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# Improved Spatial Resolution in Modeling of Nitrogen Oxide Concentrations in the Los Angeles Basin

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**ABSTRACT:** The extent to which emission control technologies and policies have reduced anthropogenic  $\text{NO}_x$  emissions from motor vehicles is large but uncertain. We evaluate a fuel-based emission inventory for southern California during the June 2021 period, coinciding with the Re-Evaluating the Chemistry of Air Pollutants in California (RECAP-CA) field campaign. A modified version of the Fuel-based Inventory of Vehicle Emissions (FIVE) is presented, incorporating 1.3 km resolution gridding and a new light-/medium-duty diesel vehicle category.  $\text{NO}_x$  concentrations and weekday–weekend differences were predicted using the WRF-Chem model and evaluated using satellite and aircraft observations. Model performance was similar on weekdays and weekends, indicating appropriate day-of-week scaling of  $\text{NO}_x$  emissions and a reasonable distribution of emissions by sector. Large observed weekend decreases in  $\text{NO}_x$  are mainly due to changes in on-road vehicle emissions. The inventory presented in this study suggests that on-road vehicles were responsible for 55–72% of the  $\text{NO}_x$  emissions in the South Coast Air Basin, compared to the corresponding fraction (43%) in the planning inventory from the South Coast Air Quality Management District. This fuel-based inventory suggests on-road  $\text{NO}_x$  emissions that are  $1.5 \pm 0.4$ ,  $2.8 \pm 0.6$ , and  $1.3 \pm 0.7$  times the reference EMFAC model estimates for on-road gasoline, light- and medium-duty diesel, and heavy-duty diesel, respectively.



**KEYWORDS:** air pollution, emission inventory, motor vehicles, satellite

## INTRODUCTION

Nitrogen oxides ( $\text{NO}_x = \text{NO} + \text{NO}_2$ ) are highly reactive air pollutants that are produced primarily from the combustion of fossil fuels. Exposure to ground-level  $\text{NO}_2$  is associated with increased mortality due to respiratory and cardiovascular diseases, and  $\text{NO}_x$  is a major precursor to tropospheric ozone ( $\text{O}_3$ ), which has negative impacts on human health and the environment.<sup>1–4</sup> Nitrogen oxides also react in the atmosphere to create acid rain and particulate matter in the form of aerosol nitrates.<sup>4</sup> To manage the human health and environmental impacts of  $\text{NO}_x$ , an improved understanding of  $\text{NO}_x$  emission sources and the distribution of  $\text{NO}_x$  in high-population areas is needed.

In the United States, the development of emissions control technologies has been driven by laws and regulations aimed at reducing air pollution, with major changes starting in 1963 with the Clean Air Act and its subsequent amendments. The primary technologies to reduce  $\text{NO}_x$  emissions from vehicles involve devices that convert  $\text{NO}_x$  emissions into water and  $\text{N}_2$ . The widespread adoption of catalytic converters has significantly decreased the  $\text{NO}_x$  emissions from gasoline vehicles since the 1990s. As a result, heavy-duty diesel trucks became the largest mobile source of  $\text{NO}_x$  emissions as light-duty gasoline emissions declined and diesel fuel sales

continued to grow;<sup>5–7</sup> these trends were reinforced due to ineffective diesel  $\text{NO}_x$  emission controls in the 1990s.<sup>8</sup> Selective catalytic reduction (SCR) systems were introduced on new heavy-duty diesel engines starting in 2010. SCR systems use a urea solution to convert  $\text{NO}_x$  to  $\text{N}_2$  and have been shown to reduce in-use  $\text{NO}_x$  emissions from heavy-duty diesel engines by >75%.<sup>9–14</sup> However, the effectiveness of these systems in reducing  $\text{NO}_x$  emissions under real-world driving conditions can be impaired at times due to factors such as extended idling and low exhaust temperature leading to inactivation of the emission control system.<sup>11,14</sup>

The extent to which adoption of more effective  $\text{NO}_x$  control technology has affected emission trends in the United States remains unclear. Field campaigns, satellite-based inventories, and chemical transport models have produced results that highlight uncertainties in  $\text{NO}_x$  emission inventories, including how much  $\text{NO}_x$  is being emitted and the extent to which  $\text{NO}_x$

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emissions have decreased in recent years. Estimates of  $\text{NO}_x$  based on the MOtor Vehicle Emission Simulator (MOVES) emission model overestimated  $\text{NO}_x$  when compared to field campaigns in the Baltimore-Washington area and in the southeast US.<sup>15–17</sup> The MOVES model also estimated larger decreases in  $\text{NO}_x$  emissions than  $\text{NO}_2$  satellite retrievals suggest.<sup>18</sup> These differences are coupled with the fact that while precursor emissions to ozone have decreased over the past several decades, ozone concentrations in urban areas have declined relatively slowly and background ozone in the northern hemisphere has been slowly increasing.<sup>19,20</sup>

