

VOC reactivity in central California: comparing an air quality model to ground-based measurements

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Abstract. Volatile organic compound (VOC) reactivity in central California is examined using a photochemical air quality model (the Community Multiscale Air Quality model; CMAQ) and ground-based measurements to evaluate the contribution of VOC to photochemical activity. We classify VOC into four categories: anthropogenic, biogenic, aldehyde, and other oxygenated VOC. Anthropogenic and biogenic VOC consist of primary emissions, while aldehydes and other oxygenated VOC include both primary anthropogenic emissions and secondary products from primary VOC oxidation. To evaluate the model treatment of VOC chemistry, we compare calculated and modeled OH and VOC reactivities using the following metrics: 1) cumulative distribution functions of NO_x concentration and VOC reactivity ($R_{OH,VOC}$), 2) the relationship between $R_{OH,VOC}$ and NO_x, 3) total OH reactivity ($R_{OH,total}$) and speciated contributions, and 4) the relationship between speciated $R_{OH,VOC}$ and NO_x. We find that the model predicts $R_{OH,total}$ to within 25–40% at three sites representing urban (Sacramento), suburban (Granite Bay) and rural (Blodgett Forest) chemistry. However in the urban area of Fresno, the model under predicts NO_x and VOC emissions by a factor of 2–3. At all locations the model is consistent with observations of the relative contributions of total VOC. In urban areas, anthropogenic and biogenic $R_{OH,VOC}$ are predicted fairly well over a range of NO_x conditions. In suburban and rural locations, anthropogenic and other oxygenated $R_{OH,VOC}$ relationships are reproduced, but calculated biogenic and aldehyde $R_{OH,VOC}$ are often poorly characterized by measurements, making evaluation of the model with available data unreliable. In central California,

30–50% of the modeled urban VOC reactivity is due to aldehydes and other oxygenated species, and the total oxygenated $R_{OH,VOC}$ is nearly equivalent to anthropogenic VOC reactivity. In rural vegetated regions, biogenic and aldehyde reactivity dominates. This indicates that more attention needs to be paid to the accuracy of models and measurements of both primary emissions of oxygenated VOC and secondary production of oxygenates, especially formaldehyde and other aldehydes, and that a more comprehensive set of oxygenated VOC measurements is required to include all of the important contributions to atmospheric reactivity.

1 Introduction

Atmospheric concentrations of volatile organic compounds (VOC) depend on primary anthropogenic and biogenic emissions, chemical formation and loss in the atmosphere (due to gaseous, aqueous and heterogeneous chemical reactions), and removal from the atmosphere due to dry and wet deposition. A wide suite of VOC with varying chemical properties is present in the atmosphere, and the chemical structures of these compounds affect reactivity (Atkinson, 2000). Once in the atmosphere, VOC oxidation leads to the formation of tropospheric ozone when in combination with sufficient concentrations of nitrogen oxides (NO_x=NO+NO₂) and sunlight. VOC oxidation also leads to the formation of secondary organic aerosol via gas-to-particle conversion (e.g., Odum et al., 1997), aqueous phase chemistry (e.g., Ervens et al., 2004), and/or oligomerization (e.g., Kalberer et al., 2004).

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Reducing anthropogenic VOC and/or NO_x precursor emissions has been the main method of controlling ozone in the United States over the past several decades (National Research Council, 1991). In some regions, such as the Los Angeles area in southern California, control of anthropogenic VOC from automobiles and industrial processes has proven effective at reducing ozone exceedances over the past three decades, indicating that VOC were the limiting factor in ozone production in this region (Milford et al., 1989; Martien and Harley, 2006). In other regions with high emissions of biogenic VOC, NO_x control has been more efficient at reducing ozone (Trainer et al., 1988; Sillman et al., 1990; Cardelino and Chameides, 1990; Han et al., 2005). In central California, differences between weekday and weekend ozone concentrations indicate that ozone production in urban regions are NO_x -saturated while more remote areas such as the southern San Joaquin Valley and the Sierra Nevada are more NO_x -sensitive (Blanchard and Fairley, 2001; Marr and Harley, 2002; Murphy et al., 2006a, b).

Certain VOC such as alkenes are more reactive than alkanes and, in the hours immediately after emission, they can contribute more to ozone formation per unit mass. As a result, VOC have been ranked according to their ozone formation potential and incremental contributions to ozone production (e.g., Carter, 1994). One element of the overall control strategy has been to use modeling studies to identify and reduce VOC with high ozone formation potential (Russell et al., 1995; Avery, 2006).

This approach requires an assessment of regional VOC reactivity and applicable control strategies, which are often performed using air quality models. Recent studies have used ground-based VOC measurements to assess VOC reactivity and validate emission inventories. These results vary by location, but studies suggest that urban alkanes and alkenes tend to be under estimated by current inventories, while aromatics such as toluene tend to be over estimated (Jiang and Fast, 2004; Choi et al., 2006; Velasco et al., 2007; Warnecke et al., 2007). This suggests that proposed control strategies targeting specific VOC species might be less effective than predicted due to errors in emissions inventories.

In our prior analysis of climate effects on ozone in California (Steiner et al., 2006), we used ground-based measurements of ozone concentrations to evaluate model performance and found that the model reproduced the spatial distributions and daily maximum ozone concentrations in the San Joaquin Valley and in the eastern portion of the San Francisco Bay area. Here, we utilize ground-based VOC and NO_x measurements to develop metrics to evaluate the representation of VOC reactivity in a regional chemical transport model. We do this by comparing calculated and modeled 1) cumulative distribution functions of NO_x and $R_{\text{OH},\text{VOC}}$, 2) the relationship between $R_{\text{OH},\text{VOC}}$ and NO_x , 3) speciated contributions to $R_{\text{OH},\text{total}}$, and 4) the relationship between speciated $R_{\text{OH},\text{VOC}}$ and NO_x . Additionally, we provide a discussion of the present day and future VOC component of ozone produc-

tion in order to understand more fully the chemical factors that may change under future climate and guide the development of air quality control strategies.

2 Methods

2.1 Model simulations

We investigate VOC reactivity for an episodic air quality simulation in central California using the Community Multiscale Air Quality model (CMAQ, Byun and Ching, 1999). We compare two five-day simulations, where the first simulation represents an ozone episode during the summer of 2000 (29/7/2000–3/8/2000). During this time period, stagnant conditions and warm temperatures led to high ozone concentrations in the southern San Joaquin Valley and in the San Francisco Bay area. Further details of this simulation, including model domain, resolution, and base case emissions are described in Steiner et al. (2006) and are described only briefly here. The model domain (96 grid cells in the east-west direction and 117 grid cells in the north-south direction) is focused on central California (approximately 34.5°N to 39°N and 118.5°W to 123°W), with a horizontal resolution of 4 km. There are 27 terrain-following vertical sigma layers from the surface to approximately 100 mb, where the first near-surface layer is approximately 20 m. Concentrations presented here are averaged over the first eleven vertical layers of the atmosphere, representing approximately 500 m above ground level. This vertical averaging was utilized to approximate an average boundary layer concentration and account for vertical and horizontal mixing.

Mesoscale meteorological simulations (MM5) are used to drive the CMAQ simulations, and were performed as part of the Central California Ozone Study (CCOS). MM5 testing over the region indicated that the model performs optimally when including the NOAA LSM, Eta boundary layer scheme, and FDDA nudging techniques (Wilczak et al., 2004). The use of the nudged simulation yields good agreement between observed and modeled wind speeds (within 0.2 m/s averaged over the full domain, and a maximum difference of 1–2 m/s) and wind direction (Wilczak et al., 2004; results available online at <http://www.etl.noaa.gov/programs/modeling/ccos/data/>). Observed and modeled boundary layer heights in the San Joaquin Valley agree within about 20%, and the model captures the spatial distribution of variable boundary layer height well compared to observations. Greater discrepancies between the observed and modeled boundary layer height are noted in the coastal regions; however, the analysis here focuses on the Central Valley and Sierra Mountains.

Anthropogenic emission estimates are based on point, area and mobile sources from gridded emission inventory data supplied by the California Air Resources Board (CARB) for a summer 1990 air pollution episode (Marr et al., 2002),

and scaled to include changes due to growth and technology to the year 2000 (Marr, 2002). The mobile portion of this inventory was further adjusted to reflect on-road and ambient measurements of pollutant emissions, the ratios of these emissions, and different activity patterns by time of day and day of week for gasoline versus diesel engines. This resulted in an increase in VOC emissions and a decrease in NO_x emissions relative to the original CARB inventory (Marr et al., 2002; Harley et al., 2005). Biogenic VOC emissions for these simulations include isoprene, monoterpenes, and methylbutenol, with emissions estimated hourly over the model domain using the BEIGIS modeling system (Scott and Benjamin, 2003).

The second simulation emulates future climate by using temperature changes estimated from a $2\times\text{CO}_2$ regional climate simulation (Snyder et al., 2002). An increase in temperature is predicted from the doubling of CO_2 , and this increase is applied to the present-day model atmospheric temperatures. We allow temperature to affect the simulation in three ways: 1) by altering the kinetics of the gas phase chemistry, 2) by increasing the atmospheric water vapor content due to the increased temperature, and 3) by increasing the emissions of biogenic VOC compounds, which generally increase with temperature. This simulation does not include a change in vegetation or land cover due to the temperature increase, which could alter biogenic VOC emissions through changes in vegetation species composition and biomass. The changes to temperature, water vapor, and biogenic VOC emissions are referred to as the “combined climate simulation” and are detailed in Steiner et al. (2006), therefore we provide only a brief summary of this simulation here. Temperature increases about 1.5°C along the coast to approximately 4°C in the Sierra Nevada Mountains, with a similar spatial impact on atmospheric moisture. The higher temperatures increase biogenic VOC emissions by 20–45%. Together, these three temperature impacts are included in a single simulation and we utilize this combined climate simulation to estimate the impact of future climate on VOC reactivity.

2.2 VOC classification

Gas phase chemistry is modeled in CMAQ using the SAPRC99 mechanism, which lumps VOC into functional groups based on structure and reactivity (Carter, 2000). For this analysis, we group VOC into four categories reflecting different primary emission sources and secondary chemistry in the atmosphere: anthropogenic, biogenic, aldehydes and other oxygenated VOC. Anthropogenic and biogenic VOC reactivity are solely from primary sources and reflect the majority of primary VOC emissions provided to the chemistry model. Aldehydes and other oxygenated VOC reactivity include a minor primary emission source plus a secondary source from oxidation of all VOC in the atmosphere.

