

COVID-19 lockdown emission reductions can explain over half of the coincident increase in global atmospheric methane

David S. Stevenson¹, Richard G. Derwent², Oliver Wild³, William J. Collins⁴

¹School of GeoSciences, The University of Edinburgh, Edinburgh EH9 3FF, UK

5 ²rdscientific, Newbury, UK

³Lancaster Environment Centre, Lancaster University, Lancaster, UK

⁴Department of Meteorology, University of Reading, Reading, UK

Correspondence to: David S. Stevenson (David.S.Stevenson@ed.ac.uk)

Abstract. Compared to 2019, measurements of the global growth rate of background (marine air) atmospheric methane rose by 5.5 ppb/yr in 2020, reaching 15.2 ppb/yr. Global atmospheric chemistry models have previously shown that reductions in nitrogen oxide (NO_x) emissions reduce levels of the hydroxyl radical (OH), and lengthen the methane lifetime. Acting in the opposite sense, reductions of carbon monoxide (CO) and non-methane volatile organic compound (NMVOC) emissions increase OH and shorten methane's lifetime. Using estimates of NO_x, CO and NMVOC emission reductions associated with COVID-19 lockdowns around the world in 2020, together with model-derived regional and aviation sensitivities of methane to these emissions, we find that NO_x emissions reductions led to a 4.3 (3.6 to 5.0) ppb/yr increase in the global methane growth rate. Reductions in CO and NMVOC emissions partly counteracted this, changing (reducing) the methane growth rate by -1.1 (-0.5 to -1.5) ppb/yr (CO) and -0.1 (0.0 to -0.3) ppb/yr (NMVOC). Uncertainties refer to ±1 standard deviation model ranges in sensitivities. Whilst changes in anthropogenic emissions related to COVID-19 lockdowns are probably not the only important factor that influenced methane during 2020, these results indicate that they have had a large impact, and that the net effect of NO_x, CO and NMVOC emissions changes can explain over half of the observed 2020 methane changes. Large uncertainties remain in both emissions changes during the lockdowns and methane's response to them; nevertheless, this analysis suggests that further research into how atmospheric composition changed over the lockdown periods will help us to interpret past methane changes and to constrain future methane projections.

1 Introduction

25 Methane is a powerful greenhouse gas and important precursor of tropospheric ozone; both are key air pollutants and short-lived climate forcers (SLCFs). Several factors in addition to rising anthropogenic methane emissions have influenced the evolution of atmospheric methane from its pre-industrial level of ~700 ppb to its present-day value of over 1900 ppb. The Intergovernmental Panel on Climate Change's Sixth Assessment Report (Szopa et al., 2021) assessed how changes in emissions of NO_x, CO, and NMVOCs have contributed to historical changes in methane, through their impacts on OH, the main sink for methane. A range of modelling studies have explored these indirect impacts on methane (e.g., Shindell et al.,

30

2005, 2009; Stevenson et al. 2013; Thornhill et al., 2021). For example, the Atmospheric Chemistry and Climate Model Intercomparison Project found that 1850-2000 increases in anthropogenic NO_x emissions had reduced year 2000 methane levels by 955 ppb, whilst growing emissions of CO and NMVOCs had increased methane by 150 ppb and 59 ppb, respectively (Table 7 of Stevenson et al., 2013). These results have quite large uncertainties (at least ±10%, based on the model range in
35 Stevenson et al. 2013), but indicate that non-methane (especially NO_x) emissions have had very significant impacts on methane. Better understanding of what controls methane and its evolution is vital for progress towards the Paris Climate Agreement target that seeks to limit warming to 1.5°C above pre-industrial levels.

Following the onset of the COVID-19 pandemic in early 2020, the trace gas composition of the global atmosphere changed substantially. Atmospheric nitrogen oxide levels reduced as surface and aviation NO_x emissions fell (Bauwens et al., 2020;
40 Cooper et al., 2022), whilst the measured growth rate of methane (CH₄) rose sharply in 2020 (Laughner et al., 2021). The observed NO_x changes are clearly linked to falls in emissions resulting from lockdowns, but the driver of the methane increases is less clear, with some studies discussing causes related to decreases in OH (e.g., Weber et al., 2020; Laughner et al., 2021) while others suggest rises in sources (e.g., Feng et al., 2022). Methane, NO_x, CO and NMVOCs are linked via the oxidising capacity of the atmosphere, specifically by the abundance of the hydroxyl (OH) radical. The response of global atmospheric
45 chemistry to the large lockdown perturbation since early 2020 provides an opportunity to explore the sensitivity of the NO_x-CO-NMVOC-OH-CH₄ system, and compare models and observations. Here we use model-derived sensitivities of global methane to NO_x, CO and NMVOC emissions, together with estimated changes in anthropogenic emissions of these species related to the COVID-19 lockdowns, to calculate estimated impacts from lockdown emissions changes on the growth rate of global methane, and compare this to observations.

50 **2 Measurements of atmospheric methane and nitrogen oxides**

Recent methane measurements from the US National Oceanographic and Atmospheric Administration (NOAA) show that the atmospheric (marine air background) methane growth rate rose sharply from 9.7 ppb/yr in 2019 to 15.2 ppb/yr in 2020, higher than any preceding annual value in the NOAA record, that started in 1984 (Dlugokencky, 2022). Many of the earlier large year-to-year jumps in methane's growth rate relate in part to variability in climate and emissions associated with El Niño
55 Southern Oscillation (ENSO), and in part because of modulation of methane's main sink, oxidation by OH (Turner et al., 2018; Zhao et al., 2020). The start of 2020 marked the onset of a La Niña that has persisted into 2022. Past La Niña's have not always shown clear links with methane's growth rate, and the influence of the current ENSO phase on methane is uncertain.

