

Elucidating secondary organic aerosol from diesel and gasoline vehicles through detailed characterization of organic carbon emissions

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Emissions from gasoline and diesel vehicles are predominant anthropogenic sources of reactive gas-phase organic carbon and key precursors to secondary organic aerosol (SOA) in urban areas. Their relative importance for aerosol formation is a controversial issue with implications for air quality control policy and public health. We characterize the chemical composition, mass distribution, and organic aerosol formation potential of emissions from gasoline and diesel vehicles, and find diesel exhaust is seven times more efficient at forming aerosol than gasoline exhaust. However, both sources are important for air quality; depending on a region's fuel use, diesel is responsible for 65% to 90% of vehicular-derived SOA, with substantial contributions from aromatic and aliphatic hydrocarbons. Including these insights on source characterization and SOA formation will improve regional pollution control policies, fuel regulations, and methodologies for future measurement, laboratory, and modeling studies.

motor vehicle emission factors | photochemical oxidation | urban air quality | volatile organic compound emissions | petroleum fuel composition

Organic aerosol (OA) in the atmosphere is detrimental to human health and represents a highly uncertain forcing of climate change (1). The use of petroleum-derived fuels is an important source of reactive gas-phase organic carbon that provides key precursors to the formation of secondary OA (SOA) and tropospheric ozone (1). Controlling these emissions from gasoline and diesel vehicles is central to air quality mitigation policies in urban areas (2). Previous work has concluded that further research is necessary to elucidate all organic sources of SOA precursors (3, 4). Significant controversy exists over the contributions of precursors from gasoline and diesel vehicles, and the relative importance of each for SOA formation remains in question, in part because of insufficient chemical characterization of fuels and emissions, and the difficulty of ambient measurements of gas-phase compounds emitted from diesel sources (1, 4–8).

In the United States, diesel fuel accounts for 21% of on-road fuel use (by volume), with off-road sources increasing total use to 28% diesel. In California, the diesel share of on-road use ranges from approximately 10% in coastal cities to more than 30% in agricultural regions (*SI Appendix, Table S1*) (2, 9, 10). Noncombusted hydrocarbons from the fuels are emitted in the exhaust of gasoline and diesel engines, and also via evaporation from gasoline vehicles and service stations. These compounds in unburned gasoline and diesel fuel dominate vehicular emissions of reactive gas-phase carbon that have the potential to form SOA (11, 12). Previous work has shown nontailpipe emissions account for ~30% of gasoline-related emissions in urban regions, but limited work exists constraining the emissions and SOA formation potential of gas-phase organic carbon from gasoline and diesel sources (13). By using extensive fuel analyses and field data from two sites that include many compounds with no previous in

situ measurements, we present the most comprehensive data to date on the chemical composition, mass distribution, emissions, and SOA formation potential of nontailpipe gasoline, gasoline exhaust, and diesel exhaust. We determine the relative importance of gasoline and diesel sources for SOA formation in, and downwind of, urban regions. We assess these results in the context of other studies during the past decade and discuss their significant implications for air pollution measurement, modeling, and control.

Results and Discussion

A total of 40 gasoline and 12 diesel fuel samples from California were collected (coincident with field data) and characterized by using several gas-chromatography methods, yielding comprehensive speciation of the “unresolved complex mixture” in diesel fuel. This was accomplished by using soft photoionization techniques, and provides unprecedented detail on the molecular identification and mass distribution of hydrocarbons in diesel fuel (14). Gasoline and diesel fuel, and thus their emissions of unburned hydrocarbons, can be classified by vapor pressure and span the volatile organic compound (VOC) range and the less volatile intermediate-volatility organic compound (IVOC) range (Fig. 1). Gasoline hydrocarbons fall mostly within the VOC range, with some aromatics extending into the IVOC range, whereas only 30% of diesel fuel hydrocarbons are in the VOC range. Diesel fuel is widely distributed across molecules containing 8 to 25 carbon atoms with a peak around 10 to 13 carbon atoms (Fig. 24). This peak is a result of aromatics and cycloalkanes, as straight and branched alkanes are evenly distributed between 10 and 20 carbon atoms. Aromatic and aliphatic hydrocarbons make up 23% and 68% of diesel fuel, respectively. By comparison, gasoline contains ~30% aromatics, with the remainder of the nonethanol fraction dominated by straight and branched alkanes with less than 10 carbon atoms (Table 1 and Fig. 24).

