



Contributions of lightning to long-term trends and inter-annual variability in global atmospheric chemistry constrained by Schumann resonance observations

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Abstract. Lightning is a significant source of nitrogen oxides ($\text{NO}_x \equiv \text{NO} + \text{NO}_2$) in the free troposphere. Variations in global lightning activity influence long-term trends (LTTs) and inter-annual variability (IAV) in tropospheric NO_x , ozone (O_3), and hydroxyl radicals (OH). However, accurately quantifying these impacts is hindered by uncertainties in representing year-to-year fluctuations in global lightning activity in models. Here, we apply Schumann resonance (SR) observations, which are sensitive to changes in global lightning activity, to better constrain inter-annual variations in lightning NO_x (LNO_x) emissions. By integrating this update into an atmospheric chemical transport model, we assess the contributions of lightning to both LTT and IAV in global atmospheric chemistry from 2013–2021. The updated parameterisation predicts an insignificant trend in global LNO_x emissions, contrasting with a significant increase of $6.4\% \text{ dec}^{-1}$ ($P < 0.05$) by the original parameterisation, reducing lightning contributions to LTTs in NO_x , O_3 , and OH. The updated simulation better aligns with satellite-observed trends in global and northern hemispheric NO_2 but further underestimates tropospheric O_3 increases. The updated parameterisation reveals twice the IAV in global LNO_x emissions but 20 % smaller IAVs in global O_3 and OH because lightning generally counteracts other sources of natural variability. A $\sim 10\%$ decline in lightning in 2020 relative to 2019 led to a $\sim 2\%$ decrease in global OH, explaining half of observed annual methane growth. These findings highlight the value of Schumann resonance observations in constraining global lightning activity, thereby enhancing our understanding of lightning's role in atmospheric chemistry.

1 Introduction

Lightning generates reactive nitrogen oxides ($\text{NO}_x \equiv \text{NO} + \text{NO}_2$) by breaking down N_2 and O_2 in the intense heat of its flash channels, making it a primary natural source of NO_x in the atmosphere (U. Schumann and Huntrieser, 2007). Although lightning accounts for only approximately 10 % of global NO_x emissions, lightning NO_x (referred to

as LNO_x) emissions have a disproportionately large impact on tropospheric ozone (O_3) and hydroxyl radicals (OH) because lightning emits NO_x directly to the free troposphere, where O_3 and OH production is more sensitive to NO_x than at the surface (Liaskos et al., 2015; Murray, 2016; Murray et al., 2021; Nussbaumer et al., 2023). For example, although LNO_x emissions are of a similar magnitude to those from biomass burning or soils, their contribution to tropospheric

O₃ is about 3 times larger (Dahmann et al., 2011). The contribution of LNO_x to global atmospheric OH abundances is more than 20 % (Murray et al., 2021).

Tropospheric O₃ and OH are important atmospheric species both chemically and environmentally. O₃ is a potent greenhouse gas in the upper troposphere (IPCC, 2021), and when transported to the surface, it acts as an air pollutant that adversely affects the health of ecosystems and populations (Fleming et al., 2018; Unger et al., 2020). OH is the main sink for a diverse array of reduced species in the atmosphere, thereby influencing their environmental and climate impacts. For instance, even a small fluctuation in global mean OH concentrations can have a large effect on the budget of methane (CH₄), the second-most important greenhouse gas after CO₂, at the inter-annual scale (Turner et al., 2017; Rigby et al., 2017). Therefore, through regulating atmospheric chemistry, changes in global lightning activity can carry substantial implications for ecosystems, human health, and the climate.

Most atmospheric chemistry transport models (CTMs) rely on sub-grid parameterisations of lightning flash rates to estimate LNO_x emissions, as fine-scale lightning physics cannot be fully captured by coarse-resolution CTMs (Murray et al., 2012). These parameterisations are often based on proxies of deep convection, such as the cloud-top height (Price and Rind, 1992) and the upward cloud ice flux (Finney et al., 2014). Cloud-top height parameterisation and its variants are commonly employed in CTMs. While the parameterised lightning flash rates align with convection patterns in simulations, they may not replicate the global spatial patterns of lightning activity (Tost et al., 2007). This limitation is improved by incorporating lightning observations from satellite-based lightning imaging sensors (LISs) and an optical transient detector (OTD) to constrain the distribution of LNO_x emission provided by the parameterisation (Murray et al., 2012).

In addition, there is no assurance that parameterised lightning flash rates can capture year-to-year variations in global lightning activity and LNO_x emissions (Clark et al., 2017). Studies have reported substantial disagreement between different parameterisations on the magnitude and even the direction of LTTs. For instance, parameterisation based on cloud-top height tends to predict an increase in global lightning activity in response to climate change, while estimations based on upward cloud ice fluxes predict the opposite outcome (Jacobson and Streets, 2009; Clark et al., 2017; Finney et al., 2018). This drawback impedes the assessment of lightning's impact on the LTTs and IAV of global tropospheric O₃ and OH budgets.

Although lightning can be observed from ground- and space-based platforms, these observations may not offer adequate global-scale constraints on changes in lightning activity. The World Wide Lightning Location Network (WWLLN), operational since 2012, is a global ground-based lightning observation network. Despite providing detailed in-

formation on individual cloud-to-ground lightning strikes, these measurements lack sensitivity to intra-cloud lightning strikes (Abarca et al., 2010) and exhibit uneven global detection efficiency that varies over years (Hutchins et al., 2012), thereby limiting their utility in long-term global investigations (Bozóki et al., 2023). On the other hand, satellite observations, including the OTD on board OrbView-1 (formerly MicroLab-1) (May 1995–February 2000; coverage between ±70° latitude) and the LISs on board the Tropical Rainfall Measuring Mission (TRMM) satellite (January 1998–March 2015; coverage between ±38° latitude) and the International Space Station (ISS) (March 2017–October 2023; coverage between ±55° latitude), provide more uniform detection. However, these satellite data records are not continuous, and their data coverage and detection efficiency vary with the instrument and observation platform (Blakeslee et al., 2020; Clark et al., 2017).