Measurements of nitrogen dioxide (NO₂) from satellite instruments and nitrogen monoxide (NO) and NO₂ from surface sites
60 show that levels of atmospheric NO_x (NO + NO₂) dramatically fell globally during 2020 (Bauwens et al., 2020; Laughner et al., 2021; Cooper et al., 2022). This was driven by COVID-19 lockdowns around the world that reduced emissions, mainly from transportation (Venter et al., 2020; Lamboll et al., 2021; Doumbia et al., 2021).

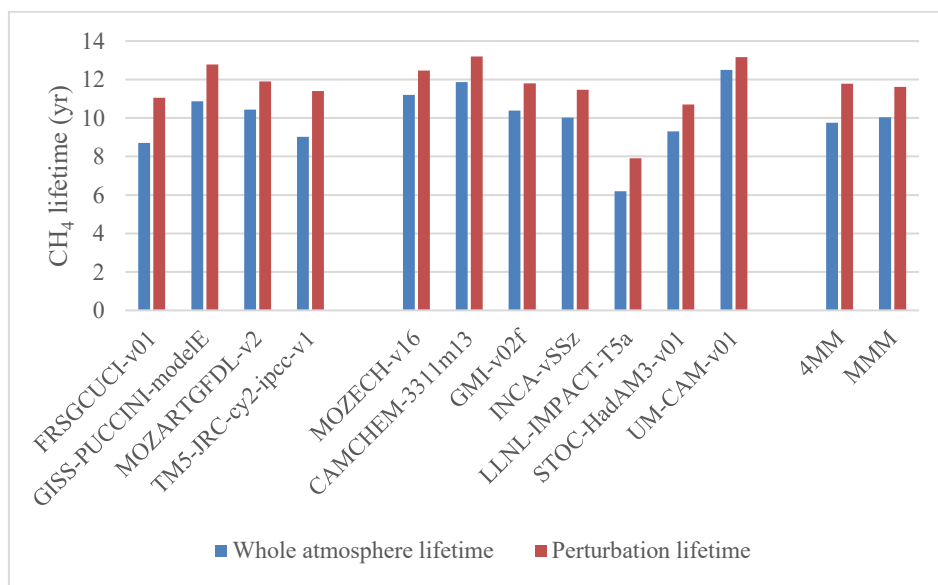
3 Sensitivity of global methane to NO_x, CO and NMVOC emissions

Global atmospheric chemistry model simulations indicate that decreases in NO_x emissions lead to reductions in OH and increases in the global methane lifetime (Prather, 1994; Derwent et al., 2001; Wild et al., 2001; Stevenson et al., 2004; Weber et al., 2020). Similarly, decreases in CO and NMVOC emissions lead to increases in OH and decreases in methane lifetime (Derwent et al., 2001; Wild et al., 2001). Although methane has an atmospheric lifetime of about 10 years, models show that its peak response occurs within a few months of the cessation of a sudden short-lived (month- or year-long) pulse of extra emissions (Derwent et al., 2001; Wild et al., 2001; Stevenson et al., 2004). This indicates that the impacts on methane from the sudden changes in emissions associated with lockdowns will have had rapid impacts on methane's growth rate.

We first illustrate the basis of our approach by describing the model experiments performed by Derwent et al. (2001), who conducted a series of simulations with the global tropospheric chemistry model STOCHEM to quantify the impact of NO_x emissions on methane. They compared a 4-year long base simulation with a perturbation simulation that was identical apart from an enhancement in NO_x emissions of magnitude 1 Tg(NO₂), added during the first month with the Northern Hemisphere surface anthropogenic NO_x emissions distribution. The extra NO_x produced a short-lived increase in OH, and this led to a rapid depletion of global methane, which peaked with a magnitude of around 0.39 Tg(CH₄) after about six months. The methane deficit then exponentially decayed with an e-folding timescale of about 12 years (the methane perturbation lifetime, τ), with methane levels returning towards their base values. Wild et al. (2001) conducted similar experiments, with year-long emissions perturbations using a different model (UCI CTM), and found very similar behaviour but with slightly larger sensitivities: 1 Tg(NO₂) from global fossil fuel sources yielded a 0.55 Tg depletion of CH₄. These studies also investigated the impact of CO and NMVOC emissions. Changes in global methane burden (Tg) are converted to changes in tropospheric mole fraction (ppb) using the total atmosphere mass of 5.113×10^9 Tg and a fill factor of 0.973 for conversion of a total atmosphere abundance to a tropospheric abundance (Prather et al., 2012). We assume the troposphere is well mixed, so surface changes will be the same as whole troposphere changes.

More recently, Fry et al. (2012) analysed results from 11 global models that took part in the Hemispheric Transport of Air Pollutants (HTAP) study in order to isolate the impacts on methane of surface NO_x, CO and NMVOC emissions from Europe (EU), North America (NA), South Asia (SA) and East Asia (EA). We utilise that ensemble of model results here; model descriptions are given in Fiore et al. (2009). Models performed a base simulation, and a series of further repeat simulations with 20% lower anthropogenic emissions for each species for each region. In addition to the 20% regional emission reduction experiments, some models also performed global 20% emission reduction experiments (Wild et al., 2012). Four models include results from all the regional and global perturbation simulations: FRSGUCI-v01, GISS-PUCCINI-modelE, MOZARTGFDL-v2, and TM5-JRC-cy2-ipcc-v1. We calculate a 'four model mean' (4MM) based on these model results. We also show results from the other models to illustrate the range of model behaviour, and show 'multi-model mean' (MMM) results from all available simulations.