Anthropogenic VOC includes ethene (ETHE), five lumped categories of alkanes (ALK1–ALK5), and two lumped cate-

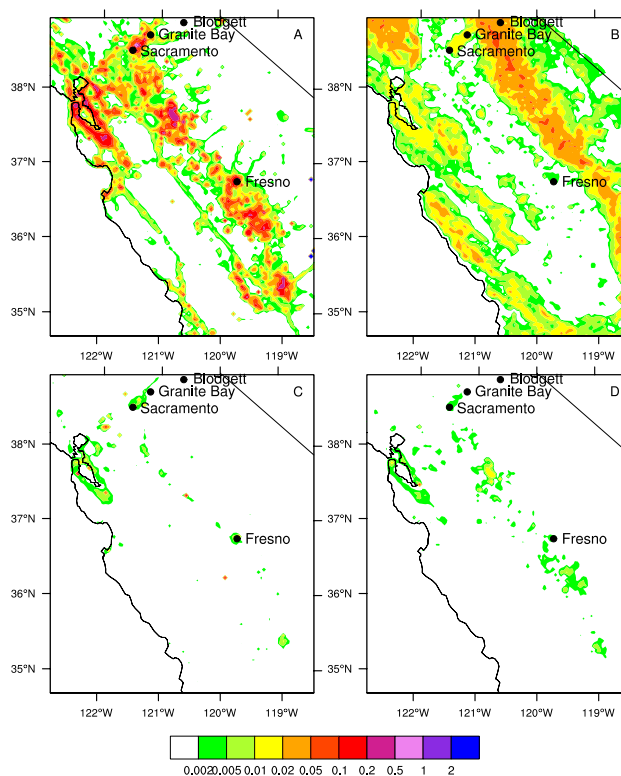


Fig. 1. Primary, present-day emissions (mol s^{-1}) for four VOC categories: (a) anthropogenic, (b) biogenic, and (c) aldehyde and (d) other oxygenated VOC. All figures display the average emission flux at 09:00 LT.

gories each for olefins (OLE1, OLE2) and aromatics (ARO1, ARO2). Model anthropogenic species are listed in Table 1 along with average morning contributions to total VOC reactivity. The modeled anthropogenic VOC category represents the total amount of anthropogenic hydrocarbon primary emissions (spatial distribution shown in Fig. 1a) based on gridded emission inventory data supplied by the California Air Resources Board and modified by Marr et al. (2002) and Harley et al. (2005). Emissions of anthropogenic VOC are dominated by higher alkanes (ALK3–ALK5) and ethene.

The second category, biogenic VOC, includes primary emissions of isoprene, monoterpenes, and methylbutenol from natural, agricultural, and urban areas estimated with the BEIGIS model (Scott and Benjamin, 2003) (Fig. 1b). Anthropogenic and biogenic VOC are emitted in similar magnitude although their spatial distributions differ, with anthropogenic VOC emitted from dense urban areas and transportation corridors and biogenic species emitted from California’s natural and crop vegetation. We note that urban locations in the northern portion of the modeling domain (e.g. the San Francisco Bay area and the Sacramento metropolitan area) still have significant amounts of biogenic VOC emissions from isoprene, while agricultural areas in the Central Valley are low in both anthropogenic and biogenic VOC emissions.

Table 1. Average morning modeled $R_{OH,VOC}$ for four VOC categories: anthropogenic, biogenic, aldehydes, and other oxygenated VOC. k_{OH} values for anthropogenic VOC reactions (alkanes, olefins and aromatics) are in $\text{ppm}^{-1} \text{min}^{-1}$ (Carter, 2000).

CMAQ Category	Description	Sacramento	Fresno	Granite Bay	Blodgett Forest
Anthropogenic		2.635	1.988	1.590	0.146
ETHENE	Ethene	0.516	0.330	0.326	0.036
ALK1	alkanes/non-aromatics, $200 < k_{OH} < 500$	0.015	0.046	0.017	0.004
ALK2	alkanes/non-aromatics, $500 < k_{OH} < 2500$	0.028	0.028	0.019	0.004
ALK3	alkanes/non-aromatics, $2500 < k_{OH} < 5000$	0.088	0.133	0.066	0.005
ALK4	alkanes/non-aromatics, $5000 < k_{OH} < 10\,000$	0.231	0.238	0.177	0.010
ALK5	alkanes/non-aromatics, $k_{OH} > 10\,000$	0.285	0.282	0.196	0.007
OLE1	Alkenes with $k_{OH} < 70\,000$, usu. terminal alkenes	0.506	0.230	0.305	0.036
OLE2	Alkenes with $k_{OH} > 70\,000$, usu. internal or disubstituted alkenes	0.364	0.215	0.017	0.016
ARO1	Aromatics with $k_{OH} < 20\,000$, usu. toluene and monoalkyl benzenes	0.131	0.104	0.101	0.011
ARO2	Aromatics with $k_{OH} > 20\,000$, usu. xylenes and polyalkyl benzenes	0.546	0.429	0.384	0.019
Biogenic		1.893	0.392	0.318	6.124
ISOP	Isoprene	1.473	0.349	0.240	2.570
TRP	Biogenic alkenes less isoprene	0.356	0.046	0.022	0.359
MBO	2-methyl-3-buten-2-ol	0.236	0.049	0.037	2.885
Aldehydes		1.642	1.492	1.506	1.071
HCHO	Formaldehyde	0.908	0.629	0.862	0.411
CCHO	Acetaldehyde and glycolaldehyde	0.354	0.444	0.323	0.309
RCHO	Lumped C3+ aldehydes	0.321	0.353	0.270	0.321
BALD	Aromatic aldehydes	0.012	0.010	0.011	0.001
GLY	Glyoxal	0.014	0.015	0.012	0.003
MGLY	Methyl glyoxal & higher α -dicarbonyls	0.033	0.041	0.028	0.026
Other Oxygenated		0.399	0.338	0.174	0.413
ACET	Acetone	0.003	0.008	0.002	0.004
MEK	Ketones with $k_{OH} < 5 \times 10^{-12} \text{ cm}^3 \text{ molec}^{-1} \text{ s}^{-1}$	0.010	0.029	0.009	0.008
MEOH	Methanol	0.002	0.003	0.001	0.001
PHEN	Phenol	0.0003	0.0002	0.0001	0.00002
CRES	Cresols	0.010	0.010	0.009	0.001
MVK	Unsaturated ketones	0.049	0.043	0.029	0.070
MACR	Methacrolein	0.154	0.050	0.026	0.105
IPRD	Unsaturated aldehydes other than acrolein and methacrolein	0.105	0.076	0.043	0.153
PROD2	Ketones with $k_{OH} < 5 \times 10^{-12} \text{ cm}^3 \text{ molec}^{-1} \text{ s}^{-1}$	0.066	0.119	0.055	0.071

The third category, aldehydes, includes primary emissions sources as well as the secondary oxidation products of anthropogenic and biogenic VOC. In the model, we consider six aldehyde categories, detailed in Table 1. Primary anthropogenic emissions of aldehydes in the model are shown in Fig. 1c and include formaldehyde (HCHO), acetaldehyde (CCHO), benzaldehyde, other C3+ aldehydes (RCHO), glyoxal (GLY) and methylglyoxal (MGLY). Primary urban emissions of aldehydes are dominated by HCHO and CCHO.

The fourth category, other oxygenated VOC, includes nine model categories (Table 1). Primary emissions (Fig. 1d) include acetone, C4+ ketones (MEK), phenol, cresol and methanol, and the dominant emissions of oxygenated species in urban regions are MEK, methanol and acetone, although aldehyde and other oxygenated VOC emissions tend to be 1–2 orders of magnitude less than anthropogenic or biogenic VOC emissions. The primary emissions of aldehydes

and other oxygenated VOC in the model are solely anthropogenic and are located near urban and industrial areas. These anthropogenically-emitted species can also be produced in the atmosphere as secondary products from the oxidation of anthropogenic and biogenic VOC. Additional other oxygenated VOC that are a result of atmospheric oxidation only include PROD2 (a lumped oxygenated VOC that represents secondary products from both biogenic and anthropogenic VOC oxidation), MVK, MACR, and IPRD (resulting primarily from biogenic VOC oxidation).

We note a few caveats to the emissions and description of the other oxygenated VOC category that suggest this VOC category may be underestimated. Several compounds grouped in the anthropogenic VOC category include primary emissions of oxygenated species such as ethanol, other alcohols, glycols, and some ethers. One example is methyl tert-butyl ether, which is included as an anthropogenic VOC

Table 2. Measured chemical species in each of the four VOC categories, ranked in order of reaction rate. List of species used to estimate calculated $R_{OH,VOC}$ at each of the following sampling locations: Sacramento PAMS (2000 data), Blodgett Forest Research Station (22/07/2001–19/09/2001), and Granite Bay (16/07/2001–17/09/2001), with average afternoon concentrations (ppb). CMAQ lumped categories for each measured species are indicated for comparison with modeled $R_{OH,VOC}$.

