

Estimates of lightning NO_x production based on high-resolution OMI NO₂ retrievals over the continental US

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Abstract. Lightning serves as the dominant source of nitrogen oxides $(NO_x = NO + NO_2)$ in the upper troposphere (UT), with a strong impact on ozone chemistry and the hydroxyl radical production. However, the production efficiency (PE) of lightning nitrogen oxides (LNO_x) is still quite uncertain (32-1100 mol NO per flash). Satellite measurements are a powerful tool to estimate LNO_x directly compared to conventional platforms. To apply satellite data in both clean and polluted regions, a new algorithm for calculating LNO_x has been developed that uses the Berkeley High-Resolution (BEHR) v3.0B NO₂ retrieval algorithm and the Weather Research and Forecasting model coupled with chemistry (WRF-Chem). LNO_x PE over the continental US is estimated using the NO2 product of the Ozone Monitoring Instrument (OMI) data and the Earth Networks Total Lightning Network (ENTLN) data. Focusing on the summer season during 2014, we find that the lightning NO₂ (LNO₂) PE is 32 ± 15 mol NO₂ per flash and 6 ± 3 mol NO₂ per stroke while LNO_x PE is $90 \pm 50 \text{ mol NO}_x$ per flash and $17 \pm 10 \text{ mol NO}_{x}$ per stroke. Results reveal that our method reduces the sensitivity to the background NO₂ and includes much of the below-cloud LNO₂. As the LNO_x parameterization varies in studies, the sensitivity of our calculations to the setting of the amount of lightning NO (LNO) is evaluated. Careful consideration of the ratio of LNO₂ to NO₂ is also needed, given its large influence on the estimation of LNO_2 PE.

1 Introduction

Nitrogen oxides (NO_x) near the Earth's surface are mainly produced by soil, biomass burning, and fossil fuel combustion, while NO_x in the middle and upper troposphere originates largely from lightning and aircraft emissions. NO_x plays an important role in the production of ozone (O_3) and the hydroxyl radical (OH). While the anthropogenic sources of NO_x are largely known, lightning nitrogen oxides (LNO_x) are still the source with the greatest uncertainty, though they are estimated to range between 2 and $8 \, \mathrm{Tg} \, \mathrm{N} \, \mathrm{yr}^{-1}$ (Schumann and Huntrieser, 2007). LNO_x is produced in the upper troposphere (UT) by O₂ and N₂ dissociation in the hot lightning channel as described by the Zel'dovich mechanism (Zel'dovich and Raizer, 1967). With the recent updates of UT NO_x chemistry, the daytime lifetime of UT NO_x is evaluated to be \sim 3 h near thunderstorms and \sim 0.5–1.5 d away from thunderstorms (Nault et al., 2016, 2017). This results in enhanced O₃ production in the cloud outflow of active convection (Pickering et al., 1996; Hauglustaine et al., 2001; DeCaria et al., 2005; Ott et al., 2007; Dobber et al., 2008; Allen et al., 2010; Finney et al., 2016). As O₃ is known as a greenhouse gas, strong oxidant, and absorber of ultraviolet radiation (Myhre et al., 2013), the contributions of LNO_x to O₃ production also have an effect on climate forcing. Finney et al. (2018) found different impacts on atmospheric composition and radiative forcing when simulating future lightning using a new upward cloud ice flux (IFLUX) method versus the commonly used cloud-top height (CTH) approach. While global lightning is predicted to increase by 5%-16% over the next century with the CTH approach (Clark et al., 2017; Banerjee et al., 2014; Krause et al., 2014), a 15 % decrease in global lightning was estimated with IFLUX in 2100 under a strong global warming scenario (Finney et al., 2018). As a result of the different effects on radiative forcing from ozone and methane, a net positive radiative forcing was found with the CTH approach while there is little net radiative forcing with the IFLUX approach (Finney et al., 2018). However, the convective available potential energy (CAPE) times the precipitation rate (P) proxy predicts a $12 \pm 5\%$ increase in the continental US (CONUS) lightning strike rate per kelvin of global warming (Romps et al., 2014), while the IFLUX proxy predicts the lightning will only increase $3.4 \% \text{ K}^{-1}$ over the CONUS. Recently, Romps (2019) compared the CAPE $\times P$ proxy and IFLUX method in cloud-resolving models. They reported that higher CAPE and updraft velocities caused by global warming could lead to the large increases in tropical lightning simulated by the CAPE $\times P$ proxy, while the IFLUX proxy predicts little change in tropical lightning because of the small changes in the mass flux of ice.

In view of the regionally dependent lifetime of NO_x and the difficulty of measuring LNO_x directly, a better understanding of the LNO_x production is required, especially in the tropical and midlatitude regions in summer. Using its distinct spectral absorption lines in the near-ultraviolet (UV) and visible (VIS) ranges (Platt and Perner, 1983), NO2 can be measured by satellite instruments like the Global Ozone Monitoring Experiment (GOME; Burrows et al., 1999; Richter et al., 2005), SCanning Imaging Absorption SpectroMeter for Atmospheric CHartographY (SCIA-MACHY; Bovensmann et al., 1999), the Second Global Ozone Monitoring Experiment (GOME-2; Callies et al., 2000), and the Ozone Monitoring Instrument (OMI; Levelt et al., 2006). OMI has the highest spatial resolution, least instrument degradation, and longest record among these satellites (Krotkov et al., 2017). Satellite measurements of NO₂ are a powerful tool compared to conventional platforms because of their global coverage, constant instrument features, and temporal continuity.

Recent studies have determined and quantified LNO_x using satellite observations. Beirle et al. (2004) constrained the LNO_x production to 2.8 (0.8–14) Tg N yr⁻¹ by combining GOME NO₂ data and flash counts from the Lightning Imaging Sensor (LIS) aboard the Tropical Rainfall Measurement Mission (TRMM) over Australia. Boersma et al. (2005) estimated the global LNO_x production of 1.1-6.4 Tg N yr⁻¹ by comparing GOME NO₂ with distributions of LNO_2 mod-

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eled by Tracer Model 3 (TM3). Martin et al. (2007) analyzed SCIAMACHY NO₂ columns with Goddard Earth Observing System chemistry model (GEOS-Chem) simulations to identify LNO_x production amounting to $6 \pm 2 \text{ Tg N yr}^{-1}$.

As these methods focus on monthly or annual mean NO₂ column densities, more recent studies applied specific approaches to investigate LNO_x directly over active convection. Beirle et al. (2006) estimated LNO_x as 1.7 (0.6– 4.7) Tg N yr⁻¹ based on a convective system over the Gulf of Mexico, using National Lightning Detection Network (NLDN) observations and GOME NO2 column densities. However, it is assumed that all the enhanced NO₂ originated from lightning and did not consider the contribution of anthropogenic emissions. Beirle et al. (2010) analyzed LNO_x production systematically using the global dataset of SCIA-MACHY NO2 observations combined with flash data from the World Wide Lightning Location Network (WWLLN). Their analysis was restricted to $30 \text{ km} \times 60 \text{ km}$ satellite pixels where the flash rate exceeded 1 flash km⁻² h⁻¹. But they found LNO_x production to be highly variable, and correlations between flash-rate densities and LNO_x production were low in some cases. Bucsela et al. (2010) estimated LNO_x production as $\sim 100-250 \text{ mol NO}_x$ per flash for four cases, using the DC-8 and OMI data during NASA's Tropical Composition, Cloud and Climate Coupling Experiment (TC4).

Based on the approach used by Bucsela et al. (2010), a special algorithm was developed by Pickering et al. (2016) to retrieve LNO_x from OMI and the WWLLN. The algorithm takes the OMI tropospheric slant column density (SCD) of NO₂ (S_{NO_2}) as the tropospheric slant column density of LNO₂ (S_{LNO₂}) by using cloud radiance fraction (CRF) greater than 0.9 to minimize or screen the lower tropospheric background. To convert the SLNO2 to the tropospheric vertical column density (VCD) of LNO_x (V_{LNO_x}), an air mass factor (AMF) is calculated by dividing the a priori S_{LNO_2} by the a priori $V_{LNO_{x}}$. The a priori $S_{LNO_{2}}$ is calculated using a radiative transfer model and a profile of LNO₂ simulated by the NASA Global Modeling Initiative (GMI) chemical transport model. The a priori V_{LNO_r} is also obtained from the GMI model. Results for the Gulf of Mexico during 2007-2011 summer yield LNO_x production of 80 ± 45 mol NO_x per flash. Since they considered NO₂ above the cloud to be LNO₂ in the algorithm due to the difficulty and uncertainty in determining the background NO₂, their AMF and derived VCD of LNO_x (LNO₂) are named AMF_{LNO_xClean} (AMF_{LNO₂Clean}) and LNO_xClean (LNO₂Clean), respectively. Note that Pickering et al. (2016) considered the two estimates of background derived from aircraft flights in the Gulf of Mexico region (3% and 33%) and subtracted the mean value (18%)from the estimated mean LNO_x production efficiency (PE) for the background bias. However, we use the original algorithm directly without correction to distinguish the effect of different AMFs on LNO_x estimation in the remainder of this paper. Unless otherwise specified, abbreviations S and V are

respectively defined as the tropospheric SCD and VCD in this paper.

