Surface and lightning sources of nitrogen oxides over the United States: Magnitudes, chemical evolution, and outflow


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We use observations from two aircraft during the ICARTT campaign over the eastern United States and North Atlantic during summer 2004, interpreted with a global 3-D model of tropospheric chemistry (GEOS-Chem) to test current understanding of regional sources, chemical evolution, and export of NO\(_x\). The boundary layer NO\(_x\) data provide top-down verification of a 50% decrease in power plant and industry NO\(_x\) emissions over the eastern United States between 1999 and 2004. Observed NO\(_x\) concentrations at 8–12 km altitude were 0.55 ± 0.36 ppbv, much larger than in previous U.S. aircraft campaigns (ELCHEM, SUCCESS, SONEX) though consistent with data from the NOXAR program aboard commercial aircraft. We show that regional lightning is the dominant source of this upper tropospheric NO\(_x\) and increases upper tropospheric ozone by 10 ppbv. Simulating ICARTT upper tropospheric NO\(_x\) observations with GEOS-Chem requires a factor of 4 increase in modeled NO\(_x\) yield per flash (to 500 mol/flash). Observed OH concentrations were a factor of 2 lower than can be explained from current photochemical models, for reasons that are unclear. A NO\(_x\)-CO correlation analysis of the fraction \(f\) of North American NO\(_x\) emissions vented to the free troposphere as NO\(_y\) (sum of NO\(_x\) and its oxidation products) shows observed \(f = 16 ± 10\%\) and modeled \(f = 14 ± 9\%\), consistent with previous studies. Export to the lower free troposphere is mostly HNO\(_3\) but at higher altitudes is mostly PAN. The model successfully simulates NO\(_x\) export efficiency and speciation, supporting previous model estimates of a large U.S. anthropogenic contribution to global tropospheric ozone through PAN export.

1. Introduction

[2] Quantifying the sources and fate of nitrogen oxides (NO\(_x\) ≡ NO + NO\(_2\)) over northern midlatitudes continent is critical for assessing anthropogenic influence on global tropospheric ozone [Pickering et al., 1992; Jacob et al., 1993; Thompson et al., 1994; Li et al., 2004]. The International Consortium on Atmospheric Transport and Transformation (ICARTT) aircraft study [Singh et al., 2006; Fehsenfeld et al., 2006], which took place in July–August 2004 over the eastern United States and the North Atlantic, provides an opportunity for this purpose. We present here a global 3-D model analysis of ICARTT observations for NO\(_x\), its chemical reservoirs, and related species including hydrogen oxide (HO\(_x\)) radicals to quantify continental NO\(_x\) sources from combustion and lightning, determine the chemical fate of NO\(_x\) in the United States boundary layer and in North American outflow, and examine the implications for ozone.

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Ozone production in the troposphere is principally limited by the supply of NOx [Chameides et al., 1992]. Fossil fuel combustion accounts for over half of the global NOx source [Intergovernmental Panel on Climate Change, 2001]. The United States has been actively reducing its summertime NOx emissions since 1998 to decrease ozone smog. The Environmental Protection Agency (EPA) NOx State Implementation Plan (SIP) Call mandated that 22 eastern states meet state-specific total NOx emissions reductions by 2003 (Phase 1) and further reduction by 2007 (Phase 2). By 2003, all 22 states had reduced NOx stationary source emissions to their Phase 1 levels. Frost et al. [2006] determined from stack sampling that power plant emissions of NOx decreased 50% between 1999 and 2003. NOx emissions from the power sector over the United States will likely continue to drop. In March 2005, the EPA issued the Clean Air Interstate Rule, which will, when fully implemented, permanently reduce NOx emissions to 60% of 2003 levels in 25 eastern states (http://www.epa.gov/interstateairquality/). The ICARTT observations offer an opportunity to check on these emission reductions.

Oxidation of NOx to HNO3, peroxyacetyl nitrate (PAN), and other minor products takes place on the order of hours in the continental boundary layer (CBL), seemingly limiting its affect on global ozone. However, the dependence of ozone production on NOx is highly nonlinear; the ozone production efficiency per unit NOx consumed (OPE) increases rapidly as the NOx concentration decreases [Liu et al., 1987]. This means that a small fraction of emitted NOx exported to the free troposphere by frontal lifting, deep convection, or boundary layer venting could lead to significant ozone production in the free troposphere over the continent or downwind [Jacob et al., 1993; Thompson et al., 1994]. Similarly, PAN (which is thermally unstable and not water-soluble) can be vented from the boundary layer and transported on a global scale at cold temperatures, eventually decomposing to release NOx as air masses subside. Quantifying the sources, chemical evolution, and export of anthropogenic NOx (and PAN) is thus critical to understanding the North American contribution to the global ozone budget. This contribution is important from the perspective of ozone as a greenhouse gas [Mickley et al., 2004] and for intercontinental transport of ozone pollution [Jacob et al., 1999; Holloway et al., 2003].

Early studies estimated that the fraction f of NOx emitted in the United States that is exported out of the CBL as NOy (sum of NOx and its oxidation products) is about 40% [Logan, 1983; Galloway et al., 1984; Kasibhatla et al., 1993]. Later work by Horowitz et al. [1998] and Liang et al. [1998] using an Eulerian 3-D model with detailed chemistry indicated values of f ranging from 25% in summer to 35% in winter. Subsequent Lagrangian analyses using NOx/CO correlations measured from aircraft in free tropospheric outflow (2–6 km), over the North Atlantic in September (North Atlantic Regional Experiment ’97) yielded much lower estimates of f, ranging from only 3% [Stohl et al., 2002] to 9 ± 14% [Parrish et al., 2004]. Li et al. [2004] reconciled these results by pointing out that the early Eulerian models had insufficient HNO3 scavenging, while the Lagrangian models underestimated background CO. They derived a consistent value f = 17–20% by both Eulerian and Lagrangian approaches for the NARE’97 period. The ICARTT study offers far more geographical coverage and chemical information in the boundary layer and the free troposphere than previous studies, enabling better constraints on the estimates of anthropogenic export and associated NOy speciation.