Alternatively, global lightning activity can be derived from measurements of Schumann resonances (SRs), which are low-frequency electromagnetic waves within the atmosphere, primarily generated by lightning activity and resonating between the surface and the ionosphere (Balser and Wagner, 1960; Schumann, 1952; Nickolaenko and Hayakawa, 2002). Despite offering limited spatial information, the SR signal is sensitive to global lightning activity, in contrast to other observations with restricted coverage and variable detection efficiencies. This global sensitivity can be attributed to the minimal attenuation of lightning-induced radio waves within this specific frequency band (Chapman et al., 1966). Long-term SR measurements are also influenced by the solar cycle, as variations in solar X-ray flux and precipitating energetic particles lead to changes in wave propagation conditions (Bozóki et al., 2021). To effectively characterise global lightning activity using SR data, it is essential to eliminate the influence of solar cycles (Bozóki et al., 2021).

In this study, we apply SR observations to constrain global lightning activity and LNO_x emissions. Using model simulations, we quantify the impact of this update on LNO_x emissions and subsequently on global atmospheric chemistry. We focus the analysis on the global and hemispheric LTTs and IAV of NO_x, O₃, and OH. We also explore the implications of lightning variability for the global budgets of tropospheric O₃ and CH₄.

2 Data and methods

2.1 Lightning observation data

We use SR measurements at the Eskdalemuir Geomagnetic Observatory (ESK) in the UK, operated by the British Geological Survey (Beggan and Musur, 2018; Musur and Beggan, 2019), between 2013 and 2021. The measurements are performed at a cadence of 100 Hz with two orthogonal horizontal induction coils. Following Bozóki et al. (2021), we apply the weighted average method to obtain the intensities of

the first three SR modes and manually exclude the disturbed days from the dataset. Such disturbed days are either of natural (e.g., local lightning activity; see Tatsis et al., 2021) or anthropogenic (e.g., nearby human activity; see Tritakis et al., 2021) origin. The removal of these data is based on the evidence that the diurnal variation in SR intensities is usually very similar within a given month (Bozóki et al., 2021; Sántori, 1996). Therefore, days with unusual diurnal variation are considered to be disturbed and are removed from the dataset during the manual sanitation process. We then derive the SR intensity, as a proxy for monthly global lightning activity, by summing the intensities of the first three resonance modes for both horizontal magnetic field components. We employ a third-order polynomial fit on the full data record to remove the influence of solar cycle variations.

To mitigate the impact of changes in the source–observer distance (Sántori et al., 2009) and day–night asymmetry (Melnikov et al., 2004), we utilise the SR intensity to parameterise the inter-annual variation in global lightning activity for each individual month. The premise is that in a given month the day–night condition does not change substantially and that the source–observer distance is the same year by year (Sántori et al., 2024). As a result, we use this SR dataset to estimate the shift in global lightning activity between, for instance, January 2013–January 2014, while it is not used to discern the changes from January 2013–February 2013. Figure 1 presents the 2013–2021 annual variation in monthly global lightning activity derived from SR observations.

We also compare SR-based global lightning activity with other global lightning observation data, including the ISS LIS data for 2018–2021, the TRMM LIS data for 2013–2014, and the WWLLN data (v2022.0.0) for 2013–2021 (Kaplan and Lau, 2021, 2022).

2.2 Lightning NO_x emissions

LNO_x emissions are computed in GEOS-Chem as the product of the lightning flash rate (F) and NO production per flash (P_{NO}) for each grid-cell column at every model time step. The column total LNO_x emissions are then vertically distributed based on Ott et al. (2010). GEOS-Chem estimates the lightning flash rate based on the cloud height parameterisation (Price and Rind, 1992), with its global distribution constrained by the climatology of OTD/LIS observations (Murray et al., 2012). This method is referred to as the original parameterisation (OP) hereafter.

To constrain the IAV of the global lightning flash rate with SR intensity, we apply a global scaling factor ($S(m, y)$), as a function of month m and year y , to the lightning flash rates computed from the OP method. Here we assume that for each individual month, the global lightning flash rate is proportional to the SR intensity, which is consistent with Boldi et al. (2018), Dyrda et al. (2014), and Heckman et al. (1998).

The scaling factor is computed as

$$S(m, y) = \frac{\overline{F_{\text{OP}}(m)}}{F_{\text{OP}}(m, y)} \frac{\text{SR}(m, y)}{\overline{\text{SR}(m)}}, \quad (1)$$

where $F_{\text{OP}}(m, y)$ is the global lightning flash rate in month m and year y computed by the OP method, $\overline{F_{\text{OP}}(m)}$ is the multi-year (2013–2021) average of $F_{\text{OP}}(m, y)$, $\text{SR}(m, y)$ is the SR intensity in month m and year y , and $\overline{\text{SR}(m)}$ is the multi-year average of $\text{SR}(m, y)$. Hereafter, this updated parameterisation is referred to as the SR method.

In addition, we also update the value of P_{NO} to be 330 mol NO per flash all over the world (Luhar et al., 2021), consistent with the global average derived from satellite NO_2 observations (Miyazaki et al., 2014; Marais et al., 2018). By comparison, the default P_{NO} value in GEOS-Chem is 500 mol NO per flash north of 35°N and 260 mol for the rest of the world (Murray et al., 2012). Such a large enhancement of P_{NO} in the northern midlatitudes relative to the rest of the world is not supported by a recent analysis of observed NO_2 vertical profiles (Horner et al., 2024). NO_2 satellite observations also indicate little difference in P_{NO} between the midlatitudes and the tropics (Allen et al., 2019; Bucsela et al., 2019).

2.3 Atmospheric chemistry simulations

We use GEOS-Chem v14.0.2 in a global simulation of atmospheric chemistry between 2013 and 2021. The model includes a detailed mechanism for O_3 – NO_x – CO – VOC –aerosol chemistry with recent updates in halogen chemistry, isoprene chemistry, and aerosol and cloud NO_x uptake (Wang et al., 2021; Bates and Jacob, 2019; Holmes et al., 2019). The simulation, conducted at a $2^\circ \times 2.5^\circ$ horizontal resolution, is driven by reanalysed meteorology from the Modern-Era Retrospective analysis for Research and Applications, version 2 (MERRA-2) (Gelaro et al., 2017).

The simulation uses anthropogenic emissions compiled from multiple global and regional inventories. This includes global emissions from the CEDS emission inventory (Hoesly et al., 2018), which are superseded by the National Emissions Inventory 2011 (NEI-2011) in the US (US EPA, 2013), the Air Pollutant Emission Inventory (APEI) in Canada, MIX-Asia v1.1 in Asia (Li et al., 2017), and DICE-Africa in Africa (Marais and Wiedinmyer, 2016). Aircraft emissions for 2019, as provided by the Aircraft Emissions Inventory Code (AEIC) (Simone et al., 2013), are adjusted annually based on the Global Aviation emissions Inventory (GAIA) (Teoh et al., 2023, 2024) to account for inter-annual changes (Lee et al., 2021).