VOC Categories and Chemical Species	$k_{OH}^a \times 10^{-12}$ (cm ³ molec ⁻¹ s ⁻¹)	PAMS ^b	Blodgett ^b	Granite Bay ^b	CMAQ
Anthropogenic VOC					
1,3-butadiene	68.3			0.026	OLE2
trans-2-pentene	67.0	0.107		0.016	OLE2
trans-2-butene	66.0	0.130		0.009	OLE2
cis-2-pentene	65.0	0.099		0.007	OLE2
2-methyl-1-butene	60.7			0.019	OLE2
styrene	58.0	0.089			OLE2
cis-2-butene	57.9	0.121			OLE2
1,3,5-trimethylbenzene	57.5	0.836			ARO2
cyclopentene	57.0			0.004	OLE2
2-methylpropene (isobutene)	52.8		0.027	0.081	OLE2
1,2,4-trimethylbenzene	37.5	0.084			ARO2
1,2,3-trimethylbenzene	32.7	0.174			ARO2
1-butene	32.2	0.144	0.038	0.034	OLE1
1-pentene	31.4	0.118		0.036	OLE1
propene	30.0	1.029		0.367	OLE1
m-xylene	23.6	1.792 ^c		0.240	ARO2
m-ethyltoluene	17.1	0.018			ARO2
p-xylene	14.3	1.792 ^c		0.133	ARO2
m-diethylbenzene	14.2				ARO2
o-xylene	13.7	0.296		0.123	ARO2
o-ethyltoluene	12.3	0.184			ARO2
n-undecane	12.9	2.021			ALK5
p-ethyltoluene	11.4	0.073			ARO2
n-decane	11.2	0.070			ALK5
methylcyclohexane	10.0	0.170			ALK5
n-nonane	9.91	0.063			ALK5
n-octane	8.60	0.086			ALK5
3-methylheptane	8.56	0.073			ALK5
2-methylheptane	8.28	0.903			ALK5
p-diethylbenzene	8.11	0.097			ARO2
3-methylhexane	7.16	0.232			ALK5
ethylbenzene	7.10	0.497	0.021	0.080	ARO1
cyclohexane	7.10	0.193			ALK5
2,3,4-trimethylpentane	7.10	0.010			ALK5
n-heptane	6.98	0.208		0.243	ALK5
methylcyclopentane	6.80 ^d	0.353			ALK4
isopropylbenzene	6.50	0.085			ARO1
toluene	6.40	0.340	0.085	0.488	ARO1
2,3-dimethylpentane	6.10	0.119			ALK5
propyne (methyl acetylene)	5.90		0.022	0.022	ALK4
2,3-dimethylbutane	5.75	0.147			ALK4
n-propylbenzene	5.70	0.187			ARO1

in the ALK3 category (Table 2) despite the oxygenated nature of the compound. However, MTBE and other similar compounds tend to have low reactivity and comprise a small portion of the total anthropogenic VOC emissions. Addi-

tionally, we have not included primary biogenic emissions of oxygenated VOC. Crops and natural vegetation may directly emit some aldehydes and other oxygenated VOC including acetone, methanol, acetaldehyde and a wide variety of other

Table 2. Continued.

Anthropogenic VOC, cont.					
3-methylpentane	5.40	0.294	0.011	0.351 ^e	ALK4
n-hexane	5.40	0.276	0.015	0.135	ALK4
3-methyl-1-butene	5.32			0.016	OLE1
2-methylpentane	5.30	0.626	0.022	0.351 ^e	ALK4
2-methylhexane	5.10	0.154			ALK5
2,4-dimethylpentane	5.00	0.091			ALK4
cyclopentane	4.92	0.122		0.041	ALK4
n-pentane	3.91	0.567	0.038	0.259	ALK4
isopentane	3.70	1.579	0.067	0.665	ALK4
methyl tert-butyl ether	2.96			1.943	ALK3
2,2-dimethylbutane	2.34	0.112			ALK4
n-butane	2.34	0.779	0.080	0.443	ALK3
isobutane	2.15	0.498	0.045	0.300	ALK3
benzene	1.17	0.136	0.227	0.152	BENZ
propane	1.07	2.406		1.489	ALK2
acetylene	0.78	1.587			ALK2
ethene	0.236	1.648			ETHE
ethane	0.236	2.017			ALK1
2,2-dimethylpropane	0.0000179			0.005	ALK2
Biogenic VOC					
d-limonene	171.0			0.031	TRP1
isoprene	103.0	1.521	0.262	0.516	ISOP
3-carene	88.0			0.010	TRP1
b-pinene	80.5		0.125		TRP1
methylbutenol	64.0		1.135		MBO
a-pinene	55.1		0.123	0.035	TRP1
Aldehydes					
hexanal	31.7 ^f			0.276	RCHO
pentanal	29.9 ^f		0.064	0.118	RCHO
acetaldehyde	15.8	1.398	1.386	1.417	CCHO
formaldehyde	9.20 ^g	2.658			HCHO
Oxygenated VOC					
3-methylfuran	93.5 ^h			0.009	
methacrolein	35.0		0.289	0.213	MACR
methyl vinyl ketone	18.8		0.331	0.309	MVK
ethanol	3.27			2.257	ALK3
methyl ethyl ketone	1.15			0.213	MEK
methanol	0.94		10.735	7.467	MEOH
acetone	0.22		2.507	2.911	ACET

^a k_{OH} values based on Atkinson (1994) unless noted otherwise.

^b Average AM (06:00–08:00 a.m. local time) concentrations over the sampling period, in ppb. PAMS data listed is an average of the four sites in the Sacramento metropolitan area.

^c PAMS stations report co-eluting m- and p-xylene as a single sum, therefore concentrations shown here are the sum of these two isomers.

^d Estimated from Kwok and Atkinson (1995).

^e Granite Bay measurements include total methylpentanes, here listed as 2-methylpentane.

^f Papagni et al. (2000)

^g Carter (2000)

^h Atkinson et al. (1989)

alcohols, ketones, esters, and ethers (Kesselmeier and Staudt, 1999; Schade and Goldstein, 2001). However, measurements and modeling efforts have yet to thoroughly quantify these species on the scale needed for regional emissions modeling, and they are not included in our biogenic VOC emissions inventory.

2.3 Ground-based measurements

We utilize ground-based measurements at four locations in central California (marked in Fig. 1). For two locations, Sacramento and Fresno, we use observations from the Photochemical Assessment Monitoring Stations (PAMS) network. At eleven sites in four different geographic regions in California (in Sacramento, Fresno, Kern and Madera counties), ground-based O₃, CO, NO_x, and VOC measurements have been collected from 1994 to the present. The VOC dataset consists of a long-term record of 55 VOC species (see Table 2) collected at 3-h intervals, including alkanes, alkenes and aromatics (data available at www.epa.gov/air/oaqps/pams). This dataset predominantly reflects the anthropogenic VOC category as defined in Sect. 2.2, with isoprene measurements at these locations representing only a portion of the biogenic VOC category. Selected PAMS sites measure a suite of carbonyl species, including formaldehyde and acetaldehyde, and we include these measurements when available. Here, we include concurrent VOC and NO_x data from the summer of 2000 for the Sacramento (four measurement sites) and Fresno (three measurement sites) metropolitan areas. While the PAMS data have been known to have several artifacts in the hydrocarbon and aldehyde samples due to the nature of canister samples (McClenney et al., 2002), we utilize this data here in the absence of in-situ VOC sampling in the region and note that these results should be reviewed with caution with respect to the carbonyl species.

A third sampling location, Granite Bay, was established in the summer of 2001 to evaluate the transport of pollutants at the eastern edge and downwind of Sacramento. NO₂ was measured from 19/07/2001–15/09/2001, and NO_x concentrations are estimated from NO₂ photolysis, measured ozone, and modeled RO₂ (Cleary et al., 2005). A suite of VOC was measured at the same location, including 28 anthropogenic, 4 biogenic, 4 aldehydes and 7 other oxygenated VOC (Millet et al., 2005). As shown in Table 2, measurements in the anthropogenic, biogenic and other oxygenated VOC categories represent a significant portion of the modeled VOC reactivity, while the aldehyde category is missing the important contribution of HCHO.

The fourth location, Blodgett Forest Research Station, is located in the Sierra Nevada mountains approximately 75 km ENE of Sacramento, and represents a rural site where concentrations and fluxes of biogenic VOC compounds have been measured during summer months since 1997 (e.g., Lamanna and Goldstein, 1999). Additional VOC concentrations were measured in 2001 in conjunction with Granite

Bay measurements. These include a limited number of anthropogenic (10 species), aldehydes (2 species), other oxygenated VOC (4 species), and the dominant biogenic VOC (4 species) (Table 2). While the number of VOC species measured at Blodgett is small compared to other sites, the measurements comprise the major biogenic species emitted and are likely representative of VOC chemistry at the site. Additionally, NO_x has been measured at the site (Day et al., 2003, 2007), and data were filtered for concurrent NO_x and VOC measurements.

2.4 R_{OH,total} and R_{OH,VOC} definitions

Here, we define the total OH reactivity (R_{OH,total}, s⁻¹) as OH loss with the following species: CO, CH₄, NO₂, anthropogenic VOC, biogenic VOC, aldehydes, and other oxygenated VOC:

$$R_{\text{OH,total}} = k_{\text{CO+OH}} [\text{CO}] + k_{\text{CH}_4+\text{OH}} [\text{CH}_4] + k_{\text{NO}_2+\text{OH}} [\text{NO}_2] \\ + \sum_{n=1, \text{navoc}} k_{n+\text{OH}} [\text{aVOC}_{n,\text{meas}}] + \sum_{n=1, \text{nbvoc}} k_{n+\text{OH}} [\text{bVOC}_{n,\text{meas}}] \\ + \sum_{n=1, \text{naldvoc}} k_{n+\text{OH}} [\text{aldVOC}_{n,\text{meas}}] + \sum_{n=1, \text{noxvoc}} k_{n+\text{OH}} [\text{oxVOC}_{n,\text{meas}}] \quad (1)$$

where navoc (nbvoc, naldvoc, noxvoc) is the number of anthropogenic (biogenic, aldehyde, other oxygenated) VOC species, $k_{n+\text{OH}}$ is the rate coefficient (cm³ molecules⁻¹ s⁻¹) and [VOC_{*n*,meas}] is the measured VOC concentration. VOC reactivity (R_{OH,VOC}) is calculated from the four VOC categories only:

$$R_{\text{OH,VOC}} = \sum_{n=1, \text{navoc}} k_{n+\text{OH}} [\text{aVOC}_{n,\text{meas}}] + \sum_{n=1, \text{nbvoc}} k_{n+\text{OH}} [\text{bVOC}_{n,\text{meas}}] \\ + \sum_{n=1, \text{naldvoc}} k_{n+\text{OH}} [\text{aldVOC}_{n,\text{meas}}] + \sum_{n=1, \text{noxvoc}} k_{n+\text{OH}} [\text{oxVOC}_{n,\text{meas}}] \quad (2)$$

Reactivities are calculated from ground-based CO, CH₄, NO₂ and VOC concentrations and kinetic rate coefficients calculated from measured temperature and pressure (Table 2).