In addition to convectively lofted anthropogenic NOy, a highly uncertain source of NOy to the upper troposphere is from lightning. Global lightning source estimates range from 1 to 25 Tg N yr−1 [Price et al., 1997; Nesbitt et al., 2000], with the most recent estimates in the range 1–6 Tg N yr−1 [Boersma et al., 2005; Beirle et al., 2006; Martin et al., 2007]. Past studies disagree on the relative importance of lightning versus convective injection of surface emissions in supplying upper tropospheric NOy [Jaeglé et al., 1998; Levy et al., 1999; Grewe et al., 1999; Jeker et al., 2000; Li et al., 2005]. Here we use the ICARTT data to examine the contribution of each.

2. ICARTT Study

ICARTT took place over eastern North America and the North Atlantic in July–August 2004. A major objective was to quantify North American sources and outflow of pollutants and climatically important species. Two principal components directed at that objective were the NOAA New England Air Quality Study/Intercontinental Transport and Chemical Transformation (NEAQS-ITCT 2004) and the NASA Intercontinental Transport Experiment–North America, Phase A (INTEX-A).

The NOAA NEAQS-ITCT 2004 campaign [Fehsenfeld et al., 2006] took place 3 July to 15 August over the NW Atlantic and the NE United States out of Portsmouth, New Hampshire (Figure 1). It used a WP-3D aircraft (ceiling ~6 km). The NASA INTEX-A campaign [Singh et al., 2006] took place 29 June to 14 August over the central and eastern United States, and the North Atlantic, from bases at Edwards (California), St. Louis (Missouri), and Portsmouth (New Hampshire). It used a DC-8 aircraft (ceiling ~12 km) with extensive vertical profiling and boundary layer mapping at 0.15–0.5 km above the local surface (Figure 1). The WP-3D remained close to New England throughout the mission and flew mostly in the boundary layer, focusing on emissions verification and chemical transformation of major urban pollution plumes. Detailed descriptions of aircraft payloads are in Fehsenfeld et al. [2006] and Singh et al. [2006].

We make use here of the following 1-min average measurements (accuracies given in parentheses for DC-8, WP-3D respectively): ozone (5%,3%); CO (5%,2.5%); CH2O (10%, DC-8 NCAR measurement only); HNO3 (15%, DC-8 only); H2O2 (20%, DC-8 only); NO (15%, 5%, DC-8 Pennsylvania State U. measurement only); NO2 (10%,8%); HNO3 (15%, 15%); PAN(15%,15%), OH(15%, DC-8 only), and HO2 (15%, DC-8 only). We estimate the concentration of NOx as the sum of NOx, PAN, and HNO3. Nitrate aerosol, organic nitrates other than PAN, and HNO4 were also measured aboard the DC-8 [Dibb et al., 2003; Huey et al., 2004; Bertram et al., 2006], but generally accounted for less than 10% of NOx in the free troposphere and the data sets were sparse.

Ventilation of the eastern United States in summer is primarily driven by the cold fronts from cyclones tracking...
eastward typically every 5 days in the 45–55°N band [Li et al., 2005]. Local convection from fair weather cumuli as well as thunderstorms is another ventilation mechanism [Parrish et al., 2004; Li et al., 2005]. During ICARTT a persistent trough along the east coast led to cyclones extending farther south than normal [Fuelberg et al., 2007], and lack of stagnation events led to a record low number of air quality violations (A. M. Thompson et al., Tropospheric ozone in eastern North America in July–August 2004: Profile views from “the summer that wasn’t,” manuscript in preparation, 2007).

3. Model Description

[11] We simulate the ICARTT observations with the GEOS-Chem global 3-D model of tropospheric chemistry (version 7.02; http://www.as.harvard.edu/chemistry/trop/geo/) driven by assimilated meteorological observations from the Goddard Earth Observing System (GEOS-4) of the NASA Global Modeling and Assimilation Office (GMAO). The model is applied to a global simulation of ozone-NO$_x$-hydrocarbon-aerosol chemistry with 120 species simulated explicitly. A general description of GEOS-Chem is given by Bey et al. [2001] and a description of the coupled oxidant-aerosol simulation as used here is given by Park et al. [2004]. Partitioning of total nitric acid between the gas and aerosol phases is calculated using the MARS-A thermodynamic equilibrium model [Binkowski and Roselle, 2003]. Emissions in the model are as described by Park et al. [2004] unless specified otherwise.

[12] Meteorological fields in the GEOS-4 data have a temporal resolution of 6 hours (3 hours for surface variables and mixing depths) and a horizontal resolution of 1° latitude by 1.25° longitude, with 55 vertical levels between the surface and 0.1 hPa (including about 16 in the troposphere and 5 in the boundary layer up to 2 km). For input to GEOS-Chem we degrade the horizontal resolution to 2° latitude by 2.5° longitude over the eastern United States in the GEOS-4 data for the ICARTT period were 1100 ± 400 m AGL. Tropopause heights were 13.3 ± 2.2 km. The cross-tropopause ozone flux is specified with the Synoz method [McLinden et al., 2000] while the cross-tropopause NO$_x$ flux is calculated from N$_2$O oxidation in the model stratosphere [Bey et al., 2001]. Global net cross-tropopause fluxes of ozone and NO$_x$ are 495 Tg O$_3$ yr$^{-1}$ and 0.5 Tg N yr$^{-1}$, respectively. The model wet deposition scheme [Liu et al., 2001] includes contributions from scavenging in convective updrafts, and rainout and washout from convective anvils and large-scale precipitation, and it allows for return to the atmosphere following evaporation. Soluble gases are taken up by liquid water on the basis of their effective Henry’s law, by ice on the basis of cocondensation or surface coverage, with species-specific retention efficiencies when droplets freeze, as described by Mari et al. [2000].