To examine contributions from each source to reactive gas-phase organic carbon in the ambient atmosphere and on-road emissions measured in a roadway tunnel, we used a chemical mass balance model with effective variance weighting on overconstrained

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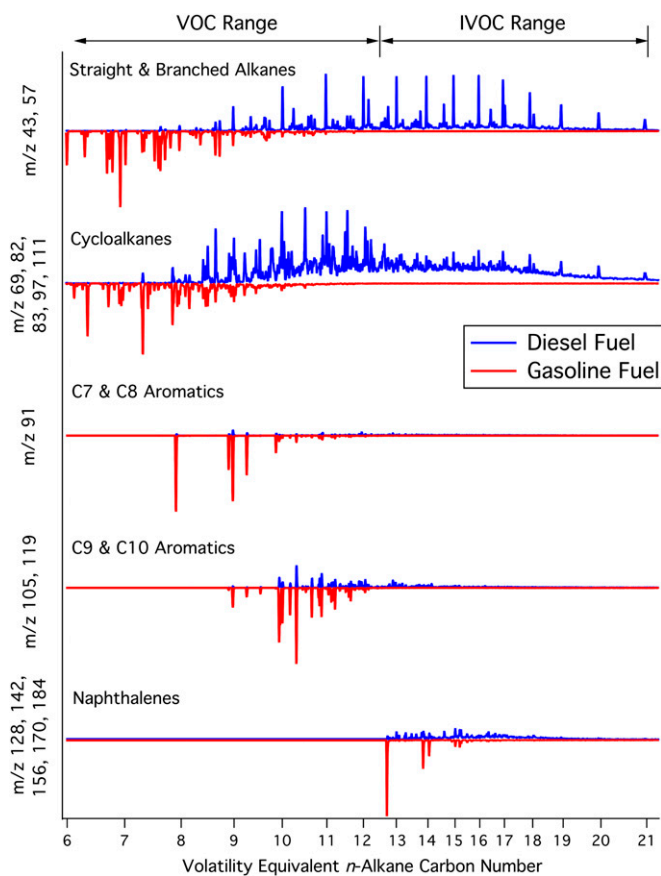


Fig. 1. Distributions of chemical classes for diesel (blue) and gasoline (red) are distinct with some overlap as shown via GC/MS for representative fuel samples. Fuels span both the VOC and IVOC volatility ranges. Chemical classes are represented by their dominant mass fragments and shown as a function of *n*-alkane carbon number.

least-squares regressions (*SI Appendix*) (15). The model uses a subset of measured compounds and capitalizes on differences in the chemical composition of sources to assess the magnitude of total noncombusted hydrocarbon emissions from each source. The source profiles used as a priori information are constructed from liquid fuel data to represent gasoline and diesel exhaust, and vapor-liquid equilibrium calculations to represent nontailpipe gasoline emissions. Equivalent chemical composition in exhaust and liquid fuel has been reported previously for gasoline and is demonstrated in this work for gasoline and diesel at both measurement sites (*SI Appendix*, Fig. S1) (16). Extensive diagnostics were used to assess model performance, including comparisons against independent compounds to confirm the model's ability to predict the behavior of reactive VOCs and IVOCs emitted by gasoline and diesel sources (*SI Appendix*, Figs. S2–S4).