The simulation also incorporates a variety of natural emissions, including fire emissions from the Global Fire Emissions Database (GFED4; van der Werf et al., 2017), biogenic emissions from the Model of Emissions of Gases and Aerosols from Nature version 2.1 (MEGAN2.1; Guenther et al., 2012), soil NO_x emissions from the Berkeley–

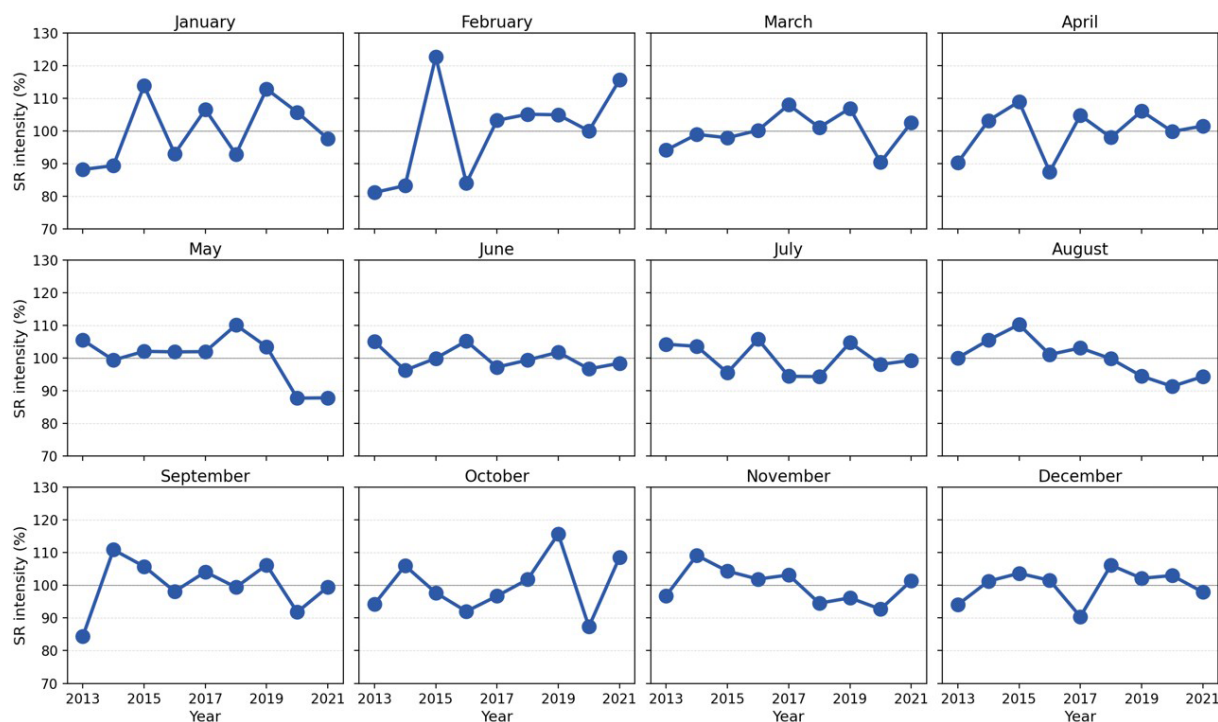


Figure 1. The inter-annual variation in global lightning activity based on SR observations in different months during 2013–2021.

Dalhousie Soil NO_x Parameterization (Hudman et al., 2012), and LNO_x emissions from either OP or SR methods.

We compare atmospheric chemistry simulations driven by OP- or SR-based LNO_x emissions, with a focus on the LTTs and IAV in NO_x , O_3 , and OH. To isolate the lightning contribution to the LTTs and IAV, we also perform a simulation driven by the multi-year (2013–2021) average LNO_x emissions and analyse the difference from simulations with annually varying LNO_x emissions. The simulation with 2013–2021 average LNO_x emissions also provides information on the contribution of non-lightning factors (e.g., fires, biogenic emissions, and meteorology) to the LTTs and IAV in NO_2 , O_3 , and OH, allowing for the decomposition of the LTTs and IAV into contributions of lightning and non-lightning factors.

3 Results and discussion

3.1 LTTs and IAV of global lightning activity and LNO_x emissions

Figure 2 presents global lightning activity and LNO_x emissions as determined by the SR method, which are compared with results from the OP method and lightning observations from the ground WLLN and satellite LIS instruments. The SR method indicates an insignificant LTT in global lightning activity and LNO_x emissions over the study period, in contrast to the OP method, which predicts an increase by $6.4\% \text{ dec}^{-1}$ ($P < 0.05$) from 2013–2021 (Fig. 2a). As a comparison, the WLLN also observes insignificant LTTs,

although the SR method and WLLN observations do not always agree on annual anomalies. Indeed, the OP method in GEOS-Chem predicts significant increases in global LNO_x emissions over an even longer period (1990–2022; Fig. 3). Although our SR data are unavailable for an earlier period, this result of the OP method is again inconsistent with TRMM LIS observations, which show no trend for 1998–2014 in the tropics (Fig. 3). These results highlight the significant uncertainties in the unconstrained OP method in estimating the LTTs of global lightning activity, demonstrating the value of applying the SR method to long-term atmospheric chemistry simulations.

We quantify the magnitude of IAV in LNO_x emissions by computing the standard deviation of detrended annual anomalies. The results show that the IAV derived from the SR method (0.23 Tg N a^{-1} or 3.3 %) is twice that derived from the OP method (0.13 Tg N a^{-1} or 1.8 %) (Fig. 2a), indicating that the IAV in LNO_x emissions is underestimated by the OP method. The magnitude of IAV in global lightning activity observed by the WLLN (0.25 Tg N a^{-1} or 3.6 %) is close to that observed by the SR observations (Fig. 2a). Consistent with our result, a previous study by Murray et al. (2013) also found that the cloud-top height parameterisation underestimates the IAV in tropical LNO_x emissions by a factor of approximately 2 compared to TRMM LIS satellite observations.