Many urban areas in North America are transitioning toward  $\text{NO}_x$ -limited ozone formation regimes, implying that  $\text{NO}_x$  emission reductions are increasingly necessary to reduce ozone.<sup>21,22</sup> The Los Angeles area is one of the largest and most populated urban areas in the United States, with air pollution problems that are linked to local topography, high volumes of vehicle traffic, and two major ports that handle nearly 30% of all imports and exports over the water in the United States.<sup>23</sup> Ozone concentrations in the region remain high despite major reductions in ozone precursors, and as a result, the South Coast Air Basin (SOCAB) was designated “extreme nonattainment” for ozone in 2018.<sup>7,24</sup>

Chemical transport models (CTMs) have been useful tools for understanding source contributions, supporting air-quality planning, and informing policy decisions. However, in order to maintain reasonable computational time and costs, most modeling studies focused on  $\text{NO}_x$  chemistry remain at a spatial resolution too coarse to see individual roadway effects. While these setups have generally been able to answer questions about the distribution and sources of  $\text{NO}_x$  on state-wide and continental levels, they have generally not been useful for resolving sharp near-source gradients, especially near major roadways. At higher spatial resolutions, there are several additional benefits, including being able to model the local effects of highways, look at neighborhood-scale differences in human health exposure, and validate models with higher-resolution satellite retrievals. While the current highest resolution satellite measurements for  $\text{NO}_2$  from the Tropospheric Monitoring Instrument (TROPOMI) include reprocessed data at a resolution of  $3.5 \text{ km} \times 5.5 \text{ km}$ ,<sup>25</sup> new instruments such as the Tropospheric Emissions: Monitoring of Pollution (TEMPO) instrument will provide even higher-resolution data on pollutant concentrations in the future.<sup>26</sup> Moving to a high-resolution inventory is in line with recent improvements in satellite spatial resolution and the increasing need to examine how changes to emissions affect the exposure of distinct communities to air pollutants.

Fuel-based inventory methods provide a complementary perspective to emission model predictions and trends inferred from satellite data, providing critical insights into how emission control technologies have impacted  $\text{NO}_x$  concentrations in heavily trafficked urban areas. To gain a better representation of the temporal and spatial patterns of  $\text{NO}_x$ , this research develops and evaluates a  $1.3 \text{ km} \times 1.3 \text{ km}$  high-resolution fuel-based inventory for Los Angeles, CA. This study evaluates this inventory as input in the Weather Research and Forecasting with Chemistry (WRF-Chem) model by comparing aircraft measurements taken during the June 2021 RECAP-CA campaign and vertically integrated tropospheric  $\text{NO}_2$  concentrations from the Sentinel-5P/TROPOMI satellite. The resulting spatial and temporal agreement with observational

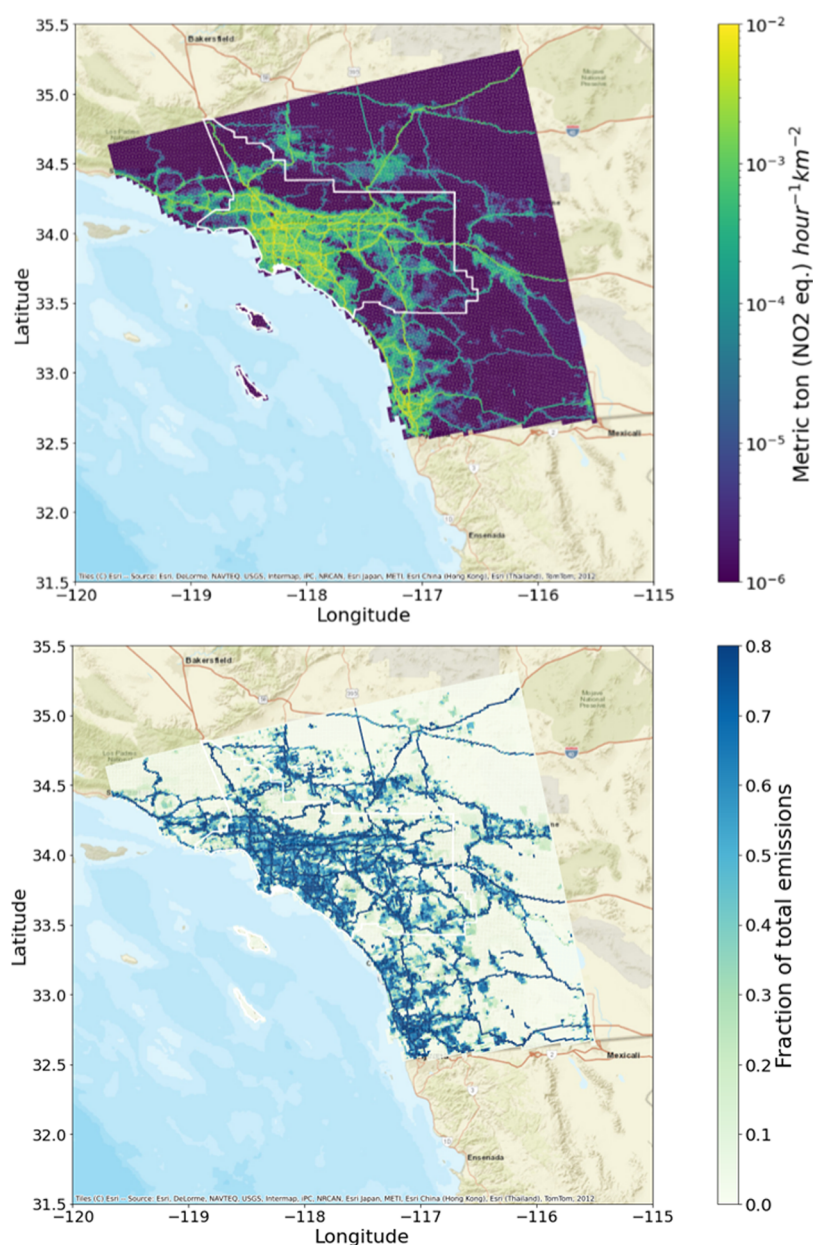
data allows us to understand the individual contributions of  $\text{NO}_x$  emission sectors in a major urban area.

