



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

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Abstract. Compared to 2019, the global growth rate of atmospheric methane rose by about 50% in 2020, reaching 15 ppb/yr. Models of global atmospheric chemistry show that reductions in nitrogen oxide (NO_x) emissions reduce levels of the hydroxyl radical, and lengthen the methane lifetime. Using estimates of NO_x emission reductions associated with COVID-19 lockdowns around the world in 2020, together with model-derived regional and sectoral sensitivities of methane to NO_x emissions, we find that NO_x emissions reductions can fully explain the observed surge in the global methane growth rate. Whilst changes in NO_x emissions are probably not the only important factor that has influenced methane since the beginning of 2020, it is clear that they are a key factor that will need to be included within any attribution study, and that they may well be the dominant driver of these recent methane changes. The major global scale changes in composition of the Earth's atmosphere measured during lockdown provide unprecedented constraints on the sensitivity of the atmospheric chemical system to changes in emissions, and are of great utility for evaluating policy-relevant models.

1 Introduction

20 Methane is a powerful greenhouse gas and important precursor of tropospheric ozone; both are key air pollutants and short-lived climate forcers (SLCFs). The 2013 Intergovernmental Panel on Climate Change assessment estimated methane's Global Warming Potential (GWP) over a 100 year time horizon to be 28 (Myhre et al., 2013); updates to its short-wave radiative forcing have increased this value by 14% (Etminan et al., 2016). Advances in our understanding of how GWP metrics relate to climate change indicate that it is not only the magnitude of emissions, but also the rates of change of SLCFs like methane that strongly influence near-term global temperature changes (Allen et al., 2018; Cain et al., 2019). These post-2013 updates increase the importance of methane and its evolution in the context of 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 has shown substantial changes. Atmospheric nitrogen oxide (NO_x) levels have reduced, whilst the measured growth rate of methane (CH₄) has risen sharply. The observed NO_x changes are clearly linked to falls in emissions resulting from lockdowns, but the



methane increases remain unexplained (e.g., Vaughan, 2021). Methane and NO_x are linked through the oxidising capacity of the atmosphere, specifically by the abundance of the hydroxyl (OH) radical. The response of global atmospheric chemistry to the large lockdown perturbation since early 2020 provides an opportunity to explore the sensitivity of the NO_x-OH-CH₄ system, and compare models and observations. Here we use model-derived sensitivities of methane to NO_x, together with
35 estimated changes in anthropogenic NO_x emissions related to the COVID-19 lockdowns, to calculate the impacts on the growth rate of global methane, and compare this to methane observations.

2 Measurements of atmospheric methane and nitrogen oxides

Recent methane measurements from the US National Oceanographic and Atmospheric Administration (NOAA) show that the atmospheric methane growth rate rose sharply from 9.9 ppb/yr in 2019 to 14.8 ppb/yr in 2020, the highest annual value in the
40 37-year NOAA record (Figure 1; Dlugokencky, 2021). 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 Southern Oscillation (ENSO), and in particular because of modulation of methane's main sink, oxidation by OH (Turner et al., 2018; Zhao et al., 2020). ENSO indices have not shown strong variations over 2019-2020 (WMO, 2021).

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

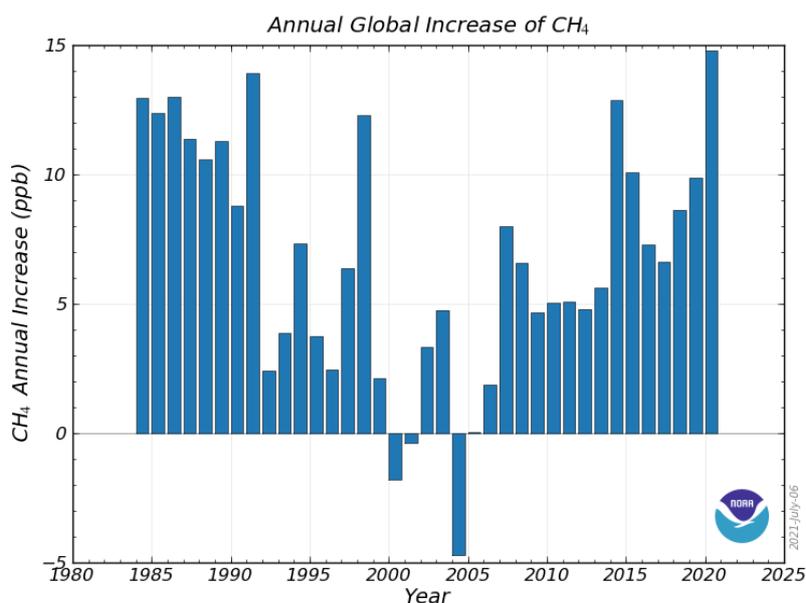


Figure 1: Global annual changes in surface atmospheric methane mole fraction (ppb) 1984-2020 (Dlugokencky, 2021).