More recently Bucsela et al. (2019) obtained an average PE of $180 \pm 100 \text{ mol NO}_x$ per flash over East Asia, Europe, and North America based on a modification of the method used in Pickering et al. (2016). A power function between LNO_x and lightning flash rate was established, while the minimum flash-rate threshold was not applied. The tropospheric NO_x background was removed by subtracting the temporal average of NO_x at each box where the value was weighted by the number of OMI pixels which meet the optical cloud pressure and CRF criteria required to be considered deep convection but have one flash or fewer instead. The lofted pollution was considered to be 15 % of total NO_x according to the estimation from DeCaria et al. (2000, 2005), and the average chemical delay was adjusted by 15 % following the $3 h LNO_x$ lifetime in the nearby field of convection (Nault et al., 2017). However, there were negative LNO_x values caused by the overestimation of the tropospheric background and stratospheric NO₂ at some locations.

On the other hand, Lapierre et al. (2020) constrained LNO₂ to $1.1 \pm 0.2 \text{ mol NO}_2$ per stroke for intracloud (IC) strokes and $10.7 \pm 2.5 \text{ mol NO}_2$ per stroke for cloud-toground (CG) strokes over the CONUS. LNO₂ per stroke was scaled to $24.2 \mod NO_x$ per flash using mean values of strokes per flash and the ratio of NO_x to NO_2 in the UT. They used the regridded Berkeley High-Resolution (BEHR) v3.0A $0.05^{\circ} \times 0.05^{\circ}$ "visible only" NO₂ VCD (V_{vis}) product which includes two parts of NO2 that can be "seen" by the satellite. The first part is the NO₂ above clouds (pixels with CRF > 0.9) and the second part is the NO₂ detected from cloud-free areas. A threshold of 3×10^{15} molecules cm⁻², the typical urban NO₂ concentration, was applied to mask the contaminated grid cells (Beirle et al., 2010; Laughner and Cohen, 2017). The main difference between Lapierre et al. (2020) and Pickering et al. (2016) is the air mass factor for lightning (AMF_{LNO_x}) implemented in the basic algorithm. In Lapierre et al. (2020), the air mass factor was used to convert S_{NO_2} to V_{vis} , while in Pickering et al. (2016) it was used to convert S_{LNO_2} to V_{LNO_x} , assuming that all S_{NO_2} is generated by lightning.

To apply the approach used by Bucsela et al. (2010), Pickering et al. (2016), Bucsela et al. (2019), and Lapierre et al. (2020) without geographic restrictions, the contamination by anthropogenic emissions must be taken into account in detail. The Weather Research and Forecasting (WRF) model coupled with chemistry (WRF-Chem) has been employed to evaluate the convective transport and chemistry in many studies (Barth et al., 2012; Wong et al., 2013; Fried et al., 2016; Li et al., 2017). Meanwhile, Laughner and Cohen (2017) showed that the OMI AMF is increased by $\sim 35\%$ for summertime when LNO₂ simulated by WRF-Chem is included in the a priori profiles to match aircraft observations. The simulation agrees with observed NO₂ profiles and the bias of AMF related to these observations is reduced to $< \pm 4\%$ for OMI viewing geometries.

In this paper, we focus on the estimation of LNO_2 production per flash (LNO_2 per flash), LNO_2 production per stroke (LNO_2 per flash), LNO_2 production per stroke (LNO_2 per stroke), and LNO_x production per stroke (LNO_x per stroke) in May–August (MJJA) 2014 by developing an algorithm similar to that of Pickering et al. (2016) based on the BEHR NO₂ retrieval algorithm (Laughner et al., 2018, 2019), but it performs better over background NO₂ sources. Section 2 describes the satellite data, lightning data, model settings, and the algorithm in detail. Section 3 explores the suitable data criteria, compares different methods, and evaluates the effect of background NO₂, cloud, and LNO_x parameterization on LNO_x production estimation. Section 4 examines the effect of different sources of the uncertainty on the results. Conclusions are summarized in Sect. 5.

2 Data and methods

2.1 Ozone Monitoring Instrument (OMI)

OMI is carried on the Aura satellite (launched in 2004), a member of the A-train satellite group (Levelt et al., 2006, 2018). OMI passes over the Equator at ~ 13:45 LT (ascending node) and has a swath width of 2600 km, with a nadir field-of-view resolution of $13 \text{ km} \times 24 \text{ km}$. Since the beginning of 2007, some of the measurements have become useless as a result of anomalous radiances called the "row anomaly" (Dobber et al., 2008; KNMI, 2012). For the current study, we used the NASA standard product v3.0 (Krotkov et al., 2017) as input to the LNO_x retrieval algorithm.

The main steps of calculating the NO₂ tropospheric VCD (V_{NO_2}) in the NASA product include the following.

- SCDs are determined by the OMI-optimized differential optical absorption spectroscopy (DOAS) spectral fit.
- 2. A corrected ("de-striped") SCD is obtained by subtracting the cross-track bias caused by an instrument artifact from the measured slant column.
- 3. The AMF for stratospheric (AMF_{strat}) or tropospheric column (AMF_{trop}) is calculated from the NO₂ profiles integrated vertically using weighted scattering weights with the a priori profiles. These profiles are obtained from GMI monthly mean profiles using 4 years (2004–2007) of simulation.
- 4. The stratospheric NO₂ VCD (V_{strat}) is calculated from the subtraction of the a priori contribution from tropospheric NO₂ and a three-step (interpolation, filtering, and smoothing) algorithm (Bucsela et al., 2013).
- 5. V_{strat} is converted to the slant column using AMF_{strat} and subtracted from the measured SCDs to yield S_{NO_2} , leading to $V_{NO_2} = S_{NO_2}/AMF_{trop}$.

Based on this method, we developed a new AMF_{LNO_x} to obtain the desired V_{LNO_x} ($V_{LNO_x} = S_{NO_2}/AMF_{LNO_x}$) by replacing the original step.

6. Details of this algorithm are discussed in Sect. 2.4.

2.2 The Earth Networks Total Lightning Detection Network (ENTLN)

The Earth Networks Total Lightning Network (ENTLN) operates a system of over 1500 ground-based stations around the world with more than 900 sensors installed in the CONUS (Zhu et al., 2017). Both IC and CG lightning flashes are located by the sensors with detection frequency ranging from 1 Hz to 12 MHz based on the electric field pulse polarity and wave shapes. Groups of pulses are classified as a flash if they are within 700 ms and 10 km. In the preprocessed data obtained from the ENTLN, both strokes and lightning flashes composed of one or more strokes are included.

Rudlosky (2015) compared ENTLN combined events (IC and CG) with LIS flashes and found that the relative flash detection efficiency of ENTLN over CONUS increases from 62.4 % during 2011 to 79.7 % during 2013. Lapierre et al. (2020) also compared combined ENTLN and the NLDN dataset with data from the LIS during 2014 and found the detection efficiencies of IC flashes and strokes to be 88 % and 45 %, respectively. Since we only use the ENTLN data in 2014 as Lapierre et al. (2020), and NLDN detection efficiency of IC pulses should be lower than 33 %, which is calculated by the data in 2016 (Zhu et al., 2016), only the IC flashes and strokes are divided by 0.88 and 0.45, respectively, while CG flashes and strokes are unchanged because of the high detection efficiency.

2.3 Model description

The present study uses WRF-Chem version 3.5.1 (Grell et al., 2005) with a horizontal grid size of $12 \text{ km} \times 12 \text{ km}$ and 29 vertical levels (Fig. 1). The initial and boundary conditions of meteorological parameters are provided by the North American Regional Reanalysis (NARR) dataset with a 3-hourly time resolution. Based on Laughner et al. (2019), 3D wind fields, temperature, and water vapor are nudged towards the NARR data. Outputs from version 4 of the Model for Ozone and Related chemical Tracers (MOZART-4; Emmons et al., 2010) are used to generate the initial and boundary conditions of chemical species. Anthropogenic emissions are driven by the 2011 National Emissions Inventory (NEI), scaled to model years by the Environmental Protection Agency annual total emissions (EPA and OAR, 2015). The Model of Emissions of Gases and Aerosol from Nature (MEGAN; Guenther et al., 2006) is used for biogenic emissions. The chemical mechanism is version 2 of the Regional Atmospheric Chemistry Mechanism (RACM2; Goliff et al., 2013) with updates from Browne et al. (2014) and Schwantes et al. (2015). In addition, lightning flash rate based on the



Figure 1. Domain and terrain height (m) of the WRF-Chem simulation with 350×290 grid cells and a horizontal resolution of 12 km.

level of neutral buoyancy parameterization (Price and Rind, 1992; Wong et al., 2013) and LNO_x parameterizations is activated (200 mol NO per flash, the factor to adjust the predicted number of flashes is set to 1, hereinafter referred to as "1 × 200 mol NO per flash"). Simulated total flash densities are higher than ENTLN observations over the southeast US and lower than observations in the north-central US (Fig. 2). The impact of these biases on LNO_x production is discussed and mitigated in Sect. 3.1 and 3.4. The bimodal profile modified from the standard Ott et al. (2010) profile (Laughner and Cohen, 2017) is employed as the vertical distribution of lightning NO (LNO) in WRF-Chem, while outputs of LNO and LNO₂ profiles are defined as the difference of vertical profiles between simulations with and without lightning.