## METHODS

**Atmospheric Model and Study Domain.** We use WRF-Chem (version 4.2.2<sup>27</sup>) to predict meteorological and air-quality-related variables for all of June 2021, using the last 3 days of May as a model spin-up period. The model was applied over two nested domains: (1) all of California and Nevada at 4 km horizontal resolution and (2) southern California at an unusually fine horizontal resolution of 1.3 km. The model includes 50 vertical levels of up to 50 hPa. Initial and boundary conditions for chemical species tracked in the model are from a parent 12 km resolution continental US simulation that used a similar model setup and covered the same period. Initial and boundary conditions for meteorological variables are from the 3 km horizontal resolution High-Resolution Rapid Refresh (HRRR) model, which features hourly assimilation of meteorological variables such as wind and temperature from in-flight commercial aircraft, as well as ground-based radar reflectivity observations of precipitation.<sup>28</sup> Several meteorological setups were tested and compared with RECAP-CA aircraft wind measurements. Using HRRR gave better model performance for wind speed and direction relative to setups using the North American Mesoscale (NAM) and Rapid Refresh (RAP) models. The chemical mechanism used in this study is a version of the Regional Atmospheric Chemistry Mechanism<sup>29</sup> with updates to account for key oxygenated volatile chemical product (VCP) emissions (RACM-ESRL-VCP) as specified in Coggon et al.<sup>29,30</sup> Chemical vertical mixing is enhanced for a low boundary layer height under polluted and fire conditions. Further details of the input data and WRF-Chem model parametrizations are included in Table S1 of the [Supporting Information](#).

**Emission Inventory.** Spatially and temporally resolved emissions from on-road vehicles in southern California were estimated and mapped using a modified version of the Fuel-based Inventory for Vehicle Emissions (FIVE).<sup>31,32</sup> The methodology used to create the FIVE inventory and temporal scaling factors can be found in further detail in Harkins et al.<sup>31</sup> Emissions are distributed spatially using 2018 data from a national database of traffic counts from the Federal Highway Administration and are scaled to June 2021 using state-wide taxable fuel sales.<sup>33,34</sup> Comparing the 2018 FHWA traffic counts to state-level traffic statistics, the traffic count data accounts for the majority of fuel use both for gasoline (68%) and for diesel fuel (77%) in California.<sup>35</sup> The remaining fuel use is mostly on small local roads rather than on major roadways and can thus be adequately distributed using census block-level population data gridded to 1.3 km resolution.<sup>36</sup> Gridded gasoline and diesel fuel consumption estimates are combined with fuel-specific emission factors (grams of pollutant emitted per kilogram of gasoline or diesel fuel burned) to calculate emissions. Emission factors for each vehicle type are determined based on regression analyses of measurements from roadside remote sensing and highway tunnel studies conducted in California.<sup>7</sup>

The major modifications to the FIVE inventory for this study include moving to 1.3 km resolution and the reapportionment of 22% of the total diesel fuel to a light-/medium-duty diesel vehicle category, separate from the existing gasoline and heavy-duty diesel vehicle categories. These light-/medium-duty diesel vehicles now have higher fuel-based  $\text{NO}_x$



**Figure 1.** Spatial distribution of average 2021 weekday  $\text{NO}_x$  emission rates from on-road vehicles in southern California (top) and the fraction of total emissions from on-road vehicles (bottom), mapped at a 1.3 km horizontal resolution. The South Coast Air Basin is outlined.

emission factors compared to heavy-duty diesel trucks.<sup>7</sup> Reasons for higher emission factors in this category may vary due to the inclusion of a wide range of vehicle types, but contributing factors include slow fleet turnover of light-duty diesel engines and less deployment of advanced emission control technology (i.e., selective catalytic reduction for  $\text{NO}_x$  control) in comparison with heavy-duty diesel trucks. Heavy-duty diesel engines have been a high state-wide priority in efforts to accelerate replacement of older (pre-2010) engines with newer and lower-emitting engines.<sup>37</sup> Diurnal and day-of-week variations in vehicular emissions are based on weigh-in-motion traffic count data, with separate temporal variation profiles for light- and heavy-duty vehicles.<sup>32</sup> We assume that the spatial distribution of light- and medium-duty diesel truck traffic matches that of light-duty gasoline vehicles and use the activity profile for heavy-duty trucks to specify temporal variations. In comparison to the existing FIVE inventory, this

reapportionment of diesel fuel to light-/medium-duty vehicles results in higher on-road  $\text{NO}_x$  emissions overall, especially on weekdays, and more emissions being attributed to light-duty vehicle spatial patterns.