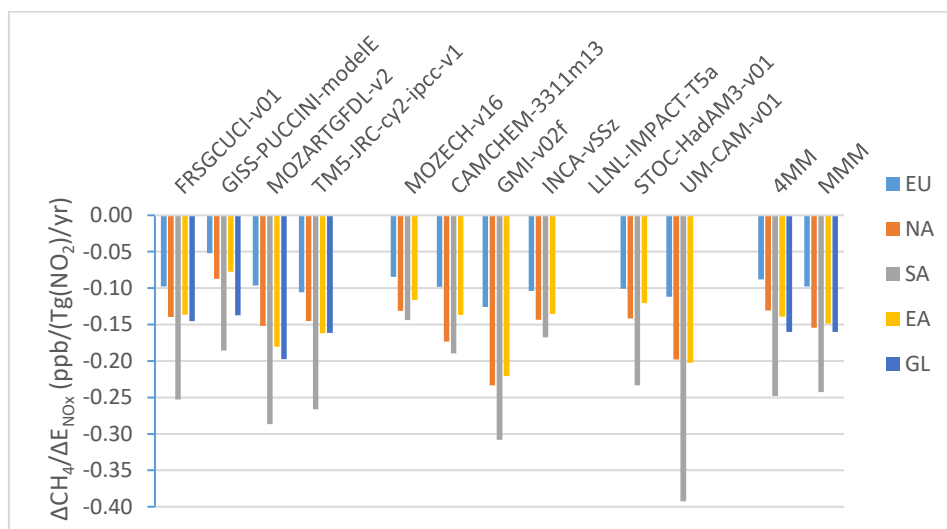
95 In the HTAP simulations, methane was fixed as a prescribed boundary condition, precluding direct diagnosis of changes in methane. However, methane changes can be diagnosed indirectly, by analysing the methane lifetime associated with the tropospheric OH sink in each run. We convert these to whole atmosphere lifetimes by assuming fixed lifetimes for methane loss to soils (160 yr) and in the stratosphere (120 yr) (Prather et al., 2012). The HTAP experiments also included a global methane perturbation simulation – allowing the methane feedback factor and perturbation lifetime to be calculated (Prather, 100 1994; Holmes, 2018). Figure 1 shows whole atmosphere and perturbation methane lifetimes for the HTAP models, with typical values of around 10 years and 12 years, respectively.



105 **Figure 1.** Whole atmosphere and perturbation methane lifetimes (years) for the HTAP models, together with the four model mean (4MM) of the core models (four models on the left), and the multi-model mean (MMM).

Differences between simulations yielded the change in methane lifetimes due to changes in regional emissions. From these changes in methane lifetime, the equilibrium change in methane was calculated; that is the change in methane that would have occurred if methane levels had been free to respond (e.g., see Stevenson et al., 2013). In model simulations where methane is not prescribed, methane adjusts towards equilibrium with an e-folding timescale given by its perturbation lifetime (Derwent et al., 2001; Wild et al., 2001; Holmes, 2018). We convert equilibrium methane changes derived from sustained changes in emissions to the equivalent methane response for a pulse of emissions for each experiment. We use each model's perturbation lifetime to calculate the fraction of the equilibrium response that would have been reached after one year; e.g., for the multi-model mean (MMM) methane perturbation lifetime of 11.6 years (Figure 1) this fraction is $(1 - e^{-1/\tau}) = 8.2\%$. This method is 110 appropriate because we compare to changes in the observed annual growth rate, and is justified by the rapid response of global methane seen in transient model simulations where methane is free to respond, and because the largest lockdown emissions? 115

perturbations occurred in the first half of 2020. We normalise results to produce global methane sensitivities per Tg of gas emitted for each HTAP region and globally for each model. Figures 2, 3 and 4 show global methane sensitivities for NO_x, CO and NMVOC emissions, respectively.



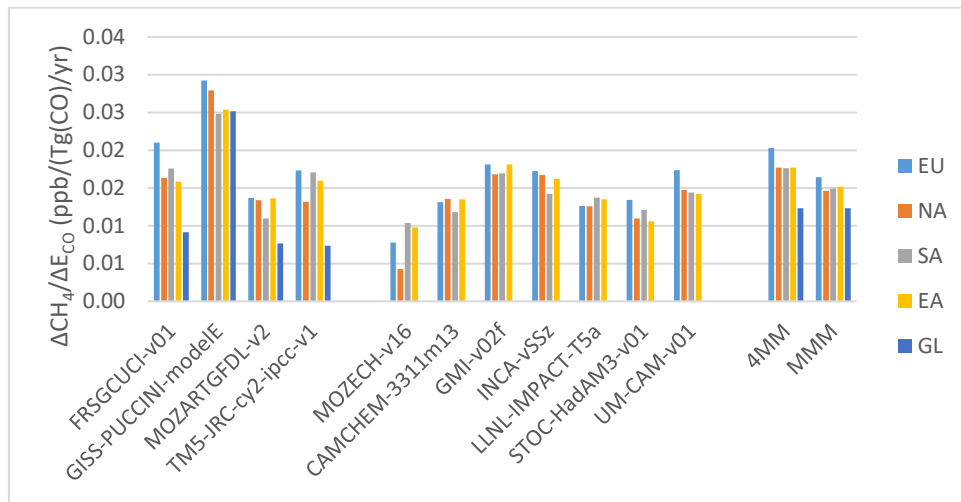
120

Figure 2. Sensitivity of global methane (ppb) to changes in surface anthropogenic NO_x emissions (Tg(NO₂)/yr) for the HTAP models for four regions (Europe, EU; North America, NA; South Asia, SA; and East Asia (EA), and globally (GL).

Global results are only available for the four core models, shown on the left of the figure. Also shown are the 4MM and MMM. There are no results for the LLNL-IMPACT-T5a model for NO_x; it is included to maintain consistency with Figures 3 and 4.

125

Figure 2 shows relatively consistent responses to NO_x emissions, with all models least sensitive to EU NO_x emissions and most sensitive to SA, with NA and EA in between. The 4MM sensitivities are slightly lower than the MMM.



130

Figure 3. As Figure 2, but methane sensitivities for changes in surface anthropogenic CO emissions (Tg(CO)/yr).

As for NO_x, Figure 3 shows relatively consistent behaviour across the models for CO, with less variation between regions, reflecting the longer lifetime of CO, which makes the location of emissions less important. The 4MM sensitivities for CO are slightly larger than the MMM values.

