

# *Halfway to doubling of CO<sub>2</sub> radiative forcing*

Article

Accepted Version

Myhre, G., Myhre, C. L., Forster, P. M. and Shine, K. P.  
ORCID: <https://orcid.org/0000-0003-2672-9978> (2017) Halfway to doubling of CO<sub>2</sub> radiative forcing. Nature Geoscience, 10 (10). pp. 710-711. ISSN 1752-0894 doi: 10.1038/ngeo3036 Available at <https://centaur.reading.ac.uk/72841/>

It is advisable to refer to the publisher's version if you intend to cite from the work. See [Guidance on citing](#).

Published version at: <http://dx.doi.org/10.1038/ngeo3036>

To link to this article DOI: <http://dx.doi.org/10.1038/ngeo3036>

Publisher: Nature Publishing Group

All outputs in CentAUR are protected by Intellectual Property Rights law, including copyright law. Copyright and IPR is retained by the creators or other copyright holders. Terms and conditions for use of this material are defined in the [End User Agreement](#).

[www.reading.ac.uk/centaur](http://www.reading.ac.uk/centaur)

**CentAUR**

Central Archive at the University of Reading

Reading's research outputs online

# Halfway to doubling of CO<sub>2</sub> radiative forcing

Gunnar Myhre<sup>1\*</sup>, Cathrine Lund Myhre<sup>2</sup>, Piers M. Forster<sup>3</sup>, Keith P. Shine<sup>4</sup>

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

<sup>2</sup>NILU-Norwegian Institute for Air Research, Kjeller, Norway

<sup>3</sup>University of Leeds, Leeds, UK

<sup>4</sup>Department of Meteorology, University of Reading, Reading, UK

\*Corresponding author ([gunnar.myhre@cicero.oslo.no](mailto:gunnar.myhre@cicero.oslo.no))

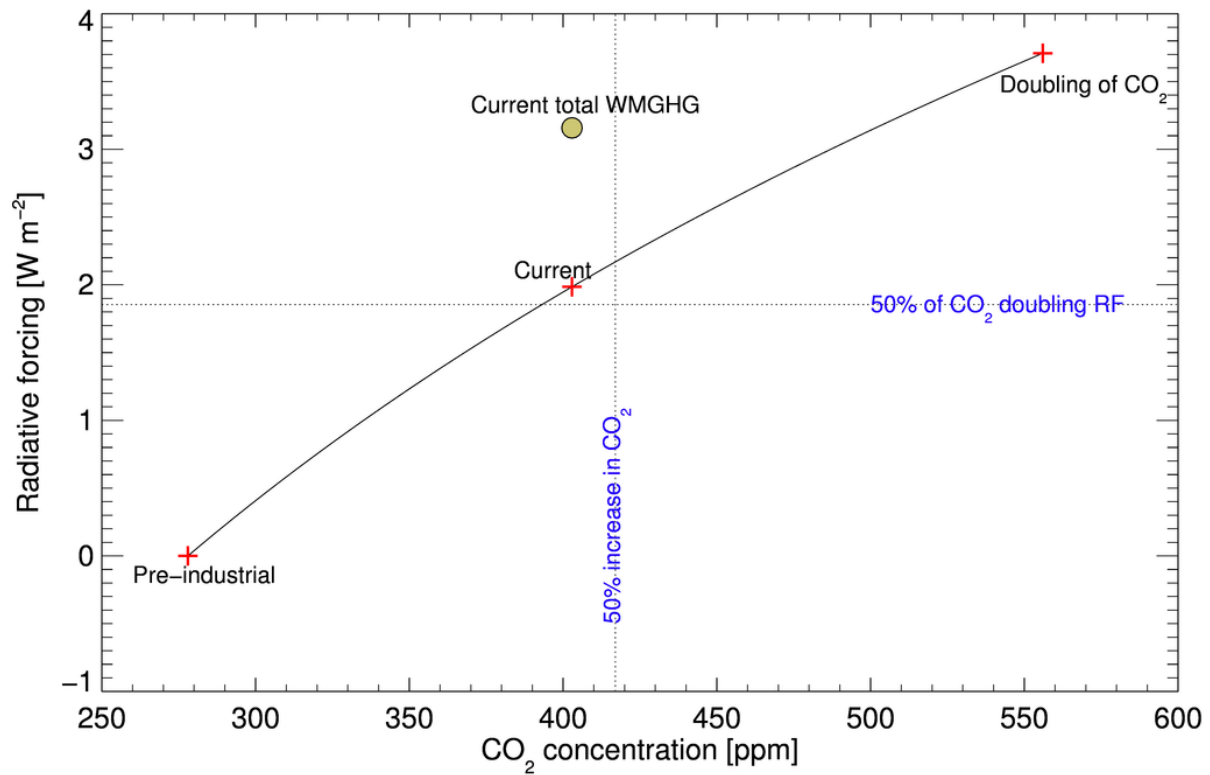
The “double CO<sub>2</sub>” experiment has become a standard experiment in climate science, and a convenient way of comparing the sensitivity of different climate models. Double CO<sub>2</sub> was first used by Arrhenius<sup>1</sup> in the 19<sup>th</sup> century and in the classic paper by Manabe and Wetherald<sup>2</sup>, published 50 years ago, which marked the start of the modern era of climate modeling. Doubling CO<sub>2</sub> now has an iconic role in climate research. The equilibrium climate sensitivity (ECS) is defined as the global-mean surface temperature change resulting from a doubling of CO<sub>2</sub><sup>3-5</sup>, which is a headline result in Intergovernmental Panel on Climate Change (IPCC) assessments. In its most recent assessment IPCC concluded that the ECS “is likely in the range 1.5 to 4.5°C”. We show that we are now halfway to doubling of CO<sub>2</sub> since pre-industrial times in terms of radiative forcing, but not in concentration.

