Evaluation of improved land use and canopy representation in BEIS v3.61 with biogenic VOC measurements in California

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9 Abstract

Biogenic volatile organic compounds (BVOC) participate in reactions that can lead to 10 11 secondarily formed ozone and particulate matter (PM) impacting air quality and climate. BVOC 12 emissions are important inputs to chemical transport models applied on local to global scales but considerable uncertainty remains in the parametrization of canopy parameterizations and 13 emission algorithms from different vegetation species. The Biogenic Emission Inventory System 14 (BEIS) has been used to support both scientific and regulatory model assessments for ozone and 15 16 PM. Here we describe a new version of BEIS which includes updated input vegetation data and canopy model formulation for estimating leaf temperature and vegetation data on estimated 17 18 BVOC. The Biogenic Emission Landuse Database (BELD) was revised to incorporate land use data from the Moderate Resolution Imaging Spectroradiometer (MODIS) land product and 2006 19 20 National Land Cover Database (NLCD) land coverage. Vegetation species data is based on the U.S. Forest Service (USFS) Forest Inventory and Analysis (FIA) version 5.1 for years from 2002 21 22 to 2013 and U.S. Department of Agriculture (USDA) 2007 census of agriculture data. This 23 update results in generally higher BVOC emissions throughout California compared with the 24 previous version of BEIS. Baseline and updated BVOC emissions estimates are used in Community Multiscale Air Quality Model (CMAQ) simulations with 4 km grid resolution and 25 evaluated with measurements of isoprene and monoterpenes taken during multiple field 26 campaigns in northern California. The updated canopy model coupled with improved land use 27

and vegetation representation resulted in better agreement between CMAQ isoprene and
monoterpene estimates compared with these observations.

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31 **1** Introduction

Volatile organic compounds (VOC) are known to contribute to ozone (O_3) and particulate matter 32 less than 2.5 microns in diameter (PM2.5) formation in the troposphere. Elevated concentrations 33 of O_3 and PM2.5 have known deleterious health effects (Bell et al., 2004; Pope and Dockery, 34 2006; Pope et al., 2006) and climate implications. Biogenic VOC (BVOC) are highly reactive and 35 36 contribute to local and continental scale O₃ and PM2.5 (Carlton et al., 2009;Chameides et al., 37 1988; Wiedinmyer et al., 2005). Terrestrial biogenic emissions are an important input to photochemical transport models which are used to quantify the air quality benefits and climate 38 impact of emission control plans. Despite the important role of BVOC in atmospheric chemistry, 39 the spatial representation of vegetation species, their emission factors, and canopy 40

41 parameterization remain highly uncertain.

42 Isoprene, a highly reactive BVOC, contributes to O₃ (Chameides et al., 1988) and influences

43 secondary organic aerosol (SOA) formation (Carlton et al., 2009). Monoterpenes and

sesquiterpenes are BVOCs known to react in the atmosphere to form SOA (Sakulyanontvittaya

45 et al., 2008). BVOC emissions are important enough to be specifically quantified for impacts on

46 O₃ and PM2.5 (Fann et al., 2013;Kwok et al., 2013;Lefohn et al., 2014). The Biogenic Emission

47 Inventory System (BEIS) (Pierce and Waldruff, 1991;Schwede et al., 2005) estimates these and

48 other BVOC species and has been used extensively to support scientific (Carlton and Baker,

49 2011;Fann et al., 2013;Kelly et al., 2014;Simon et al., 2013;Wiedinmyer et al., 2005) and

regulatory (U.S. Environmental Protection Agency, 2010, 2011, 2012b, a) model applications.