135

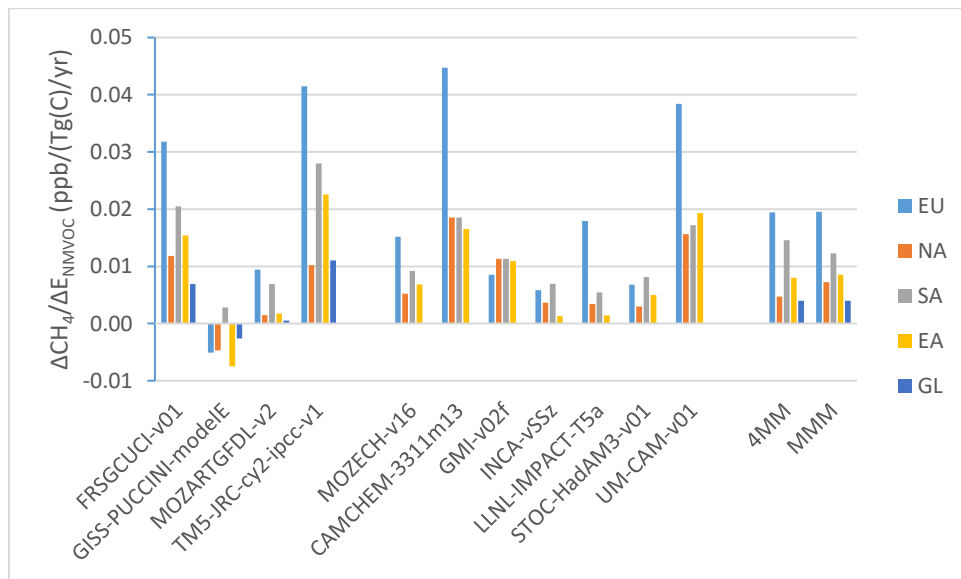


Figure 4. As Figure 2, but methane sensitivities for changes in surface anthropogenic NMVOC emissions (Tg(C)/yr).

140 Figure 4 shows more divergence in model response to NMVOC emissions, with one model (GISS-PUCCINI-modelE) displaying an opposite sensitivity to the other models (apart from for SA emissions), and some models showing quite large

sensitivities, whilst others are small. This probably reflects differing methods of representing NMVOCs in each model, in terms of both the number of species, grouping together of species, and the sophistication of their oxidation chemistry. Somewhat fortuitously, the 4MM and MMM are similar.

145 The HTAP experiments used 2001 as their base year, prescribing global methane to be 1760 ppb, and each model used their own best estimates of global 2001 emissions. In 2020, surface level background global mean methane was ~1870 ppb, and emissions of NO_x, CO and NMVOCs had changed relative to 2001. Sensitivities of methane to emissions derived from the HTAP results will differ somewhat from those that would be found if 2020 conditions were used, and this represents an important caveat to our results. However, these differences are unlikely to be substantial, and no more up-to-date multi-model
150 study of the impacts of regional NO_x, CO and NMVOC emissions on methane has been published to date, so it represents our best source of information in the literature.

Similar model simulations have calculated the sensitivity of methane to aviation NO_x emissions. Wild et al. (2001) and Stevenson et al. (2004) conducted pulse experiments adding NO_x using the global aviation NO_x emission distribution, and found a peak impact on global methane of about 2.5-2.6 Tg (equivalent to mole fractions of 0.88-0.92 ppb) for a 1 Tg(NO₂)
155 emission perturbation. Stevenson and Derwent (2009) also found spatial variation in sensitivity for aviation NO_x, with the more sensitive regions tending to have lower background NO_x levels. The most up-to-date study of aviation NO_x is Lee et al. (2021), who assessed multi-model results using sustained emissions changes, similarly to the HTAP study. Lee et al. (2021) report (their Table 3) a methane radiative forcing sensitivity to aviation NO_x emissions of -15.8 mW m⁻² (Tg(N) yr⁻¹)⁻¹. We convert this to a methane mole fraction sensitivity to NO_x emissions using the relationship between changes in mole fraction
160 and radiative forcing given by Myhre et al. (1998), and then, using a similar methodology to that described above, to the equivalent response for a pulse of emissions. This yields a sensitivity of methane to a pulse change in aviation NO_x emissions of 1.12 ppb (CH₄)/Tg(NO₂) yr⁻¹, similar to, but slightly higher than results from the earlier studies. Lee et al. (2021) also report a 95% likelihood range on the radiative forcing sensitivity, which translates to a standard deviation of 0.21 ppb (CH₄)/Tg(NO₂) yr⁻¹, which we take to be a representative uncertainty for the mole fraction sensitivity to aviation NO_x emissions.

165 **4 COVID-19 lockdown impacts on emissions**

Lamboll et al. (2021) compiled estimates of the impact of COVID-19 lockdowns on global anthropogenic NO_x, CO and NMVOC emissions, as monthly mean time series, with spatial resolution 0.5° latitude by 0.5° longitude. We use these data to calculate the difference in surface NO_x emissions between 2019 (pre-lockdown) and 2020 for the four HTAP regions, as well as globally, and hence for the 'Rest of the World' (ROW) region (i.e. everywhere beyond the four HTAP regions). The annual
170 reduction in global surface NO_x emissions from 2019 to 2020 was about 19.38 Tg(NO₂), or 15%. Lamboll et al. (2021) also compiled data on aviation emissions, estimating a global reduction of about 0.83 Tg(NO₂), or 25%. Global and regional annual changes in NO_x, CO and NMVOC emissions are summarised in Table 1.

	NO _x Tg(NO ₂)	CO Tg(CO)	NMVOC Tg(C)
Global surface emissions (GL)	-19.38	-73.38	-15.65
Europe (EU)	-2.65	-6.09	-1.71
North America (NA)	-2.55	-7.49	-1.56
South Asia (SA)	-3.78	-16.76	-4.34
East Asia (EA)	-4.40	-24.58	-2.41
Rest of the World (ROW)	-6.00	-18.46	-5.63
Global aviation	-0.83	-	-

Table 1. Changes in global and regional annual anthropogenic emissions from 2019 to 2020, assumed to be associated with COVID-19 lockdowns. Derived from data in Lamboll et al. (2021).

