# **1** Supplementary Material

### 2 The SSP greenhouse gas concentrations and their extensions to 2500

Malte Meinshausen<sup>1,2,3</sup>, Zebedee Nicholls<sup>1,2</sup>, Jared Lewis<sup>1</sup>, Matthew J. Gidden<sup>4,5</sup>, Elisabeth Vogel<sup>1,2</sup>,
Mandy Freund<sup>1,6</sup>, Urs Beyerle<sup>7</sup>, Claudia Gessner<sup>7</sup>, Alexander Nauels<sup>1,5</sup>, Nico Bauer<sup>3</sup>, Josep G. Canadell<sup>8</sup>,
John S. Daniel<sup>9</sup>, Andrew John<sup>1,10</sup>, Paul Krummel<sup>11</sup>, Gunnar Luderer<sup>3</sup>, Nicolai Meinshausen<sup>12</sup>, Stephen A.
Montzka<sup>13</sup>, Peter Rayner<sup>2,1</sup>, Stefan Reimann<sup>14</sup>, Steven J. Smith<sup>15</sup>, Marten van den Berg<sup>16</sup>, Guus J.M.
Velders<sup>17,18</sup>, Martin Vollmer<sup>14</sup>, Hsaing Jui (Ray) Wang<sup>19</sup>

- 8
- 9 <sup>1</sup> Climate & Energy College, The University of Melbourne, Parkville, Victoria, Australia
- 10 <sup>2</sup> School of Earth Sciences, The University of Melbourne, Parkville, Victoria, Australia
- <sup>11</sup> <sup>3</sup> Potsdam Institute for Climate Impact Research (PIK), Potsdam, Germany
- 12 <sup>4</sup>IIASA Institute for Applied Systems Analysis, Laxenburg, Austria
- <sup>5</sup>Climate Analytics, Berlin, Germany
- <sup>6</sup> Marine and Atmospheric Research, CSIRO, Hobart, Tasmania, Australia
- <sup>15</sup> <sup>7</sup> Institute for Atmospheric and Climate Science, Swiss Federal Institute of Technology, Zurich (ETH Zurich), Switzerland
- 16 <sup>8</sup> Global Carbon Project, CSIRO Oceans and Atmosphere, Canberra, ACT, Australia
- <sup>9</sup>NOAA, Earth System Research Laboratory, Chemical Sciences Division, Boulder, Colorado, USA
- 18 <sup>10</sup> Department of Infrastructure Engineering, The University of Melbourne, Parkville, Victoria, Australia
- 19 <sup>11</sup>CSIRO Oceans and Atmosphere, Aspendale, Victoria, Australia
- 20 <sup>12</sup> Seminar for Statistics, Swiss Federal Institute of Technology (ETH Zurich), Zurich, Switzerland.
- 21 <sup>13</sup>NOAA, Earth System Research Laboratory, Global Monitoring Division, Boulder, Colorado, USA
- <sup>14</sup> Empa, Laboratory for Air Pollution/Environmental Technology, Swiss Federal Laboratories for Materials Science and
- 23 Technology, Dübendorf, Switzerland
- 24 <sup>15</sup> Joint Global Change Research Institute, Pacific Northwest National Laboratory, College Park, MD, USA
- 25 Potsdam, Germany
- 26 <sup>16</sup> PBL Netherlands Environmental Assessment Agency, the Netherlands
- <sup>17</sup> National Institute for Public Health and the Environment (RIVM), Bilthoven, Netherlands
- <sup>18</sup> Institute for Marine and Atmospheric Research Utrecht (IMAU), Utrecht University, Utrecht, The Netherlands
- <sup>19</sup> School of Earth and Atmospheric Sciences, Georgia Institute of Technology, Atlanta, GA 30332-0340, USA
- 30
- 31 Correspondence to: M. Meinshausen (malte.meinshausen@unimelb.edu.au)

32

# **33 1 Supplementary Material**

Note: All data and all scenario and individual gas visualisation for the SSP GHG projections under this study are available at greenhousegases.science.unimelb.edu.au
The supplementary material contains a comparison of our GHG concentration projections with those that are part of the IIASA SSP database as initial values using MAGICC6. Efforts are underway to update the IIASA database values with the GHG

concentrations presented here. Also, this supplementary presents additional figures, such as those related to the methane gas
 cycle and nitrous oxide gas cycles.

45

34

#### 46 1.1 Comparison to earlier SSP scenario data in IIASA database

Several Integrated Assessment Models (IAM) and the IIASA emission scenario database include the MAGICC6 model to produce atmospheric concentrations, radiative forcing and temperatures. Here, we compare our CMIP6 GHG concentration projections using the updated MAGICC7.0 model, with its CMIP6 default settings, to the concentration projections within the IIASA database, both the concentration projections from the harmonized and non-harmonized emission scenarios. The harmonisation process adjusted the various IAM group's emission scenarios to common starting values in 2015 (Gidden et al., 2018). In the case of CO<sub>2</sub>, the difference between the harmonized and non-harmonized MAGICC6 concentration projections is generally small – given that most IAMs' recent historical CO<sub>2</sub> emission assumptions were relatively close to each other from the start (see dashed and dash-dotted line in panel a of Supplementary Figure 4). A slightly more pronounced upward adjustment for the higher scenarios is due to the shift from the MAGICC6 default version to the MAGICC7 default version used in this study, predominantly due to the more sensitive carbon cycle setting used here. As illustration of this difference, we consider the non-harmonised SSP5-8.5 scenario from the REMIND-MAgPIE modelling group. Upper range 2100 CO<sub>2</sub> concentrations for the official CMIP6 recommendations resulting from this study are 1135 ppm, whereas the IIASA database lists 1089 ppm for the non-harmonized SSP5-8.5 scenario from the REMIND-MAgPIE modelling group (Supplementary Figure 4).