Seasonally, the SR method finds that the magnitudes of IAVs in LNO_x emissions are $0.028 \text{ Tg N month}^{-1}$ in DJF

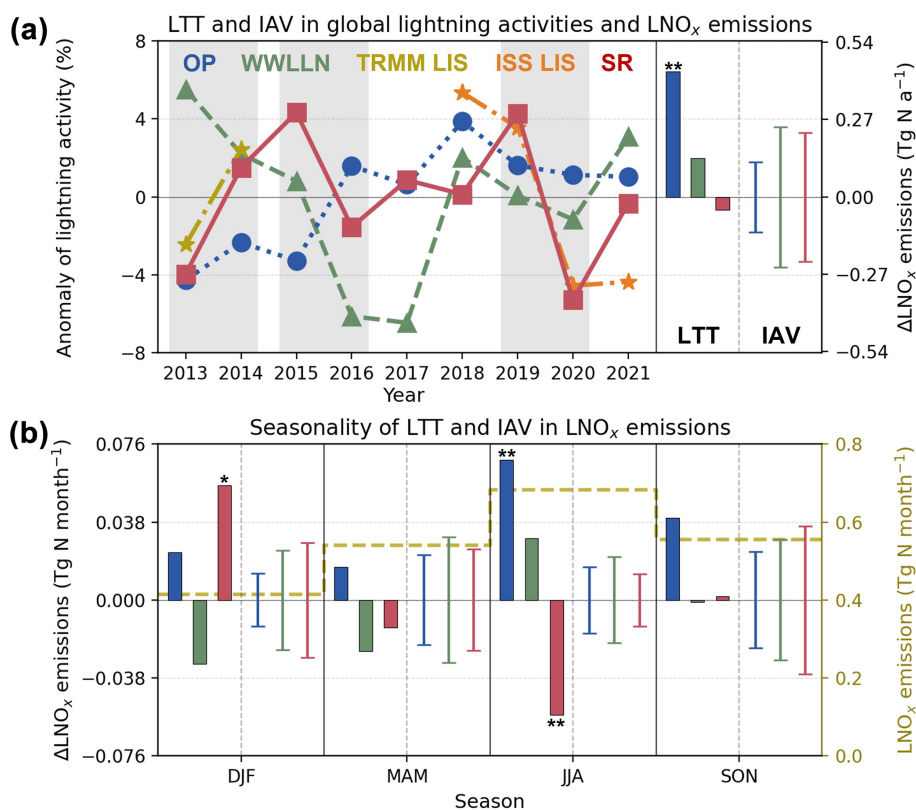


Figure 2. (a) Annual anomalies in global lightning activities and LNO_x emissions during 2013–2021 based on the OP method and observations from SR, WWLLN, the TRMM LIS, and the ISS LIS. Grey shading indicates years with large anomalies (> 5%) in global lightning activities and LNO_x emissions based on SR observations. LTTs and IAV of the annual anomaly time series are shown to the right. (b) Seasonal LTTs and IAV in global LNO_x emissions based on the OP method and observations from SR and WWLLN during 2013–2021. The dashed yellow lines show global mean LNO_x emissions in different seasons. LTT values are in units of Tg N a⁻¹ dec⁻¹ in (a) and Tg N month⁻¹ dec⁻¹ in (b). ** indicates significant trends ($P < 0.05$) and * moderately significant trends ($0.05 \leq P < 0.1$). IAV values represent the standard deviation of annual anomalies after the LTT is removed.

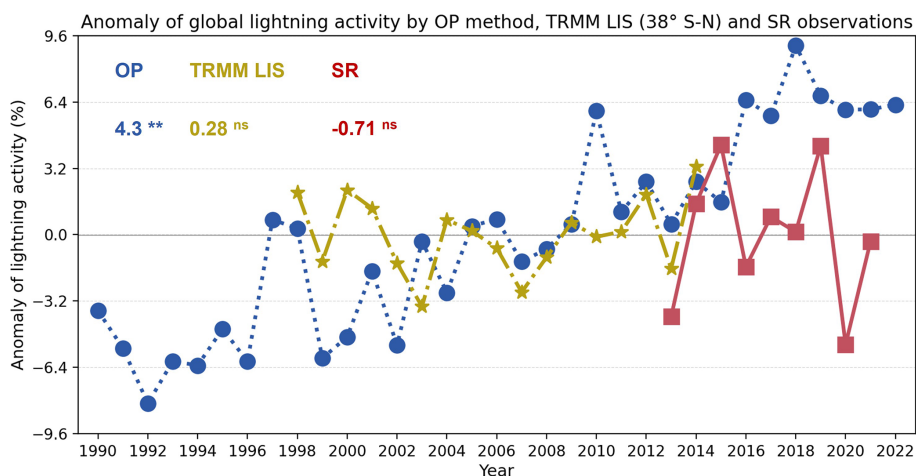


Figure 3. Anomaly of global lightning activity based on the OP method, TRMM LIS (38° S–38° N), and SR observations. The linear trends (% dec⁻¹) are shown in the upper-left corner. ** denotes significant trends ($P < 0.05$), * moderately significant trends ($0.05 \leq P < 0.1$), and ns insignificant trends.

(austral summer) and $0.013 \text{ Tg N month}^{-1}$ in JJA (boreal summer) (Fig. 2b), which translates to relative IAVs of 6.7 % and 1.8 % for the two seasons, respectively (average LNO_x emissions are 0.42 and $0.73 \text{ Tg N month}^{-1}$ in DJF and JJA, respectively) (Fig. 2b). Because of the dominant occurrence of lightning in the summer hemisphere, this result based on the SR method indicates a factor of 3.7 greater relative IAVs in LNO_x emissions over the Southern Hemisphere than over the Northern Hemisphere. In contrast, the OP method predicts similar levels of relative IAVs in both hemispheres (Fig. 2b).

In terms of year-to-year anomalies, the SR method shows large increases in global lightning activity and LNO_x emissions during 2013–2015, 2018–2019, and 2020–2021 and decreases in 2015–2016 and 2019–2020 (Fig. 2a). These large year-to-year anomalies are not captured by the OP method (Fig. 2a). Although the SR method and WWLLN observations show consistent increases for 2020–2021 and decreases for 2015–2016 and 2019–2020, they disagree on year-to-year changes during 2013–2015, 2017–2018, and 2018–2019, probably because of variable global detection efficiency by the ground WWLLN (Hutchins et al., 2012). The SR results agree well with satellite-based LIS observations for the increase during 2013–2014 and the decrease during 2019–2020, while satellite-based observations are unavailable for 2015–2017. Noteworthily, SR and ISS LIS observations conformably show that global lightning activity decreased by $\sim 10\%$ from 2019–2020 (Fig. 2a). Such a large inter-annual deviation is unprecedented in the history of LIS observation records (Fig. 3), which is in accordance with previous finds by Dunn et al. (2022; see Fig. SB2.1b).