51 BVOC emissions are highly variable among different types of vegetation, therefore the

52 representation of vegetative coverage is critically important for accurate spatial distribution of

53 emissions. Northern California has a large gradient in high isoprene emitting vegetation

- 54 extending from the Sacramento valley eastward toward the Sierra Nevada (Dreyfus et al.,
- 55 2002;Karl et al., 2013;Misztal et al., 2014). Many counties in California have been designated as
- 56 non-attainment of both the 8-hr O₃ and PM2.5 National Ambient Air Quality Standards
- 57 (NAAQS). Recent field studies measuring BVOC concentrations in this area provide a unique

opportunity to evaluate photochemical model estimated BVOC ambient concentrations using an 58 existing (BEIS version 3.14) and updated version of BEIS (version 3.61) and input vegetation 59 data. Ground measurements of BVOC concentrations were made during the Carbonaceous 60 Aerosols and Radiative Effects Study (CARES) campaign in an urban area (Sacramento) and at a 61 site downwind from Sacramento (Cool, CA) that is located near vegetation known for high 62 isoprene emissions (Zaveri et al., 2012). The Biosphere Effects on Aerosols and Photochemistry 63 Experiment (BEARPEX) 2009 campaign provides BVOC measurements at a remote location in 64 the Sierra Nevada foothills to the east of Sacramento and Cool (Beaver et al., 2012), an area of 65 high monoterpene emitting vegetation. 66 In this manuscript, BVOC emissions estimated with the existing, version 3.14 (Schwede et al., 67

2005), and updated version of BEIS, version 3.61, are input to the Community Multiscale Air 68 69 Quality (CMAQ) photochemical transport model (Hutzell et al., 2012;Byun and Schere, 70 2006; Foley et al., 2010) and estimated BVOC ambient concentrations are compared to surface 71 observations at these field campaigns in central and northern California. Canopy coverage and 72 vegetation species data has been updated with the United States Forest Service Forest Inventory 73 and Analysis (FIA) version 5.1 database and 2006 United States Geological Survey National 74 Land Cover Database (NLCD) using more spatially explicit techniques for tree species 75 allocation. BEIS 3.61 has been updated with new a canopy model of leaf temperature for 76 emissions estimation. Canopy leaf temperature estimates are also compared with infrared skin 77 temperature measurements over a grass canopy made at Duke Forest. BVOC estimates from the Model of Emissions of Gases and Aerosols from Nature (MEGAN) (Guenther et al., 2012) are 78 also input to CMAQ and model predictions are compared with field study measurements to 79 provide additional context for BEIS updates. 80

81

82 2 Methods

83 2.1 Land Cover & Vegetation Speciation

BEIS 3.14 used the BELD 3 landuse dataset relied on combined U.S. county level USDA-USFS

85 Forest Inventory and Analysis (FIA) vegetation speciation circa 1992 information with the 1992

- USGS landcover information (Kinnee et al., 1997). A new land cover dataset (BELD 4)
- 87 integrating multiple data sources has been generated at 1 km resolution covering North America.

- Landuse categories are based on the 2001 to 2011 National Land Cover Dataset (NLCD), 2002
- and 2007 USDA census of agriculture county level cropping data, and Moderate Resolution
- 90 Imaging Spectroradiometer (MODIS) satellite products where more detailed data was
- 91 unavailable.
- 92 Fractional tree canopy coverage is based on the 30 m resolution 2001 NLCD canopy coverage
- 93 (http://nationalmap.gov/landcover.html: Homer et al., 2004) and land cover is based on 30 m
- 94 resolution 2006 NLCD Land Cover data. The 2001 canopy data was used because there was no
- 95 canopy product developed for the 2006 NLCD. Land cover for areas outside the conterminous
- 96 United States is based on 500 m MODIS land cover data for 2006
- 97 (https://lpdaac.usgs.gov/products/modis_products_table; MCD12Q1) using the International
- 98 Geosphere Biosphere Programme classification.
- 99 Vegetation speciation is based on multiple data sources. Tree species are based on 2002 to 2013
- 100 Forest Inventory and Analysis (FIA) version 5.1 and crop species information is based on 2002
- and 2007 USDA census of agriculture data. The FIA includes approximately 250,000
- representative plots of species fraction data that are within approximately 75 km of one another
- 103 in areas identified as forest by the NLCD tree canopy coverage. USDA census of agriculture data
- is available on a State and County level only and has been used to refine the agricultural classes
- to the NLCD agricultural land use categories.
- 106 FIA version 5.1 location data has been degraded to enhance landowner privacy in accordance
- 107 with the Food Security Act of 1985 (O'Connell et al., 2012). The provided locations are accurate
- 108 within approximately 1.6 km with most plots being within 0.8 km of the reported coordinates
- and have accurate State and County identification codes (O'Connell et al., 2012). BELD 3 FIA
- 110 vegetation specie fractions were aggregated to county level based on national above ground
- biomass estimates for deciduous, pine, juniper, fir, and hemlock species. In the BELD 4 data set,
- 112 FIA plot level forest biomass (kg/ha) and specific leaf area (g/m^{-2}) were estimated using the
- allometric scaling methods of Jenkins et al. (2003) and Chojnacky et al. (2014). Plot level tree
- biomass estimates were corrected for sampled bole biomass and scaled to a per hectare bases
- following O'Connell et al. (2012). The plot level total and foliage biomass estimates are then
- 116 extrapolated to the continental United States by spatial kriging using the plots longitude, latitude
- and elevation as predictors and weighted by the NLCD canopy fraction. If elevation was not