The greenhouse effect due to change in CO<sub>2</sub> – quantified using calculations of radiative forcing – follows, to a good approximation, a logarithmic dependence on the ambient concentration in the atmosphere over the last 1000 years<sup>6</sup>. Due to this relationship between radiative forcing and CO<sub>2</sub> concentration, the radiative forcing due to a doubling of CO<sub>2</sub> is approximately independent of background levels. A doubling of CO<sub>2</sub> is estimated by IPCC to cause a radiative forcing of 3.7 W m<sup>-2</sup>. Recent detailed radiative transfer calculations arrived at a similar estimate<sup>7</sup>. The uncertainties are small for the radiative forcing due to CO<sub>2</sub>; uncertainties associated with spectroscopic parameters that underpin forcing calculations are estimated to be less than 1% in a recent study<sup>8</sup>, with overall uncertainties assessed to be 10%<sup>6</sup> (with 90% confidence). Forcing estimates of doubling of CO<sub>2</sub> from global climate models have the same best estimate as the IPCC value<sup>6</sup>, even though these models include rapid atmospheric adjustments, which modify the forcing calculated using a radiative transfer model.

It is timely to assess where we are now, relative to a doubling. The global-mean CO<sub>2</sub> abundance in 2016 was 403 ppm according to global observations<sup>9</sup> which is less than 50% higher than the pre-industrial CO<sub>2</sub> concentration of 278 ppm. However, due to the logarithmic forcing relationship, a halfway to doubling of CO<sub>2</sub>, in terms of radiative forcing, has now been reached. Figure 1a illustrates that this halfway point happened at 393 ppm, which was reached in 2012. A halfway to doubling in the CO<sub>2</sub> concentration is 417 ppm and will be reached before 2025 with current CO<sub>2</sub> growth rates. Hence, at CO<sub>2</sub> concentrations between of 393 and 417 ppm we are more than a halfway to CO<sub>2</sub> doubling in radiative forcing, but not in concentration (Figure 1a).