2.4 Method for deriving AMF

The V_{LNO_r} near convection is calculated according to

$$V_{LNO_x} = \frac{S_{NO_2}}{AMF_{LNO_x}},\tag{1}$$

where S_{NO_2} is the OMI-measured tropospheric slant column NO₂, and AMF_{LNO_x} is a customized lightning air mass factor. The concept of AMF_{LNO_x} was also used in Beirle et al. (2009) to investigate the sensitivity of satellite instruments to freshly produced lightning NO_x. In order to estimate LNO_x, we define the AMF_{LNO_x} as the ratio of the "visible" modeled NO₂ slant column to the total modeled tropospheric LNO_x vertical column (derived from the a priori NO and NO₂ profiles, scattering weights, and cloud radiance fraction):

$$AMF_{LNO_x} = \frac{(1-f_r) \int_{\rho_{surf}}^{\rho_{to}} w_{clear}(p) NO_2(p) \, dp}{\int_{\rho_{surf}}^{\rho_{to}} w_{cloudy}(p) NO_2(p) \, dp},$$
(2)

where f_r is the cloud radiance fraction (CRF), p_{surf} is the surface pressure, p_{tp} is the tropopause pressure, p_{cloud} is the cloud optical pressure (CP), w_{clear} and w_{cloudy} are respectively the pressure-dependent scattering weights from



Figure 2. Comparison between total flash densities from ENTLN and WRF-Chem during MJJA 2014.

the TOMRAD lookup table (Bucsela et al., 2013) for clear and cloudy parts, and $NO_2(p)$ is the modeled NO_2 vertical profile. Details of these standard parameters and calculation methods are given in Laughner et al. (2018). $LNO_x(p)$ is the LNO_x vertical profile calculated by the difference of vertical profiles between WRF-Chem simulations with and without lightning.

Please note that the CP is a reflectance-weighted pressure retrieved by the collision-induced O₂-O₂ absorption band near 477 nm (Acarreta et al., 2004; Sneep et al., 2008; Stammes et al., 2008). For a deep convective cloud with lightning, the CP lies below the geometrical cloud top, which is much closer to that detected by thermal infrared sensors, such as CloudSat and the Aqua Moderate Resolution Imaging Spectrometer (MODIS) (Vasilkov et al., 2008; Joiner et al., 2012). Hence, much of the tropospheric NO₂ measured by OMI lies inside the cloud rather than above the cloud top. In the following, "above cloud" or "below cloud" is relative to the cloud pressure detected by OMI. The sensitivity study of Beirle et al. (2009) compared the chemical composition from the cloud bottom to that of the cloud top and revealed that a significant fraction of the NO₂ within the cloud originating from lightning can be detected by the satellite. This valuable cloud pressure concept has been applied not only in the LNO_x research but also in the cloud slicing method of deriving the UT O_3 and NO_x (Ziemke et al., 2009, 2017; Choi et al., 2014; Strode et al., 2017; Marais et al., 2018). As discussed in Pickering et al. (2016), the ratio of V_{LNO_2} seen by OMI to V_{LNO_x} is partly influenced by p_{cloud} . The effects of LNO₂ below the cloud will be discussed in Sect. 3.4.

To compare our results with those of Pickering et al. (2016) and Lapierre et al. (2020), we calculate their

AMF_{LNO_xClean} and AMF_{NO₂Vis}, respectively:

$$AMF_{LNO_{x}}Clean = \frac{(1-f_{r})\int_{p_{surf}}^{p_{tp}} w_{clear}(p)LNO_{2}(p) dp}{\int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p)LNO_{2}(p) dp}, \qquad (3)$$
$$(1-f_{r})\int_{p_{surf}}^{p_{tp}} w_{clear}(p)NO_{2}(p) dp$$

$$AMF_{NO_2Vis} = \frac{+f_r \int_{\rho_{cloud}}^{\rho_{tp}} w_{cloudy}(p) NO_2(p) \, dp}{(1 - f_g) \int_{\rho_{surf}}^{\rho_{tp}} NO_2(p) \, dp}, \qquad (4)$$
$$+f_g \int_{\rho_{cloud}}^{\rho_{tp}} NO_2(p) \, dp$$

where f_g is the geometric cloud fraction and LNO₂(*p*) is the modeled LNO₂ vertical profile. Besides these AMFs, another AMF called AMF_{LNO₂Vis} is developed for later comparison.

$$AMF_{LNO_2}v_{is} = \frac{(1-f_r)\int_{p_{surf}}^{p_{tp}} w_{clear}(p)NO_2(p) \, dp}{+f_r \int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p)NO_2(p) \, dp} + f_g \int_{p_{surf}}^{p_{tp}} LNO_2(p) \, dp$$
(5)

A full definition list of the used AMFs is shown in Appendix A.

2.5 Procedures for deriving LNO_x

 V_{LNO_x} is re-gridded to $0.05^{\circ} \times 0.05^{\circ}$ grids using the constant value method (Kuhlmann et al., 2014). Then, it is analyzed in $1^{\circ} \times 1^{\circ}$ grid boxes with a minimum of 50 valid $0.05^{\circ} \times 0.05^{\circ}$ grids to minimize the noise. The main procedures of deriving LNO_x are as follows.

CRFs (CRF ≥ 70%, CRF ≥ 90%, and CRF = 100%) and CP ≤ 650 hPa are various criteria of deep convective clouds for OMI pixels (Ziemke et al., 2009; Choi et al., 2014; Pickering et al., 2016). The effect of different CRFs on the retrieved LNO_x is explored in Sect. 3.2. Furthermore, another criterion of cloud fraction (CF) is applied to the WRF-Chem results for the successful simulation of convection. The CF is defined as the maximum cloud fraction calculated by the Xu–Randall method between 350 and 400 hPa (Xu and Randall, 1996; Strode et al., 2017). This atmospheric layer (between 350 and 400 hPa) avoids any biases in the simulation of high clouds. We choose CF ≥ 40% suggested by Strode et al. (2017) to determine cloudy or clear for each simulation grid.

Besides cloud properties, a time period and sufficient flashes (or strokes) are required for fresh LNO_x to be detected by OMI. The time window (t_{window}) is the hours prior to the OMI overpass time. t_{window} is limited to 2.4 h by the mean wind speed at pressure levels 500-100 hPa during OMI overpass time and the square root of the $1^{\circ} \times 1^{\circ}$ box over the CONUS (Lapierre et al., 2020). Meanwhile, 2400 flashes per box and 8160 strokes per box per 2.4 h time window are chosen as sufficient for detecting LNO_x (Lapierre et al., 2020). These criteria will result in a low bias in the PE results, as Bucsela et al. (2019) found that the PE is larger at small flash rates, which are discarded here. Since our study focuses on developing a new AMF and compares results with other works using similar lightning thresholds (Lapierre et al., 2020; Pickering et al., 2016), we will only discuss results based on the strict criteria in the main text. For comparisons between the criterion of 2400 flashes per box and that of one flash per box, scatter diagrams using different lightning criteria are presented in Appendix B.

To ensure that lightning flashes are simulated successfully by WRF-Chem, the threshold of simulated total lightning flashes (TL) per box is set to 1000, which is fewer than that used by the ENTLN lightning observation, considering the uncertainty of lightning parameterization. In view of other NO₂ sources in addition to LNO₂, the ratio of modeled lightning NO₂ above cloud (LNO₂Vis) to modeled NO₂ above cloud (NO₂Vis) is defined to check whether enough LNO₂ can be detected by OMI. The ratio \geq 50 % indicates that more than half of the NO_x above the cloud must have an LNO_x source.

Finally, the NO₂ lifetime due to oxidation should be taken into account. As estimated by Nault et al. (2017), the lifetime (τ) of NO₂ in the near field of convections is ~ 3 h. The initial value of NO₂ is solved by Eq. (6) as

$$NO_2(0) = NO_2(OMI) \times e^{0.5t/\tau},$$
(6)

where NO₂(0) is the moles of NO₂ emitted at time t = 0, NO₂(OMI) is the moles of NO₂ measured at the OMI overpass time, and 0.5*t* is the half cross grid time, which is 1.2 h, assuming that lightning occurred at the center of each 1° × 1° box. For each grid box, the mean LNO_x vertical column is obtained by averaging V_{LNO_x} values from all regridded $0.05^{\circ} \times 0.05^{\circ}$ pixels in the box. This mean value is converted to moles of LNO_x using the dimensions of the grid box. Two methods are applied to estimate the seasonal mean LNO₂ per flash, LNO_x per flash, LNO₂ per stroke, and LNO_x per stroke:

- 1. summation method, dividing the sum of LNO_x by the sum of flashes (or strokes) in each $1^\circ \times 1^\circ$ box in MJJA 2014;
- 2. linear regression method, applying the linear regression to daily mean values of LNO_x and flashes (or strokes).

3 Results

3.1 Criteria determination

To determine the suitable criteria from the conditions defined in Sect. 2.5, six different combinations are defined (Table 1) and applied to the original data with a linear regression method (Table 2).