In addition to on-road vehicle emissions, the emission inventory used in this study also includes fuel-based emissions from off-road engines used in agricultural and construction equipment.<sup>31,38,39</sup> Also included are emissions from oil and natural gas production,<sup>40</sup> power plants based on Continuous Emission Monitoring System data,<sup>41</sup> and other point and area sources from the 2017 National Emission Inventory scaled to 2021 based on activity factors.<sup>31,42</sup> Ocean-going vessel emissions and emissions from all sources in Mexico were from the Copernicus Atmosphere Monitoring Service (CAMS) inventory.<sup>43</sup> Biogenic emissions are based on the Biogenic Emissions Inventory System (BEIS) v3.14 model.<sup>44,45</sup> Consistent with previous modeling work in Los Angeles,<sup>46</sup>

Table 1. Estimates of NO<sub>x</sub> Emissions from On-Road Vehicles in Southern California<sup>a</sup>

	vehicle category			on-road total
	gasoline	LD + MD diesel <sup>b</sup>	HD diesel <sup>b</sup>	
fuel burned (t/day)	41,763 ± 2506	2295 ± 298	6015 ± 782	
EF NO <sub>x</sub> (g/kg)	2.0 ± 0.6	20.2 ± 3.4	9.2 ± 4.6	
avg emissions (t/day)	87 ± 27	49 ± 10	62 ± 32	198 ± 69
WD <sup>c</sup> emissions (t/day)	89	60	76	224
WE <sup>c</sup> emissions (t/day)	82	20	26	127
EMFAC (t/day)	59	17	48	124
fuel-based/EMFAC <sup>d</sup>	1.5 ± 0.4	2.8 ± 0.6	1.3 ± 0.7	1.6 ± 0.5

<sup>a</sup>Emission estimates for the South Coast Air Basin, including Los Angeles, Orange County, and portions of Riverside and San Bernardino Counties. <sup>b</sup>Diesel vehicle weight categories are light-, medium-, and heavy-duty (LD, MD, and HD). <sup>c</sup>Emissions for average weekday (WD) and weekend (WE) conditions. <sup>d</sup>Ratio of fuel-based emission inventory (weighted average of WD and WE values) from this study with corresponding estimates for summer 2021 from the most current version of the EMFAC model.<sup>59</sup>

BEIS emissions for isoprene and monoterpenes from the urban land cover type were updated based on Scott and Benjamin.<sup>47</sup>

**Aircraft and Surface Monitor Data.** Model predictions were compared to vertical profile NO<sub>2</sub> measured during aircraft flights over the Los Angeles basin that took place during the summer of 2021 as part of the RECAP-CA (Re-Evaluating the Chemistry of Air Pollutants in California) field campaign.<sup>48,49</sup> Relevant flights occurred at midday hours on three weekend days (June 6, 12, and 19) and six weekdays (June 1, 4, 10, 11, 18, and 21). NO<sub>x</sub> was measured at 5 Hz using a thermal dissociation laser-induced fluorescence (TD-LIF) instrument, and a detailed overview of the instrument can be found in Thornton et al. and Day et al.<sup>50,51</sup> Instrument calibration details and methodology used during this campaign can be found in Zhu et al.<sup>49</sup> Here, we report only data from the stacked racetrack patterns (see Figure S1), where the plane flew 4–6 different altitudes' layer stacked on top of one another within the planetary boundary, which were designed to measure vertical concentration profiles. The aircraft data used here were split into flights that occurred closer to the coast (west/central LA) and at locations further inland (east basin) where temperatures were higher. Only altitude bins with greater than five observation points were used in the comparison. The aircraft data were matched with corresponding model predictions using a nearest neighbor method, averaging the aircraft data every 30 s and pulling comparison data for the nearest model grid point.

In addition to verifying the vertical profiles against aircraft observations, the South Coast Air Quality Management District (AQMD) monitoring network was used for comparison to verify the diurnal patterns of NO<sub>2</sub> in the model.<sup>46</sup> The NO<sub>2</sub> data for 24 sites located in the South Coast Air Basin were averaged for each hour of the day on weekdays and weekends. The quantity of NO<sub>2</sub>\* (NO<sub>2</sub>\* = NO<sub>2</sub> + PAN + alkyl nitrates + HONO + 2\*N<sub>2</sub>O<sub>5</sub>) was used from the model for comparison to the measured NO<sub>2</sub> due to the conversion of other nitrogenous species in addition to NO<sub>2</sub> by the molybdenum converter installed within standard chemiluminescent NO<sub>x</sub> analyzers.<sup>52,53</sup>