[13] The simulations are conducted for July–August 2004 and are initialized with a 12-month spin-up simulation. Comparison with observations uses model output sampled for the aircraft flight tracks and times. The comparisons exclude fresh pollution plumes as diagnosed by NO$_x$/NO$_y$ > 0.4 mol mol$^{-1}$ or (if NO$_x$ is not available) NO$_y$ > 4 ppbv and altitude <3 km; biomass burning plumes as diagnosed by HCN > 500 pptv or CH$_3$CN > 225 pptv or flight logs; and stratospheric air as diagnosed by ozone/CO > 1.25 mol mol$^{-1}$. These filters exclude 8%, 22% (fresh pollution plumes); 5%, 8% (biomass burning plumes); and 6%, 0% (stratospheric air) of the data for the DC-8 and WP-3D, respectively.

[14] A major focus of our work is to use the ICARTT observations to evaluate U.S. NO$_x$ emission estimates. We will show simulations with “standard” emissions based on a priori information from the standard version 7.02 of the GEOS-Chem model, and “improved” emissions that reflect the constraints from the ICARTT observations. These emissions for the ICARTT period (1 July to 15 August 2004) are summarized in Table 1. Standard fossil and biofuel emis-

Figure 1. Flight tracks for (left) INTEX-NA and (right) ITCT 2k4 aircraft campaigns (1 July to 15 August 2004) shaded by altitude above ground level (AGL). Shaded areas define regions used in comparisons between model and observations: south (medium grey), northeast (light grey), and Midwest (dark grey).
ions in the United States are from the EPA 1999 National Emission Inventory (NEI99). They amount to 0.79 Tg N for 1 July to 15 August and 6.2 Tg N annually, with distribution shown in Figure 2 (left). Transportation accounts for 35%, industry 17%, power generation 26%, and other sources 22% (mostly off-road vehicles). We show in section 5 that, this 1999 inventory overestimates the boundary layer NO$_x$ concentrations observed in ICARTT, consistent with 1999–2004 reduction of NO$_x$ emissions from power plants and industry by 50% driven by the NO$_x$ SIP Call [Frost et al., 2006]. These reductions result in the improved inventory of Table 1 and Figure 2 (right). The ICARTT data also show that U.S. anthropogenic CO emissions in the NEI99 inventory (94 Tg CO yr$^{-1}$) are at least 30% too high, as discussed below, and we make this adjustment in the improved inventory.

[15] The lightning source of NO$_x$ in GEOS-Chem is computed locally in deep convection events with the scheme of Price and Rind [1992] that relates number of flashes to convective cloud top heights, and the vertical distribution from Pickering et al. [1998] (55–75% above 8 km). Implementation in GEOS-Chem is as described by Wang et al. [1998]. It yields $2.7 \times 10^6$ flashes yr$^{-1}$ in the GEOS-4 fields for 2004. The standard GEOS-Chem simulation assumes a global source of 4.7 Tg N yr$^{-1}$ to match observational constraints in the tropics [Martin et al., 2002], which results in a global mean NO yield of 125 mol per flash averaged over intracloud (IC) and cloud-to-ground (CG) flashes (see Wang et al. [1998] for the model partitioning of flashes between IC and CG, and the NO yield difference between these two types of flashes). The resulting U.S. emissions for the ICARTT period are 0.067 Tg N for the contiguous United States and coastal waters (130–70$^\circ$W, 25–50$^\circ$N), concentrated over the southwest and along the Gulf of Mexico. As we will see in section 6, successful simulation of the ICARTT upper tropospheric NO$_x$ observations in GEOS-Chem requires a factor of 4 increase in this source along with its upward extension to the tropopause, and this is included in the improved inventory.

[16] Extensive and persistent boreal forest fires took place in Alaska and NW Canada in summer 2004; in contrast, there were no significant fires in Siberia. We use the daily biomass burning inventory of Turquety et al. [2007] for North American fires during ICARTT. This inventory was constructed by combining daily area burned reports from government agencies and hot spots detected from space by the MODIS instrument with estimates of fuel loadings and emission factors depending on the type of ecosystem burned. The resulting North American fire emissions for 1 July to 15 August are 20 Tg CO and 0.3 Tg N of NO$_x$. The inventory for CO has been successfully evaluated against MOPITT columns [Turquety et al., 2007]. Short-lived hydrocarbons emitted from fires drive fast conversion

### Table 1. NO$_x$ Emissions in the Contiguous United States for 1 July to 15 August 2004$^a$

<table>
<thead>
<tr>
<th>Source Type</th>
<th>Standard Magnitude, Tg N</th>
<th>Improved Magnitude, Tg N</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fuel</td>
<td>0.79</td>
<td>0.62$^a$</td>
</tr>
<tr>
<td>Biomass burning$^a$</td>
<td>0.01</td>
<td>0.006</td>
</tr>
<tr>
<td>Fertilizer</td>
<td>0.03</td>
<td>0.03</td>
</tr>
<tr>
<td>Soils</td>
<td>0.09</td>
<td>0.09</td>
</tr>
<tr>
<td>Aircraft</td>
<td>0.021</td>
<td>0.021</td>
</tr>
<tr>
<td>Lightning$^d$</td>
<td>0.068</td>
<td>0.27</td>
</tr>
<tr>
<td>Total</td>
<td>1.01</td>
<td>1.03</td>
</tr>
</tbody>
</table>

$^a$The “standard” emissions are from the standard version 7.02 of the GEOS-Chem model. The “improved” emissions reflect the constraints from the ICARTT observations.