180

5 Impacts of reduced lockdown emissions on global methane

To calculate an approximate impact of the lockdown emission reductions on global methane, we simply multiply the regional/aviation sensitivities and emissions changes and sum over the globe. To calculate ROW contributions, we assume that the global sensitivity values can be linearly constructed from the four regions and the ROW, weighting each region by its emissions.

185

Figure 5 shows calculated regional contributions to the global methane growth rate from changes in surface NO_x emissions for each of the HTAP models, together with the 4MM and MMM values. Equivalent results for CO and NMVOCs are shown in Figures 6 and 7, respectively. Table 2 summarises the regional and aviation components for all emissions, using results from the 4MM.

We find that reduced NO_x emissions during lockdown increased the methane growth rate in total by 4.3 ± 0.7 ppb/yr (4MM; a slightly larger impact of 4.4 ± 0.8 ppb/yr is found for the MMM). South Asia is the largest contributing HTAP region, although this is exceeded by the impact from NO_x emissions changes from outside the four HTAP regions. Aviation NO_x is also an important contributor. Reduced CO emissions partly counteracted this positive impact on the methane growth rate, with an overall impact of -1.1 ± 0.5 ppb/yr (4MM; a slightly smaller impact of -1.0 ± 0.4 ppb/yr is found for the MMM). East Asia, followed by South Asia, are the largest contributing regions. Reduced NMVOC emissions had a small additional effect in the

195

same sense as CO, but about one order of magnitude smaller, and with a large uncertainty. The overall impact from NMVOC was -0.13 ± 0.15 ppb/yr (4MM; very similar value for MMM: -0.13 ± 0.10 ppb/yr).

We find a net total impact on methane of 3.0 ± 0.8 ppb/yr (4MM; 3.2 ± 0.9 ppb/yr MMM), with the largest contributing region overall being ROW, followed by South Asia.

200

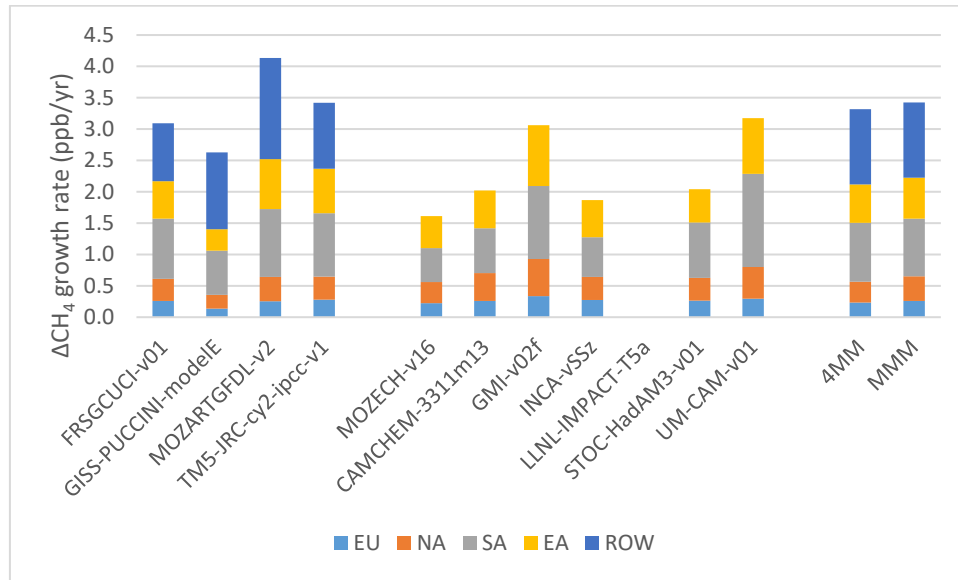


Figure 5. Calculated changes in global methane growth rate from changes in surface NO_x emissions during the 2020 lockdown, for each of the HTAP models. Also shown are values for the mean of the four core models (shown on left) (4MM) that reported results for all simulations, together with multi-model mean (MMM) results based on all available models.

205

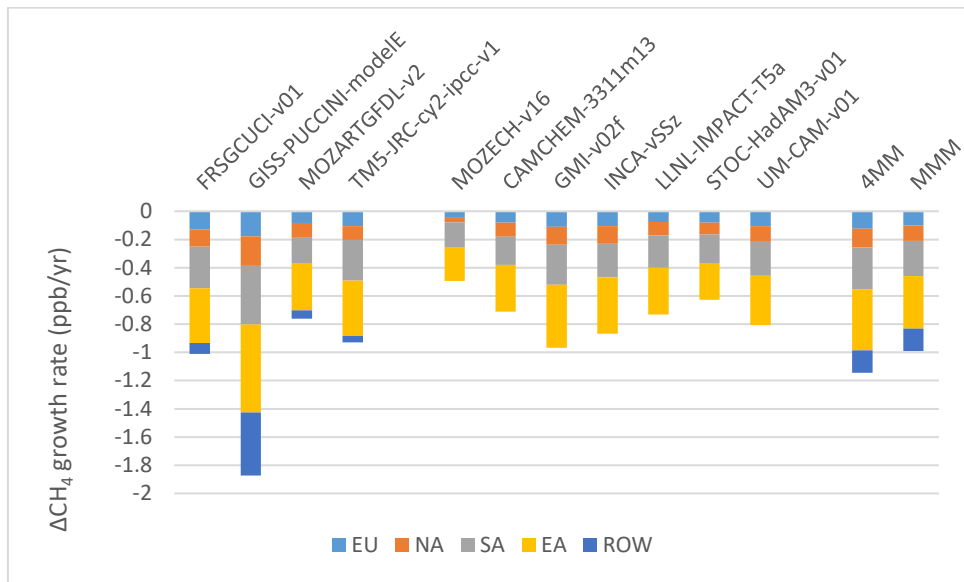


Figure 6. As Figure 5, but for CO emissions.