Emission factors for noncombusted gas-phase organic carbon in exhaust were determined to be $0.38 \pm 0.11 \text{ gC}\cdot\text{L}^{-1}$ for gasoline and $0.86 \pm 0.25 \text{ gC}\cdot\text{L}^{-1}$ for diesel vehicles, which are consistent with values calculated by using California's emissions model for the same period (17). With respect to contributions of noncombusted hydrocarbons from gasoline and diesel exhaust, diesel accounted for 24% at the tunnel study in a coastal city, compared with 56% in the urban center of an agricultural region. Accounting for differences in emission factors and fuel densities, this is consistent with on-road fuel sales data in both regions—11% and 33% diesel fuel by volume, respectively (*SI Appendix*, Table S1) (10).

To assess the importance of gasoline and diesel sources for SOA in urban areas, we calculated bulk SOA yields for all three sources and compared them in context of our emission factors and source contributions. Data on SOA yields are limited for many of the hydrocarbons; the mass fraction of diesel, gasoline, and nontailpipe gasoline emissions that have unknown yields are 66%, 25%, and 7%, respectively. Thus, we modeled high- NO_x SOA yields by using published data (where available) and an estimation of yields and uncertainties for unknown values based on best estimates from various plausible scenarios (Fig. 2*B* and *SI Appendix*, Fig. S5).

For the same mass of unburned fuel emissions reacted, diesel exhaust forms 6.7 ± 2.9 times more SOA than gasoline exhaust (bulk SOA yields of 0.15 ± 0.05 and $0.023 \pm 0.007 \text{ }\mu\text{gSOA}\cdot\mu\text{g}^{-1}$, respectively). Considering differences in emission factors, diesel exhaust is expected to form 15 times more SOA than gasoline per liter of fuel burned. For populated regions with 10% to 30% diesel fuel use, this implies that diesel exhaust is responsible for two to seven times more SOA than gasoline exhaust (Fig. 3). Nontailpipe gasoline emissions were 39% to 77% lower than gasoline exhaust emissions and produce negligible SOA as a result of a substantially lower yield (0.0024 ± 0.0001).

Our methods also allowed us to examine the most important chemical classes and mass distribution of SOA formation. The vast majority of SOA from gasoline sources is a result of its aromatic content, whereas diesel SOA is predicted to be $47 \pm 7\%$ from aliphatics, with the remainder from aromatics (Fig. 2*B* and Table 1).

Regional estimates of daytime SOA concentrations from diesel and gasoline using our model results and calculated SOA yields are consistent with independent positive matrix factor analysis results for aromatic and aliphatic SOA from fossil fuel combustion in the San Joaquin Valley using aerosol MS (AMS) and Fourier transform IR spectroscopy (FTIR) measurements. Based on our model results, we expect an average of $1.3 \pm 0.4 \text{ }\mu\text{gOA}\cdot\text{m}^{-3}$ from motor vehicles compared with average $1.0\text{-}\mu\text{m}$ particulate matter concentrations of 1.8 to $2.1 \text{ }\mu\text{g OA}\cdot\text{m}^{-3}$ from FTIR and AMS data, respectively (18). These independent data also support the predominance of diesel SOA in the San Joaquin Valley, as young aerosol (O:C ratios of 0.27–0.36) was 58% aliphatic and 42% aromatic (18).

SOA models have made considerable progress by using a parameterization known as the volatility basis set to estimate contributions from unmeasured intermediate and semivolatiles compounds (5, 19). Together with traditional explicit models for individual hydrocarbons in the VOC range, models are better able to predict the magnitude of observed SOA, but not all temporal patterns or physical/chemical characteristics (3, 19, 20). Here we evaluate the inclusion of SOA precursors in these models and their distribution in gasoline and diesel exhaust. Aromatics with single or multiple rings have rightfully received considerable attention historically, but their distribution between gasoline and diesel emissions has been relatively unexplored. Gasoline exhaust dominates emissions of C_7 and C_8 aromatics. C_9 aromatic content is four times greater in gasoline than in diesel, and there are nearly equivalent amounts of C_{10} aromatics. For an urban region with 15% diesel fuel use, this implies that gasoline emits more than 90% of the C_9 aromatics and 75% of the C_{10} aromatics. Gasoline SOA from C_9 and C_{10} aromatics represent 26% and 14% of total SOA from gasoline, respectively, and C_{9-11} aromatics represent 5% of SOA from diesel exhaust (Table 1). Emissions of naphthalene and similar small polycyclic aromatic hydrocarbons (PAHs) are shared by gasoline and diesel vehicles, but represent only a minor contribution to potential SOA formation as a result of their minor weight fractions in the fuels (Fig. 2 and *SI Appendix*, Tables S9 and S10).