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3 Sensitivity of global methane to NO_x emissions

Global atmospheric chemistry model simulations indicate that decreases in NO_x emissions lead to reductions in OH and increases in the methane lifetime (Prather, 1994; Derwent et al., 2001; Wild et al., 2001; Stevenson et al., 2004; Weber et al., 2020). Multi-model studies have calculated methane effects for NO_x emissions from specific world regions (Fry et al., 2012) and the aviation sector (Lee et al., 2021). Although methane has an atmospheric lifetime of about 10 years, the 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 NO_x emissions (Derwent et al., 2001; Wild et al., 2001; Stevenson et al., 2004).

Derwent et al. (2001) conducted a series of experiments 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 at 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), with methane levels returning towards their base values. Wild et al. (2001) conducted similar experiments, with year-long perturbations using a different model (UCI CTM), and found slightly larger sensitivities: 1 Tg(NO₂) from global fossil fuel sources yielded a 0.55 Tg depletion of CH₄. Fry et al. (2012) analysed multi-model experiments from the Hemispheric Transport of Air Pollutants (HTAP) study that isolated the impacts on methane of surface NO_x emissions from Europe, North America, and South and East Asia. For each region, Fry et al. (2012) compared a base simulation to one with 20% lower anthropogenic NO_x emissions from that region. These simulations had methane fixed as a prescribed boundary condition, but diagnosed the change in methane lifetime associated with changes in OH. From these changes in methane lifetime, the equilibrium change in methane was calculated, that is the change in methane that would have been achieved if methane levels had been free to respond (e.g., see Stevenson et al., 2013). In model simulations with methane not prescribed, it adjusts towards equilibrium with an e-folding time given by the methane perturbation lifetime (τ) (Holmes, 2018). To convert equilibrium methane changes derived from sustained changes in emissions to the equivalent response for a pulse of emissions, we use the perturbation lifetime to calculate the fraction of the equilibrium response that would have been reached after one year; for a methane perturbation lifetime of 12.4 years (Holmes, 2018) this fraction is $(1 - e^{-1/\tau}) = 7.7\%$.

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 emissions distribution, and found a peak impact on methane of about 2.5-2.6 Tg for a 1 Tg(NO₂) emission perturbation. Lee et al. (2021) assessed multi-model results for aviation NO_x emissions using sustained changes, which we convert, as described above, to the equivalent response for a pulse of emissions, and find a similar sensitivity.



Table 1 summarises results from all these existing studies. For surface NO_x emissions reductions of 1 Tg(NO₂) over one year, models find peak increases in global methane burden of about 0.5 Tg, with about five times higher sensitivities for reductions in aviation NO_x. There is significant variation in sensitivity between regions for surface NO_x emissions, with South Asia about twice as sensitive as North America and East Asia, and three times as sensitive as Europe (Fry et al., 2012). 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.

NO _x emission region/sector	$\Delta\text{CH}_4/\Delta E_{\text{NO}_x}$ Tg(CH ₄)/Tg(NO ₂) yr ⁻¹	2020-2019 ΔE_{NO_x} Tg(NO ₂) yr ⁻¹	ΔCH_4 Tg(CH ₄)
<i>Surface emissions</i>			
Global	-0.55 ^a	-19.38	
N. Hemisphere	-0.39 ^b	-16.72	
S. Hemisphere	-1.1 ^b	-2.66	2.93
Europe	-0.28 ^c	-2.65	0.74
N. America	-0.47 ^c	-2.55	1.20
E. Asia	-0.44 ^c	-4.40	1.94
S. Asia	-0.88 ^c	-3.78	3.33
NH minus 4 regions	-0.39 ^b	-3.34	1.30
<i>Aviation emissions</i>			
Global	-2.6 ^a	-0.83	
Global	-2.5 ^d	-0.83	
Global	-2.3 ^e	-0.83	1.91
Various 10° x 10° model grid-boxes	-1.9 to -15 ^f		

90 **Table 1: Sensitivity of changes in the global methane burden (ΔCH_4 ; units Tg(CH₄)) to changes in NO_x emissions (ΔE_{NO_x} ; units Tg(NO₂) yr⁻¹) from several modelling studies, calculated for a variety of surface and aviation emissions from different regions. Also shown are COVID-19 lockdown impacts on NO_x emissions (Tg(NO₂) yr⁻¹) between 2019 and 2020 from Lamboll et al. (2021) for regions and global aviation, and the corresponding contributions to the change in global methane burden (Tg(CH₄); values only given for contributions used in Section 4). References for the sensitivity values: (a) Wild et al. (2001); (b) Derwent et al. (2001); (c) Fry et al. (2012); (d) Stevenson et al. (2004); (e) Lee et al. (2021); and (f) Stevenson and Derwent (2009).**



4 COVID-19 lockdown impacts on NO_x emissions

Lamboll et al. (2021) compiled estimates of the impact of COVID-19 lockdowns on global anthropogenic NO_x 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 for the Northern and Southern Hemispheres. The annual 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%. Regional changes in NO_x emissions are summarised in Table 1.