$^b$Improved fuel emissions include 0.28 Tg N from transport, 0.17 Tg N from power generation and industry, and 0.17 Tg N from other fuel use including industrial solvents, aircraft takeoff and landing, residential fossil fuel, residential biofuel, and all other anthropogenic sources.

$^c$North American fires during ICARTT were mainly outside the contiguous United States; Alaska and Canada each contributed 0.14 Tg N.

$^d$Contiguous United States and coastal waters (130–70$^\circ$W, 25–50$^\circ$N).
of NOx to PAN, slowing down ozone formation [Jacob et al., 1992] but this is inadequately represented in GEOS-Chem where the fastest-reacting pyrogenic hydrocarbon is propene. In the improved emission inventory we release 80% of the biomass burning NOx as PAN.

4. Hydrogen Oxide Radicals and Reservoir Species

[17] Hydrogen oxide radicals (HOx \equiv OH + peroxy radicals) and their reservoirs (H2O2, CH2O, HNO4) largely determine the photochemical environment for NOx oxidation and ozone production. Figure 3 shows simulated and observed mean vertical distributions of OH, HO2, H2O2, CH2O, and HNO4 for the ensemble of INTEX-A flights. Observed HO2 and H2O2 decrease with altitude, reflecting the decrease in water vapor [Heikes, 1992], while observed OH increases with altitude, reflecting the positive relationship of the OH/HO2 ratio with NO [Jaegle et al., 2000]. CH2O decreases with altitude, largely reflecting the boundary layer source from oxidation of isoprene [Millet et al., 2006]. HNO4, a thermally unstable molecule that is an important HOx reservoir in the upper troposphere [Jaegle et al., 2000], has an 8–9 km peak of 62 pptv.

[18] OH and HO2 in the standard simulation match observations in the lowest km, but overestimate at higher altitudes by 60% and 30% respectively. Nominal measurement accuracy for both is 15% (X. Ren et al., HOx observation and model comparison during INTEX-NA 2004, unpublished manuscript, 2007, hereinafter referred to as Ren et al., unpublished manuscript, 2007). Increasing the lightning NOx source in the improved simulation decreases HO2 in the troposphere while increasing OH. This improves the HO2 simulation but worsens that of OH. Similar HOx simulation biases in comparison with these observations have been found in a box photochemical model constrained with the ensemble of concurrent ICARTT observations [Olson et al., 2005; Ren et al., unpublished manuscript, 2007]. They must reflect either instrument error or a fundamental flaw in current understanding of HOx photochemistry. In what follows we take the model OH at face value but we also discuss the effect of possible model bias.

[19] The model simulates in general well the concentrations of the HOx reservoirs H2O2, CH2O, and HNO4 (Figure 3). H2O2 is unbiased in the free troposphere, but is overestimated below 3 km by \sim 30%. A possible cause is the model conversion of HO2 to H2O2 in aerosols [Martin et al., 2003], which recent measurements find to be much lower than previously estimated [Thornton and Abbatt, 2005]. The CH2O simulation, discussed in detail by Millet et al. [2006], matches observations closely and is largely insensitive to changes in OH (which affect sources and sinks in the same direction). HNO4 is well matched in the standard simulation but is overestimated by 30% when the lightning source is increased.

5. Boundary Layer NOx and CO Over the United States

[20] Figure 4 compares simulated and observed mean vertical distributions of CO, NOx, PAN, HNO3, and ozone concentrations for the ensemble of DC-8 and WP-3D flights. Observed CO, NOx, PAN, and HNO3 are elevated below 3 km, reflecting anthropogenic sources (and also biogenic sources for CO). The model with standard emis-

Figure 3. Mean vertical profiles of OH, HO2, H2O2, HCHO and HNO4 concentrations. Observations from the DC-8 aircraft (thick black) are compared to model results from the standard (dashed red) and improved (thick red) simulations. Horizontal bars are standard deviations on the observations. Here and in subsequent figures, the ICARTT observations have been filtered to remove urban plumes, biomass burning plumes, and stratospheric air as described in the text. Model results are sampled along the flight tracks at the time of flights and gridded to the model resolution. Number of gridded observations per 1-km bin is shown. Modifications to emissions from the standard to the improved simulation include a four-fold increase in the lightning source, upward extension of the lightning source to the tropopause height, and a 50% reduction of the NEI99 CO transport and NOx point source emissions.
emissions is too high for NO\textsubscript{x} and CO. Parrish [2006], using urban ambient measurements of the CO/NO\textsubscript{x} ratio along with fuel sales data, previously found that the on-road vehicular source in the NEI99 inventory is accurate for NO\textsubscript{x}, but 50\% too high for CO. Applying this correction to the on-road vehicular CO source, which represents 60\% of total CO emissions in the United States according to NEI99, decreases the mean CO bias below 1 km from 35 ppbv to 14 ppbv in the WP-3D data and from 26 to 13 ppbv for the DC-8 data (Figure 4). Further source reduction would be needed to match the boundary layer observations for CO.