3.2 Lightning-driven LTTs of global atmospheric chemistry

Figure 4 shows the contribution of LNO_x emissions, as determined by the OP or SR methods, to global LTTs and IAV in key atmospheric chemicals (NO_x , O_3 , and OH) based on GEOS-Chem simulations. The contribution of lightning is further compared to that of other influencing factors. The stable LNO_x emissions predicted by the SR method result in insignificant lightning-induced LTTs in NO_x , O_3 , and OH during 2013–2021, both globally and in two hemispheres (Figs. 4a–c, S1a–c, and S2a–c in the Supplement). In contrast, the OP method, with increases in LNO_x emissions, leads to substantial lightning-induced LTTs in these species. These trends are statistically significant for NO_x ($2.1\% \text{ dec}^{-1}$), O_3 ($1.0\% \text{ dec}^{-1}$), and OH ($1.4\% \text{ dec}^{-1}$) in the Northern Hemisphere and for NO_x ($1.9\% \text{ dec}^{-1}$) globally (Figs. S1a–c and 4a).

When all factors are accounted for, the GEOS-Chem model based on the SR method simulates a strong decrease in global ($-5.4\% \text{ dec}^{-1}$) and northern hemispheric ($-10\% \text{ dec}^{-1}$) NO_x concentrations during 2013–2021 (Figs. 4a and S1a), driven primarily by a reduction

in global anthropogenic emissions, especially from China (Wang et al., 2022). However, if the simulation is based on the OP method, these negative trends are reduced to $-3.8\% \text{ dec}^{-1}$ globally and $-8.3\% \text{ dec}^{-1}$ in the Northern Hemisphere (Figs. 4a and S1a). The sharper decrease predicted by the SR simulation is in better agreement with the trends in tropospheric NO_2 columns from satellite observations during the period (Fig. 5a and b), though the simulation still does not fully capture the inter-annual anomalies, possibly due to errors in non-lightning factors. This difference between the OP- and SR-based simulations highlights the critical role of accurately accounting for lightning in better interpreting long-term trends in global background NO_x levels and estimating the expected effects of emission reduction policies for improving air quality.

The GEOS-Chem simulation using SR LNO_x emissions estimates $\sim 2.5\% \text{ dec}^{-1}$ increases for global and hemispheric tropospheric O_3 (Figs. 4b, S1b, and S2b). These trends are $\sim 30\%$ greater when the simulation is based on the OP method. This result suggests that non-lightning factors are major drivers of the simulated increases in tropospheric O_3 , while lightning is a non-negligible contributor. Previous studies have reported that CTMs tend to underestimate the observed long-term increase in tropospheric O_3 (Christiansen et al., 2022; Wang et al., 2022) over a longer time frame (since the 1990s) than the one in our current study. Their results are based on LNO_x emissions by the OP method, which predicts accelerated increases in global lightning activity in the last 3 decades ($4.3\% \text{ dec}^{-1}$ increase during 1991–2000, $6.9\% \text{ dec}^{-1}$ during 2001–2010, and $7.9\% \text{ dec}^{-1}$ during 2011–2020), inconsistent with the SR intensity in our study and satellite observations for the earlier period (Fig. 3). This result implies that after adjusting the LTTs in LNO_x emissions with SR observations, the model underestimation of O_3 trends should be even more pronounced, underscoring the gap in our understanding of tropospheric O_3 trends.

3.3 Lightning-driven IAV in atmospheric oxidants and implications for the global methane budget

Figures 4b and c show that the simulation using SR LNO_x emissions exhibits greater magnitudes of lightning-induced IAVs in global average O_3 and OH concentrations compared to the simulation using OP LNO_x emissions. This is due to the larger IAV in LNO_x emissions predicted by the SR method (Fig. 2a). However, when all factors are accounted for, the IAVs in global O_3 and OH are more than 20 % smaller in the SR simulation compared to the OP simulation. The smaller all-factor IAVs in the SR simulation arise from a negative correlation between the lightning and non-lightning contributions ($r \approx -0.4$), indicating that lightning serves as a balancing factor that reduces the IAVs of O_3 and OH globally. Conversely, in the OP simulation, the lightning contributions positively correlate with the non-lightning contribu-