210

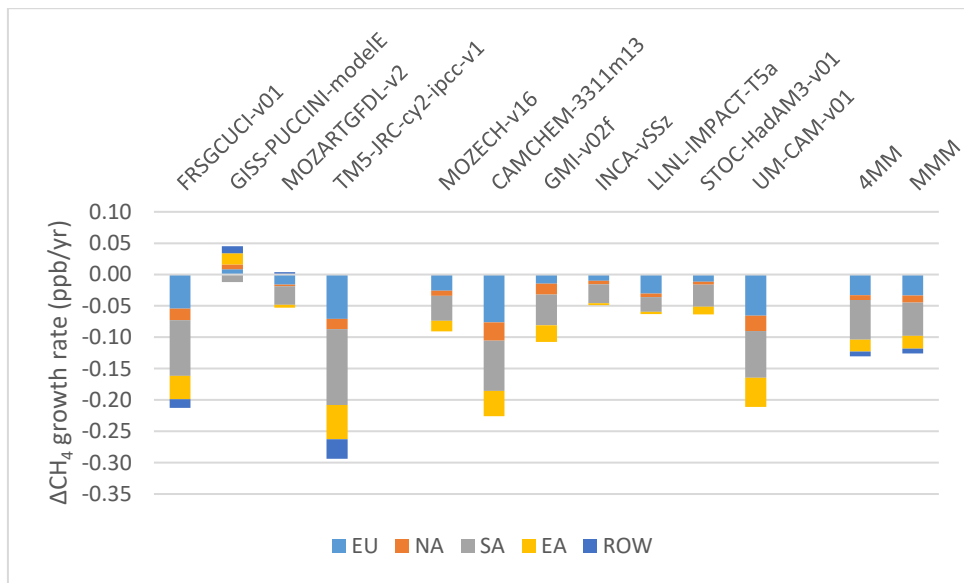


Figure 7. As Figure 5, but for NMVOC emissions.

215

	NO _x	CO	NM VOC	Total
Europe	0.23 ± 0.06	-0.12 ± 0.04	-0.03 ± 0.04	0.08 ± 0.08
N America	0.33 ± 0.08	-0.13 ± 0.05	-0.01 ± 0.01	0.19 ± 0.09
S Asia	0.94 ± 0.17	-0.30 ± 0.09	-0.06 ± 0.05	0.58 ± 0.20
E Asia	0.61 ± 0.20	-0.44 ± 0.13	-0.02 ± 0.03	0.16 ± 0.24
ROW	1.20 ± 0.30	-0.16 ± 0.19	-0.01 ± 0.02	1.04 ± 0.36
Aviation	0.93 ± 0.18			0.93 ± 0.18
Total	4.25 ± 0.66	-1.14 ± 0.50	-0.13 ± 0.15	2.97 ± 0.84

220 **Table 2.** Summary of impacts on the 2020 global methane growth rate (ppb/yr) relative to 2019 due to COVID-19 lockdown
emission reductions based on 4MM results.

6 Discussion and Conclusions

225 These model-derived results can be compared to the observed increase in methane growth rate from 2019 to 2020 of 5.5 ppb/yr,
and suggest that lockdown emission changes in NO_x, CO and NMVOCs can explain 54-58% of this increase. Uncertainties
estimated from the standard deviation of the HTAP (Fry et al., 2012) and aviation NO_x (Lee et al. 2021) model's sensitivity
results are about ±30%. No uncertainty estimate is included here for the magnitude of lockdown emissions changes, which is
probably similar in magnitude. Refinements to this relatively simply derived estimate will need to account for several
additional complications. The emission changes have temporal structure (Lamboll et al., 2021), as do the sensitivities of
230 methane to NO_x, CO and NMVOCs, and these will interact. One study has reported a reduction in lightning during 2020
(Vasquez, 2022), which may contribute much like reductions in aircraft NO_x. The regional sensitivities derived here are based
on emissions changes with the spatial distributions and base magnitudes of the 2001 anthropogenic emissions, rather than a
2020 emissions baseline and the actual changes during lockdown, so the real sensitivities are likely to be slightly different.
Detailed modelling of the lockdown period is starting to explore these effects (Weber et al., 2020; Miyazaki et al., 2021). There
235 is also spatio-temporal structure in the observed methane changes (e.g., Laughner et al., 2021; Feng et al., 2022) that will yield
further information. There are undoubtedly several other factors, in addition to changes in anthropogenic NO_x, CO and
NMVOC emissions that influenced methane during 2020. Nevertheless, it seems likely that the dramatic reductions in these
emissions, especially NO_x, brought about by the COVID-19 lockdowns can explain a large component of the surge in methane
growth rate seen during 2020. These influences on methane related to changes in OH need to be carefully accounted for in any
240 attribution study that attempts to explain the recent observed dramatic changes in methane.

Author contributions

DSS wrote the text and performed the main analysis. OW and WJC performed additional analysis and commented on the text. RGD commented on the text.

Competing interests

245 The authors declare that they have no conflict of interest.

Code/data availability

Original data used here are all freely available in the cited references.

Acknowledgements

This work was partly supported by the Natural Environment Research Council (NE/S009019/1) and the Royal Society
250 (IES\R3\193183). We acknowledge all the modellers who contributed results to the HTAP Phase 1 study: without those results, this work would not have been possible. Jize Jiang is thanked for his technical help with the analysis.