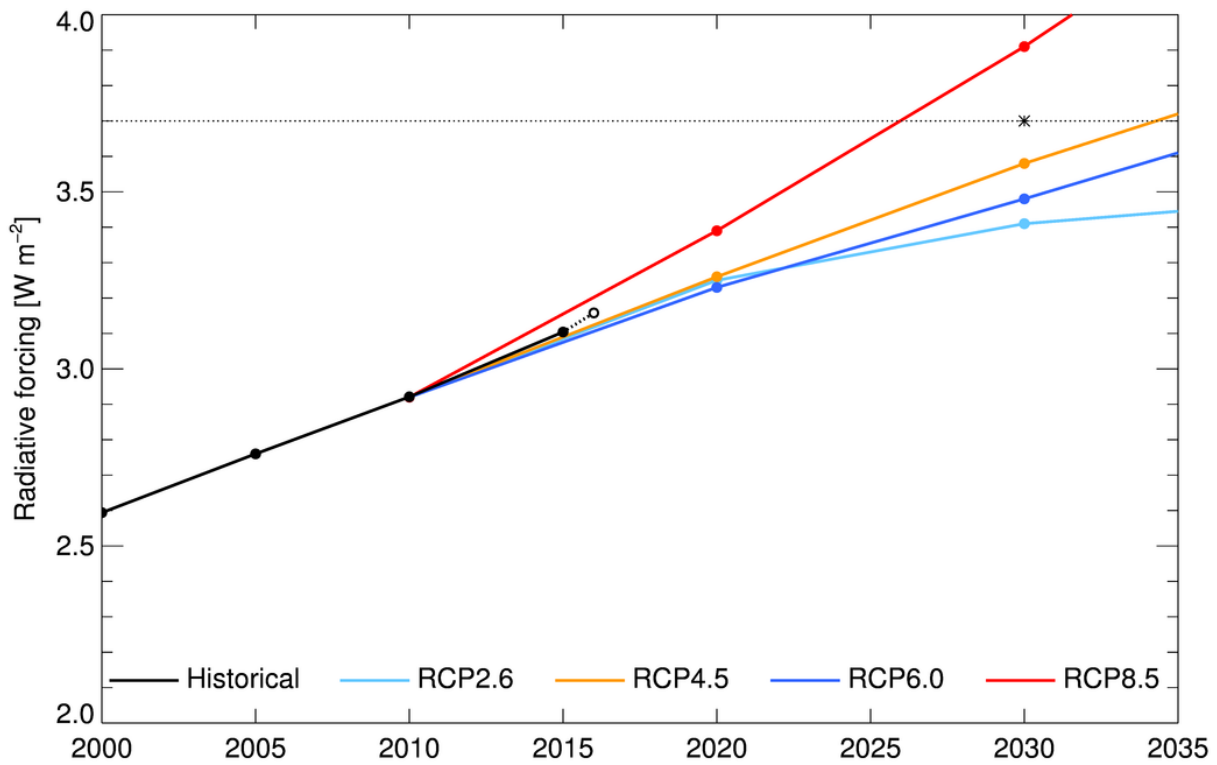
Climate change over the industrial era is caused by several anthropogenic climate drivers in addition to CO<sub>2</sub>, including other atmospheric gases and aerosols and changes to the land surface<sup>6</sup>. Increases in concentrations of well-mixed greenhouse gases (WMGHGs) other than CO<sub>2</sub> (notably CH<sub>4</sub>, N<sub>2</sub>O and halocarbons) contribute to a stronger greenhouse effect. The combined radiative forcing from all WMGHGs is 3.1 W m<sup>-2</sup> in 2015 (Figure 1b) and hence in CO<sub>2</sub>-equivalent forcing terms, is 84% of the way to a doubling. This value includes a recent estimate of methane's radiative forcing which incorporated its absorption of solar radiation; this update resulted in an increase in the 1750-2011 CH<sub>4</sub> forcing from 0.48 (the value in IPCC fifth assessment<sup>6</sup>) to 0.61 W m<sup>-2</sup> <sup>7</sup>. This increase is, in radiative forcing terms, close to the increase in CO<sub>2</sub> concentration over the 5 year period from 2010 to 2015. Consequently, we estimate that total WMGHG radiative forcing will be equivalent to doubling of CO<sub>2</sub>, with present growth rates, by around 2030 (Figure 1b). This is almost 5 years earlier than is estimated without the update to the CH<sub>4</sub> forcing. Aerosols generally cool the Earth and have historically countered much of this additional WMGHG forcing. The total anthropogenic forcing is expected to be close to the CO<sub>2</sub>-only forcing, but aerosols add uncertainty<sup>6</sup>. Nevertheless, in terms of radiative forcing we are more than half way to a doubling of CO<sub>2</sub>.