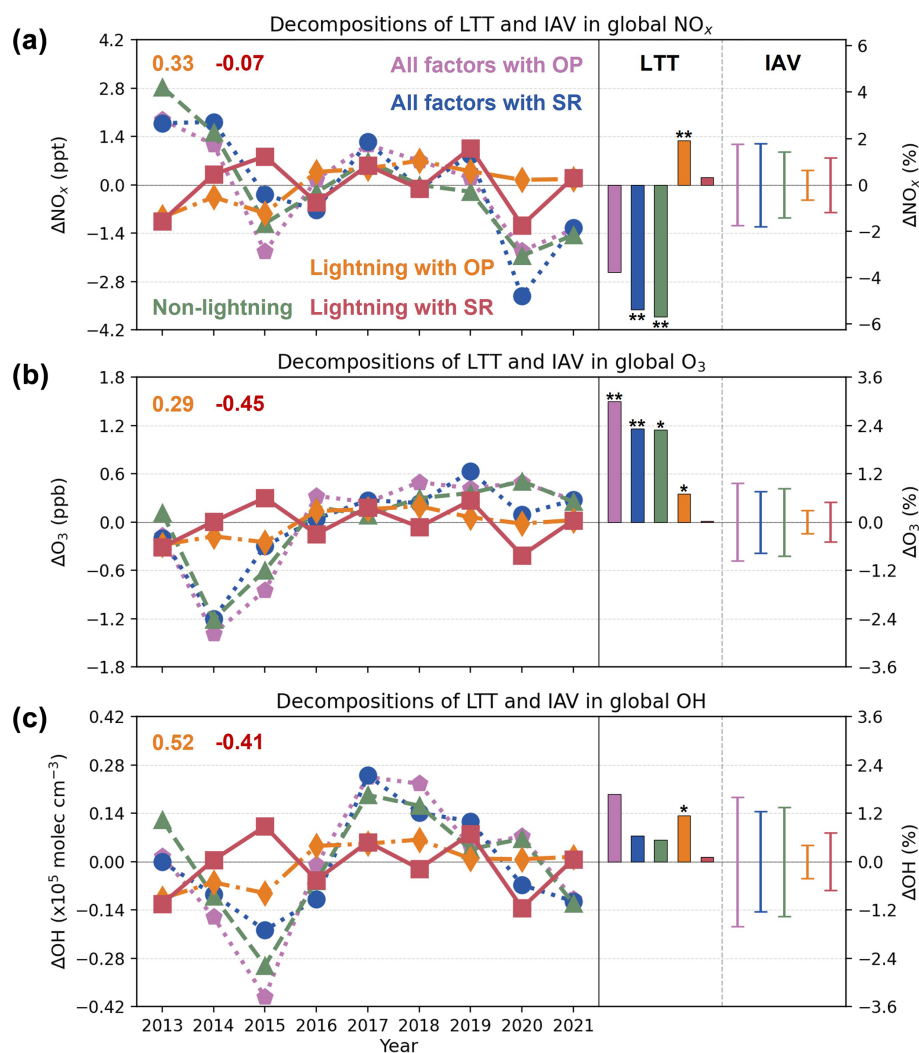


Figure 4. Annual anomalies, LTTs, and IAV of global NO_x (a), O_3 (b), and OH (c) and their decompositions into lightning and non-lightning factors during 2013–2021. LTT values are reported in units of $\% \text{ dec}^{-1}$. ** denotes significant trends ($P < 0.05$) and * moderately significant trends ($0.05 \leq P < 0.1$). IAV values represent the standard deviation of annual anomalies after LTTs are removed. Shown in the upper-left corner of each panel are correlation coefficients between the annual anomalies (with LTTs removed) of lightning (OP: orange; SR: red) and non-lightning contributions.

tions ($r = 0.3$ for O_3 and 0.5 for OH), thereby amplifying the IAVs of O_3 and OH (Fig. 4b and c).

The year-to-year variations in global OH concentrations can impact the annual growth rates of atmospheric methane, as the OH radical is the main sink of methane in the atmosphere. NOAA observations indicate high annual methane growth rates in 2014, 2015, 2020, and 2021 (Fig. 6), although the underlying causes are not yet fully understood (Feng et al., 2023; Stevenson et al., 2022; Yin et al., 2021; Zhang et al., 2021, 2023). Figure 6 quantifies the contribution of lightning-induced OH variations to annual methane growth rates from 2013–2021, based on the simulation using SR LNO_x emissions. The influence of other factors, including direct methane emissions and OH variations due to non-

lightning drivers, is deduced from the difference between observed growth rates and lightning contributions.

Based on the simulation using SR emissions, lightning-driven OH variations account for $0.7\text{--}3.0 \text{ ppb a}^{-1}$ year-to-year anomalies in methane growth rates, with positive contributions in 2015–2016, 2017–2018, and 2019–2020 and negative contributions in other years (Fig. 6). In most cases, the influence of lightning counteracts that of other factors, suppressing changes in methane growth rates, as seen in 2013–2014, 2015–2016, and 2020–2021. However, there are also cases where lightning works in conjunction with other factors. Notably, in 2019–2020, both lightning and other factors contribute to an accelerated methane growth. As mentioned in Sect. 3.1, global lightning declined by $\sim 10\%$

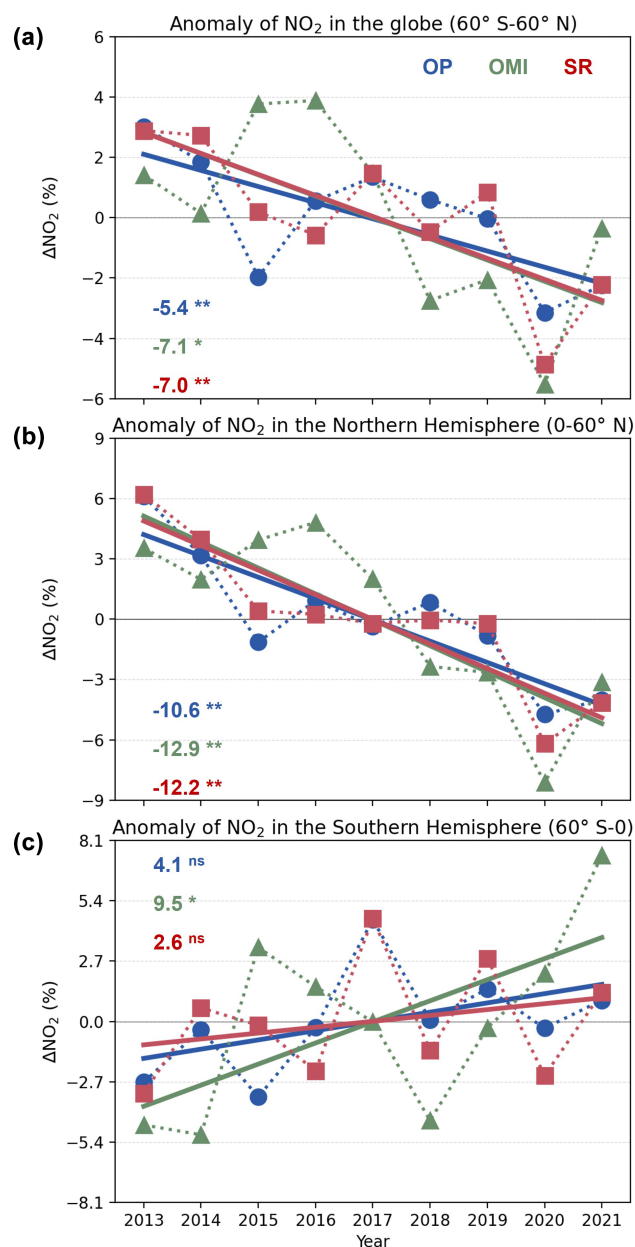


Figure 5. The anomaly of tropospheric NO₂ vertical column densities averaged over the globe (a), the Northern Hemisphere (b), and the Southern Hemisphere (c) during 2013–2021 for OP and SR simulations and OMI satellite observations. The linear trends (% dec⁻¹) are inset. ** denotes significant trends ($P < 0.05$), * moderately significant trends ($0.05 \leq P < 0.1$), and ^{ns} insignificant trends. Following Qu et al. (2021), we filter the data using quality flags reported in the OMI data product. We exclude data affected by the row anomalies and retain those with a cloud fraction < 0.2 , surface albedo < 0.3 , solar zenith angle $< 75^\circ$, and view zenith angle $< 65^\circ$.

from 2019–2020. Our analysis indicates that the reduction in global lightning activity can generate a $\sim 2\%$ decrease in OH sinks and explain $\sim 54\%$ of the 2020 surge in atmospheric methane. Current theories have not accounted for the impact of abnormal lightning activity on the 2020 methane spike (Feng et al., 2023; Peng et al., 2022; Stevenson et al., 2022; Chen et al., 2025).