References

- Bauwens, M., Compernelle, S., Stavrakou, T., Müller, J.-F., van Gent, J., Eskes, H., et al.: Impact of coronavirus outbreak on NO₂ pollution assessed using TROPOMI and OMI observations. *Geophys. Res. Lett.*, 47, e2020GL087978,
255 <https://doi.org/10.1029/2020GL087978>, 2020.
- Cooper, M.J., Martin, R.V., Hammer, M.S. et al.: Global fine-scale changes in ambient NO₂ during COVID-19 lockdowns. *Nature* 601, 380–387, <https://doi.org/10.1038/s41586-021-04229-0>, 2022.
- Derwent, R.G., Collins, W.J., Johnson, C.E., and Stevenson, D.S.: Transient Behaviour of Tropospheric Ozone Precursors in a Global 3-D CTM and Their Indirect Greenhouse Effects, *Climatic Change* 49, 463–487,
260 <https://doi.org/10.1023/A:1010648913655>, 2001.
- Dlugokencky, E.: NOAA/GML Trends in Atmospheric Methane, https://gml.noaa.gov/ccgg/trends_ch4/, accessed 03/08/2022, 2022.
- Doumbia, T., C. Granier, Elguindi N., Bouarar I., Darras S., Brasseur G., Gaubert B., Liu Y., Shi X., Stavrakou J., Tilmes S., Lacey F., Deroubaix A., Wang T. 2021 Changes in global air pollutant emissions during the Covid-19 pandemic: a dataset for
265 atmospheric chemistry modeling *Earth Syst. Sci. Data*, 4191–4206 <https://doi.org/10.5194/essd-13-4191-2021>, 2021.

- Feng, L., Palmer, P. I., Parker, R. J., Lunt, M. F., and Boesch, H.: Methane emissions responsible for record-breaking atmospheric methane growth rates in 2020 and 2021, *Atmos. Chem. Phys. Discuss.* [preprint], <https://doi.org/10.5194/acp-2022-425>, in review, 2022.
- Fiore, A. M., et al.: Multimodel estimates of intercontinental source-receptor relationships for ozone pollution, *J. Geophys. Res.*, 114, D04301, doi:10.1029/2008JD010816, 2009.
- Fry, M. M., et al.: The influence of ozone precursor emissions from four world regions on tropospheric composition and radiative climate forcing, *J. Geophys. Res.*, 117, D07306, doi:10.1029/2011JD017134, 2012.
- Holmes, C. D.: Methane feedback on atmospheric chemistry: Methods, models, and mechanisms. *Journal of Advances in Modeling Earth Systems*, 10, 1087– 1099. <https://doi.org/10.1002/2017MS001196>, 2018.
- Lamboll, R. D., Jones, C. D., Skeie, R. B., Fiedler, S., Samset, B. H., Gillett, N. P., Rogelj, J., and Forster, P. M.: Modifying emissions scenario projections to account for the effects of COVID-19: protocol for CovidMIP, *Geosci. Model Dev.*, 14, 3683–3695, <https://doi.org/10.5194/gmd-14-3683-2021>, 2021.
- Laughner, Joshua L., Jessica L. Neu, David Schimel, Paul O. Wennberg, Kelley Barsanti, Kevin W. Bowman, Abhishek Chatterjee, Bart E. Croes, Helen L. Fitzmaurice, Daven K. Henze, Jinsol Kim, Eric A. Kort, Zhu Liu, Kazuyuki Miyazaki, Alexander J. Turner, Susan Anenberg, Jeremy Avice, Hansen Cao, David Crisp, Joost de Gouw, Annmarie Eldering, John C. Fyfe, Daniel L. Goldberg, Kevin R. Gurney, Sina Hasheminassab, Francesca Hopkins, Cesunica E. Ivey, Dylan B. A. Jones, Junjie Liu, Nicole S. Lovenduski, Randall V. Martin, Galen A. McKinley, Lesley Ott, Benjamin Poulter, Muye Ru, Stanley P. Sander, Neil Swart, Yuk L. Yung, Zhao-Cheng Zeng, Societal shifts due to COVID-19 reveal large-scale complexities and feedbacks between atmospheric chemistry and climate change, *Proceedings of the National Academy of Sciences Nov 2021*, 118 (46) e2109481118; DOI: 10.1073/pnas.2109481118, 2021.
- Lee, D.S., D.W. Fahey, A. Skowron, M.R. Allen, U. Burkhardt, Q. Chen, S.J. Doherty, S. Freeman, P.M. Forster, J. Fuglestedt, A. Gettelman, R.R. De León, L.L. Lim, M.T. Lund, R.J. Millar, B. Owen, J.E. Penner, G. Pitari, M.J. Prather, R. Sausen, and L.J. Wilcox: The contribution of global aviation to anthropogenic climate forcing for 2000 to 2018, *Atmospheric Environment*, Volume 244, 117834, <https://doi.org/10.1016/j.atmosenv.2020.117834>, 2021.
- Miyazaki, K., K. Bowman, T. Sekiya, M. Takigawa, J. L. Neu, K. Sudo, G. Osterman, and H. Eskes: Global tropospheric ozone responses to reduced NOx emissions linked to the COVID-19 worldwide lockdowns, *Science Advances*, 7, 24, eabf7460 DOI:10.1126/sciadv.abf7460, 2021.
- Myhre, G., E. Highwood, K. Shine, and F. Stordal: New estimates of radiative forcing due to well mixed greenhouse gases, *Geophys. Res. Lett.*, 25(14), 2715– 2718, doi:10.1029/98GL01908, 1998.
- Prather, M.J.: Lifetimes and Eigenstates in Atmospheric Chemistry, *Geophys. Res. Lett.*, 21, 801-804, 1994.
- Prather, M. J., Holmes, C. D., and Hsu, J.: Reactive greenhouse gas scenarios: Systematic exploration of uncertainties and the role of atmospheric chemistry, *Geophys. Res. Lett.*, 39, L09803, doi:10.1029/2012GL051440, 2012.
- Shindell, D.T., G. Faluvegi, N. Bell, and G.A. Schmidt: An emissions-based view of climate forcing by methane and tropospheric ozone. *Geophys. Res. Lett.*, 32, L04803, doi:10.1029/2004GL021900, 2005.