Modeled reactivities are determined from integrated reaction rates calculated by CMAQ, which estimates the loss rate of each species that reacts with OH integrated over each hour:

$$\text{OH}_{\text{loss}}(t + \Delta t) = \text{OH}_{\text{loss}}(t) + \int_t^{t+\Delta t} k_{n+\text{OH}} [\text{OH}] [n] dt \quad (3)$$

$$\text{VOC}_{\text{loss}}(t + \Delta t) = \text{VOC}_{\text{loss}}(t) + \int_t^{t+\Delta t} k_{n+\text{OH}} [\text{OH}] [\text{VOC}] dt \quad (4)$$

where OH_{loss} and VOC_{loss} is the integrated reaction rate of each species (ppb hr⁻¹), Δ*t* is the integration time of the chemical solver, $k_{n+\text{OH}}$ is the kinetic rate constant of species

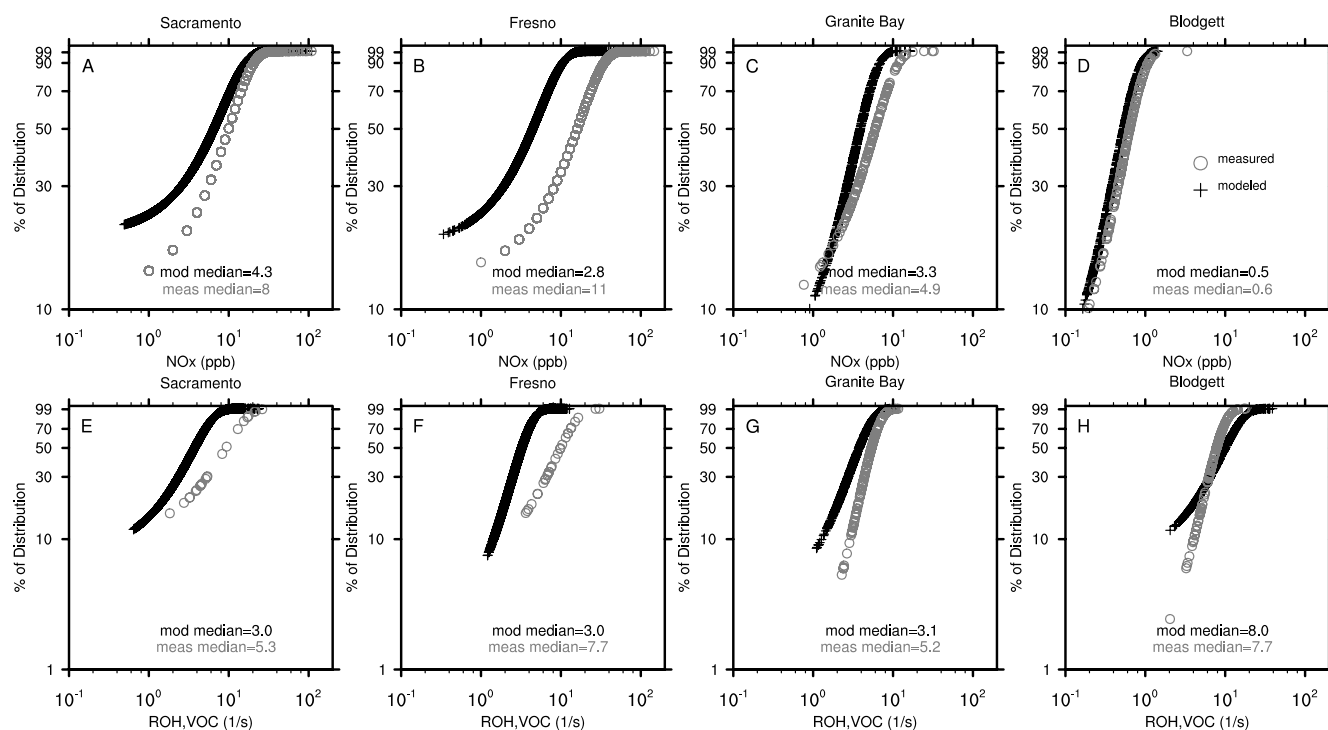


Fig. 2. Measured (gray circles) versus modeled (black crosses) cumulative distribution functions for (a) Sacramento NO_x, (b) Fresno NO_x, (c) Granite Bay NO_x, (d) Blodgett NO_x, and (e) Sacramento R_{OH,VOC}, (f) Fresno R_{OH,VOC}, (g) Granite Bay R_{OH,VOC}, and (h) Blodgett R_{OH,VOC}.

n with the OH radical, and [OH] and $[n]$ represent concentrations at time t (Byun and Ching, 1999). For OH loss, n includes CO, CH₄, NO₂, and the four VOC categories. For VOC loss, n includes only the four VOC categories. This representation provides an accurate method of assessing the loss rates of each species with OH. In order to compare the model loss rates with observations in California and elsewhere, we assume that the modeled OH concentrations are relatively stable over the timescale of the hour of integration and calculate the modeled OH and VOC reactivity as:

$$R_{\text{OH,total}} = \frac{\text{OH}_{\text{loss}}}{[\text{OH}]} \quad (5)$$

$$R_{\text{OH,VOC}} = \frac{\text{VOC}_{\text{loss}}}{[\text{OH}]} \quad (6)$$

For R_{OH,VOC}, we calculate this quantity as the sum of individual species for each of the four VOC categories, considering only daytime modeled and calculated reactivity calculations, when OH dominates radical reactivity with VOC. We note the definitions of R_{OH,total} and R_{OH,VOC} are closely connected, as changes in concentrations of CO, NO₂ and CH₄ will impact OH concentrations and in turn the rate of VOC reactions. Therefore, R_{OH,VOC} is a subset of R_{OH,total}.

3 Results and discussion

We use ground-based measurements of NO₂, CO, and VOC to calculate R_{OH,total} and R_{OH,VOC} at four locations in central California. We compare calculated and modeled reactivities using four metrics: 1) cumulative distribution functions of NO_x concentrations and R_{OH,VOC}, 2) the relationship between R_{OH,VOC} and NO_x concentration, and 3) speciated contributions to R_{OH,total}. Additionally, at the Sacramento location we evaluate relationships between speciated R_{OH,VOC} and NO_x. These comparisons evaluate the model performance in order to gain insight into the fast chemistry and the importance of VOC sources in central California.

3.1 Cumulative distribution functions of R_{OH,VOC} and NO_x

To determine if the model is reproducing conditions similar to those estimated by ground-based measurements, we evaluate the cumulative distribution functions (CDFs) of NO_x and R_{OH,VOC}. Comparing measured versus modeled CDFs can indicate if the model is able to reproduce the general mode of NO_x and R_{OH,VOC} in the atmosphere. The CDFs shown in Fig. 2 are based on summertime (01/06/2000–01/09/2000) daytime (6 a.m.–6 p.m. local time) data at the four measurement locations described in Sect. 2.4, including two urban locations (Sacramento and Fresno), one suburban location (Granite Bay), and one rural location (Blodgett Forest). Re-

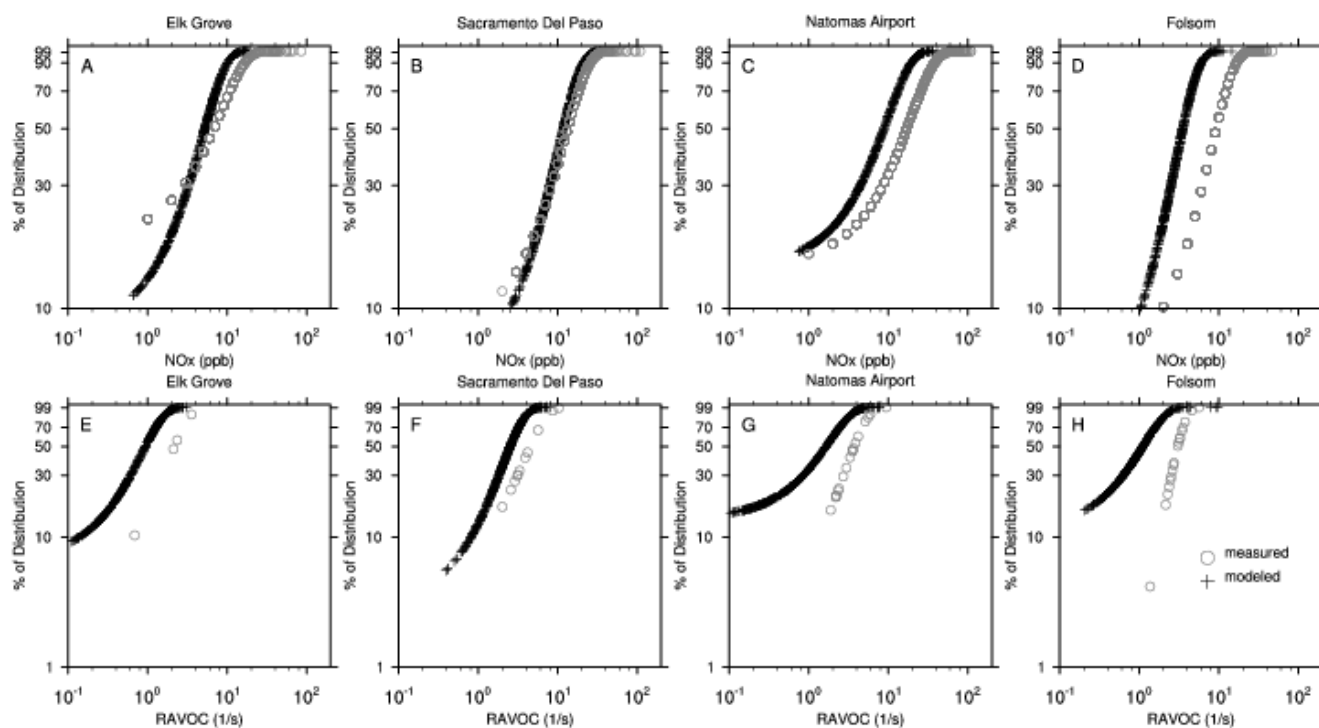


Fig. 3. Measured (gray circles) versus modeled (black crosses) cumulative distribution functions for four Sacramento locations (**a**) upwind location, Elk Grove NO_x , (**b**) near-urban location, Sacramento Del Paso Manor NO_x , (**c**) near-urban location, Natomas Airport NO_x , (**d**) downwind location, Folsom NO_x , (**e**) Elk Grove R_{AVOC} , (**f**) Sacramento Del Paso Manor R_{AVOC} , (**g**) Natomas Airport R_{AVOC} , and (**h**) Folsom R_{AVOC} .

activities calculated from Granite Bay and Blodgett forest are based on measurements conducted during 2001.