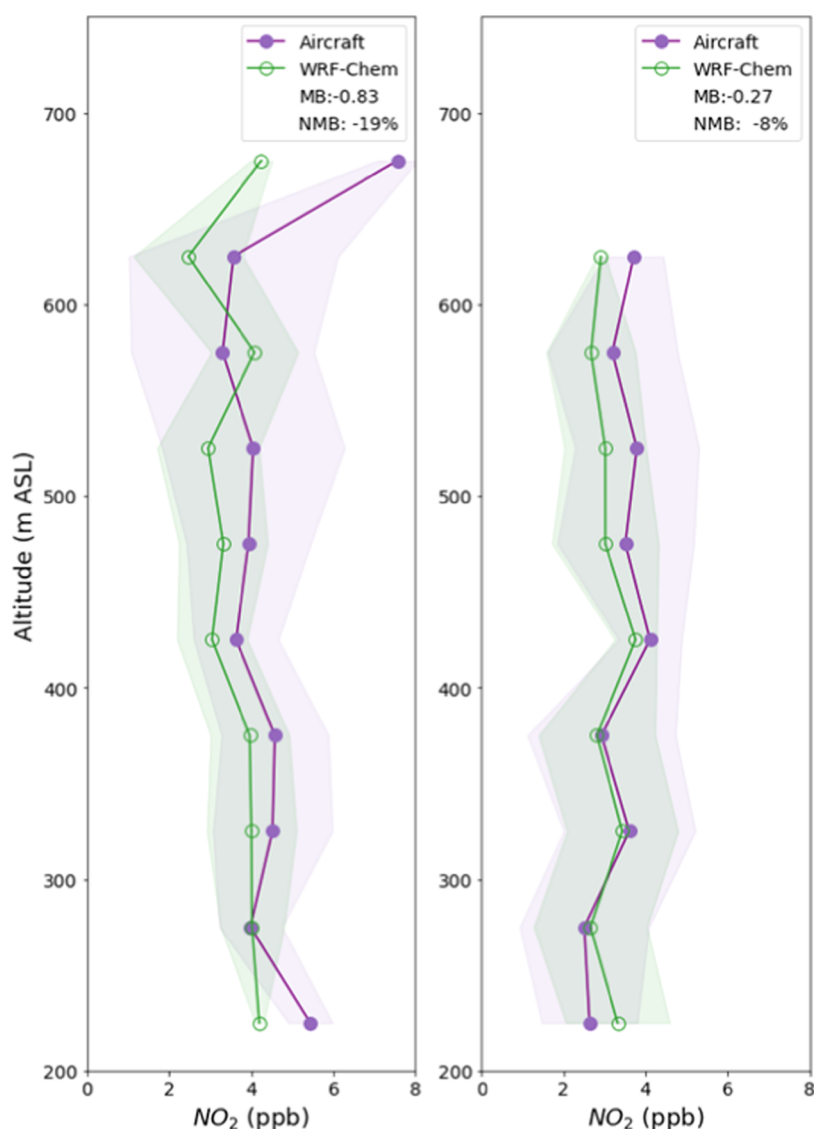
**Satellite Observations.** TROPOMI satellite measurements are used in this study to evaluate tropospheric column-integrated model predictions of NO<sub>2</sub> concentrations. We use the latest NO<sub>2</sub> product reprocessed to a spatial resolution of 3.5 km × 5.5 km from the Sentinel-5P Products Algorithm Laboratory (SP5-PAL) for June 2021.<sup>25,54</sup> We used this reprocessed data product to create average June tropospheric NO<sub>2</sub> columns using methods described by Li et al., as

well as separate weekday and weekend average tropospheric NO<sub>2</sub> columns for the study period.<sup>55</sup> Pixels with cloud cover, snow, or ice, or with otherwise problematic retrievals, are filtered out to reduce uncertainties by using only data where the quality flag ≥ 0.75.<sup>56</sup> NO<sub>2</sub> vertical profiles require an assumption for the air mass factor (AMF), which may affect the ability to compare tropospheric NO<sub>2</sub> columns between the model and satellite data. In order to eliminate biases introduced by a priori profile assumptions, the NO<sub>2</sub> vertical profiles used in this study are calculated using averaging kernels and the AMF was calculated from WRF-Chem, rather than using the AMF derived from the a priori TMS model.<sup>57</sup> This approach has been shown to increase the satellite NO<sub>2</sub> concentrations over urban areas by 20% on average.<sup>55</sup> Even accounting for these improvements, there is still an expectation of a low bias of −20% in TROPOMI tropospheric NO<sub>2</sub> columns over polluted cities in comparison to observations based on studies compared to ground-based Pandora measurements.<sup>54,55,58</sup>

