, United Kingdom and New York, NY, USA, 2013), pp. 571-657.
- 306 <sup>18</sup> Haywood, J. and Schulz, M., Causes of the reduction in uncertainty in the anthropogenic  
307 radiative forcing of climate between IPCC (2001) and IPCC (2007). *Geophysical Research*  
308 *Letters* **34** (2007).
- 309 <sup>19</sup> Boucher, O. and Haywood, J., On summing the components of radiative forcing of climate  
310 change. *Climate Dynamics* **18**, 297-302 (2001).

311 20 Forest, C. E. et al., Quantifying uncertainties in climate system properties with the use of  
312 recent climate observations. *Science* **295**, 113-117 (2002).  
313 21 Knutti, R., Stocker, T. F., Joos, F., and Plattner, G. K., Constraints on radiative forcing and  
314 future climate change from observations and climate model ensembles. *Nature* **416**, 719-723  
315 (2002).  
316 22 Peters, G. P. et al., COMMENTARY: The challenge to keep global warming below 2 degrees C.  
317 *Nature Climate Change* **3**, 4-6 (2013).  
318 23 Murphy, D. M., Little net clear-sky radiative forcing from recent regional redistribution of  
319 aerosols. *Nature Geoscience* **6**, 258-262 (2013).  
320 24 Shindell, D. T., Inhomogeneous forcing and transient climate sensitivity. *Nature Climate*  
321 *Change* **4**, 274-277 (2014).  
322 25 Forster, P. M. et al., Evaluating adjusted forcing and model spread for historical and future  
323 scenarios in the CMIP5 generation of climate models. *Journal of Geophysical Research-*  
324 *Atmospheres* **118**, 1139-1150 (2013).  
325 26 Meinshausen, M. et al., Greenhouse-gas emission targets for limiting global warming to 2  
326 degrees C. *Nature* **458**, 1158-1162 (2009).  
327 27 Prather, M. et al., *IPCC 2013: Annex II: Climate System Scenario Tables*, in *Climate Change*  
328 *2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment*  
329 *Report of the Intergovernmental Panel on Climate Change*, edited by T. F. Stocker et al.  
330 (Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013), pp.  
331 1395-1445.  
332 28 van Vuuren, D. P. et al., The representative concentration pathways: an overview. *Climatic*  
333 *Change* **109**, 5-31 (2011).  
334 29 Skeie, R. B. et al., Anthropogenic radiative forcing time series from pre-industrial times until  
335 2010. *Atmospheric Chemistry and Physics* **11**, 11827-11857 (2011).  
336 30 Shindell, D. T. et al., Radiative forcing in the ACCMIP historical and future climate simulations.  
337 *Atmospheric Chemistry and Physics* **13**, 2939-2974 (2013).

338

339

340

341 Correspondence and requests for material should be addressed to GM

342 ([gunnar.myhre@cicero.oslo.no](mailto:gunnar.myhre@cicero.oslo.no))

343

#### 344 **Acknowledgements**

345 GM was supported from the Norwegian Research Council project SLAC (208277)

346

#### 347 **Author contributions**

348 GM, FMB, DS initiated the study with additional contribution on the design on the study from

349 PF and OB. GM, OB, FMB, PF, and DS performed the analysis and wrote the paper.

350