In Sacramento during the daytime, the NO_x CDF (Fig. 2a) indicates that the model predicts less NO_x than observed at the lower end of the NO_x concentrations, while the Fresno results indicate that the model predicts less NO_x at all times and over the full width of the distribution (Fig. 2b). Mean daytime NO_x concentrations are less than those measured (Sacramento measured daytime mean $\text{NO}_x=10$ ppb, modeled mean $\text{NO}_x=7$ ppb; Fresno measured daytime mean $\text{NO}_x=23.5$ ppb, modeled mean $\text{NO}_x=5$ ppb). A portion of this daytime discrepancy is likely due to the NO_2 chemiluminescence measurement technique employed at the EPA PAMS monitoring locations. This technique is well known to include PAN and other higher oxides of nitrogen in the reported NO_2 and thus to have a high bias (e.g., Winer et al., 1974). Recent measurements using current commercial implementations of the technique side by side with instruments that specifically measure NO_2 show that high biases of 20–50% occur routinely (Dunlea et al., 2007; Steinbacher et al., 2007).

Other possibilities for measured and modeled discrepancies include the vertical averaging technique, the impacts of dilution due to meteorological parameters, and the location of urban monitoring stations. Vertical averaging is not a sig-

nificant factor, as comparisons of concentrations in the surface layer only (surface to 20 m) only impact the top 90% of the distribution, and indicate that the model still underestimates the majority of the distribution in urban locations (results not shown). Because measured and modeled boundary layer heights are similar in Sacramento (daily maxima approximately 500–600 m) and Fresno (daily maxima approximately 600–800 m) during the simulation time period and daytime wind speeds and wind directions agree fairly well (Wilczak et al., 2004; data available at <http://www.etl.noaa.gov/programs/modeling/ccos/data/>), dilution is not likely a strong factor in the differences in the measured and modeled distribution functions.

Past studies indicate that some PAMS measurement locations are likely too close to mobile sources to be regionally representative (e.g., Murphy et al., 2006a), leading to elevated measurements that should not be reproduced in the relatively large size of the model grid cell (4 km \times 4 km). There are several PAMS sites at the two urban locations, and a portion of the measured versus modeled distribution functions could be influenced by the averaging area of the model grid cells around these different locations. To evaluate the impact of spatial averaging, we examine the NO_x and anthropogenic $R_{\text{OH,VOC}}$ (R_{AVOC}) at four individual locations in Sacramento and compare the observations to the model grid cells in the

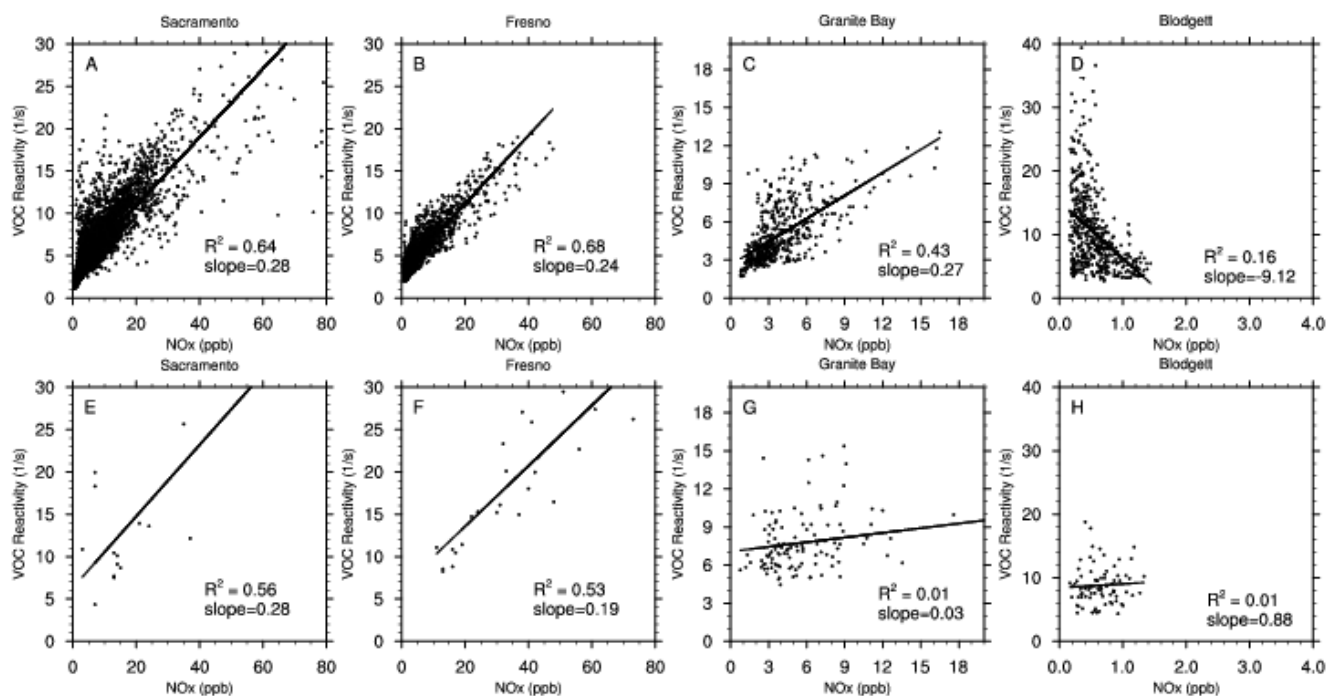


Fig. 4. Total $R_{OH,VOC}$ (s^{-1}) as a function of NO_x concentrations (ppb), as modeled (a–d) and measured (e–h) in four locations: Sacramento (a, e), Fresno (b, f), Granite Bay (c, g), and Blodgett Forest (d, h). See Table 1 for VOC included in these calculations.

immediate vicinity (Fig. 3). The four PAMS sites represent upwind (Type 1 sites), near urban (Type 2 sites), and downwind locations (Type 3 sites). Elk Grove (Fig. 3a and e) corresponds to an upwind location, while Sacramento Del Paso (Fig. 3b and f) and Natomas Airport (Fig. 3c and g) represent locations near maximum emissions in the urban area, and Folsom (Fig. 3d and h) is located downwind of the urban area and near the location of maximum ozone. The site locations relative to the urban area (e.g., upwind or downwind) do not have a strong impact on the NO_x and R_{AVOC} CDFs. In the upwind location of Elk Grove, the model captures the median NO_x accurately but does not capture the minimum or the maximum NO_x concentrations. At the Del Paso urban site (Fig. 3b) the model accurately reproduces the measured CDF while at Natomas Airport (a second urban location) the model consistently predicts less NO_x than observed. At the downwind site, Folsom, the model predicts less NO_x and R_{AVOC} than observed.

Because these factors do not show a significant impact, this suggests that the discrepancy between measured and modeled NO_x distributions are due to differences in anthropogenic emissions. Similar to the NO_x comparisons in Sacramento and Fresno, modeled $R_{OH,VOC}$ is also less than observed throughout the entire distribution (Fig. 2e and f), suggesting an underestimate in modeled anthropogenic VOC emissions as well. Overall, the CDFs indicate that the model underestimates emissions of NO_x and anthropogenic VOC throughout the Sacramento and Fresno regions.

At Granite Bay, a suburban Sacramento location at the same longitude as Folsom but located approximately 12 km north, a different NO_2 measurement technique is used (Laser Induced Fluorescence (LIF); Thornton et al., 2000; Cleary et al., 2002). Here, the CDFs indicate that observed NO_x concentrations are fairly well represented by the model in the bottom 50% of the distribution (similar mode, with modeled median = 3.3 ppb and measured median = 4.9 ppb), with increasing differences at the high end (model less than observations, Fig. 2c). A similar NO_x distribution mode is predicted by the model at Granite Bay as at Folsom, but measured NO_x is lower at Granite Bay than at Folsom. This fact indicates that in the Sacramento region, the overestimate of NO_2 by the PAMS monitoring network is one of the dominant sources of the model-measurement difference, but that also some underestimate of emissions by the model is also contributing. For the $R_{OH,VOC}$ distribution (Fig. 2g), the model predicts less $R_{OH,VOC}$ than indicated by measurements, particularly at the lower end of the distribution. This is due in part to the greater contribution of biogenic VOC to calculated $R_{OH,VOC}$ than predicted by the model (see further discussion below in Sect. 3.3).

At Blodgett Forest, a location also with LIF NO_2 measurements, measured and modeled NO_x concentration distributions are similar (Fig. 2d), while the mode and extremes of the $R_{OH,VOC}$ data are significantly different (Fig. 2h). This is likely due to greater values of modeled biogenic $R_{OH,VOC}$, which exhibits a higher reactivity over the narrow range of NO_x concentrations. Further discussion on the contributions

of individual VOC categories to $R_{OH,total}$ and $R_{OH,VOC}$ is included in Sects. 3.3 and 3.4 below.

3.2 The relationship between $R_{OH,VOC}$ and NO_x

Another approach to evaluating modeled VOC is to compare the functional relationships of $R_{OH,VOC}$ versus NO_x with observations (Fig. 4). In Sacramento (Fig. 4a and e), the measurements and model both span a similar range of NO_x concentrations (up to 80 ppb) and $R_{OH,VOC}$ (up to 30 s^{-1}). The increase in $R_{OH,VOC}$ with NO_x predicted by the model is close to the observations (slope of 0.28 for both the model and measurements with similar R^2 values), however the sparse data in Sacramento limits the robustness of this result. In Fresno (Fig. 4b and f), NO_x concentrations and $R_{OH,VOC}$ range up to 60 ppb and 30 s^{-1} , respectively, for both measured and modeled data, with similar slopes of approximately $0.2\text{ s}^{-1}/\text{ppb}$. This indicates increasing model emissions to match observed CDFs would need to be from a source that emits NO_x and $R_{OH,VOC}$ in the same proportion as observed. Because biogenic VOC emissions in Fresno are relatively small, this suggests an anthropogenic source that co-emits NO_x and VOC, including some industrial processes and motor vehicle emissions. Another possibility in the Central Valley is agricultural emissions, which can also emit both NO_x and VOC. Our simulations do not include agricultural NO_x emissions, and only traditional biogenic VOC emissions (isoprene and monoterpenes) from crops are included in the biogenic inventory despite knowledge that crops emit a wide variety of biogenic oxygenated VOC (e.g., Kesselmeier and Staudt, 1999).