A daily search of the NO₂ product for coincident ENTLN flash (stroke) data results in 99 (102) valid days under the CRF90_ENTLN condition. Taking the flash-type ENTLN data as an example, the number of valid days decreases from 99 to 81 under the CRF90_ENTLN_TL1000_ratio50 condition, while LNO_x per flash increases from 52.1 ± 51.1 to 54.5 ± 48.1 mol per flash. The result is almost the same as that under the CRF90 ENTLN TL1000 condition, which is without the condition of more than half of the above-cloud NO_x having an LNO_x source. Although this indicates the criterion of TL works well, it is better to include the ratio criterion in case there are some exceptions in the different AMF methods. Since $CF \ge 40\%$ leads to a sharp loss of valid numbers and production, it is not a suitable criterion. Instead the CRF criteria are used. Finally, coincident ENTLN data, TL \geq 1000, and ratio \geq 50 % are chosen as the thresholds to explore the effects of three different CRF conditions (CRF \geq 70%, CRF \geq 90%, and CRF = 100%) on LNO_x production (Table 3). Apart from the fewer valid days under higher CRF conditions (CRF > 90% and CRF = 100 %), LNO_x per flash increases from 35.7 ± 36.8 to 54.5 \pm 48.1 mol per flash and decreases again to 20.8 \pm 37.4 mol per flash while LNO_x per stroke enhances from 4.1 ± 3.9 to 7.0 ± 4.8 mol per stroke and drops again to 2.6 ± 4.0 mol per stroke (Table 3), as the CRF criterion increases from 70 % to 90 % and to 100 %. When the CRF increases from 90 % to 100 %, the LNO_x PE decreases because of the higher lightning density with less LNO_x (not shown). The increment of LNO_x PE caused by the CRF increase from 70 % to 90 % is opposite to the result of Pickering et al. (2016). This is an effect of the consideration of NO2 contamination transported from the boundary layer in our method. Although enhanced NO_x is often observed in regions with CRF > 70 % (Pickering et al., 2016), the following analysis will be based on the criterion of CRF \geq 90 % considering the contamination by low and midlevel NO₂ and comparisons with the results of Pickering et al. (2016) and Lapierre et al. (2020).

3.2 Comparison of LNO_x production based on different AMFs

Lapierre et al. (2020) derived LNO₂ production based on the BEHR NO₂ product. In order for our results to be comparable with those of Pickering et al. (2016) and Lapierre et al. (2020), we choose NO₂ instead of NO_x to derive production per flash (production efficiency, PE). In Fig. 3, time series of NO₂Vis, LNO₂Vis, LNO₂, and LNO₂Clean production per

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Abbreviations	Full form (source)
CRF	Cloud radiance fraction (OMI)
CP	Cloud optical pressure (OMI)
CF	Cloud fraction (WRF-Chem)
TL	Total lightning flashes (WRF-Chem)
Ratio	Modeled LNO ₂ Vis / modeled NO ₂ Vis (WRF-Chem)
CRFα_ENTLN	$CRF \ge \alpha + ENTLN$ flashes (strokes) ≥ 2400 (8160) (ENTLN)
$CRF\alpha_CF40_ENTLN$	$CRF \ge \alpha + ENTLN$ flashes (strokes) $\ge 2400 (8160) + CF \ge 40 \%$
$CRF\alpha_ENTLN_TL1000$	$CRF \ge \alpha + ENTLN$ flashes (strokes) $\ge 2400 (8160) + TL \ge 1000$
CRFα_CF40_ENTLN_TL1000	$CRF \ge \alpha + ENTLN$ flashes (strokes) $\ge 2400 (8160) + CF \ge 40 \% + TL \ge 1000$
CRFa_ENTLN_TL1000_ratio50	$CRF \ge \alpha + ENTLN$ flashes (strokes) $\ge 2400 (8160) + TL \ge 1000 + ratio \ge 50 \%$
CRFα_CF40_ENTLN_TL1000_ratio50	$CRF \ge \alpha + ENTLN$ flashes (strokes) $\ge 2400 (8160) + CF \ge 40\% + TL \ge 1000 + ratio \ge 50\%$
CRFa_ENTLN1(3.4)_TL1_ratio50	$CRF \ge \alpha + ENTLN$ flashes (strokes) ≥ 1 (3.4) + $TL \ge 1 + ratio \ge 50 \%$

 α has three options: 70 %, 90 %, or 100 %.

Table 2. LNO_x production efficiencies for different combinations of criteria defined in Table 1.

Condition ¹	ENTLN data type ²	LNO_x per flash or LNO_x per stroke	R value	Intercept (10 ⁶ mol)	Days ³
CRF90_ENTLN	Flash	52.1 ± 51.1	0.20	0.21	99
CRF90_CF40_ENTLN	Flash	84.2 ± 31.5	0.54	-0.04	70
CRF90_ENTLN_TL1000	Flash	61.9 ± 49.1	0.27	0.33	83
CRF90_CF40_ENTLN_TL1000	Flash	63.4 ± 52.9	0.38	0.26	38
CRF90_ENTLN_TL1000_ratio50	Flash	54.5 ± 48.1	0.25	0.39	81
CRF90_CF40_ENTLN_TL1000_ratio50	Flash	90.0 ± 65.0	0.46	0.15	32
CRF90_ENTLN	Stroke	6.7 ± 4.1	0.31	0.23	102
CRF90_CF40_ENTLN	Stroke	10.3 ± 3.6	0.55	0.08	79
CRF90_ENTLN_TL1000	Stroke	7.5 ± 5.1	0.29	0.38	94
CRF90_CF40_ENTLN_TL1000	Stroke	8.6 ± 6.2	0.39	0.27	46
CRF90_ENTLN_TL1000_ratio50	Stroke	7.0 ± 4.8	0.29	0.42	93
CRF90_CF40_ENTLN_TL1000_ratio50	Stroke	8.9 ± 7.0	0.39	0.31	40

¹ These conditions are defined in Table 1. ² The thresholds of ENTLN data are 2400 flashes per box and 8160 strokes per box during the period of 2.4 h before OMI overpass time. ³ The number of valid days with specific criteria in MJJA 2014.

day over CONUS are plotted for MJJA 2014 with the criterion of CRF \geq 90 % and a flash threshold of 2400 flashes per 2.4 h. LNO₂ PEs are mostly in the range from 20 to 80 mol per flash. LNO₂Vis PEs are smaller than LNO₂ PEs, which contain LNO2 below clouds. The simulation of GMI in Pickering et al. (2016) indicated that 25 % - 30 % of the LNO_x column lies below the CP, while the ratio in our WRF-Chem simulation is 56 ± 20 %. The effect of cloud properties on LNO_x PE will be discussed in more detail in Sect. 3.4. Generally, the order of estimated daily PEs is LNO₂Clean > LNO₂ $> NO_2 Vis > LNO_2 Vis$. The percent difference in the estimated PE (Δ PE) between NO₂Vis and LNO₂Vis indicates a certain amount of background NO₂ exists above clouds. Overall, the tendency of that ΔPE is consistent with another ΔPE between NO₂Vis and LNO₂Clean. When the region is highly polluted (ΔPE between NO₂Vis and LNO₂Vis is larger than 200%), PEs based on NO₂Vis and LNO₂Clean are significantly overestimated. In other words, NO₂Vis and LNO₂Clean are more sensitive to background NO₂. The extent of the overestimation of NO₂Vis is larger than that of LNO₂Clean in highly polluted regions, while it is usually opposite in most regions.

Figure 4 shows the linear regression for ENTLN data versus NO₂Vis, LNO₂Vis, LNO₂, and LNO₂Clean with the same criteria as shown in Fig. 3. LNO₂Clean PE (the largest slope) is $25.2 \pm 22.3 \text{ mol NO}_2$ per flash with a correlation of 0.25 and $2.3 \pm 2.1 \text{ mol NO}_2$ per stroke with a correlation of 0.22. As shown in Fig. 3, positive percent differences between NO₂Vis PE and LNO₂Clean PE occur much less often than negative differences. As a result, NO₂Vis PE (17.1 ± 17.2 mol NO₂ per flash and $0.4 \pm 1.0 \text{ mol NO}_2$ per stroke) is smaller than LNO₂Clean PE using the linear regression method.

In order to compare our result with that of Lapierre et al. (2020), we tried to remove the CP \leq 650 hPa, TL \geq 1000, and ratio \geq 50% conditions from criteria. But, our result

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CRF (%)	ENTLN data type ¹	LNO_x per flash or LNO_x per stroke	R value	Intercept (10 ⁵ mol)	Days ²
70	Flash	35.7 ± 36.8	0.21	4.91	85
90	Flash	54.5 ± 48.1	0.25	3.90	81
100	Flash	20.8 ± 37.4	0.13	5.67	71
70	Stroke	4.1 ± 3.9	0.21	5.16	96
90	Stroke	7.0 ± 4.8	0.29	4.16	93
100	Stroke	2.6 ± 4.0	0.14	5.41	82

Table 3. LNO_x production efficiencies for different thresholds of CRF with coincident ENTLN data, TL \geq 1000, and ratio \geq 50 %.

¹ The thresholds of ENTLN data are 2400 flashes per box and 8160 strokes per box during the period of 2.4 h before OMI overpass time. ² The number of valid days with specific criteria in MJJA 2014.



Figure 3. (a) Time series of NO₂Vis, LNO₂Vis, LNO₂, and LNO₂Clean production per day over the CONUS for MJJA 2014 with CRF \geq 90 % and a flash threshold of 2400 flashes per 2.4 h. (b) Time series of the percent differences between NO₂Vis and LNO₂Vis and the percent differences between NO₂Vis and LNO₂Clean with CRF \geq 90 %. The value of the black dot on 23 August (not shown) is 1958 %.

based on daily summed NO₂Vis values $(3.8 \pm 0.5 \text{ mol per stroke})$ is still larger than the value of 1.6 ± 0.1 mol per stroke mentioned in Lapierre et al. (2020). This may be caused by the different version of the BEHR algorithm, as Lapierre et al. (2020) used BEHR v3.0A and our algorithm is based on BEHR v3.0B (Laughner et al., 2019). The input of S_{NO₂} in both versions is from the NASA standard product v3, and the major improvements of BEHR v3.0B are listed below.