61 For CH<sub>4</sub>, the overall concentration differences are somewhat more pronounced. For example, the effect of the scenario harmonisation for the SSP5-8.5 scenario (compare red dashed and dash-dotted lines in panel b of Supplementary Figure 4) is 62 a substantial downward adjustment. See also Figure 7b in Gidden et al. (2019). This is partly offset by an upward adjustment 63 64 that results from using the new MAGICC7.0 CH<sub>4</sub> cycle calibration (section). The MAGICC7.0 CH<sub>4</sub> gas cycle generally results in an upward adjustment of the projections from the harmonized emission dataseries, whereas the harmonization process itself 65 66 resulted in both upwards and downwards adjustments. Similarly, for N<sub>2</sub>O (panel c in Supplementary Figure 4), the updated 67 gas cycle leads in slight upward adjustments of N<sub>2</sub>O timeseries, whereas the harmonisation process resulted in both up and 68 downwards adjustments.

69

M. Meinshausen et al. - Supplementary Material "The SSP greenhouse gas concentrations and their extensions to 2500"

#### 70 1.2 Supplementary Figures - Captions

71

72	Supplementary Figure 1 – Calibration of MAGICC7's methane gas cycle to concentrations, lifetimes and natural emissions from Holmes
73	et al. (2013), varying 10 of MAGICC's gas cycle parameters with the goodness of fit being a weighted mean squared error across the four
74	variables (shown columns) and scenarios. MAGICC's CH4 projections are emission driven from year 1900. The constant natural emissions
75	assumed in MAGICC7 are calculated over the budgeting period from 1994 to 2004, whereas Holmes' et al. natural emission assumptions
76	have a slight time-variation.

77

Supplementary Figure 2 – Calibration of MAGICC's N<sub>2</sub>O concentration projections for the four RCPs and the concentrations projected by Prather et al. (2012) with natural N<sub>2</sub>O emissions in left column of panels, the N<sub>2</sub>O concentrations in the middle column of panels and the total N<sub>2</sub>O lifetime in the right column of panels. Each row of panels refers to one of the RCP scenarios, with RCP2.6, RCP4.5, RCP6.0 and RCP8.5 shown in order from top to bottom.

82

Supplementary Figure 3 - One-year (2050) and cumulative emissions (2018-2050) and their relationship with mid-century concentrations
 for methane (panels a, b) and CO<sub>2</sub> (panels c, d) across the SSP, RCP and 475 other scenarios from the IPCC SR1.5 database. The considered
 scenarios are the same as in Figure 9.

86

87 Supplementary Figure 4 - Comparison of final CMIP6 recommendation datasets for CO<sub>2</sub> (panel a), CH<sub>4</sub> (panel b) and N<sub>2</sub>O (panel c)
 88 concentrations with earlier derivations of concentrations using MAGICC6.

89

M. Meinshausen et al. - Supplementary Material "The SSP greenhouse gas concentrations and their extensions to 2500"

90

### 91 References

- Gidden, M. J., Fujimori, S., van den Berg, M., Klein, D., Smith, S. J., van Vuuren, D. P., and Riahi, K.: A methodology and
   implementation of automated emissions harmonization for use in Integrated Assessment Models, Environmental Modelling &
   Software, 105, 187-200, https://doi.org/10.1016/j.envsoft.2018.04.002, 2018.
- Gidden, M. J., Riahi, K., Smith, S. J., Fujimori, S., Luderer, G., Kriegler, E., van Vuuren, D. P., van den Berg, M., Feng, L.,
  Klein, D., Calvin, K., Doelman, J. C., Frank, S., Fricko, O., Harmsen, M., Hasegawa, T., Havlik, P., Hilaire, J., Hoesly, R.,
  Horing, J., Popp, A., Stehfest, E., and Takahashi, K.: Global emissions pathways under different socioeconomic scenarios for
  use in CMIP6: a dataset of harmonized emissions trajectories through the end of the century, Geosci. Model Dev., 12, 14431475, 10.5194/gmd-12-1443-2019, 2019.
- Holmes, C. D., Prather, M. J., Sovde, O. A., and Myhre, G.: Future methane, hydroxyl, and their uncertainties: key climate
   and emission parameters for future predictions, Atmospheric Chemistry and Physics, 13, 285-302, 2013.
- Prather, M. J., Holmes, C. D., and Hsu, J.: Reactive greenhouse gas scenarios: Systematic exploration of uncertainties and the
   role of atmospheric chemistry, Geophysical Research Letters, 39, 2012.
- 104
- 105
- 106

- . . .

### 116 Supplementary Figures



Supplementary Figure 1



Supplementary Figure 2



Supplementary Figure 3



Supplementary Figure 4