## RESULTS AND DISCUSSION

Figure 1 shows the spatial distribution of NO<sub>x</sub> emissions from on-road vehicles for the 1.3 km resolution southern California modeling domain. NO<sub>x</sub> emissions are the highest in the urban center, with lower emissions in sparsely populated rural and mountainous areas. This pattern follows from the emissions mapping methodology, which relies on vehicle traffic counts that indicate high traffic densities within urban areas. Sharp spatial gradients in vehicle emissions near major highways are clearly apparent in the top panel of Figure 1, not only in rural areas but also within densely populated urban areas. While high NO<sub>x</sub> emissions in and around downtown LA are mostly due to high volumes of light-duty vehicle traffic, elsewhere along major highways such as I-5 (running to the northwest of LA) and I-15 (running to the northeast of LA toward Las Vegas), the majority of the NO<sub>x</sub> emissions are due to heavy-duty diesel trucks. Even though emissions from on-road vehicles decrease on weekends primarily due to the steep weekend decrease in diesel vehicle activity, vehicles overall are still the dominant source of NO<sub>x</sub> emissions, responsible for 52% of the NO<sub>x</sub> on weekends versus 62% of the NO<sub>x</sub> on weekdays. The fraction of the total emissions from the on-road sector on weekdays can be seen in the bottom panel of Figure 1.

More details concerning NO<sub>x</sub> emissions from on-road vehicles are presented in Table 1. We estimate that average daily NO<sub>x</sub> emissions for the South Coast Air Basin were 198 ±



**Figure 2.** Vertical profiles of measured (RECAP flights) and modeled NO<sub>2</sub> concentrations over Western (left panel) and Eastern Los Angeles (right panel), binned over 100 m intervals. Shaded bands indicate 1 standard deviation from the mean values.

69 t/day during our June 2021 study period, which is  $1.6 \pm 0.5$  times the corresponding summer-season estimate from the California EMFAC model. Our estimates of NO<sub>x</sub> emissions are comparable for heavy-duty diesel and higher for gasoline and light-/medium-duty diesel categories, with the largest relative difference (a ratio of  $2.8 \pm 0.6$ ) for light- and medium-duty diesel vehicles. As shown in Table 1, on-road vehicle emissions are lower on weekends than on weekdays, as expected, due mainly to the decreased activity and emissions from diesel trucks. The overall decrease in vehicular NO<sub>x</sub> emissions on weekends is about 100 t/day, which is a 43% reduction relative to baseline weekday conditions. A potential cause of the lower emissions in the EMFAC model in comparison to the fuel-based inventory is in how emission factor trends are derived. The modified FIVE inventory used in this study uses on-road remote-sensing data trends, resulting in emission factors that are higher overall and are decreasing more slowly in comparison to laboratory-measured emission factors used in other inventories. The large sample size of remote-sensing data is better equipped to capture the effects of individual vehicles with ineffective or nonfunctioning emission control systems.

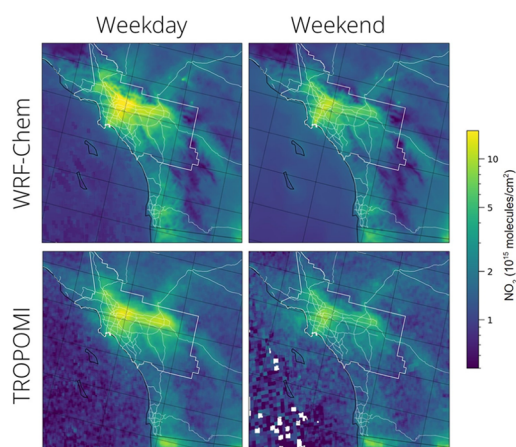
While the remote-sensing studies capture driving conditions that reflect typical vehicle activity patterns, they do not capture start-up and idling conditions, leaving room for actual average emission factors to be even higher. In the case of heavy-duty vehicles, selective catalytic reduction (SCR) systems do not operate effectively at low temperatures, leading to potential underestimates of mobile source emissions in urban areas when idealized SCR performance is assumed.

Figure 2 shows modeled and measured NO<sub>2</sub> concentrations as a function of altitude above sea level; values are binned within 100 m intervals of altitude. The plots are separated into two regions, Western LA and Eastern LA, in order to eliminate spatial biases. Flight tracks showing the specific locations of these measurements are shown in Figure S1. Normalized mean biases for NO<sub>2</sub> are -19 and -8% for the western and eastern portions of Los Angeles, respectively. Vertical profiles are in good agreement at most altitudes, especially below 600 m. In Eastern LA, the NO<sub>2</sub> concentrations are closer to the observations near the surface and diverge slightly with an increasing altitude. In Western LA, we see that generally the model and observation of NO<sub>2</sub> match very closely, especially

between 250 and 600 m. Despite slight underestimation in the model compared to the aircraft observations overall, these results indicate that the vertical representation of  $\text{NO}_2$  in the model closely resembles the  $\text{NO}_2$  measured during the field campaign during the same period. An accurate vertical profile in the model is critical to the calculation of tropospheric  $\text{NO}_2$  columns from the satellite, as the AMF used in the calculation is revised to use the  $\text{NO}_2$  vertical profile shape from the WRF-Chem simulation and averaging kernels.