We examined the compounds included in SOA models and found that 20% to 30% of the SOA formed from gasoline

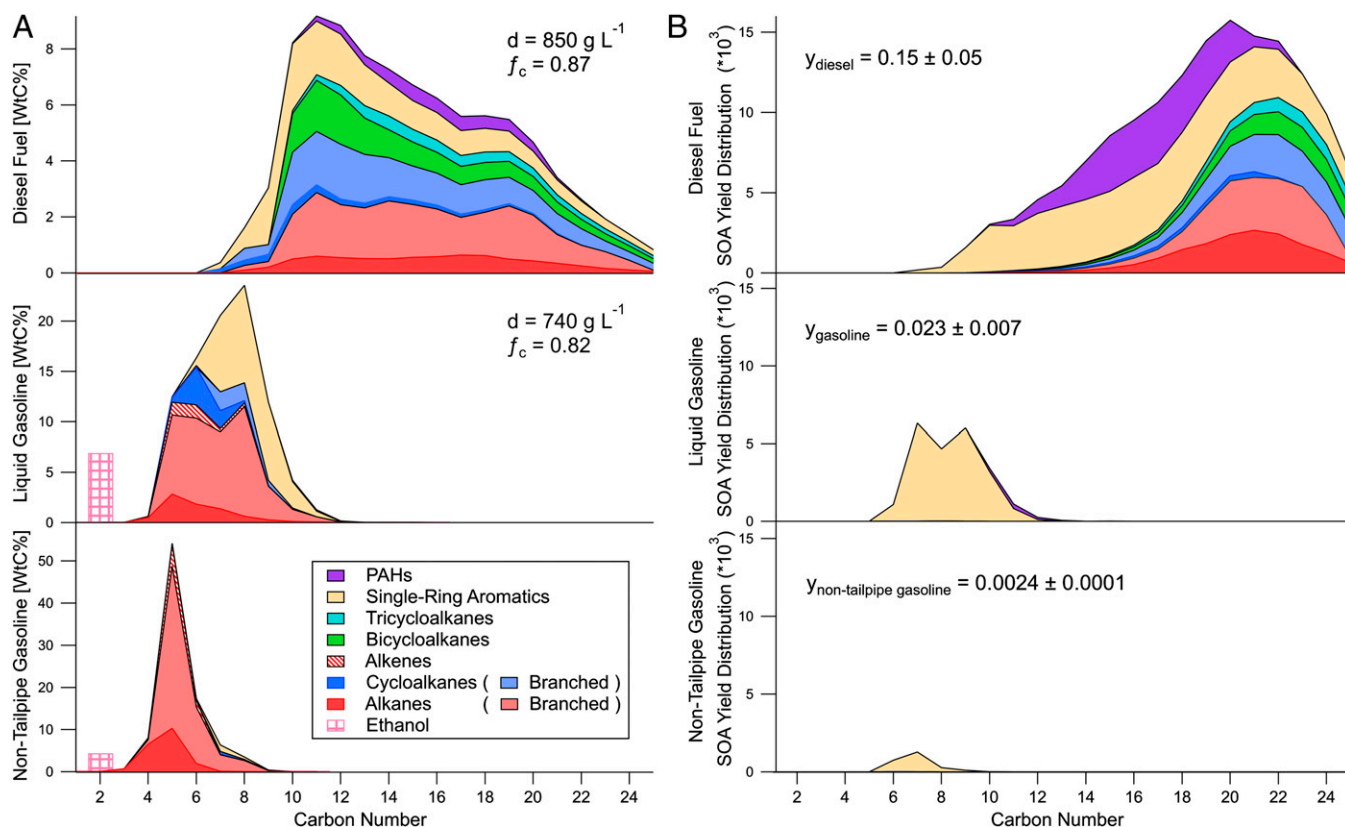


Fig. 2. Distribution of mass (A) and SOA formation potential (in $\mu\text{g SOA}\cdot\mu\text{g}^{-1}$; B) in diesel and gasoline fuel (representative of exhaust) and nontailpipe gasoline emissions. Distributions in A and B are colored by chemical class. Fuel properties (density, carbon fraction) and bulk SOA yields (at an organic particle loading of $10\ \mu\text{g}\cdot\text{m}^{-3}$) are superposed on A and B, respectively. Predicted SOA from gasoline exhaust is much lower than diesel and dominated solely by aromatic content, whereas diesel SOA is produced from a mix of aromatic and aliphatic compounds. A distribution of the SOA potential uncertainties is provided in *SI Appendix, Fig. S5*.

exhaust was not included in recent urban studies (21–23) (*SI Appendix*). Given the contributions of C_{9-11} aromatics to SOA formation from gasoline and diesel vehicles, it is important that they are better represented in explicit traditional SOA models or the extension of volatility basis set modeling to include the 10^7 and $10^8\ \mu\text{g}\cdot\text{m}^{-3}$ effective saturation concentration (C^*) bins that fall in the VOC range (*SI Appendix, Fig. S6*) (5, 19, 22). For recent urban studies, we scaled up traditional compound-explicit SOA models (without the volatility basis set) to include the missing 20% to 30% of gasoline SOA and contributions from diesel