Figure 5 shows the geographical distribution of median ratios of simulated to observed NO\textsubscript{x} concentrations in the lowest 2 km for the standard simulation. The largest overestimate is in the Midwest (median ratio of 1.88), where the contribution from power plants to the NO\textsubscript{x} source is largest (Figure 2). The improved simulation with 50\% decrease in point source NO\textsubscript{x} emissions decreases the median ratio to 1.28 in the Midwest. We also find an overestimate over the northeast in the standard simulation, though significantly weaker (median ratio of 1.26). In contrast, Martin et al. [2006] find the NEI99 inventory to be underestimated in the New York City metropolitan area when comparing GEOS-Chem standard model results with mean WP-3D profiles and SCIAMACHY NO\textsubscript{2} satellite observations over and downwind of the city. This difference in conclusions appears to reflect the use of median versus mean NO\textsubscript{x} concentrations as comparison metrics, and the exclusion of fresh pollution plumes in our analysis. Assessment of NEI99 emission bias is ambiguous in the northeast but it is robust in the Midwest.

6. Upper Tropospheric NO\textsubscript{x} Over the United States

[22] NO\textsubscript{x} concentrations observed in ICARTT show a decrease from the boundary layer to the free troposphere, but then a sharp rise with altitude above 6 km (Figure 4). Mean concentrations reach 0.55 ± 0.36 ppbv at 8–12 km altitude, higher than in the boundary layer. The NO/NO\textsubscript{x} molar ratio averages 75\% both in the observations and the model for the 8–12 km region (all data are for daytime). Observed PAN shows a broad maximum at 6–10 km. In contrast, HNO\textsubscript{3} is depleted in the free troposphere because of scavenging during uplift. Mean ozone increases with altitude from 50 ppbv near the surface to 75 ppbv at 8 km. [23] The standard model greatly underestimates the upper tropospheric NO\textsubscript{x} enhancement; the discrepancy increases with altitude from a factor of 3 at 8 km to a factor of 5 at 11 km. Simulated PAN is too low by ~30\% while HNO\textsubscript{3} is well simulated. Ozone is too low by 10 ppbv throughout the free troposphere.

[24] Measurements of upper tropospheric NO\textsubscript{x} from previous aircraft campaigns over the United States indicate lower concentrations than observed in ICARTT. Jaeglé et al. [1998] report mean NO concentrations from the SUCCESS campaign out of Kansas (April–May 1996) of 0.030 ± 0.022 ppbv for 8–10 km and 0.061 ± 0.045 ppbv for 10–12 km. Ridley et al. [1994] report mean NO
concentrations of 0.2 ± 0.1 ppbv over New Mexico during ELCHEM (July–August 1989), even though convection was frequently targeted. SONEX observations in October–November 1997 over Maine and Atlantic Canada indicate mean NO concentrations between 6–12 km of 0.1 ppbv (all data) and 0.23 ppbv (convective outflow) [Crawford et al., 2000]. On the other hand, measurements from commercial aircraft by Brunner et al. [2001] over the east coast of the United States in 1995–1997 (NOXAR program) showed mean upper tropospheric NO concentrations of 0.60 ppbv in spring, 0.41 ppbv in summer, and 0.23 ppbv in fall and winter, consistent with the observations here. Brunner et al. [2001] attributed the high spring-summer values to a combination of lightning and convective injection of pollution [also see Jeker et al., 2000]. As we show below, lightning was the dominant factor in ICARTT.

Figure 6 (left) shows the observed mean spatial distribution of upper tropospheric NO concentrations. Values exceeding 1 ppbv extend over much of the southeast and Midwest. Deep convective injection of boundary layer pollution cannot explain these high values since the NO mixing ratio above 8 km is greater than that in the boundary layer (Figure 4). Aircraft emissions cannot provide an explanation either because the geographical distribution does not match the aircraft corridor along the eastern United States and North Atlantic [Gauss et al., 2006], and in any case these emissions are fairly well constrained from atmospheric measurements [Meijer et al., 2000] and repre-

Figures 5. Median simulated-to-observed NO\textsubscript{x} concentrations in the lowest 2 km using the NEI99 v1 inventory.

![Figure 5](image_url)

Figure 6. Mean upper tropospheric NO\textsubscript{x} concentrations (8–12 km) during ICARTT (July–August 2004). (left) Observations mapped on the 2° × 2.5° GEOS-Chem model grid are compared to (right) model values for the improved simulation. The improved simulation has a factor of four increase in the United States emissions relative to the standard GEOS-Chem version.
sent only a small source of upper tropospheric NO\textsubscript{x} (Table 1). Brunner et al. [2001] previously argued that an aircraft source for upper tropospheric NO\textsubscript{x} would be inconsistent with their observed spring/summer maximum of NO\textsubscript{x} concentrations.

Lightning provides the best explanation for the elevated NO\textsubscript{x} in the upper troposphere during ICARTT. Figure 7 (left) shows National Lightning Detection Network (NLDN) mean lightning flash rates for 1 July to 15 August. The NLDN data (>100 sites in the continental United States) were collected by Vaisala (http://www.vaisala.com) and supplied to us by the Global Hydrology Resource Center at NASA Marshall Space Flight Center. The NLDN network measures only cloud-to-ground lightning flashes, and intracloud flashes are estimated to be about 3 times that amount [Boccippio et al., 2001]. Cloud-to-ground flash detection efficiencies are >90% over the continental United States and degrade rapidly offshore and beyond U.S. borders. Figure 7 shows heaviest lightning in the Gulf of Mexico region and secondary maxima in the Midwest. We see substantial coincidence in Figures 6 and 7 between the observed geographical distribution of lightning and that of upper tropospheric NO\textsubscript{x}.