At Granite Bay (Fig. 4c and g), daytime measured and modeled NO_x concentrations range from 0–18 ppb and $R_{OH,VOC}$ ranges from 0– 15 s^{-1} . Here the model predicts a stronger linear trend of NO_x and $R_{OH,VOC}$ (slope=0.43 and $R^2=0.27$), while the measurements indicate little to no trend (slope=0.03 and $R^2=0.01$). At Blodgett Forest, NO_x concentrations are low (<2 ppb) at this site in both the model and the measurements. The model predicts $R_{OH,VOC}$ approximately twice that of the measurements and shows that $R_{OH,VOC}$ and NO_x are anti-correlated, which is expected at a rural location. Further discussion of these differences is presented in Sects. 3.3 and 3.4.

3.3 The diurnal cycle of $R_{OH,total}$

We next look at the contributions of individual species to modeled $R_{OH,total}$ over the daytime diurnal cycle (Fig. 5). During the daytime, VOC play a dominant role while NO_2 loss dominates at night. In the urban locations (Sacramento and Fresno; Fig. 5a and b), the dominant daytime $R_{OH,total}$ is due to anthropogenic VOC and aldehydes. For these two VOC categories, $R_{OH,total}$ peaks during the early morning rush hour, and a rise in the contribution of the secondary VOC categories (aldehydes and other oxygenated

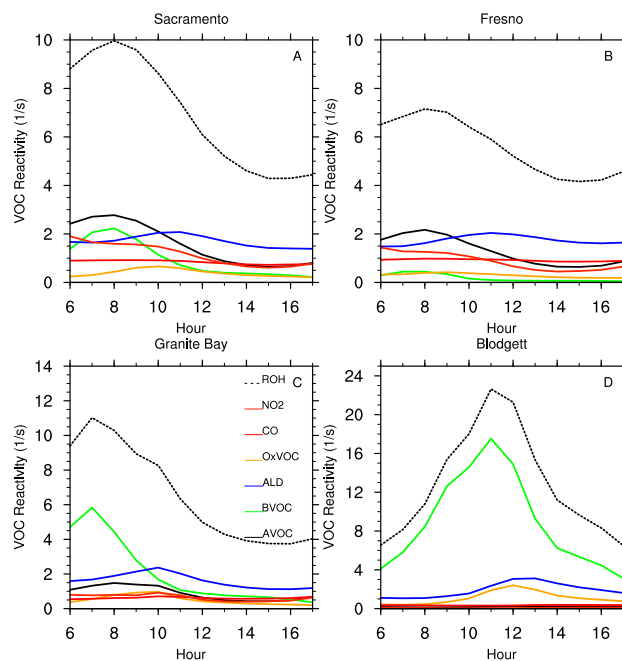


Fig. 5. Average daytime (LT) modeled diurnal cycle of $R_{OH,total}$ (black dashed line) and reactivities of six species at four locations: (a) Sacramento, (b) Fresno, (c) Granite Bay, and (d) Blodgett Forest. Species contributing to $R_{OH,total}$ include NO_2 (brown solid), CO (red solid), anthropogenic VOC (black solid), biogenic VOC (green solid), aldehydes (blue solid), and other oxygenated VOC (orange solid). CH_4 (not shown) contributes a constant value of 0.3 s^{-1} to R_{OH} .

VOC species) occurs about an hour or two later. In Sacramento, biogenic VOC play a role equal to aldehydes in the model, while biogenic VOC are of minor importance in Fresno. The remaining three categories (CO, CH_4 and other oxygenated VOC) tend to contribute less than 1 s^{-1} to $R_{OH,total}$.

In Granite Bay (Fig. 5c), $R_{OH,total}$ also peaks during the morning rush hour when photochemistry is accelerated, predominantly due to biogenic VOC. Biogenic $R_{OH,VOC}$ also peaks at 6 s^{-1} , a value greater than that of anthropogenic VOC or aldehydes. We note that the model does not reproduce the second diurnal peak in reactivity observed at Granite Bay (Murphy et al., 2006b), and this could be due to the lack of afternoon emissions in the model (discussed below with reference to Fig. 6) or poor representation of mixing and chemistry during the evening boundary layer transition. Additionally, the diurnal cycle of modeled biogenic $R_{OH,VOC}$ does not reflect the emissions pattern, which peaks at midday due to temperature and radiation patterns. The accuracy of the model related to this peak in biogenic $R_{OH,VOC}$ is discussed in greater detail in the discussion below.

At the rural Blodgett location (Fig. 5d), biogenic VOC dominates the reactivity, leading to a midday peak of

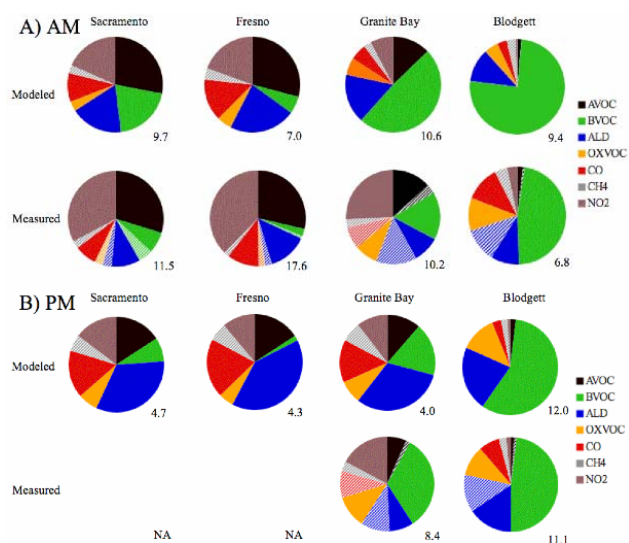


Fig. 6. Modeled and calculated percent contribution to $R_{OH,total}$ (s^{-1}) for the four central California locations during (a) morning hours (6–8 a.m. local time) and (b) afternoon hours (1–3 p.m. local time). $R_{OH,total}$ includes contributions from four VOC categories (anthropogenic, biogenic, aldehydes, and other oxygenated VOC), CO, CH₄ and NO₂. Calculated contributions explained in the text and Tables 1 and 2. $R_{OH,total}$ for each location is noted at the lower right hand of each pie chart. Hatched regions indicate estimated contributions.

$R_{OH,total} = 17 s^{-1}$, corresponding to the time of maximum biogenic VOC flux. Oxidation products of these biogenic VOC (included in the aldehyde and other oxygenated VOC categories) are the second greatest contributors, reaching a maximum approximately one to two hours after the biogenic $R_{OH,VOC}$ peak. In comparison with calculated diurnal variation in reactivity at Blodgett (Murphy et al., 2006b), the model produces a much stronger midday peak in reactivity ($17 s^{-1}$) compared to an average afternoon reactivity observed by Murphy et al. (2006b) of $4\text{--}5 s^{-1}$. Because the biogenic VOC model is tuned to Blodgett measurements, modeled fluxes of well-characterized biogenic VOC such as methylbutenol compare well with measurements (Steiner et al., 2007), and similar chemical species are included in the measurements and model due to the large amount of data available for Blodgett forest. Therefore, the differences are likely due to errors in the model representation of oxidation rates or mixing rates.

Measurements at the PAMS sites are insufficient to assess the full modeled $R_{OH,total}$ diurnal cycle, but there is sufficient data to compare the model and measurements during the morning and afternoon time periods. In some cases, we estimate the contribution of important species that are not included in the measurements. The contribution of CH₄ to $R_{OH,total}$ is estimated for both modeled and calculated $R_{OH,total}$ based on a concentration of 1800 ppbv. For the

PAMS measurement sites (Sacramento and Fresno), the measurements include a large number of anthropogenic hydrocarbons, accounting for many of the reactive species and accounting for a large portion of the modeled lumped anthropogenic categories. However, some important biogenic compounds (terpenes and MBO), aldehyde (RCHO), and other oxygenated VOC that can yield a significant contribution to the total are not measured (see Table 1). For these species, we use model reactivities to estimate their contribution to $R_{OH,total}$, as indicated by the hatched regions of Fig. 6. For Granite Bay and Blodgett, the anthropogenic measurements are not as complete as those provided by the PAMS stations. Therefore, we include model estimates of the top two contributors to anthropogenic $R_{OH,VOC}$ (ethene and ARO₂), which are not measured at these locations. We also include model estimates of two important aldehydes (HCHO and RCHO), which are unmeasured yet can contribute significantly to $R_{OH,VOC}$. The detailed measurements of biogenic and other oxygenated species at Granite Bay and Blodgett include most of the important reactive species for these two categories (Table 2).

Calculated and modeled $R_{OH,total}$ for the four locations and the relative contributions of individual species are compared at 06:00–08:00 LT (Fig. 6a) and 13:00–15:00 LT (Fig. 6b). The morning time period has the greatest number of VOC measurements, yet also coincides with the early-morning break up of the atmospheric boundary layer and the strongest anthropogenic emissions, which is when comparison of model and observations is most complicated. In the morning in Sacramento (Fig. 6a), modeled $R_{OH,total}$ is about 17% less than $R_{OH,total}$ estimated from ground-based measurements. The main difference is due to NO₂, where the measurements indicate a contribution of twice that of the model (17% of $R_{OH,total}$ in the model versus 33% based on the measurements). As previously discussed, measurement techniques may lead to an overestimate of the calculated $R_{OH,total}$ from NO₂ at the PAMS sites (see Sect. 3.1), but the majority of this difference appears to be due to an underestimate of emissions. Calculated (or estimated) and modeled CH₄ and CO compares well and contributes about 3% and 8%, respectively, to R_{OH} . For the VOC component, calculated $R_{OH,VOC}$ is about 60% of $R_{OH,total}$ while the model predicts a greater contribution of 70%. Anthropogenic $R_{OH,VOC}$ contributes about 30% to $R_{OH,total}$ in both the measurements and the model, indicating that the PAMS measurements of anthropogenic $R_{OH,VOC}$ are well described by the model at Sacramento in the morning. The discrepancy in $R_{OH,VOC}$ is due to the model predicting a greater contribution of biogenic VOC and aldehydes than are observed. Biogenic $R_{OH,VOC}$ contributes about 20% to $R_{OH,total}$ in the model and about 12% in the measurements, even when including estimates of modeled biogenic VOC species not measured, such as terpenes and MBO. Aldehydes contribute 13% to calculated $R_{OH,total}$ and 17% to the modeled $R_{OH,total}$. The minor contribution (3%) of the higher order aldehydes estimated from

the model (RCHO) indicates that the aldehyde measurements at this location (HCHO and CCHO) are fairly representative of the modeled aldehyde species at this location. Other oxygenated VOC measurements are not available for Sacramento, although they contribute a relatively small fraction in the model (4%). During the afternoon (Fig. 6b), incomplete measurements do not allow for a comparison, but the modeled NO₂ fraction increases along with a decrease in the primary emissions reactivity (anthropogenic and biogenic) and an increase in the secondary VOC reactivity (aldehydes and other oxygenated VOC).