(assuming 15% diesel fuel use) and calculated a fivefold increase in modeled SOA from vehicular exhaust. Such an inclusion dramatically improves model closure, which has typically underestimated SOA in urban regions by 80% to 90% (19), but additional contributions from other sources of SOA precursors remain critical to model all observed SOA. Further chamber and modeling studies on SOA yields of aromatics with nine or more carbon atoms are important to reduce uncertainties in the SOA-forming potential of gasoline and diesel exhaust emissions and their overall contribution to SOA in urban regions. Additional

Table 1. Distribution of mass and SOA potential by chemical class for diesel exhaust, gasoline exhaust, and nontailpipe gasoline

Compound class	Weight by carbon, wtC%			Potential SOA formation, wt%		
	Diesel exhaust	Gasoline exhaust	Non-tailpipe gasoline	Diesel exhaust	Gasoline exhaust	Non-tailpipe gasoline
Total aliphatic	68 ± 8	58 ± 2	85 ± 4	47 ± 4	0.38 ± 0.07	0.9 ± 0.4
Straight-chain alkanes	7 ± 1	7.7 ± 0.3	20 ± 1	11 ± 2	0.09 ± 0.003	0.02 ± 0.001
Branched alkanes	23 ± 2	40 ± 1	60 ± 3	14 ± 2	0.12 ± 0.003	0.13 ± 0.01
Cycloalkanes (single straight alkyl chain)	2.5 ± 0.2	4.3 ± 0.1	1.03 ± 0.04	1.2 ± 0.3	0.13 ± 0.07	0.7 ± 0.4
Cycloalkanes [branched/multiple alkyl chain(s)]	18 ± 2	6.2 ± 0.3	5.0 ± 0.2	11 ± 2	0.04 ± 0.02	0.05 ± 0.03
Bicycloalkanes	13 ± 1	0	0	6 ± 1	0	0
Tricycloalkanes	4.8 ± 0.6	0	0	4 ± 1	0	0
Single-ring aromatics	19 ± 2	29 ± 1	2.7 ± 0.1	36 ± 9	96 ± 22	99 ± 6
Polycyclic aromatic compounds	4 ± 2	0.32 ± 0.02	0.0003	17 ± 8	3.2 ± 0.9	0.01 ± 0.01
Alkenes (straight, branched, cyclic)	0	3.6 ± 0.1	7.4 ± 0.3	0	0	0
Ethanol	0	6.9 ± 0.5	4.4 ± 0.4	0	0	0

The wt% by total mass for each source can be found in the *SI Appendix, Table S2*.