The comparison of the diurnal patterns between the model and AQMD surface observations for the June 2021 study period is shown in Figure S2. This comparison suggests that the diurnal variations in modeled  $\text{NO}_2^*$  are consistent with observations, but there is some negative bias in the modeled  $\text{NO}_2^*$  concentrations during nighttime hours on the weekends. The analysis performed at a 4 km resolution yielded the same conclusions.

Figure 3 compares spatial distributions of modeled (WRF-Chem) tropospheric column-integrated  $\text{NO}_2$  concentrations



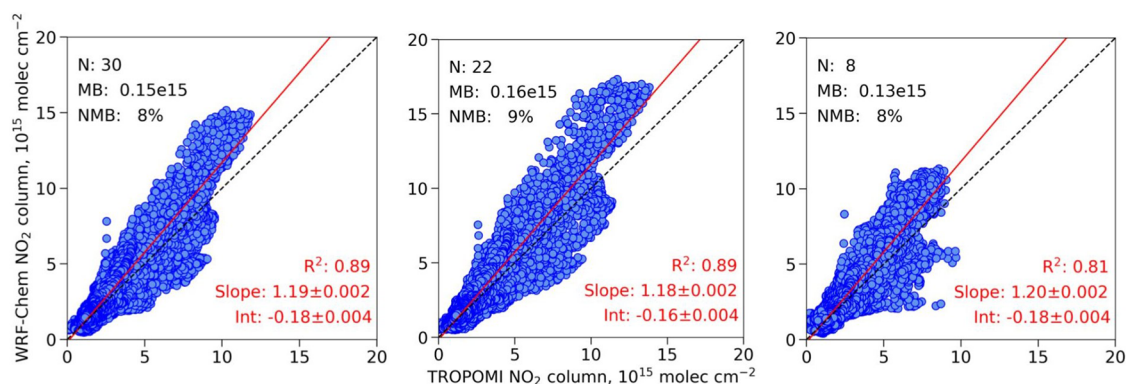
**Figure 3.** Average tropospheric  $\text{NO}_2$  columns over southern California for June 2021 were predicted using the WRF-Chem model (top row) with comparisons to TROPOMI satellite data (bottom row). Separate results are shown for weekdays (left panels) and weekends (right panels). The boundaries of the South Coast Air Basin and major highways are shown in white.

with the corresponding TROPOMI satellite-derived values. Both model and satellite data show high weekday  $\text{NO}_2$  columns over downtown LA extending south toward Long

Beach as well as extending further inland to the east as far as San Bernardino. In both cases, there is significantly lower  $\text{NO}_2$  on the weekends, especially apparent within the South Coast Air Basin along the corridors east and south of downtown LA. Elevated  $\text{NO}_2$  concentrations due to traffic along major highways running through rural areas are captured well, notably on Interstate highways I-5 (heading north toward San Francisco and Sacramento) and I-15 (heading northeast toward Las Vegas) (see Figure S3 of the Supporting Information for interstate locations). The main difference between modeled and observed  $\text{NO}_2$  in the southern California domain is that, overall, WRF-Chem has higher  $\text{NO}_2$  concentrations in the downtown region, with maximum  $\text{NO}_2$  concentrations slightly offset to the northeast compared to what the TROPOMI satellite observations show (Figure S4). These differences are slightly emphasized on weekdays in comparison to weekends.

Figure 4 shows scatterplots of modeled tropospheric  $\text{NO}_2$  columns versus corresponding satellite-derived values. The lower  $\text{NO}_2$  values that prevail on weekends are apparent in the more limited range of the data in the rightmost panel of Figure 4. The regression-derived coefficients are similar for all three plots, with near-zero intercepts and slopes of 1.18–1.20. The model explains a high fraction of the observed variance in  $\text{NO}_2$  columns, although the value of  $R^2$  decreases somewhat from 0.89 on weekdays to 0.81 on weekends. Mean normalized bias for the model relative to satellite data is +8% for all days in June, with similar values for the subsets of weekdays only and weekend days only. The finding of slight overestimation in the model compared to the satellite data is in contrast with the finding of slight underestimation compared to the aircraft observations. The differences between the model and the satellite are all within the range of and directionally consistent with findings of previous studies that suggest a negative bias of  $\sim 20\%$  in satellite-derived observations of  $\text{NO}_2$  columns in urban areas.<sup>47</sup>

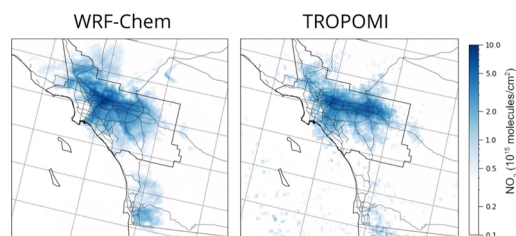
The direct comparison between model and satellite  $\text{NO}_2$  columns at 4 km resolution over the larger California domain can be seen in Figure S5 of the Supporting Information. Model performance for our study period at a 4 km resolution for California is comparable to model performance at a 1.3 km resolution for southern California within the same domain, as shown in Figure S6 of the Supporting Information. At 4 km resolution, we see an  $R^2$  of 0.83 with a slope of 1.14 for the entire California/Nevada domain and an  $R^2$  of 0.91 and a slope



**Figure 4.** Orthogonal distance regression between modeled and satellite-derived tropospheric  $\text{NO}_2$  columns over southern California in June 2021 for all days (left panel), weekdays only (middle panel), and weekends only (right panel).

of 1.09 for the southern California domain. While the results have similar statistics, the 1.3 km output has the added benefit of resolving sharp gradients in NO<sub>2</sub> over major cities and highways, which is in line with what can be resolved by the satellite.