In the morning in Fresno (Fig. 6a), the model underestimates $R_{OH,total}$ by a factor of 2–3 compared to ground-based estimates. A large portion of this difference is due to NO₂ and, as in Sacramento, the modeled contribution (18%) is about half that of the calculated portion (38%). Calculated and modeled CO and CH₄ also have similar contributions as in Sacramento (11–14% and 2–3%, respectively). VOC contributions to $R_{OH,total}$ are about 50% based on measurements and about 60% in the model. Anthropogenic VOC contributions to $R_{OH,total}$ are similar in the calculated and modeled results (~30%), while the model predicts slightly greater biogenic (6% modeled vs. 4% calculated), aldehyde (22% modeled vs. 15% calculated), and other oxygenated VOC (5% in the model) contributions. In general, missing measurements of biogenic (terpenes and MBO; green hatched regions) and aldehydes (RCHO; blue hatched regions), as described above, contributed very little to total estimates of biogenic and aldehyde species. Overall in the morning at Fresno, the relative contributions of different types of VOC to $R_{OH,total}$ are similar in the model and observations. This fact coupled with the observation that total reactivity and NO_x are both low by about the same amount suggests that the model emissions should be substantially increased for both NO_x and VOC in this region. VOC measurements are not available during the afternoon at Fresno, but similar to Sacramento, the model predicts a smaller contribution of primary VOC reactivity while secondary VOC (aldehydes and other oxygenated VOC) reactivity increases.

Direct measurements of $R_{OH,total}$ have been performed in several other urban locations in North America. In Nashville, $R_{OH,total}$ was calculated in the range of 5–25 s⁻¹, with a median value of 11 s⁻¹ (Kovacs et al., 2003); measurements range from 15–25 s⁻¹ with a median of 19 s⁻¹ in New York City (Ren et al., 2003). Other studies have estimated $R_{OH,total}$ from VOC and NO_x measurements to range from 2–20 s⁻¹ (median=4 s⁻¹) for Nashville, and 1.5–10 s⁻¹ (median=4 s⁻¹) for New York City (Kleinman et al., 2005). The $R_{OH,total}$ we estimate from Sacramento and Fresno (6–20 s⁻¹) are consistent with those of other urban areas in the United States.

At Granite Bay (Fig. 6a), the calculated (10.6 s⁻¹) and modeled (10.2 s⁻¹) morning $R_{OH,total}$ is very similar. As with Sacramento and Fresno, however, the model predicts a much smaller contribution of NO₂ to the $R_{OH,total}$, the mea-

surements show that the $R_{OH,total}$ due to NO₂ is 2.6 s⁻¹ while the model calculates 0.8 s⁻¹. Anthropogenic VOC components agree well in the morning (about 16% in the measurements and 13% in the model), and including additional anthropogenic VOC (black hatched regions) does not make a significant impact on $R_{OH,VOC}$. The model predicts a greater fraction of the total reactivity is biogenic VOC, ~50% compared to the 20% in the measurements. This suggests that the morning biogenic VOC reactivity is much larger than that observed. Calculated and modeled aldehyde reactivities are similar, contributing about 23% and 17% to R_{OH} , respectively. Calculated CCHO reactivity at Granite Bay is greater than that predicted by the model, leading to a greater aldehyde reactivity based on measurements. In the afternoon at Granite Bay (Fig. 6b), the model predicts $R_{OH,total}$ that is a factor of two smaller than the measurements. The greatest discrepancies for the calculated and modeled reactivities at Granite Bay can be attributed to NO₂, biogenic VOC, and aldehyde components. Again NO₂ is underestimated, although not as significantly as in the morning hours. Aldehyde chemistry in the model tends to follow the changes in anthropogenic and biogenic reactivity (increases as the anthropogenic and biogenic reactivity decreases), but this is not reflected in the measurements. The model predicts more biogenic reactivity than aldehyde $R_{OH,VOC}$ in the morning, while the measurements indicate that aldehyde $R_{OH,VOC}$ dominates over biogenic $R_{OH,VOC}$.

At Blodgett Forest in the morning (Fig. 6a), calculated and modeled $R_{OH,total}$ values are within 40% (6.8 s⁻¹ calculated, 9.4 s⁻¹ modeled). Contributions from NO₂ (1–3%), CO (4–12%), and CH₄ (3–4%) are small compared to that from VOC, which contribute about 80% based on measurements and 90% in the model. Anthropogenic VOC plays a small role in $R_{OH,total}$ (2–3%). At Blodgett, the differences in $R_{OH,total}$ stem primarily from discrepancies in the biogenic category. The model predicts about 75% of the $R_{OH,total}$ is due to biogenic VOC in the morning, while the measurements show this fraction is about 50%. In the afternoon (Fig. 5b), calculated and modeled $R_{OH,total}$ and relative VOC contributions are within 10% of each other, again with an overestimate of biogenic reactivity in the model compared to observations. One possible reason for this difference could be due to the biogenic emissions model, which predicts primary isoprene fluxes from Blodgett forest that are not observed at the site. These emissions, which are nearly equal to MBO emissions, could be contributing to the higher reactivity in the model. Another possibility is that fast chemistry within the forest canopy could react away many of the highly reactive terpenes and this fast chemistry cannot be reproduced by the model. Both of these factors could contribute to the over prediction of modeled $R_{OH,VOC}$.

To summarize, the urban reactivity comparison suggests VOC and NO_x emissions are underestimated in the model, yet the relative proportions of the various reactivity sources to $R_{OH,total}$ are approximately correct. Analysis of the mod-

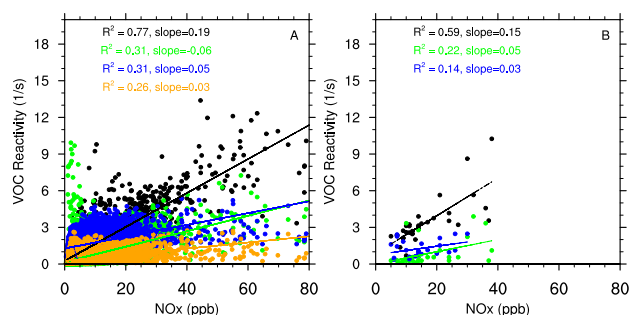


Fig. 7. Speciated $R_{OH,VOC}$ (s^{-1}) as a function of NO_x concentrations (ppb) in Sacramento, as (a) modeled and (b) calculated. The contribution of the four VOC categories to the $R_{OH,VOC}$, including anthropogenic VOC (black), biogenic VOC (green), aldehydes (blue), and other oxygenated VOC (orange). For the calculated Sacramento $R_{OH,VOC}$, biogenic $R_{OH,VOC}$ is based on isoprene concentrations only, and aldehydes are based on HCHO and CCHO concentrations only. See Table 1 for a full list of VOC species included in each category.

eled and observed VOC contributions to $R_{OH,total}$ in the two urban locations show that NO_2 and anthropogenic VOC are the two largest factors. The main categories of VOC contribution to $R_{OH,total}$ in urban areas are anthropogenic VOC (~28%), followed by aldehydes (17–22%), biogenic VOC (6–20%), and other oxygenated VOC (4–6%). Modeled biogenic VOC contribute more to OH reactivity in Sacramento (20%) than Fresno (6%), and this is reflected in the measurements.

At Granite Bay, the model and measurements agree well on $R_{OH,total}$ in the morning but not in the afternoon, while the opposite is true at Blodgett Forest. In both locations, the discrepancies are due to the relative contributions of aldehydes and biogenic VOC. At Granite Bay, modeled biogenic VOC is greater than calculated in the morning, but the converse is true in the afternoon. At Blodgett Forest, the model consistently predicts greater biogenic VOC than that calculated from measurements. These differences are indicative of errors in chemistry, as we have shown the emissions are consistent with observations.

3.4 The relationship between speciated $R_{OH,VOC}$ and NO_x

More insight into where the model is accurately describing the relationships between NO_x and VOC reactivity can be found by comparing each of the four VOC categories versus NO_x (Fig. 7). In the urban locations, we expect that anthropogenic VOC and NO_x will be positively correlated, as both emissions have primary sources from transportation and chemical removal largely occurs downwind. In Sacramento, the slopes of correlation of anthropogenic VOC reactivity with NO_x are $0.15 s^{-1}/ppb NO_x$ for the observations and $0.19 s^{-1}/ppb NO_x$ for the model. We note that some of the difference in slope could be due to the positive bias in the

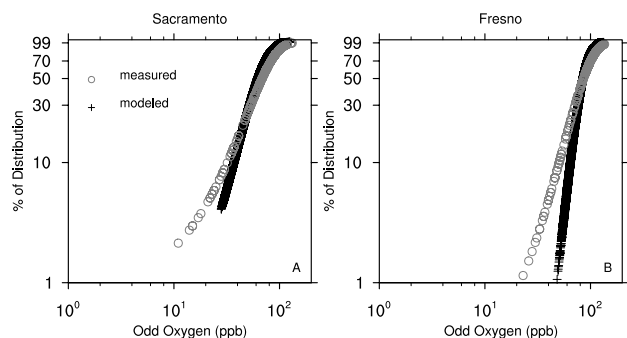


Fig. 8. Measured (gray circles) versus modeled (black crosses) cumulative distribution functions for (a) Sacramento odd oxygen (O_3+NO_2) and (b) Fresno odd oxygen. Ozone and NO_2 data from PAMS stations represents daytime data during the time period of the simulation (29/07/2000–03/08/2000).