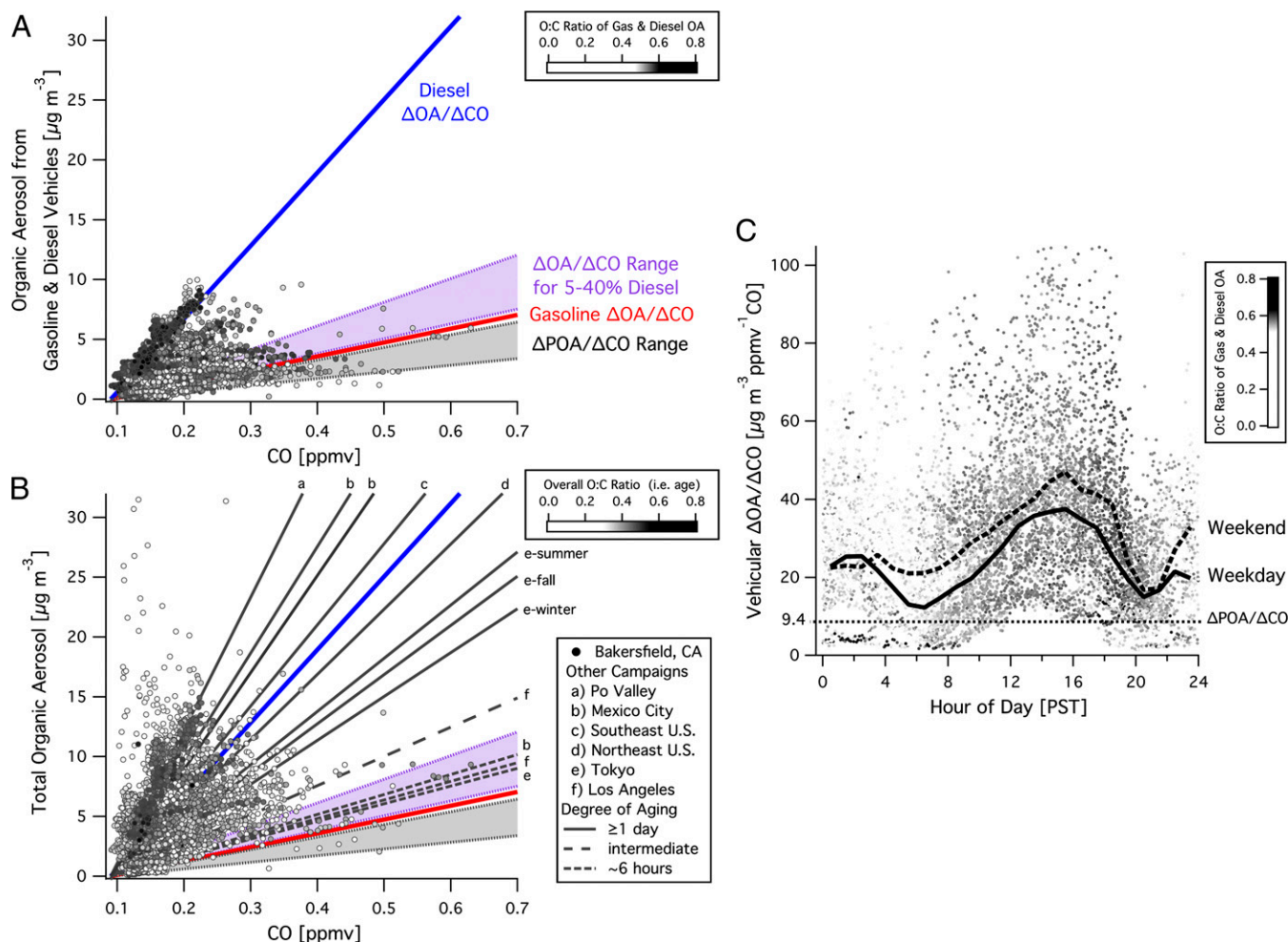


Fig. 4. Comparisons of OA vs. CO show the behavior of primary and secondary OA in the atmosphere and are used to examine vehicular OA and total OA in the San Joaquin Valley (Bakersfield, CA) and numerous other urban sites (6, 18, 21, 25–29). Photochemical aging increases $\Delta\text{OA}/\Delta\text{CO}$ ratios and is represented by increased O:C ratios shaded in each panel. (A) Best estimates for $\Delta\text{OA}/\Delta\text{CO}$ ratios expected for pure gasoline and diesel emissions are added to a $\Delta\text{POA}/\Delta\text{CO}$ value of $9.4 \mu\text{g m}^{-3} \text{ppmv}^{-1} \text{CO}$ to account for primary OA and shown with a range of $\Delta\text{POA}/\Delta\text{CO}$ values (21, 30). Vehicular OA is determined from AMS factor analysis and observations are well constrained at Bakersfield with the exception of the most aged air parcels, whose $\Delta\text{OA}/\Delta\text{CO}$ ratios are greater than expected for the mix of gasoline and diesel use. (B) Predicted $\Delta\text{OA}/\Delta\text{CO}$ slopes for a range of fuel mixtures ranging from 5% to 40% diesel agree with observations of relatively young aerosol in urban areas and vehicular OA at Bakersfield. Observed $\Delta\text{OA}/\Delta\text{CO}$ ratios increase with degree of aging and/or the influence of other SOA precursor sources that do not emit CO, which are prominent at Bakersfield and sites a–c. (C) Weekday and weekend diurnal averages of vehicular $\Delta\text{OA}/\Delta\text{CO}$ show greater ratios in the afternoon and over the weekend as a result of increased photochemical aging. Ratios are calculated with a 90 ppbv CO background, and SDs are shown in *SI Appendix, Fig. S8*.

discern organic SOA precursor contributions from gasoline vs. diesel given the variability in Los Angeles measurements (*SI Appendix, Fig. S9*) (6).