3.4 Comparison of year-to-year anomalies to regional observations

The integration of SR observations into the OP parameterisation results in differences of several percent in the annual average global background concentrations of NO_x, O₃, and OH. While these differences between the SR and OP methods can have substantial implications for the global budgets of these species, systematic assessment using atmospheric chemical observations is challenging due to the impact of other factors across multiple scales. Here, we present two cases characterised by strong lightning-driven year-to-year anomalies, allowing us to compare simulations driven by OP and SR parameterisations to regional atmospheric observations.

In the first case, we analyse the change in high-altitude NO₂ concentrations over South America between September–November of 2019 and 2020, comparing the changes simulated with OP and SR LNO_x emissions with those observed by the Tropospheric Monitoring Instrument (TROPOMI). The time frame is chosen due to the substantial decrease in global lightning activity detected by SR observations (Fig. S3a in the Supplement). South America is chosen because the continent is a lightning hotspot (Fig. S3b) and, unlike other continental hotspots, experiences lower interference by confounding factors such as fire (e.g., Africa and Indonesia) and anthropogenic (e.g., North America) emissions (see Supplement for detailed discussion).

Using the cloud-slicing technique (Marais et al., 2021; Marais and Roberts, 2020) on TROPOMI observations (version 01-03-02, accessible through the Sentinel-5P Pre-Operations Data Hub), we retrieved mean NO₂ concentrations in the upper troposphere (300–100 hPa), where the impact of lightning is greatest (Fig. S3c), at a $2^\circ \times 2.5^\circ$ resolution for September–November of 2019 and 2020 (Fig. S4 in the Supplement). The 3-month cloud-sliced satellite observations over South America are averaged and compared to GEOS-Chem simulations. The analysis reveals a change in NO₂ concentrations of -3 (-8 to 3) ppb in the upper troposphere over South America, as observed by TROPOMI, which aligns closely with the simulation driven by SR LNO_x emissions (-5.5 (-11 to 0) ppb) (Fig. 7a). In contrast, the simulation driven by OP LNO_x emissions results in an increase in NO₂ concentrations (9 (4 – 14) ppb) (uncertainties shown in the brackets are estimated with bootstrapping; the same applies hereafter).

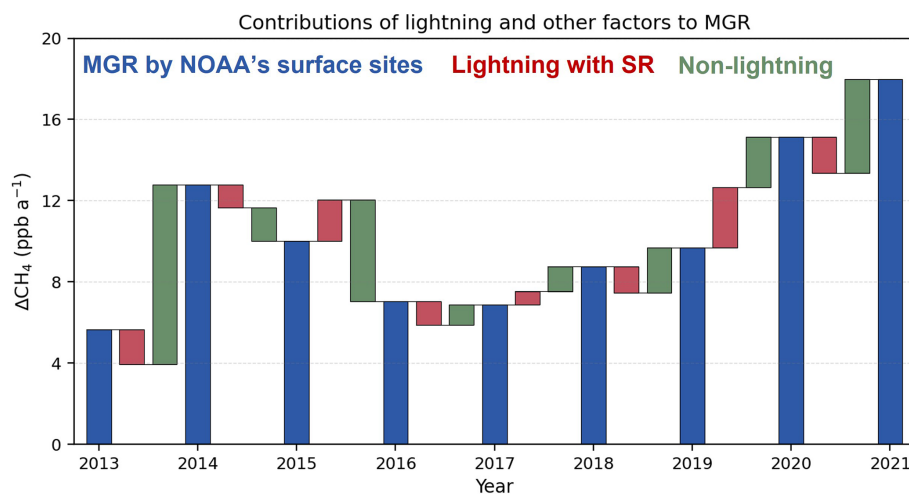


Figure 6. Contributions of lightning-driven variations in the methane sink by OH oxidation to the annual atmospheric methane growth rate (MGR) from 2013–2021 based on SR observations. The gap between the observed year-to-year anomalies in MGR and the simulated lightning contribution is attributed to non-lightning factors.

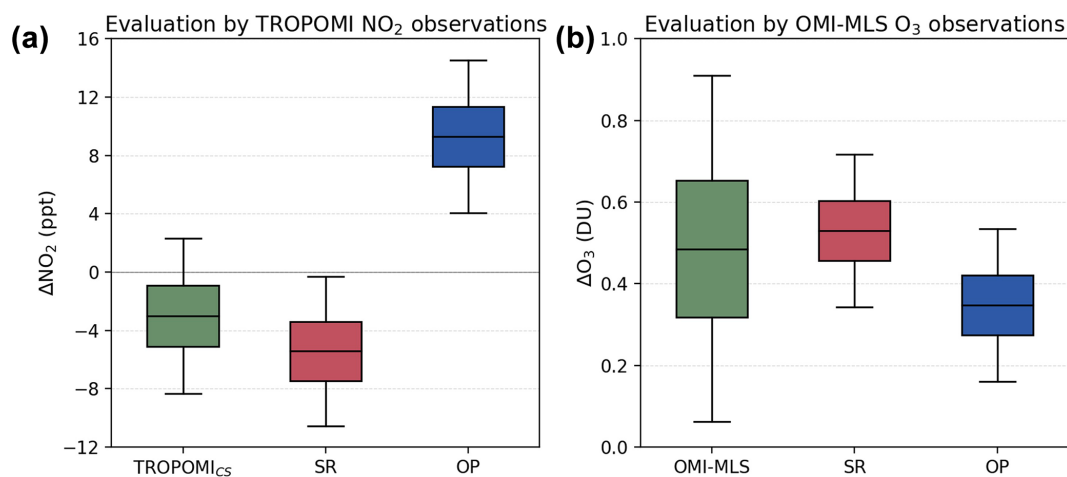


Figure 7. (a) Anomalies of NO_2 concentrations in the upper troposphere (300–100 hPa) of northern South America (20°S – 10°N and 80 – 40°W) in September–November 2020 relative to 2019 from the TROPOMI NO_2 observations by the cloud-slicing technique (green), the GEOS-Chem simulation with the SR method (red), and the GEOS-Chem simulation with the OP method (blue). (b) Anomalies of tropospheric O_3 in the North Atlantic (0 – 60°N and 100°W – 30°E) in 2015 relative to 2014 from the OMI-MLS O_3 observations (green), the GEOS-Chem simulation with the SR method (red), and the GEOS-Chem simulation with the OP method (blue). Uncertainties are calculated by 1 000 000 Monte Carlo simulations.