2004 was not an anomalous year for lightning over the United States, as shown in Figure 8 with total NLDN lightning flash counts for July–August 2000–2005. There is a jump in lightning flashes from 2001 to 2002 which reflects an upgrade in detection. The comparable years in the observations are thus 2003–2005. GEOS-Chem lightning counts computed using consistent GEOS-4 meteorology for 2000–2005 also show relatively little interannual variability in total lightning over the contiguous United States. We find in the model that lightning flash rates over the eastern United States in 2004 are everywhere within 20% of the 1995–2005 mean.

We can make a rough estimate of lightning emissions over the United States during ICARTT by using NLDN flash rates, multiplied by 4 to account for intracloud flashes [Boccippio et al., 2001], and assuming a 500 mol NO\textsubscript{x} per flash production rate derived from the mean peak NLDN flash production rate derived from the mean peak NLDN current (L. E. Ott et al., Production of lightning NO\textsubscript{x} and its vertical distribution calculated from 3-D cloud-scale transport model simulations, manuscript in preparation, 2007, hereinafter referred to as Ott et al., manuscript in preparation, 2007). This yields an emission of 0.45 Tg N, a factor of 7 above the standard GEOS-Chem simulation (Table 1). The model captures the maximum along the Gulf Coast but is too low offshore and over the Midwest (Figure 7). We tried to improve the model distribution of lightning with alternate lightning parameterizations based on cloud mass flux or convective precipitation [Allen and Pickering, 2002], but the cloud mass flux parameterization did not capture the Gulf maximum while the convective precipitation scheme did not capture the lightning distribution over land.

The global lightning source of NO\textsubscript{x} in the standard GEOS-Chem simulation is 4.7 Tg N yr\textsuperscript{-1} from 2.7 × 10\textsuperscript{9} flashes, which corresponds to a global mean average of 125 mol per flash, a factor of 4 below the Ott et al. (manuscript in preparation, 2007) estimate. Correction to the Ott et al. (manuscript in preparation, 2007) NO\textsubscript{x} yield would imply a factor of 4 increase in the GEOS-Chem lightning source over the United States. Figure 6 (right) shows the resulting mean 8–12 km NO\textsubscript{x} concentrations, successful over the south where lightning flashes are correctly simulated, but still showing discrepancies in the Midwest due to insufficient lightning. Generalization of the Ott et al. (manuscript in preparation, 2007) yield to the northern midlatitudes band (1.5 Tg N) yields a GEOS-Chem lightning source of 5.8 Tg N yr\textsuperscript{-1}, consistent with Martin et al. [2006] who found that a northern midlatitudes lightning source of 1.6 Tg N yr\textsuperscript{-1} provides the best fit to SCIAMACHY NO\textsubscript{2} satellite observations.

On a global scale though, a NO\textsubscript{x} yield of 500 mol per flash would appear to lead to an excessive lightning source. The OTD-LIS v1.0 gridded satellite lightning climatology produced by the NASA LIS/OTD Science Team (Principal Investigator, Hugh J. Christian, NASA/Marshall Space Science Center at NASA Marshall Space Flight Center) reduced the LIS/OTD Science Team (Principal Investigator, Hugh J. Christian, NASA/Marshall Space Flight Center) provided the best fit to SCIAMACHY NO\textsubscript{2} satellite observations.
Flight Center), available from the Global Hydrology Resource Center (http://ghrc.msfc.nasa.gov), yields $1.5 \times 10^{9}$/yr. Combining a NO$_x$ yield of 500 mol/flash with this global estimate would imply a lightning source of 10.5 Tg N yr$^{-1}$, which seems too high on the basis of constraints from satellite observations [Boersma et al., 2005; Martin et al., 2007; Sauvage et al., 2007] and tropical ozonesondes [Martin et al., 2002]. While physical mechanisms responsible are not well understood, recent observational evidence suggests that NO$_x$ yields per flash are lower in tropical than in midlatitude storms [Huntrieser et al., 2006].

The lifetime of NO$_x$ in the upper troposphere is a major uncertainty in scaling the lightning source to match the ICARTT observations. As shown in section 4, the model OH concentration in the upper troposphere is a factor of 2 higher than observed, for reasons that are unclear. Figure 9 shows simulated and observed frequency distributions of NO$_x$ concentrations at 8–12 km on the $2^\circ \times 2.5^\circ$ model grid. If the model lifetime were too short because of excessive OH, then one might expect the variability in the model to be larger than observed. Figure 9 shows that this is not the case, adding some support to the simulated NO$_x$ lifetime.

The large lightning source inferred from the ICARTT observations has important implications for tropospheric ozone. Li et al. [2005] found that a semipermanent upper level anticyclone above the southern United States in summer allows ozone buildup in the upper troposphere by trapping convectively lifted precursors and lightning NO$_x$. Cooper et al. [2006] confirmed the resulting ozone maximum by analysis of ozonesonde data during ICARTT, and found an associated 11–14 ppbv contribution to ozone from lightning. Figure 4 shows that the improved simulation with increased lightning largely corrects the upper tropospheric ozone bias in the standard simulation (reduction in surface anthropogenic NO$_x$ emissions in the improved simulation decreases upper tropospheric ozone by only 1–2 ppbv). The residual bias appears due to insufficient lighting generation

![Cloud-to-Ground Lightning Flashes (July-August)](image)

**Figure 8.** Cloud-to-ground lightning flash counts in July–August 2000–2005 over the United States. National Lightning Detection Network flash counts (grey bars) are compared to modeled flash counts in GEOS-Chem derived using GEOS-4 meteorology for the domain 130–70°W, 25–50°N (grey and white bars). Model flash counts are divided uniformly by a factor of 4 to account for intracloud lightning [Boccippio et al., 2001]. The jump in NLDN data from 2001 to 2002 reflects an upgrade in detection.