PAMS NO_2 as discussed in Sect. 3.1. There are no significant correlations of NO_x and the other categories of VOC in the observations or the model.

3.5 Implications for ozone chemistry

Based on the above comparisons between measured and modeled urban chemistry, the under estimation of NO_x and VOC emissions inventories are notable and likely have implications for ozone prediction in central California. Our prior central California ozone study (Steiner et al., 2006) compared modeled afternoon ozone concentrations versus ground-based observations, and found that the near-surface ozone daily maxima were reproduced by in the model in the central Valley. In the Sacramento region, however, modeled ozone maximum concentrations were about 10–20 ppb less than those observed, and this discrepancy was attributed to poor meteorological representation of upslope flow along the Sierra. In Fresno, measured concentrations of ozone show similar maxima to those modeled (90–100 ppb). Results from this study indicate that inaccurate emissions inventories could also contribute to these differences in ozone concentrations.

In both Sacramento and Fresno, we compare measured and modeled odd oxygen (O_3+NO_2) (Fig. 8). During the simulation time period, the model roughly captures the median of odd oxygen in both Sacramento and Fresno. However, the model over predicts the lower 10% of the distribution in both locations by about 5 ppb, and under estimates the top 10% by about 10–20 ppb. This behavior is not supported by the distribution functions of NO_x concentrations and NO_x and VOC reactivity (Fig. 2), which indicated that emissions were under predicted by the model at these locations over most of the distribution. This suggests that other factors are important for local ozone, including the mixing depths, large-scale meteorology, and the transport of ozone formed in other urban areas such as San Francisco. Because these distributions

are based on ozone concentrations and not local ozone production, it is impossible to discern the difference between locally produced and transported ozone at this time. However, this does suggest that merely increasing NO_x and VOC emissions to address the emission estimate discrepancies may not improve ozone prediction in central California.

3.6 Present day and future spatial distributions of ROH,VOC

Over the model domain, ROH,VOC reaches up to 16 s^{-1} in the morning hours and is greatest near urban areas and regions with high biogenic VOC emissions (Fig. 9a). ROH,VOC outside regions with high anthropogenic and biogenic VOC emissions tend to be low ($<4\text{ s}^{-1}$). The contribution of each VOC category to total modeled reactivity varies with emissions geography. Anthropogenic and biogenic VOC reactivities are located near emission sources (Fig. 9b and c), with anthropogenic ROH,VOC occurring near urban centers and biogenic ROH,VOC dominating in the mountainous and vegetated regions of the domain. However, an interesting point in this simulation is the importance of aldehyde and other oxygenated species with respect to ROH,VOC throughout the model domain (Fig. 9d). Overall, the aldehyde and other oxygenated ROH,VOC accounts for at least 40% of ROH,VOC . In agricultural regions of the Central Valley, where measurements are unavailable, the oxygenated VOC fraction represents up to 90% of total reactivity with minor contributions from primary anthropogenic and biogenic emissions. As shown in Figs. 5 and 6, the model indicates that aldehydes play an important role in the ROH,total in urban areas, accounting for 30–40% of ROH,total in the afternoon. This indicates that aldehyde compounds produced by secondary formation in the atmosphere play a significant role in the cycling of HO_x in all regions, even those with high primary emissions of anthropogenic and biogenic VOC.

For the climate change simulation described in Sect. 2, we examine changes in ROH,VOC under future climate conditions. Warmer temperatures and a moister atmosphere can alter photochemical reaction rates and VOC emission profiles, affecting total and speciated reactivities in the region. The change in morning average ROH,total is almost completely composed of ROH,VOC , which increases over almost all land points in the model domain (Fig. 9e). The largest increases in ROH,VOC occur in the Sierra Nevada Mountains, where ROH,VOC increases up to 5 s^{-1} ($\sim 33\%$) due to an increase in biogenic VOC emissions (Fig. 9g) and subsequent reactions (Fig. 9h). In urban areas such as the eastern portion of the San Francisco Bay area, ROH,VOC increases by about $1\text{--}2\text{ s}^{-1}$ (12–15%).

In regions of high reactivity, the increase in ROH,VOC is due to an increase in biogenic, aldehyde and other oxygenated reactivities. We note that the magnitude of these changes should be considered in light of our previous results in Sect. 3, which indicate that the model is predicting more biogenic and aldehyde VOC reactivity than that esti-

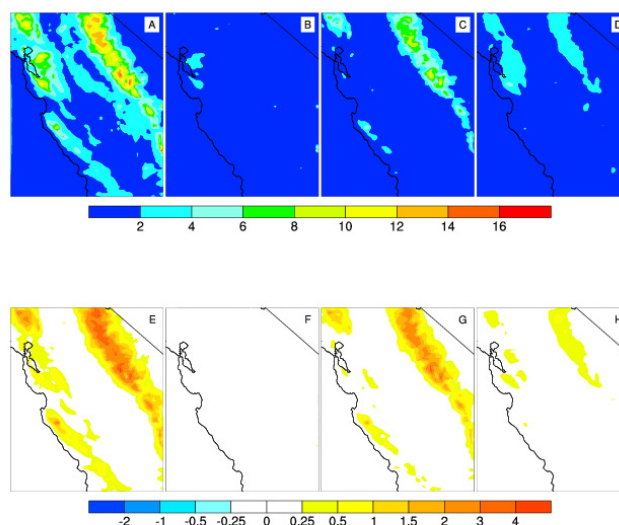


Fig. 9. Present-day and future morning reactivity (s^{-1}). Top panel is present-day reactivity for (a) ROH,VOC , including four VOC categories (anthropogenic, biogenic, aldehydes, and other oxygenated VOC) and (b) anthropogenic VOC, (c) biogenic VOC, (d) aldehydes. Bottom panel is the change to reactivity under future climate conditions for (e) ROH,VOC , (f) anthropogenic VOC, (g) biogenic VOC and (h) aldehydes. Positive values reflect an increase in total reactivity under future climate.

mated from ground-based measurements. However, despite uncertainties in model reactivities, these results indicate that climate impacts on anthropogenic VOC chemistry are small compared to impacts on biogenic, aldehyde and other oxygenated VOC species. Additionally, in urban areas with significant biogenic VOC, changes in biogenic emissions will likely have more impact on ROH,VOC than anthropogenic species. If anthropogenic emissions were to change in a future emissions scenario, then the relative impact of their reactivity will likely change as well.

4 Conclusions and implications

Comparisons of calculated and modeled ROH,total and ROH,VOC indicate that the model generally predicts ROH,total within 25–40% of the observations at three of the four locations (Sacramento, Granite Bay and Blodgett Forest). At the fourth location, Fresno, the model under predicts both NO_x and ROH,VOC emissions by a factor of 2–3. At Sacramento and Fresno, we suspect that the ROH,total differences are in part due to the inadequacies of current NO_2 measurements used in regulatory networks, which can overestimate NO_2 and hence ROH,total substantially (e.g., Dunlea et al., 2007). Because meteorological parameters such as boundary layer height, and wind speeds and directions are represented fairly well during the simulation time period (Wilczak et al., 2004), the representation of the measured and mod-

eled cumulative distribution functions indicate that the emissions inventory in urban locations underestimates both NO_x and anthropogenic VOC emissions. This result is consistent with other studies that indicate that emission inventories are underestimated with respect to on-road emissions (Parrish, 2006; Warneke et al., 2007). Despite these apparent underestimates of ozone precursors, the model predicts a cumulative distribution function for ozone that is similar to the measurements, indicating that simply increasing emissions will likely not improve ozone predictions. Suburban and rural locations show slightly better agreement, likely in part due to direct measurements of NO_2 and in part to the reduced importance of anthropogenic emissions inventories.

The fraction of $\text{R}_{\text{OH, total}}$ from VOC is well-represented at all four locations in central California. The model captures the relative relationships between anthropogenic, biogenic, aldehyde and other oxygenated $\text{R}_{\text{OH, VOC}}$ with NO_x , however some locations have notable exceptions. In urban areas, anthropogenic and biogenic $\text{R}_{\text{OH, VOC}}$ are predicted reasonably over a range of NO_x conditions. In suburban and rural locations, anthropogenic, aldehyde and other oxygenated $\text{R}_{\text{OH, VOC}}$ relationships are reproduced, but biogenic $\text{R}_{\text{OH, VOC}}$ is difficult to emulate, and this may be due to improperly modeled chemical mixing and reactions in the atmospheric boundary layer. Aldehydes make a significant contribution to modeled $\text{R}_{\text{OH, VOC}}$ at all locations, but are difficult to validate based on the limited number of oxygenated VOC measurements.

Under predicted future climate conditions, higher temperatures will affect physical and chemical properties of the atmosphere, increasing $\text{R}_{\text{OH, VOC}}$ over the model domain about 30% near regions of increased biogenic VOC emissions. Our results indicate that biogenic and oxygenated VOC will continue to contribute to radical and ozone production in the future in California, and these impacts will be greatest in the regions where biogenic emissions increase due to increasing temperature. This indicates that anthropogenic VOC control may be less effective in some regions under future scenarios, although this is strongly dependent on the fate of future anthropogenic VOC emissions.

We find that oxygenated VOC play an important role in VOC reactivity throughout central California. Other studies (Yoshino et al., 2006) have found observed $\text{R}_{\text{OH, total}}$ can be greater than modeled R_{OH} , and that this missing OH sink may be due to reaction with oxygenated VOC. In the Bay Area, up to 50% of the modeled $\text{R}_{\text{OH, VOC}}$ is due to oxygenated species and this amplifies local HO_x production cycles. Oxygenated compounds dominate $\text{R}_{\text{OH, VOC}}$ in the San Joaquin Valley, where primary emissions of anthropogenic and oxygenated species are currently estimated to be low. However, emissions of aldehydes and other oxygenated VOC are currently not well quantified from either anthropogenic or biogenic sources. Detailed anthropogenic and biogenic oxygenated VOC emission inventories could improve the emission inputs to regional air quality models. These compounds

are often not measured as part of ground-based measurement campaigns, and they are needed to verify emission inventories, reactivity in the atmosphere, and impacts on gas and particle phase chemistry. More measurements of a suite of oxygenated VOC would provide an additional means of assessing their importance in regional chemistry under present and future climate conditions.

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