- 1. The profile (v3.0B) closest to the OMI overpass time was selected instead of the last profile (v3.0A) before the OMI overpass.
- 2. The AMF uses a variable tropopause height as opposed to the fixed 200 hPa tropopause.

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3. The surface pressure is now calculated according to Zhou et al. (2009).

The detailed log of changes is available at https: //github.com/CohenBerkeleyLab/BEHR-core/blob/master/ Documentation/Changelog.txt (last access: 20 March 2020). Note that Lapierre et al. (2020) used the monthly NO₂ profile. While the daily profile is used in our study and the interval of our outputs from WRF-Chem is 30 min, which is more frequent than 1 h in the BEHR daily product, the AMF could be affected by different NO₂ profiles. In view of these factors, we compare different methods based on our data to minimize these effects.

Meanwhile, LNO₂ PE (18.7 \pm 18.1 mol per flash and 2.1 \pm 1.8 mol per stroke) is between LNO₂Clean PE and NO₂Vis

PE, which coincides with the daily results in Fig. 3. Furthermore, the LNO_x PE based on the linear regression of daily summed values, the same method used in Pickering et al. (2016), is 114.8 ± 18.2 mol per flash (or 17.8 ± 2.9 mol per stroke), which is larger than 91 mol per flash in Pickering et al. (2016), possibly due to the differences in geographic location, lightning data, and chemistry model.

The mean and standard deviation of LNO₂ PE under $CRF \ge 90\%$ using the summation method is 46.2 ± 35.1 mol per flash and 9.9 ± 8.1 mol per stroke, while LNO_x PE is 125.6 ± 95.9 mol per flash and 26.7 ± 21.6 mol per stroke (Fig. 5). The LNO₂ PE and LNO_x PE are both higher in the southeast US (denoted by the red box in Fig. 5, 25-37° N, 75-95° W), consistent with Lapierre et al. (2020) and Bucsela et al. (2019). Compared with Fig. 3, Fig. 6a and b present some large differences between NO₂Vis PE and LNO₂Vis PE, which are consistent with what we expect for polluted regions. Meanwhile, the differences between LNO₂ PE and NO_2V is PE depend on background NO_2 , the strength of updraft, and the profile. The negative differences are caused by background NO₂ carried by the updraft while parts of the below-cloud LNO₂ result in LNO₂ PE higher than NO₂Vis PE (Fig. 6c). Figure 6d shows that the ratio of LNO_2V is to LNO₂ ranges from 10 % to 80 %. This may be caused by the height of the clouds and the profile of LNO₂. If the CP is near 300 hPa, the ratio should be smaller because of the coverage of clouds. While peaks of the LNO₂ profile are below the CP, the ratio would also be smaller. Therefore, a better understanding of the LNO₂ profile and LNO_x below clouds is required.

3.3 Effects of tropospheric background on LNO_x production

With respect to the LNO₂ production, the patterns in Fig. 6 indicate the improvement of our approach is different in polluted and clean regions. To simplify the quantification, we select six grids with similar NO₂ profiles ($\sim 100 \text{ pptv}$) above the cloud with CRF = 100 %. These grid boxes contain the polluted and clean cities denoted by stars and triangles in Fig. 6a, respectively. Then, the differences between AMFs are dependent on fewer parameters.

$$AMF_{LNO_2} = \frac{\int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p)NO_2(p) \, dp}{\int_{p_{surf}}^{p_{tp}} LNO_2(p) \, dp}$$
(7)

$$AMF_{NO_2Vis} = \frac{\int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p)NO_2(p) dp}{\int_{p_{cld}}^{p_{tp}} NO_2(p) dp}$$
(8)

$$AMF_{LNO_2Clean} = \frac{\int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p) LNO_2(p) \, dp}{\int_{p_{surf}}^{p_{tp}} LNO_2(p) \, dp}$$
(9)

Figure 7 compares the mean profiles of NO_2 , background NO_2 and background NO_2 ratio in polluted and clean grids. Generally, the profiles of the ratio of background NO_2 to total NO_2 are C shaped because UT LNO_2 concentrations are

higher than UT background NO_2 concentrations. However, the ratio profile in Fig. 7e has one peak between the cloud pressure and tropopause as background NO_2 increases and LNO_2 decreases. Besides, the percentage of UT background NO_2 in polluted regions is steady and higher than that in clean regions.

Table 4 presents the relative changes among three methods in six cities. The difference between AMF_{LNO2} (Eq. 7) and AMF_{LNO2}Clean (Eq. 9) $\int_{p_{\text{cloud}}}^{p_{\text{tp}}} w_{\text{cloudy}}(p) \operatorname{NO}_2(p) \, \mathrm{d}p$ is the numerator: and $\int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p) LNO_2(p) dp$. When the ratio of LNO₂ is higher or the region is cleaner, the relative difference is smaller (e.g. 5.0%-12.0%, Fig. 7d-f). The largest relative difference (46.3%) occurs when the ratio of background NO_2 is continuously high in the UT (Fig. 7c). As a result, our approach is less sensitive to background NO2 and more suitable for convective cases over polluted locations. In contrast, production estimated by our method is larger than that based on NO₂Vis due to the LNO₂ below the cloud. When the cloud is higher, in particular the peak of the LNO profile is lower than the cloud (Fig. 7b). The relative difference is larger (121.2%) because more LNO₂ can not be included in the NO₂Vis, which has been discussed in Sect. 3.2. The relative change between AMF_{LNO₂Clean} (Eq. 9) and AMF_{NO₂Vis} (Eq. 8) depends on $\int_{p_{cloud}}^{p_{tp}} w_{cloudy}(p) LNO_2(p) dp / \int_{p_{surf}}^{p_{tp}} w_{cloudy}(p) LNO_2(p) dp,$ which is also affected by cloud, not the background NO_2 . The largest relative change (153.8%) occurs at New Orleans, which has the lowest cloud pressure and consequently the smallest visible column.

3.4 Effects of cloud and LNO_x parameterization on LNO_x production

Figure 8a presents the daily distribution of CP and the ratio of LNO₂Vis to LNO₂ during MJJA 2014 with the criteria defined in Sect. 3.1 under CRF \geq 90 %. Since the ratio of LNO₂Vis to LNO₂ decreases from 0.8 to 0.2 as the cloud pressure decreases from 600 to 300 hPa, NO₂Vis PE is smaller than LNO₂ PE in relatively clean areas as shown in Fig. 4. Apart from LNO₂Vis, the LNO₂ PE is also affected by CP. For LNO₂ PEs larger than 30 mol per stroke, the CPs are all smaller than 550 hPa (Fig. 8b). However, smaller LNO₂ PEs (< 30 mol per stroke) occur on all levels between 650 and 200 hPa. Because of the limited number of large LNO₂ PEs and lightning data, we cannot derive the relationship between LNO₂ PE and cloud pressure or other lightning properties at this stage. Because the CP only represents the development of clouds, the vertical structure of flashes can not be derived from the CP values only. As discussed in several previous studies, the flash channel length varies and depends on the environmental conditions (Carey et al., 2016; Mecikalski and Carey, 2017; Fuchs and Rutledge, 2018). Davis et al. (2019) compared two kinds of flash: normal flashes and anomalous flashes. Because updrafts are stronger and flash



Figure 4. (a) Daily NO₂Vis, LNO₂Vis, LNO₂, and LNO₂Clean versus ENTLN total flash data. (b) Same as (a) but for strokes. (c) Daily LNO_x Vis and LNO_x versus total flashes. (d) Same as (c) but for strokes.

rates are higher in anomalous storms, UT LNO_x concentrations are larger in anomalous than normal polarity storms. In general, normal flashes are coupled with an upper-level positive charge region and a midlevel negative charge region, while anomalous flashes are opposite (Williams, 1989). It is not straightforward to estimate the error resulting from the vertical distribution of LNO_x . There are mainly two methods of distributing LNO_x in models: LNO_x profiles (postconvection) in which LNO_x has already been redistributed by convective transport and LNO_x production profiles (preconvection) made before the redistribution of convective transport (Allen et al., 2012; Luo et al., 2017). However, given the similarity of results compared to other LNO_x studies, we believe that our $1^\circ \times 1^\circ$ results based on postconvective LNO_x profiles are sufficient for estimating average LNO_x production.

The LNO production settings in WRF-Chem varied in different studies. Zhao et al. (2009) set a NO_x production rate of 250 mol NO per flash in a regional-scale model, while Bela et al. (2016) chose 330 mol NO per flash used by Barth et al. (2012). Wang et al. (2015) assumed approximately 500 mol NO per flash, which was derived by a cloud-scale chemical transport model and in-cloud aircraft observations (Ott et al., 2010). To illustrate the im-

pact of LNO_x parameterization on LNO_x estimation, we apply another WRF-Chem NO₂ profile setting $(2 \times \text{ base})$ flash rate, 500 mol NO per flash, hereinafter referred to as " $2 \times 500 \text{ mol NO per flash"}$) to a priori profiles and evaluate the changes in AMF_{LNO2}, AMF_{LNOx}, LNO₂ PE, and LNO_x PE. For the linear regression method (Fig. 9), LNO₂ PE is 29.8 ± 20.5 mol per flash, which is 59.4 % larger than the basic one (18.7 \pm 18.1 mol per flash). Meanwhile, LNO_x PE (increasing from 54.5 ± 48.1 mol per flash to 88.5 ± 61.1 mol per flash) also depends on the configuration of LNO production in WRF-Chem. The comparison between Figs. 4 and 9 shows that LNO₂Clean PE and LNO₂ PE are more similar while LNO₂ PE and NO₂Vis PE present the same tendency. It remains unclear as to whether the NO-NO₂-O₃ cycle or other LNO_x reservoirs account for the increment of LNO_x PE. This would need detailed source analysis in WRF-Chem and is beyond the scope of this study.