73 a



74

75 b



76

**Figure 1:** Radiative forcing due to CO<sub>2</sub> and all well-mixed greenhouse gases (WMGHG). **a**, The CO<sub>2</sub> radiative forcing shown as a function of its global-mean abundance calculated using the IPCC forcing expressions<sup>6</sup>. Dotted lines are for a 50% increase in concentration (vertical) and radiative forcing (horizontal). **b**, Radiative forcing for all WMGHGs using the IPCC forcing expressions<sup>6</sup>, except for CH<sub>4</sub> where a stronger forcing, based on recent detailed calculations, is used<sup>7</sup>. Historical values are based on observed concentrations. Radiative forcing for CO<sub>2</sub>, N<sub>2</sub>O and halocarbons for the 2000-2010 period and future scenarios are from IPCC<sup>10</sup>. CH<sub>4</sub> concentrations are from IPCC<sup>10</sup>. For year 2015 the global annual mean concentrations of CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O are from NOAA<sup>9</sup>, and for halocarbons the relative increase since 2010 are from the Arctic Zeppelin observatory. Preliminary data for 2016 is included<sup>9</sup>, which may be subject to small changes. Growth in WMGHG radiative forcing in the 2010-2016 period is 0.04 W m<sup>-2</sup> yr<sup>-1</sup>; the asterisk shows the date at which the total WMGHG forcing equals a CO<sub>2</sub> doubling by extrapolating this trend.

91  
92  
93  
94  
95  
96  
97  
98  
99  
100  
101  
102  
103  
104  
105  
106  
107  
108  
109  
110  
111  
112  
113  
114  
115  
116  
117

<sup>1</sup> Arrhenius, S. *Philos. Mag. J. Sci.* **41**, 237–276 (1896).  
<sup>2</sup> Manabe, S. and Wetherald, R. T. *J. Atmos. Sci.* **24**, 241-259 (1967).  
<sup>3</sup> Forster, Piers M. *Annual Review of Earth and Planetary Sciences* **44**, 85-106 (2016).  
<sup>4</sup> Roe, G. H. and Baker, M. B. *Science* **318**, 629-632 (2007).  
<sup>5</sup> Collins, M. et al. in *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by T.F. Stocker, D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley (Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013), pp. 1029–1136.  
<sup>6</sup> Myhre, G. et al. in *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by T. F. Stocker et al. (Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013), pp. 659-740.  
<sup>7</sup> Etminan, M., Myhre, G., Highwood, E. J., and Shine, K. P. *Geophys. Res. Lett.* **43**, 12614-12623 (2016).  
<sup>8</sup> Mlynckzak, Martin G. et al. *Geophys. Res. Lett.* **43**, 5318-5325 (2016).  
<sup>9</sup> Blunden, J. and Arndt, D.S. *Bull. Amer. Meteor. Soc.*, **98**, Si–S277 (2017).  
<sup>10</sup> Prather, M. et al. in *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by T. F. Stocker et al. (Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013), pp. 1395-1445.