In the second case, we examine changes in tropospheric O_3 concentrations over the North Atlantic Ocean in 2014–2015. Russo et al. (2023) highlighted that annual anomalies in tropospheric O_3 over this region are closely linked to lightning activity, but this variability is not well captured by atmospheric chemical models. One such instance is the substantial increase in tropospheric ozone by 0.49 (0.1–0.9) DU measured by the OMI-MLS satellite product between 2014 and 2015, shown in Fig. S5 in the Supplement (Russo et al., 2023). In comparison, our GEOS-Chem simulation with OP LNO_x emissions underestimates this increase (0.35 (0.2–0.5) DU), while the simulation with SR LNO_x emis-

sions agrees better with observed ozone changes (0.53 (0.3–0.7) DU) (Fig. 7b). However, we note that the difference between the OP and SR results is not significant if the uncertainties are accounted for (Fig. 7b), indicating that this case represents weak evidence supporting the SR method.

4 Conclusions

In this study, we updated the parameterisation of LNO_x emissions in the GEOS-Chem model by incorporating SR observations, which offer strong global sensitivity to lightning.

This update aimed to better constrain the LTTs and IAV in global lightning activity and hence improve the simulation of global atmospheric chemistry.

The updated global LNO_x emissions showed insignificant LTTs from 2013–2021, while the original results predicted a significant increase by 6.4 % dec^{−1} ($P < 0.05$). As a comparison, the WWLLN also observed no significant trend. Indeed, the original method in GEOS-Chem indicated significant increases in global LNO_x emissions over an even longer period (1990–2022). Although our SR data were unavailable for an earlier period, this result of the original method was again inconsistent with TRMM LIS observations, which showed no trend for 1998–2014 in the tropics ($\pm 38^\circ$ latitude). Meanwhile, the updated LNO_x emissions had an IAV twice as high as that of the original result, which was close to that by WWLLN observations. These results highlighted the significant uncertainties in the unconstrained method in estimating the LTTs and IAV of global lightning, demonstrating the value of applying the SR method to long-term atmospheric chemistry simulations.

Compared to the original simulation, the simulation with updated LNO_x emissions found smaller contributions of lightning to the LTTs in NO_x, O₃, and OH. This led to sharper decreases in simulated global and northern hemispheric NO_x concentrations during 2013–2021, in better agreement with the trends in tropospheric NO₂ columns from satellite observations. Previous research has reported that CTMs tended to underestimate the observed long-term increase in tropospheric O₃ (Christiansen et al., 2022; Wang et al., 2022). Our study showed that the increase in tropospheric O₃ from the original simulation was $\sim 30\%$ greater than in the updated results, implying that the updated simulation further underestimated tropospheric O₃ trends and highlighting the gap in our understanding of tropospheric O₃.

Moreover, lightning-induced anomalies in O₃ and OH were negatively correlated with those caused by non-lightning factors (e.g., wildfires, biogenic emissions, and meteorology). This suggests that lightning serves as a counterbalancing factor, reducing the IAV of O₃ and OH, an opposite feature shown in the original simulation. Our analysis showed that the change in global OH concentration due to IAV in LNO_x emissions was a substantial contributor to the inter-annual variations in methane growth rates. Notably, SR and ISS LIS observations consistently found that 2020 global lightning fell by $\sim 10\%$ relative to 2019, in line with Dunn et al. (2022), which might be the biggest year-to-year change in nearly 3 decades. It resulted in a $\sim 2\%$ drop in OH levels, representing $\sim 54\%$ of the growth of atmospheric methane in this year. Therefore, our study also suggests that the variability in lightning activity and LNO_x emissions needs to be taken into account when making the global atmospheric methane budget.

Code and data availability. We used lightning observation data, including ISS LIS data (<https://doi.org/10.5067/LIS/ISS/LIS/DATA109>, Lang, 2022), TRMM LIS data (<https://doi.org/10.5067/LIS/LIS/DATA201>, Blakeslee, 1998), WWLLN data (v2022.0.0) (<https://doi.org/10.5281/zenodo.6007052>, Kaplan and Lau, 2021, 2022), and Schumann-resonance-derived monthly lightning variation data (shown in Fig. 1). The source code of GEOS-Chem (v14.0.2) is freely available through <https://doi.org/10.5281/zenodo.7383492> (The International GEOS-Chem User Community, 2022). The tropospheric NO₂ observation data are available at <https://doi.org/10.5067/Aura/OMI/DATA2017> (Krotkov et al., 2019). The MGR data from NOAA's surface sites can be downloaded from <https://doi.org/10.15138/P8XG-AA10> (Lan et al., 2022). We used the cloud-slicing algorithm introduced by Marais et al. (2021) and the corresponding Python code (<https://doi.org/10.5281/zenodo.3979211>, Marais and Roberts, 2020a) and subsequent improvements in the code (<https://doi.org/10.5281/zenodo.4058442>, Marais and Roberts, 2020b). The OMI-MLS observation data can be downloaded from https://acd-ext.gsfc.nasa.gov/Data_services/cloud_slice/ (Ziemke et al., 2006).

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Author contributions. YuzZ and XW conceptualised the study. XW led the formal analysis and simulated GEOS-Chem under the supervision of YuzZ. XW, YuzZ, TB, RL, XX, SZ, RW, YujZ, and SS were responsible for the methodology. XW, YuzZ, and TB provided resources. XW visualised the study. XW wrote the original draft, and YuzZ and TB reviewed and edited the paper.

Competing interests. The contact author has declared that none of the authors has any competing interests.

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