Figure 10 shows the average percentage changes in AMF_{LNO_2} , AMF_{LNO_x} , LNO_2 , and LNO_x between retrievals using profiles based on 1×200 and 2×500 mol NO per flash. These results were obtained by averaging data over MJJA 2014 based on the method described in Sect. 2.5 with the criterion of CRF $\geq 90\%$. The effects on LNO₂ and LNO_x



Figure 5. (**a**, **c**) Maps of $1^{\circ} \times 1^{\circ}$ gridded values of mean LNO_x and LNO₂ production per flash with CRF $\ge 90\%$ for MJJA 2014. (**b**, **d**) Same as (**a**) and (**c**) except for strokes. The southeastern US is denoted by the red box in panels (**a**)–(**d**).

Table 4. The percent changes in the estimated production when using different methods based on the same a priori profiles.

	City*	(LNO ₂ Clean – LNO ₂)/LNO ₂	(LNO ₂ – TropVis)/TropVis	(LNO ₂ Clean-TropVis)/TropVis
	Lansing	24.2 %	49.5 %	85.6%
Polluted	New Orleans	13.3 %	121.2 %	153.8 %
	Orlando	46.3 %	37.5 %	101.3 %
	Huron	12.0 %	56.4 %	75.2 %
Clean	Charles Town	12.0 %	82.2 %	104.1 %
	Tarboro	5.0 %	86.0 %	95.3 %

* Locations are denoted in Fig. 6a.

retrieval from increasing LNO profile values show mostly the same tendency: smaller AMF_{LNO_2} and AMF_{LNO_x} lead to larger LNO₂ and LNO_x, but the changes are regionally dependent. This is caused by the nonlinear calculation of AMF_{LNO_2} and AMF_{LNO_x} . As the contribution of LNO₂ increases, both the numerator and denominator of Eq. (2) increase. Note that the LNO₂ accounts for a fraction of NO₂ above the clouds. The magnitude of an increasing denominator could be different than that of an increasing numerator, resulting in a different effect on the AMF_{LNO2} and AMF_{LNOx}. As mentioned in Zhu et al. (2019), the lightning densities in the southeast US might be overestimated using the 2×500 mol NO per flash setting and the same lightning parameterization as ours. Fortunately, the AMFs and esti-



Figure 6. (a) Mean (MJJA 2014) NO₂ tropospheric column. Polluted cities are denoted by stars (Lansing, New Orleans, and Orlando) while clean cities are denoted by triangles (Huron, Charles Town, and Tarboro). (b) The differences of the estimated mean production efficiency between NO₂Vis and LNO₂Vis with CRF \ge 90 %. (c) The same differences as (b) but between LNO₂ and NO₂Vis. (d)The ratio of LNO₂Vis to LNO₂.

mated LNO₂ change little in that region. Because the southeast US has the highest flash density (Fig. 2), the NO₂ in the numerator of AMF is dominated by LNO₂. Both the SCD and VCD will increase when the model uses higher LNO₂. In other words, the sensitivity to the LNO setting decreases and the relative distribution of LNO₂ matters.

Figure 11 shows the comparison of the mean LNO and LNO₂ profiles in two specific regions where the $2 \times$ 500 mol NO per flash setting leads to lower and higher LNO₂ PEs, respectively. The first one (Fig. 11a) is the region (36– 37° N, 89–90° W) containing the minimal negative percent change in LNO₂ (Fig. 10c). The second one (31–32° N, 97– 98° W), Fig. 11b, has the largest positive percent change in LNO₂ (Fig. 10c). Although the relative distributions of mean LNO and LNO₂ profiles are similar in both regions, the magnitude differs by a factor of 10. This phenomenon implies that the performance of lightning parameterization in WRF-Chem is regionally dependent, and an unrealistic profile could appear in the UT. Although this sensitivity analysis is false in some regions, it allows the calculation of an upper limit on the NO₂ due to LNO and LNO₂ profiles. As discussed in Laughner and Cohen (2017), the scattering weights are uniform under cloudy conditions and the sensitivity of NO₂ is nearly constant with different pressure levels because of the high albedo. However, the relative distribution of LNO₂ within the UT should be taken carefully into consideration. If the LNO₂/NO₂ above the cloud is large enough (Fig. 11a), the AMF_{LNO₂} is largely determined by the ratio of LNO₂Vis to LNO₂, which is related to the relative distribution. When the condition of high LNO₂/NO₂ is not met, both relative distribution and ratio are important (Fig. 11b).

To clarify this, we applied the same sensitivity test of different simulating LNO amounts for all four methods mentioned in Sect. 2.4: LNO₂, LNO₂Vis, LNO₂Clean, and NO₂Vis (Fig. 12). Note that the threshold for CRF is set to 100% to simplify Eq. (2) to Eq. (7). The overall differences of LNO₂Clean and NO₂Vis are smaller than those of LNO₂ and LNO₂Vis. Comparing the numerator and denom-



Figure 7. Comparisons of mean WRF-Chem NO₂ and background NO₂ profiles in six grids with CRF $\geq 100\%$ on specific days during MJJA 2014. The (**a**), (**b**), and (**c**) data are selected from polluted regions (stars in Fig. 6a) while the (**d**), (**e**), and (**f**) data are from clean regions (triangles in Fig. 6a). The green dashed lines are the mean ratio profiles of background NO₂ to total NO₂. The zoomed figures show the profiles from the cloud pressure to the tropopause. The titles present the mean productions based on three different methods mentioned in Sect. 2.4.



Figure 8. Kernel density estimation of the (a) daily ratio of LNO_2V is to LNO_2 and (b) daily LNO_2 production efficiency versus the daily cloud pressure measured by OMI with $CRF \ge 90\%$ for MJJA 2014. The kernel density estimation was generated by kdeplot in the Python package named seaborn.

inator in the equations, it is clear why the impact of different simulating LNO amounts is smaller in Fig. 12c and d. For LNO₂Clean and NO₂Vis, both the SCD and VCD will increase (decrease) when more (less) LNO₂ or NO₂ presents. The difference between Fig. 12a and b is the denominator: the total tropospheric LNO₂ vertical column and visible

LNO₂ vertical column, respectively. As a result, the negative values in Fig. 12a are caused by the part of LNO₂ below the cloud. The uncertainty of retrieved LNO₂ and LNO_x PEs is driven by this error, and we conservatively estimate this to be ± 13 % and ± 25 %, respectively.



Figure 9. Same as Fig. 4 except for the 2×500 mol NO per flash configuration.

4 Uncertainty analysis

The uncertainties of the LNO₂ and LNO_x PEs are estimated following Pickering et al. (2016), Allen et al. (2019), Bucsela et al. (2019), Laughner et al. (2019) and Lapierre et al. (2020). We determine the uncertainty due to BEHR tropopause pressure, cloud radiance fraction, cloud pressure, surface pressure, surface reflectivity, profile shape, profile location, V_{strat}, the detection efficiency of lightning, t_{window} , and LNO₂ lifetime numerically by perturbing each parameter in turn and re-retrieving the LNO₂ and LNO_x with the perturbed values (Table 5).

The GEOS-5 monthly tropopause pressure, which is consistent with the NASA standard product, is applied instead of the variable WRF tropopause height to evaluate the uncertainty (6% for LNO₂ PE and 4% for LNO_x PE) caused by the BEHR tropopause pressure. The cloud pressure bias is given as a function of cloud pressure and fraction by Acarreta et al. (2004), implying an uncertainty of 32%, the most likely uncertainty in the production analysis, for LNO₂ PE and 34% for LNO_x PE. The resolution of GLOBE terrain height data is much higher than the OMI pixel, and a fixed scale height is assumed in the BEHR algorithm. As a result, Laughner et al. (2019) compared the average WRF surface

pressures to the GLOBE surface pressures and arrived at the largest bias of 1.5 %. Based on the largest bias, we vary the surface pressure (limited to less than 1020 hPa), and the uncertainty can be neglected.

The error in cloud radiance fraction is transformed from cloud fraction using

$$\sigma = 0.05 \cdot \left. \frac{\partial f_{\rm r}}{\partial f_{\rm g}} \right|_{f_{\rm g,pix}},\tag{10}$$

where f_r is the cloud radiance fraction, f_g is the cloud fraction, and $f_{g,pix}$ is the cloud fraction of a specific pixel. We calculate $\partial f_r / \partial f_g$ under $f_{g,pix}$ by the relationship between all binned f_r and f_g with the increment of 0.05 for the each specific OMI orbit. Considering the relationship, the error in cloud fraction is converted to an error in cloud radiance fraction of 2 % for the LNO₂ and LNO_x PEs.

The accuracy of the 500 m MODIS albedo product is usually within 5 % of albedo observations at the validation sites, and those exceptions with low-quality flags have been found to be primarily within 10 % of the field data (Schaaf et al., 2011). Since we use the bidirectional reflectance distribution function (BRDF) data directly, rather than including a radiative transfer model, 14 % Lambertian equivalent reflectivity



Figure 10. Average percent differences in (a) $\text{AMF}_{\text{LNO}_2}$, (b) $\text{AMF}_{\text{LNO}_x}$, (c) LNO_2 , and (d) LNO_x with $\text{CRF} \ge 90\%$ over MJJA 2014. Differences between profiles are generated by 2×500 and 1×200 mol NO per flash.