reported at the plot then elevation was supplied by a digital elevation model from WRF. Kriging

119 was done in 140 by 140 km windows with a 50% overlap to address regional differences in

spatial gradients. A buffer that extended beyond this window was determined by a

semivariogram. Similarly, tree species biomass information was kriged with the additional

122 constraint of the NLCD land use categories (deciduous, evergreen or mixed forest) applied as

weights.

124 The fractional species composition of the NLCD canopy coverage was then calculated and the 125 FIA 5.1.6 species were aggregated to the BELD 4 species (Table S1 and Figure S1). The NLCD 126 land cover defines trees as greater than 5 m tall, forest refers to greater than 20% canopy coverage, with deciduous forests have more than 75% foliage shed in winter and evergreen 127 forests have more than 75% of foliage retained in winter (http://www.mrlc.gov/nlcd06_leg.php). 128 129 These tolerances were used constraining the kriging processes. Total kriged biomass estimates 130 were quantitatively evaluated against the independent estimates of (Blackard et al., 2008). 131 Species specific data in BELD 4 were qualitatively evaluated against the range maps of 132 (Critchfield and Little, 1966) and (Little Jr, 1971, 1976). This kriging approach provides an estimate of vegetation speciation for land cover categories where information is not readily 133 134 available such as urban, grassland, and shrublands. While this kriging approach may provide better spatial estimates of biomass and vegetation type for most areas of the continental United 135 136 States, it is possible that small areas with vegetation and biomass dramatically different than the 137 surrounding region (e.g. some urban areas) will likely need further refinement.

138

139 2.2 Biogenic Emissions

140 MEGAN and BEIS are both used to support regional to continental scale O₃ and PM2.5 photochemical model applications (Carlton and Baker, 2011). Both modeling systems estimate 141 emissions based on vegetation type, meteorological variables, and canopy characteristics 142 (Carlton and Baker, 2011). MEGAN and BEIS both estimate BVOC emissions following the 143 144 empirical algorithm initially developed by Guenther et al (2006). The emission factors between 145 MEGAN and BEIS differ as MEGAN uses emission factors for 16 different global plant functional types (Guenther et al. 2012) while BEIS uses species or species group specific 146 emission factors where available and MODIS plant function types where no species specific data 147

is available, see section 2.1. The canopy models between BEIS and MEGAN also differ.

- 149 MEGAN uses a five layer canopy model where leaf temperature is iteratively solved for each
- 150 layer by adjusting the MEGAN modeled latent, sensible heat fluxes, and outgoing long wave
- radiation to minimize the incoming and outgoing energy balance for the modeled leaf (equation
- 152 1). BEIS approximates the leaf temperature for sun and shaded layers of the canopy from the
- surface energy and momentum balance in the meteorological model as detailed in section 2.3.
- 154 These models have been evaluated against BVOC measurements in the central United States
- 155 (Carlton and Baker, 2011) and Texas (Warneke et al., 2010) but little evaluation of both models
- 156 has been done for California. BEIS version 3.14 provides a baseline for comparison of BEIS
- 157 version 3.61 that includes enhancements described here.