![DC-8 NO$_x$ 8 - 12 km](image)

**Figure 9.** Frequency distribution of NO$_x$ concentrations at 8–12 km altitude during ICARTT. NO$_x$ observations (line) are compared to results from the improved GEOS-Chem simulation with increased lightning source (grey bars). The frequency distribution is shown as the number of occurrences along the aircraft flight tracks averaged on the $2^\circ \times 2.5^\circ$ model grid.
in the Midwest (Figure 7). We find in the model that lightning enhances upper tropospheric ozone by about 10 ppbv, consistent with the analysis of Cooper et al. [2006].

7. Chemical Evolution and Export of U.S. NO\textsubscript{x} Emissions

[33] In this section we use the ICARTT data to estimate the export of anthropogenic NO\textsubscript{x} from the U.S. boundary layer to the free troposphere and the speciation of this NO\textsubscript{x}. Figure 10 shows the simulated and observed NO\textsubscript{x} speciation below 2 km. Here and from now on, model results are from the improved simulation with reduced fuel NO\textsubscript{x} emissions and increased lightning (Table 1), and spatial patterns are discussed in the context of the regions of Figure 1. The median observed NO\textsubscript{x}/NO\textsubscript{y} ratio at 0–2 km is 18% in the northeast/Midwest, 17% in the south, and 7% offshore. The dominant component of NO\textsubscript{y} in all three regions is HNO\textsubscript{3}, with medians of 62% in the northeast/Midwest and 84% offshore. The mean PAN/NO\textsubscript{x} ratio is 1.0 in the northeast/Midwest and 1.5 in the south. PAN is favored in the south, despite high temperatures, likely because of large isoprene emission [Horowitz et al., 1998]. The model is remarkably successful at reproducing these fractions and patterns.

[34] We can estimate the export efficiency $f$ of NO\textsubscript{x} from the North American boundary layer, following the approach
of Parrish et al. [2004], by viewing CO as an inert tracer and comparing the enhancement ratio $\Delta NO_y/\Delta CO$ in North American pollution outflow to the anthropogenic CO/NO$_x$ molar emission ratio $R$:

$$f = R \cdot \alpha \frac{\Delta NO_y}{\Delta CO}$$

(1)

where $\alpha$ is a correction factor to account for the CO source from boundary layer oxidation of nonmethane hydrocarbons, particularly biogenic isoprene [Chin et al., 1994]. Derivation of the export efficiency following (1) requires estimates of $R$, $\alpha$, and the background NO$_y$ and CO concentrations to which the enhancements $\Delta$ are referenced. By adopting constant values for these variables, as discussed by Li et al. [2004], we obtain a simple observationally based diagnostic of export. We use background concentrations of 95 ppbv CO [Li et al., 2004] and 100 pptv NO$_x$ [Parrish et al., 2004], $\alpha = 1.2$ [Chin et al., 1994], and $R = 5.9$ mol mol$^{-1}$ from our improved GEOS-Chem simulation for the United States east of 100°W. $R$ varies little by region between the Midwest (5.6), south (5.9), and northeast (6.1).

[35] We apply equation (1) to every anthropogenic pollution plume observed by the DC-8 and WP-3D between 2.5 and 6.5 km as defined by a CO enhancement $\Delta CO > 30$ ppbv. Plumes above 6.5 km are ignored because of the low NO$_y$ interference, and we also exclude biomass burning plumes diagnosed from nitrate data (section 2). Anthropogenic pollution plumes defined in this manner represent 11% of the combined INTEX-A and ITCT2k4 data at 2.5–6.5 km. From these data we find a mean NO$_y$ export efficiency $f = 16 \pm 10\%$ to the free troposphere with mean composition of 13% NO$_x$, 40% PAN, and 47% HNO$_3$ for the exported NO$_y$. Sampling the model along the ICARTT flight tracks shows a similar mean value and variability, i.e., $f = 14 \pm 9\%$ with mean composition of 9% NO$_x$, 42% PAN, and 49% HNO$_3$. Previous studies using aircraft data for North American outflow in NARE’97 [Li et al., 2004; Parrish et al., 2004] and Asian outflow in TRACE-P [Koike et al., 2003; Miyazaki et al., 2003] similarly found $f$ values in the range 10–20%.

[36] Segregation of the outflow observations by altitude shows that $f$ is highest and most variable at 2.5–3.5 km (18 ± 11%), where HNO$_3$ dominates the NO$_y$ export fraction (54% on average). The model in that altitude range shows a similar value of $f$ (15 ± 10%) and HNO$_3$ fraction (55%). Parrish et al. [2004] proposed that this shallow venting is due to fair weather cumulus breaking through the afternoon boundary layer. In GEOS-Chem, scavenging of soluble species in a wet convective updraft takes place with an e-folding vertical length scale of 2 km [Liu et al., 2001], allowing significant escape from shallow convection to the lower free troposphere. At higher altitudes we find that PAN becomes the principal component of exported NO$_y$ above 4 km, both in the observations and in the model, reflecting the scavenging of HNO$_3$. The export efficiency $f$ decreases with altitude in the model and observations reflecting HNO$_3$ scavenging during uplift, except between 5.5–6.5 km in the model, where lightning may begin to affect the calculation. The observed mean $f$ is 12 ± 6% at 3.5–6.5 km with speciation of 14% NO, 41% PAN, and 45% HNO$_3$. The corresponding mean $f$ in the model is 13 ± 7% with speciation of 9% NO, 48% PAN, and 43% HNO$_3$. This dominance of PAN in free tropospheric continental outflow of NO$_y$ has been previously observed in aircraft campaigns downwind of North America [Parrish et al., 2004] and Asia [Miyazaki et al., 2003]. It leads to efficient ozone production far downwind of the continent when the PAN subsides and decomposes back to NO$_x$ [Hudman et al., 2004; Li et al., 2004].