Table 5. Uncertainties for the estimation of LNO_2 per flash, LNO_x per flash, LNO_2 per stroke, and LNO_x per stroke.

Туре	Perturbation	LNO ₂ per flash ⁵	LNO _x per flash ⁵	LNO ₂ per stroke ⁵	LNO_x per stroke ⁵
BEHR tropopause pressure ¹	NASA product tropopause	6	4	6	4
Cloud radiance fraction ¹	$\pm5\%$	2	2	2	2
Cloud pressure ²	Variable	32	34	32	34
Surface pressure ¹	±1.5 %	0	0	0	0
Surface reflectivity ¹	±17 %	0	0	0	0
LNO ₂ profile ¹	2×500 mol NO per flash	13	25	13	25
Profile location ¹	Quasi-Monte Carlo	0	1	0	1
Lightning detection efficiency ³	IC: ±16 %, CG: ±5 %	15	15	15	15
$t_{\rm window}^3$	2–4 h	10	10	8	8
LNO_x lifetime ³	2–12 h	24	24	24	24
V _{strat} ⁴	-	10	10	10	10
Systematic errors in slant column ⁴	_	5	5	5	5
Tropospheric background ⁴	_	10	10	10	10
NO/NO ₂ ⁴	$20\% \pm 15\%$	0	15	0	15
Net	-	49	56	48	56

 $PE_{uncertainty} = (Error_{rising perturbed value} - Error_{lowering perturbed value})/2, where Error_{perturbed value} = (PE_{perturbed value} - PE_{original value})/PE_{original value} - 1 Laughner et al. (2019). ² Acarreta et al. (2004). ³ Lapierre et al. (2020). ⁴ Allen et al. (2019) and Bucsela et al. (2019). ⁵ Uncertainty (%).$



Figure 11. LNO and LNO₂ profiles with different LNO settings in (a) the region containing the minimal negative percent change in LNO₂ and (b) the region containing the largest positive percent change in LNO₂ when the LNO setting is changed from 1×200 to 2×500 mol NO per flash, averaged over MJJA 2014. The profiles using 1×200 (2×500) mol NO per flash are shown in blue (red) lines. Solid (dashed) green lines are the mean ratio of LNO₂ to NO₂ with 1×200 (2×500) mol NO per flash.

(LER) error and 10% uncertainty are combined to get a perturbation of 17% (Laughner et al., 2019). The uncertainty due to surface reflectivity can be neglected with the 17% perturbation.

As discussed at the end of Sect. 3.4, another setting of LNO₂ (2 × 500 mol NO per flash) is applied to determine the uncertainty of the lightning parameterization and the vertical distribution of LNO in WRF-Chem. Differences between the two profiles lead to an uncertainty of 13% and 25% in the resulting PEs of LNO₂ and LNO_x. Another sensitivity test allows each pixel to shift by -0.2, 0, or $+0.2^{\circ}$ in the directions of longitude and latitude, taking advantage of the high-resolution profile location in WRF-Chem. The resulting uncertainty of LNO_x PE is 1%, including the error of transport and chemistry by shifting pixels.

Compared to the NASA standard product v2, Krotkov et al. (2017) demonstrated that the noise in V_{strat} is 1×10^{14} cm⁻². Errors in polluted regions can be slightly larger than this value, while errors in the cleanest areas are typically significantly smaller (Bucsela et al., 2013). We estimated the uncertainty of the V_{strat} component and the slant column errors to be 10% and 5%, respectively, following Allen et al. (2019).

Based on the standard deviation of the detection efficiency estimation over the CONUS relative to LIS, ENTLN detection efficiency uncertainties are \pm 16% for total and IC flashes and strokes. Due to the high detection efficiency of CG over the CONUS, the uncertainty is estimated to be \pm 5% (Lapierre et al., 2020). It is found that the resulting uncertainty of detection efficiency is 15% in the production analysis. We have used the t_{window} of 2.4 h for counting ENTLN flashes and strokes to analyze LNO₂ and LNO_x production. Because t_{window} derived from the ERA5 reanalysis can not represent the variable wind speeds, a sensitivity test is performed which yields an uncertainty of 10% for production per flash and 8% for production per stroke using t_{window} of 2 and 4h. Meanwhile, the lifetime of UT NO_x ranges from 2 to 12 h depending on the convective location, the methyl peroxy nitrate and alkyl, and multifunctional nitrates (Nault et al., 2017). The lifetime (τ) of NO₂ in Eq. (6) is replaced by 2 and 12 h to determine the uncertainty as 24% due to lifetime. This is comparable with the uncertainty (25%) caused by lightning parameterization for the LNO_x type.

Recent studies revealed that the modeled NO/NO₂ ratio departs from the data in the SEAC⁴RS aircraft campaign (Travis et al., 2016; Silvern et al., 2018). Silvern et al. (2018) attributed this to the positive interference on the NO₂ measurements or errors in the cold-temperature NO–NO₂–O₃ photochemical reaction rate. We assign a 20% bias with $\pm 15\%$ uncertainty to this error considering the possible positive NO₂ measurement interferences (Allen et al., 2019; Bucsela et al., 2019) and estimate the uncertainty to be 15% for LNO_x PE.

In addition, the estimation of LNO_x PE also depends on the tropospheric background NO₂. In our method, the main factors affecting this factor are the emissions inventory and



Figure 12. Average percent differences in (a) LNO_2 , (b) LNO_2Vis , (c) LNO_2Clean , and (d) NO_2Vis with CRF = 100% over MJJA 2014.

the amount of transported NO₂. For the emissions inventory, the sources of uncertainty are assumptions, methods, input data, and calculation errors. As a result, the uncertainties for different species or pollutants related to NO₂ are different and the U.S. EPA also does not publish the quantified uncertainty measures because the parties that submit emissions estimates to the U.S. EPA are not asked to include quantitative uncertainty measurements or estimates (EPA, 2015). For the simulated convective transport, Li et al. (2018) compared the cloud-resolving simulations with these based on convective parameterization and pointed out that the convective transport was weaker in the parameterization. But, we believe that the ratio condition (LNO₂Vis/NO₂Vis > 50 %) should reduce these two kinds of uncertainty, and we assume an uncertainty of 10 %, which is less than the 20 % assigned in Allen et al. (2019) and Bucsela et al. (2019).

The overall uncertainty is estimated as the square root of the sum of the squares of all individual uncertainties in Table 5. The net uncertainty is 48 % and 56 % for the LNO₂ type and LNO_x type, respectively. The mean LNO₂ per flash, LNO_x per flash, LNO₂ per stroke, and LNO_x per stroke based on the linear regression and summation method are 32 mol per flash, 90 mol per flash, 6 mol per stroke, and 17 mol per stroke. Applying the corresponding uncertainty to these mean values, we arrive at 32 ± 15 mol LNO₂ per flash, $90 \pm 50 \text{ mol LNO}_x$ per flash, $6 \pm 3 \text{ mol LNO}_2$ per stroke, and $17 \pm 10 \text{ mol LNO}_x$ per stroke. This is in the range of current literature estimates ranging from 33 to 500 mol LNO_x per flash (Schumann and Huntrieser, 2007; Beirle et al., 2010; Bucsela et al., 2010). Bucsela et al. (2010) estimated LNO_x PE of 100–250 mol per flash, which is higher than but overlaps with our estimate. Pickering et al. (2016) estimated LNO_x PE to be 80 ± 45 mol per flash for the Gulf of Mexico. This is 50 % smaller than our flash-based results over the CONUS, if we use the same linear regression method which is based on the daily summed values instead of daily mean values. Note that the criteria defined in Sect. 3.1 lead to many missing data over the Gulf of Mexico; thus it is actually a comparison between different regions. For the strokebased results, Lapierre et al. (2020) found lower LNO₂ PE of 1.6 ± 0.1 mol per stroke, and the difference is caused by the different version of BEHR algorithm and several settings as mentioned in Sect. 3.2. Bucsela et al. (2019) inferred an average value of $200 \pm 110 \text{ mol} (122 \% \text{ larger than our results})$ of LNO_x produced per flash over the North America, this is related to the different algorithm, lightning data, and lightning thresholds.

5 Conclusions

In this study, a new algorithm for retrieving LNO_2 (LNO_x) from OMI, including LNO₂ (LNO_x) below cloud, has been developed for application over active convection. It works in both clean and polluted regions because of the consideration of tropospheric background pollution in the definition of AMFs. It uses specific criteria combined with several other conditions (sufficient CRF, coincident ENTLN data, TL \geq 1000, and ratio \geq 50 %) to ensure that the electrically active regions are detected by OMI and simulated by WRF-Chem successfully. We conducted an analysis on $1^{\circ} \times 1^{\circ}$ daily boxes in MJJA 2014 and obtained the seasonal mean LNO_2 and LNO_x production efficiencies over the CONUS. Considering all the uncertainties (Table 5) and applying the summation and regression methods, the final mean production efficiencies are estimated to be 32 ± 15 mol LNO₂ per flash, $90\pm50 \text{ mol LNO}_x$ per flash, $6\pm3 \text{ mol LNO}_2$ per stroke, and $17 \pm 10 \text{ mol LNO}_x$ per stroke.