5 Impacts of reduced NO_x emissions on global methane

To calculate an approximate impact of the NO_x emission reductions on global methane, we simply multiply the regional/aviation sensitivities and emissions changes and sum over the globe. Table 1 shows calculated regional and aviation components of the methane change. We calculate a total methane burden change of 13.36 Tg(CH₄), comprising 11.44 Tg(CH₄) from surface NO_x changes, with a further 1.92 Tg(CH₄) from aircraft. The more sensitive regions (South Asia, the Southern Hemisphere) and aviation make proportionally larger contributions to the total methane change. We convert the overall change in global methane burden (Tg) to a change 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). This yields a global mean increase in tropospheric methane mole fraction of 4.9 ppb associated with the NO_x reductions. Since the troposphere is well mixed, this is also the change at the surface.

6 Discussion and Conclusions

This model-derived estimate of the extra methane expected due to the reductions in NO_x emissions exactly matches the observed extra growth in methane seen during lockdown from 2019 to 2020 (4.9 ppb), suggesting that the NO_x changes can account for all or most of the observed methane changes.

Refinements to this simple estimate will need to account for several additional complications. The NO_x emission changes have temporal structure (Lamboll et al., 2021), as do the sensitivities of methane to NO_x (e.g., Stevenson et al., 2004), and these will interact. In addition, we have ignored any spatial variations in aircraft emissions, but Stevenson and Derwent (2009) found that NO_x emissions into cleaner environments had larger effects; there are also likely spatial variations within the large regions we have used for the surface NO_x emissions. Detailed modelling of the lockdown period is starting to explore these effects (Weber et al., 2020; Miyazaki et al., 2021). There is also spatio-temporal structure in the observed methane changes that will yield further information. There are undoubtedly several other factors, in addition to changes in NO_x, that influenced methane during 2020. Nevertheless, it seems likely that the dramatic reductions in NO_x emissions brought about by the COVID-19 lockdowns can explain a large component of the surge in methane growth rate seen since early 2020.



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

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

References

- Allen, M. R., Shine, K. P., Fuglestvedt, J. S., Millar, R. J., Cain, M., Frame, D. J., and Macey, A.H.: A solution to the misrepresentations of CO₂-equivalent emissions of short-lived climate pollutants under ambitious mitigation. *npj Clim. Atmos. Sci.* 1, 16, <https://doi.org/10.1038/s41612-018-0026-8>, 2018.
- 135 Bauwens, M., Compornolle, S., Stavrou, 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, <https://doi.org/10.1029/2020GL087978>, 2020.
- Cain, M., Lynch, J., Allen, M.R. et al.: Improved calculation of warming-equivalent emissions for short-lived climate pollutants. *npj Clim. Atmos. Sci.* 2, 29, <https://doi.org/10.1038/s41612-019-0086-4>, 2019.
- 140 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, <https://doi.org/10.1023/A:1010648913655>, 2001.
- Dlugokencky, E.: NOAA/GML Trends in Atmospheric Methane, https://gml.noaa.gov/ccgg/trends_ch4/, accessed 25/06/2021, 2021.
- 145 Etminan, M., G. Myhre, E. J. Highwood, and K. P. Shine: Radiative forcing of carbon dioxide, methane, and nitrous oxide: A significant revision of the methane radiative forcing, *Geophys. Res. Lett.*, 43, 12,614–12,623, doi:10.1002/2016GL071930, 2016.
- 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.
- 150 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.



- 155 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
160 ozone responses to reduced NO_x emissions linked to the COVID-19 worldwide lockdowns, *Science Advances*, 7, 24, eabf7460
DOI:10.1126/sciadv.abf7460, 2021.
- Myhre, G., D. Shindell, F.-M. Bréon, W. Collins, J. Fuglestedt, J. Huang, D. Koch, J.-F. Lamarque, D. Lee, B. Mendoza, T. Nakajima, A. Robock, G. Stephens, T. Takemura and H. Zhang: Anthropogenic and Natural Radiative Forcing. In: *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the*
165 *Intergovernmental Panel on Climate Change* [Stocker, T.F., D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013.
- 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
170 role of atmospheric chemistry, *Geophys. Res. Lett.*, 39, L09803, doi:10.1029/2012GL051440, 2012.
- 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
from aircraft NO_x emissions: Mechanisms and seasonal dependence, *J. Geophys. Res.*, 109, D17307,
175 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,
S., Sudo, K., Nagashima, T., Josse, B., Cionni, I., Righi, M., Eyring, V., Conley, A., Bowman, K. W., Wild, O., and Archibald,
180 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.
- 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,
185 2018.
- Vaughan, A.: A mysterious rise in methane levels is sparking global warming fears, *New Scientist*,
<https://www.newscientist.com/article/mg25033350-700-a-mysterious-rise-in-methane-levels-is-sparking-global-warming-fears/> [accessed 14th July 2021], 2021.



- 190 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 forcings following emission reductions related to the COVID-19 pandemic. Geophysical Research
Letters, 47, e2020GL090326. <https://doi.org/10.1029/2020GL090326>, 2020.
- 195 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.
- 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,
200 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.