Nonvehicular anthropogenic and biogenic sources also lead to elevated $\Delta\text{OA}/\Delta\text{CO}$ ratios with higher slopes occurring in regions with large nonvehicular sources, such as Mexico City, the Southeast United States, and the Po Valley of Italy (Fig. 4B). $\Delta\text{OA}/\Delta\text{CO}$ ratios in the San Joaquin Valley span a broad range of values observed at other sites and the importance of other SOA sources is supported by elevated $\Delta\text{OA}/\Delta\text{CO}$ ratios in aged air masses and episodic contributions of low O:C OA from other sources (Fig. 4B and *SI Appendix, S10*) (6, 21, 25–29).

Our expanded measurement capabilities for gasoline and diesel compounds in the liquid fuels and the ambient atmosphere produce a more complete picture of SOA formation from motor vehicles. We provide the ability to predict emissions of SOA precursors and SOA formation that is consistent with fuel use data and ambient measurements. SOA from diesel sources outweighs gasoline contributions, and other sources provide sig-

nificant precursors in many urban regions. The inclusion of our insights will allow for the development of more effective pollution control policies and inform the design of future studies in the ambient atmosphere, laboratory experiments, and modeling efforts.

Materials and Methods

In situ gas-phase organic carbon measurements were made by using an automated sampling system coupled to a gas chromatograph and mass spectrometer at the Caldecott Tunnel in Oakland, CA, and in Bakersfield, CA, as part of the California at the Nexus of Air Quality and Climate Change (CalNex) campaign. Offline analyses used gas chromatography and MS to analyze the composition of gasoline and diesel fuels. Estimates of bulk SOA yields for each source and SOA yields for precursors without established yields were calculated via a Monte Carlo analysis by using all appropriate laboratory and modeling data. OA in Bakersfield was measured in situ with AMS and offline via FTIR analysis of filter samples. *SI Appendix* provides further detail on the instrumentation, analytical methods, and results presented in this work.

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