8. Conclusions

[37] The ICARTT study in summer 2004 provided extensive observations of reactive nitrogen (NO$_y$) species over the eastern United States and western North Atlantic, from the surface to 12 km altitude. We interpreted these observations with a global 3-D model of tropospheric chemistry (GEOS-Chem) to place constraints on the sources, chemical evolution, and export of NO$_y$ from North America.

[38] ICARTT observations in the continental boundary layer provide top-down verification of the recent decrease in stationary NO$_x$ emissions in the eastern United States mandated by the NO$_x$ SIP Call. Model simulation of NO$_y$ in ICARTT indicates that the latest comprehensive national emission inventory done for 1999 (NEI 99) is too high over the Midwest by almost a factor of 2. This is consistent with the 50% reduction in stationary sources from 1999 to 2004 inferred from power plant smokestack monitoring [Frost et al., 2006] and amounts to a 22% decrease in U.S. anthropogenic NO$_y$ emissions, to 0.62 Tg N for the 1 July to 15 August ICARTT period. GEOS-Chem emissions of anthropogenic NO$_y$ during that same period were 0.98 Tg N for east Asia and 0.53 Tg N for Europe.

[39] Observed NO$_x$ concentrations in ICARTT show a sharp rise above 6 km with mean concentrations reaching 0.55 ± 0.36 ppbv at 8–12 km. These values are higher than observed in the U.S. boundary layer and much higher than observed in the upper troposphere on previous U.S. aircraft campaigns (SONEX, SUCCESS, ELCHEM), though consistent with previous NOXAR measurements from commercial aircraft [Brunner et al., 2001]. A close correspondence is observed between the spatial distribution of upper tropospheric NO$_y$ during ICARTT and the lightning flash frequency observed by the National Lightning Detection Network (NLDN), thus identifying lightning as the dominant source for the observed NO$_x$. The NLDN data also show that 2004 was not an anomalous year for lightning over the United States.

[40] Using NLDN flash rates for 1 July to 15 August, scaled up by a factor of 4 to account for intracloud flashes [Boccioppio et al., 2001], and assuming a 500 mol NO$_y$/flash production rate following Ott et al. (manuscript in preparation, 2007), we deduce a lightning NO$_y$ emission of 0.45 Tg N for that period over the United States and adjacent coastal areas. This is a factor of 7 higher than in the standard GEOS-Chem simulation, which uses a yield of 125 mol NO$_y$/flash for a global lightning NO$_y$ source of 4.7 Tg N yr$^{-1}$. NO$_y$

[41] We can reproduce the upper tropospheric NO$_y$ observed over the southern United States in ICARTT by increasing the lightning NO$_y$ yield in the model by a factor of four to the Ott et al. (manuscript in preparation, 2007) value. This also provides a successful simulation of the
observed frequency distribution of upper tropospheric NO$_x$, and corrects a 5–10 ppbv low bias in the simulation of ozone in the free troposphere in the model. The factor of four increase is probably not extrapolatable globally as the resulting global lightning source in the model (19 Tg N yr$^{-1}$) would be too high relative to observational constraints [Boersma et al., 2005; Martin et al., 2007]. Recent observations suggest that midlatitude storms have higher NOx yields than tropical storms [Huntrieser et al., 2006].

[42] Uncertainty in OH concentrations in the upper troposphere remains an important issue for interpreting quantitatively the observed NO$_x$ concentrations in terms of an implied lightning NO$_x$ source. Simulated upper tropospheric OH concentrations in GEOS-Chem are about a factor of 2 higher than observed in ICARTT, and the same bias is found in box photochemical model calculations constrained with the aircraft observations [Olson et al., 2005; Ren et al., unpublished manuscript, 2007]. As discussed by Olson et al. [2005] and also by Spivakovskyy et al. [2000], this discrepancy is beyond what one might expect from standard error propagation in a photochemical model, and points either to instrument error or to some fundamental flaw in understanding of upper tropospheric NO$_x$ chemistry that would in turn affect the simulated NO$_x$ lifetime. A model decrease of OH by a factor of 2 would correspondingly decrease the required increase in the lightning NO$_x$ source.

[43] Speciation of NO$_x$ in the U.S. boundary layer is about 20% NO, 20% PAN, and 60% HNO$_3$, both in the observations and the model. The PAN fraction is larger in the south than elsewhere, likely reflecting high isoprene emissions. We estimated the export efficiency $f$ and related speciation of NO$_x$ out of the U.S. boundary layer with a Lagrangian analysis of NO$_x$-CO correlations in the free troposphere, following the approach of Parrish et al. [2004]. For the ICARTT data at 2.5–6.5 km altitude we find $f = 16 \pm 10$% in the observations and 14 ± 8% in the model, consistent with previous studies of North American and Asian outflow [Li et al., 2004; Parrish et al., 2004; Koike et al., 2003; Miyazaki et al., 2003]. The highest export efficiency is in the lower free troposphere but is then mostly HNO$_3$ venting from shallow convection. We find that PAN is the dominant component of exported NO$_x$ in pollution plumes above 3.5 km, consistent with previous studies of Asian outflow. The successful simulation of export of North American NO$_x$ offers confidence in current model estimates of North American influence on the global NO$_x$ and ozone budgets [Li et al., 2004].

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