To further evaluate the emission inventory, we turn to a more in-depth consideration of weekday–weekend differences in the NO<sub>2</sub> columns. Figure 5 shows weekday–weekend



**Figure 5.** Weekday–weekend difference plots for tropospheric NO<sub>2</sub> columns for the WRF-Chem model and TROPOMI satellite data.

difference plots of NO<sub>2</sub> columns for the model and for the satellite data. The spatial patterns of areas showing large weekend decreases in these plots are similar. The model shows larger weekend NO<sub>2</sub> decreases northwest of downtown along the I-5 Interstate highway through the San Fernando Valley, while the satellite data indicates larger weekend NO<sub>2</sub> decreases further inland, extending to the east as far as San Bernardino. The traffic count data underlying the fuel-based inventory used in this study are for 2018 and may not adequately reflect expansions in warehousing, truck traffic, and associated NO<sub>x</sub> emissions that have recently occurred in inland portions of the LA basin.<sup>60,61</sup>

An additional emission sensitivity case was considered in a model run with all on-road vehicle emissions zeroed out completely. Emissions from other source categories were left unchanged. As shown in Figure S7, the spatial pattern in the resulting weekday–weekend difference plot for this case without the on-road sector does not match either the observed spatial distribution or the magnitude of weekend NO<sub>2</sub> decreases shown in Figure 5. We conclude that locations that exhibit large weekend NO<sub>2</sub> decreases (i.e., from Downtown LA east to San Bernardino) are those where on-road vehicle emissions make large absolute and relative contributions to tropospheric NO<sub>2</sub> columns.

Current emission inventories used for air-quality planning purposes suggest that on-road vehicle emissions are no longer the dominant source of NO<sub>x</sub> emissions in southern California. In contrast, as shown in Table 2, the emission inventory in this

**Table 2.** Comparison of NO<sub>x</sub> Emission Inventories for the South Coast Air Basin

source sector	fuel-based inventory June 2021		south coast air-quality management district, summer 2022 <sup>a</sup>	
	NO <sub>x</sub> (t/day)	fraction of total	NO <sub>x</sub> (t/day)	fraction of total
on-road	198 ± 69	66%	103	43%
area + off-road	79	26%	116	49%
point	23	8%	20	8%
total	300		239	

<sup>a</sup>Ocean-going vessels removed to match included sectors of fuel-based inventory.

study has 25% higher NO<sub>x</sub> emissions overall when compared to the planning inventory, with the dominant contribution coming from on-road sources. More specifically, the present study includes higher emissions from on-road vehicles (198 ± 69 vs 103 tons/day) that account for 66% of the anthropogenic NO<sub>x</sub> emissions. The planning inventory assigns greater importance to area and off-road mobile sources, which together account for a higher fraction (49 vs 26%) of total NO<sub>x</sub> emissions compared to the present study.

Air-quality model results, satellite observations, and their corresponding weekday–weekend differences are consistent with a larger-than-expected on-road vehicular source of NO<sub>x</sub> emissions in southern California. Major efforts to update fleets to newer engine models with advanced emission control systems have improved heavy-duty diesel, and values from the EMFAC model are consistent with and within the uncertainty of those from the fuel-based inventory evaluated here. Still, actual emissions from heavy-duty diesel may be even higher than estimated due to ineffective NO<sub>x</sub> control during idling and low load driving conditions. Total NO<sub>x</sub> emission estimates are higher in this study for on-road vehicles in comparison to the EMFAC inventory owing to higher emissions from gasoline vehicles and light- and medium-duty diesel vehicles. Contributing factors to these differences may include slow absolute progress in reducing fleet-average NO<sub>x</sub> emission factors for gasoline vehicles since 2010 (see Figure 2 in Yu et al., 2021) and slow fleet turnover with relatively high in-use NO<sub>x</sub> emission factors for light- and medium-duty diesel trucks.

## ■ ASSOCIATED CONTENT

### Supporting Information

The emission inventory used in this study is available at the following URL: <https://csl.noaa.gov/groups/csl7/measurements/2021sunvex/emissions/>. The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.est.3c06158>.

Summary of the WRF-Chem model setup and input data, flight paths of aircraft, diurnal pattern comparison with AQMD sensors, key interstate freeways in the Los Angeles domain, absolute difference plots between model and satellite, average tropospheric NO<sub>2</sub> columns at 4 km resolution, orthogonal distance regression between model and satellite NO<sub>2</sub> at 4 km resolution for the California/Nevada domain and the southern California domain, weekday–weekend difference of the on-road sector and combined off-road, area, and point sectors (PDF)

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## Notes

The authors declare no competing financial interest.

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