Compared with Lapierre et al. (2020), we find that the LNO₂ production could be larger when the below-cloud LNO₂ is taken into account, especially for the high clouds. Meanwhile, if the method of Pickering et al. (2016) is applied without the background NO₂ correction, the derived LNO_x production efficiency is similar to ours in clean regions or regions with a high LNO₂ concentration above the cloud, but it could be overestimated by more than 18 % in polluted regions. Finally, implementing profiles generated with different model settings of lightning $(1 \times 200 \text{ and } 2 \times 500 \text{ mol NO})$ per flash), we find that the larger LNO production setting leads to 62 % larger retrieval of LNO_x on average despite some regionally dependent effects caused by the nonlinear calculation of AMF. Both the ratio of the tropospheric LNO₂ above the cloud to the total tropospheric LNO₂ and the ratio of LNO₂ to NO₂ cause different comprehensive effects due to the nonlinear calculation of AMF_{LNO₂} and AMF_{LNO_x}.

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Since other regions, like China and India, have much more NO₂ pollution than the CONUS, it is necessary to consider the background NO₂ in detail. These analyses will be complemented by the recently launched satellite instrument (TROPOspheric Monitoring Instrument, TROPOMI) (Veefkind et al., 2012; Boersma et al., 2018; Griffin et al., 2019) and Lightning Mapping Imager (LMI) on the new-generation Chinese geostationary meteorological satellite Fengyun-4 (Min et al., 2017; Yang et al., 2017; Zhang et al., 2019). Future work investigating the flash channel length and more detailed lightning parameterization in WRF-Chem would greatly benefit LNO_x estimation. Applying current methods in future studies may enhance the accuracy of LNO_x production at both local and global scales.

Appendix A: AMF definitions used in this study

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$$AMF_{LNO_2} = \frac{(1 - f_r) \int_{p_{surf}}^{p_{top}} w_{clear}(p) NO_2(p) \, dp}{\int_{p_{surf}}^{p_{top}} LNO_2(p) \, dp}$$
(A1)

$$AMF_{LNO_x} = \frac{(1-f_r) \int_{p_{surf}}^{p_{tp}} w_{clear}(p) NO_2(p) \, dp}{\int_{p_{surf}}^{p_{tp}} w_{cloudy}(p) NO_2(p) \, dp}$$
(A2)

Here f_r is the cloud radiance fraction, p_{surf} is the surface pressure, p_{tp} is the tropopause pressure, p_{cloud} is the cloud optical pressure (CP), w_{clear} and w_{cloudy} are respectively the pressure-dependent scattering weights from the TOMRAD lookup table (Bucsela et al., 2013) for clear and cloudy parts, and NO₂(p) is the modeled NO₂ vertical profile. LNO₂(p) and LNO_x(p) are respectively the LNO₂ and LNO_x vertical profiles calculated by the difference of vertical profiles between WRF-Chem simulations with and without lightning.

$$AMF_{LNO_{2}Clean} = \frac{(1-f_{r})\int_{P_{cloud}}^{P_{tp}} w_{clear}(p)LNO_{2}(p) dp}{\int_{P_{cloud}}^{P_{tp}} w_{cloudy}(p)LNO_{2}(p) dp}$$
(A3)

$$(1-f_r)\int_{p_{surf}}^{p_{tp}} w_{clear}(p) NO_2(p) dp$$

$$AMF_{NO_2Vis} = \frac{+f_r \int_{\rho_{cloud}}^{\rho_{recloud}} w_{cloudy}(\rho)NO_2(\rho) d\rho}{(1-f_g) \int_{\rho_{surf}}^{\rho_{tp}} NO_2(\rho) d\rho + f_g \int_{\rho_{cloud}}^{\rho_{tp}}}$$
(A4)

$$AMF_{NO_xVis} = \frac{\frac{(1-f_r)\int_{p_{surf}}^{p_{top}} w_{clear}(p)NO_2(p) \, dp}{+f_r \int_{p_{cloud}}^{p_{top}} w_{cloudy}(p)NO_2(p) \, dp}}{\frac{(1-f_g)\int_{p_{surf}}^{p_{top}} NO_x(p) \, dp + f_g \int_{p_{cloud}}^{p_{top}}}{NO_r}}$$
(A5)

$$\begin{array}{c} \operatorname{NO}_{X}(p) \, \mathrm{d}p \\ (1-f_{\mathrm{r}}) \int_{P_{\mathrm{surf}}}^{P_{\mathrm{p}}} w_{\mathrm{clear}}(p) \operatorname{NO}_{2}(p) \, \mathrm{d}p \\ + f_{\mathrm{r}} \int_{n_{\mathrm{res}}}^{p_{\mathrm{p}}} w_{\mathrm{cloudy}}(p) \operatorname{NO}_{2}(p) \, \mathrm{d}p \end{array}$$

$$AMF_{LNO_2Vis} = \frac{f_{SJ}p_{cloud} cloudy(p) ero_2(p) e_p}{(1-f_g) \int_{p_{surf}}^{p_{tp}} LNO_2(p) dp} + f_g \int_{p_{cloud}}^{p_{tp}} LNO_2(p) dp$$
(A6)

Here f_g is the geometric cloud fraction and NO_x(*p*) is the modeled NO_x vertical profile.

Table A1. Simple forms of abbreviations for AMFs.

Abbreviations	Numerator ¹	Denominator ²
AMF _{LNO2} AMF _{LNO2} Vis AMF _{LNO2} Clean AMF _{NO2} Vis AMF _{LNOx}	$\begin{array}{c} S_{NO_2} \\ S_{NO_2} \\ S_{LNO_2} \\ S_{NO_2} \\ S_{NO_2} \\ S_{NO_2} \\ S_{NO_2} \end{array}$	VLNO2 VLNO2Vis VLNO2 VNO2Vis VLNOx VLNOX

¹ The part of simulated VCD seen by OMI. ² The simulated VCD.

Appendix B: LNO_x production based on lower lightning thresholds

While we used 2400 flashes per box and 8160 strokes per box per 2.4 h time window for detecting LNO_x , here we show results obtained when using one flash per box and 3.4 strokes per box in the same time window. We note that the WRF total lightning threshold is also reduced to one flash per box, but we keep the ratio condition unchanged. Briefly, the condition is CRF90_ENTLN1(3.4)_TL1_ratio50 as shown in Table 1.

Similarly, the order of estimated daily PEs is LNO₂Clean > LNO₂ > NO₂Vis > LNO₂Vis (Fig. B1). Compared with Fig. 4, the LNO₂ per flash and LNO_x per flash are larger while PEs based on stroke data are smaller. Considering the additional boxes of fewer lightning counts, differences in the daily mean flashes and NO_x result in different PEs, and the relationship presents more like the power function as mentioned in Bucsela et al. (2019).

Instead of using the nonlinear regression of power function

$$y = \alpha x^{\beta}, \tag{B1}$$

where x is flashes or strokes and y is NO_2 or NO_x , we take the logarithm of both sides and apply the linear regression to data:

$$\log_{10} y = \log_{10} \alpha + \beta \log_{10} x.$$
 (B2)

As expected, the linear regression based on logarithmized data performs better in this situation and yields $\alpha = 38$ kmol and $\beta = 0.3$ for LNO_x per flash (Fig. B2). Since we use the unbinned data (flashes not divided into many groups), we compare our results with Bucsela et al. (2019) based on the same kind of data ($\alpha = 10.3$ kmol and $\beta = 0.42$). The large difference of α is related to the method of estimating LNO_x, different lightning data (WWLLN and ENTLN), and different regions (northern midlatitudes and CONUS). Note that the resolution (13 km × 24 km) of OMI could weaken the signal of LNO_x. We believe the phenomenon of higher production efficiency as flash rate decreases (Fig. B3) could be explored in much more detail with higher-resolution data like the TROPOMI data.



Figure B1. Linear regressions with $CRF \ge 90\%$ and a flash threshold of one flash per box or 3.4 strokes per box per 2.4 h. (a) Daily NO₂Vis, LNO₂Vis, LNO₂, and LNO₂Clean versus ENTLN total flash data. (b) Same as (a) but for strokes. (c) Daily LNO_xVis and LNO_x versus total flashes. (d) Same as (c) but for strokes.



Figure B2. Same as Fig. B1 but using log-log axes.



Figure B3. (a) Daily LNO_{*x*} production efficiencies versus ENTLN total flash data, with $CRF \ge 90\%$ and a flash threshold of one flash per box. (b) Same as (a) but for strokes.

Code and data availability. The retrieval algorithm used in Sect. 2.4 is available https://github.com/ at zxdawn/BEHR-LNOx (last access: 20 March 2020; https://doi.org/10.5281/zenodo.3553426, Zhang and Laughner, 2019). The WRF-Chem model output and LNO_x product are available upon request to Xin Zhang (xinzhang1215@gmail.com). MOZART-4 global model output is available at https: //www.acom.ucar.edu/wrf-chem/mozart.shtml (last access: 20 March 2020).

Author contributions. YY directed the research and RvdA, XZ, and YY designed the research with feedback from the other co-authors; RvdA and XZ developed the algorithm; JLL provided guidance and supporting data on the ENTLN data; XZ performed simulations and analysis with the help of YY, RvdA, QC, XK, SY, JC, CH, and RS; YY, RvdA, JLL, and XZ interpreted the data and discussed the results. XZ drafted the manuscript with comments from the coauthors; JLL, RvdA, and YY edited the manuscript.

Competing interests. The authors declare that they have no conflict of interest.

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