- 300 Shindell, D.T., G. Faluvegi, D.M. Koch, G.A. Schmidt, N. Unger, and S.E. Bauer: Improved attribution of climate forcing to emissions. *Science*, 326, 716-718, doi:10.1126/science.1174760, 2009.
- Stevenson, D.S. and R.G. Derwent: How does the location of aircraft nitrogen oxide emissions affect their climate impact? *Geophys. Res. Lett.*, 36, L17810, doi:10.1029/2009GL039422, 2009.
- Stevenson, D. S., Doherty, R. M., Sanderson, M. G., Collins, W. J., Johnson, C. E., and Derwent, R. G.: Radiative forcing
305 from aircraft NO_x emissions: Mechanisms and seasonal dependence, *J. Geophys. Res.*, 109, D17307, doi:10.1029/2004JD004759, 2004.
- Stevenson, D. S., Young, P. J., Naik, V., Lamarque, J.-F., Shindell, D. T., Voulgarakis, A., Skeie, R. B., Dalsoren, S. B., Myhre, G., Berntsen, T. K., Folberth, G. A., Rumbold, S. T., Collins, W. J., MacKenzie, I. A., Doherty, R. M., Zeng, G., van Noije, T. P. C., Strunk, A., Bergmann, D., Cameron-Smith, P., Plummer, D. A., Strode, S. A., Horowitz, L., Lee, Y. H., Szopa,
310 S., Sudo, K., Nagashima, T., Josse, B., Cionni, I., Righi, M., Eyring, V., Conley, A., Bowman, K. W., Wild, O., and Archibald, A.: Tropospheric ozone changes, radiative forcing and attribution to emissions in the Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP), *Atmos. Chem. Phys.*, 13, 3063–3085, <https://doi.org/10.5194/acp-13-3063-2013>, 2013.
- Szopa, S., V. Naik, B. Adhikary, P. Artaxo, T. Berntsen, W.D. Collins, S. Fuzzi, L. Gallardo, A. Kiendler-Scharr, Z. Klimont,
315 H. Liao, N. Unger, and P. Zanis, 2021: Short-Lived Climate Forcers. In *Climate Change 2021: The Physical Science Basis. Contribution of Working Group I to the Sixth Assessment Report of the Intergovernmental Panel on Climate Change* [Masson-Delmotte, V., P. Zhai, A. Pirani, S.L. Connors, C. Péan, S. Berger, N. Caud, Y. Chen, L. Goldfarb, M.I. Gomis, M. Huang, K. Leitzell, E. Lonnoy, J.B.R. Matthews, T.K. Maycock, T. Waterfield, O. Yelekçi, R. Yu, and B. Zhou (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, pp. 817–922, doi:10.1017/9781009157896.008.
- 320 Thornhill, G. D., Collins, W. J., Kramer, R. J., Olivíe, D., Skeie, R. B., O'Connor, F. M., Abraham, N. L., Checa-Garcia, R., Bauer, S. E., Deushi, M., Emmons, L. K., Forster, P. M., Horowitz, L. W., Johnson, B., Keeble, J., Lamarque, J.-F., Michou, M., Mills, M. J., Mulcahy, J. P., Myhre, G., Nabat, P., Naik, V., Oshima, N., Schulz, M., Smith, C. J., Takemura, T., Tilmes, S., Wu, T., Zeng, G., and Zhang, J.: Effective radiative forcing from emissions of reactive gases and aerosols – a multi-model comparison, *Atmos. Chem. Phys.*, 21, 853–874, <https://doi.org/10.5194/acp-21-853-2021>, 2021.
- 325 Turner, A.J., I. Fung, V. Naik, L. W. Horowitz, R. C. Cohen: Modulation of hydroxyl variability by ENSO in the absence of external forcing, *Proceedings of the National Academy of Sciences*, 115 (36) 8931-8936; DOI: 10.1073/pnas.1807532115, 2018.
- Vasquez, K.: Cleaner pandemic air led to reduced lightning strikes worldwide, *Eos*, 103, <https://doi.org/10.1029/2022EO220048>, 2022.
- 330 Venter, Z.S., K. Aunan, S. Chowdhury, J. Lelieveld: COVID-19 lockdowns cause global air pollution declines, *Proceedings of the National Academy of Sciences*, 117 (32) 18984-18990; DOI: 10.1073/pnas.2006853117, 2020.

- Weber, J., Shin, Y. M., Staunton Sykes, J., Archer-Nicholls, S., Abraham, N. L., & Archibald, A. T.: Minimal climate impacts from short-lived climate forcers following emission reductions related to the COVID-19 pandemic. *Geophysical Research Letters*, 47, e2020GL090326. <https://doi.org/10.1029/2020GL090326>, 2020.
- 335 Wild, O., Fiore, A. M., Shindell, D. T., Doherty, R. M., Collins, W. J., Dentener, F. J., Schultz, M. G., Gong, S., MacKenzie, I. A., Zeng, G., Hess, P., Duncan, B. N., Bergmann, D. J., Szopa, S., Jonson, J. E., Keating, T. J., and Zuber, A.: Modelling future changes in surface ozone: a parameterized approach, *Atmos. Chem. Phys.*, 12, 2037–2054, <https://doi.org/10.5194/acp-12-2037-2012>, 2012.
- Wild, O., Prather, M.J., and Akimoto, H.: Indirect long-term global radiative cooling from NO_x emissions, *Geophys. Res. Lett.*, 28, 1719-1722, 2001.
- 340 World Meteorological Organization: State of the Global Climate 2020 (WMO-No. 1264), <https://public.wmo.int/en/our-mandate/climate/wmo-statement-state-of-global-climate>, accessed 14-07-21, 2021.
- Zhao, Y., Saunio, M., Bousquet, P., Lin, X., Berchet, A., Hegglin, M. I., Canadell, J. G., Jackson, R. B., Deushi, M., Jöckel, P., Kinnison, D., Kirner, O., Strode, S., Tilmes, S., Dlugokencky, E. J., and Zheng, B.: On the role of trend and variability in the hydroxyl radical (OH) in the global methane budget, *Atmos. Chem. Phys.*, 20, 13011–13022, <https://doi.org/10.5194/acp-20-13011-2020>, 2020.
- 345