BEIS version 3.61 estimates emissions for 33 volatile organic compounds, carbon monoxide, and 158 159 nitric oxide. Table 1 shows the complete list of compounds estimated by BEIS with mapping to 160 contemporary gas phase chemical mechanisms SAPRC07T and CB6. BEIS estimates isoprene, 161 14 unique monoterpene compounds, and total sesquiterpenes. In addition, emissions are 162 estimated for 16 other volatile organic compounds and an aggregate group of other unspeciated VOC. All biogenic VOC emissions are a function of leaf temperature while only isoprene, 163 164 methanol, and MBO are a function of both leaf temperature and photosynthetically activated radiation (PAR). All species emissions have small indirect impacts from PAR via the canopy 165 166 module.

167 Inputs to BEIS include normalized emissions for each vegetation species, gridded vegetation 168 species, temperature, and PAR. Temperature and PAR can be provided from prognostic meteorological models such as WRF or other sources such as satellite products (Pinker and 169 170 Laszlo, 1992; Pinker et al., 2002) or ambient measurements. The BELD 4 database contains vegetation specie information for 275 different vegetation categories (Table S1). Table 2 shows 171 172 emission rates for each emitted compound by aggregated vegetation type to illustrate variability 173 in emissions. The variability in BEIS emission rates is greater than MEGAN 2.1 (Guether et al. 174 2012) due to the more detailed representation of vegetation species. These vegetation types 175 include 20 MODIS and 21 NLCD land cover categories, and 20 different types of crops both irrigated and non-irrigated (40 total). The remaining categories include tree species, much of 176 177 which are broadleaf (e.g. oak) and needle leaf (e.g. fir) species. A gridded file indicating leaf-on 178 based on the 2009 modeled meteorology, bioseasons file, is also provided as input to BEIS. In

179 the future leaf out and leaf fall dates will be matched with LAI data. Plant genus type LAIs for 180 summer and winter are estimated following Kinnee et al. (1997). However, it is unlikely the 181 current simple leaf-on parameterization will impact typical regulatory assessments since elevated O₃ and PM2.5 organic carbon events often happen outside the spring and fall seasons. 182 For various sensitivity studies presented here, BEIS 3.14 is applied with BELD 3 vegetation 183 data, WRF temperature, and both WRF and satellite derived estimates of PAR. BEIS 3.61 is 184 185 applied similarly but with BELD 3 and BELD 4 vegetation data to isolate the impact of the 186 updates to the canopy model. Note, that the BEIS BVOC emission factors were the same in these 187 BEIS 3.14 and 3.61 simulations. A gridded 0.5 by 0.5 degree resolution satellite estimate of PAR from 2009 was processed to match the model domain specifications and input to both BEIS and 188 MEGAN. The satellite estimates are based on the GEWEX Continental Scale International 189 190 Project and GEWEX Americas Prediction Project Surface Radiation Budget 191 (www.atmos.umd.edu/~srb/gcip/cgi-bin/historic.cgi; Pinker et al., 2002). MEGAN version 2.1 192 (Guenther et al., 2014;Guenther et al., 2012) with version 2011 North America Leaf Area Index 193 and Plant Functional Type (Guenther et al., 2014) was applied with WRF estimated temperature and PAR and also with satellite derived PAR. 194

195

2.3 Canopy Model – Leaf temperature update

BEIS 3.61 includes a two layer canopy model. Layer structure varies with light intensity and 197 solar zenith angle. Both layers of the canopy model include estimates of sunlit and shaded leaf 198 199 area based on solar zenith angle and light intensity, direct and diffuse solar radiation, and leaf 200 temperature. BEIS 3.14 previously used 2 m temperature to represent canopy temperature for 201 emissions estimation even though BVOC emission factors are typically based on leaf temperature (Niinemets et al., 2010). The canopy model has been updated to use land surface 202 203 physics from the Weather and Research Forecasting model and air-surface exchange algorithms 204 from the CMAQ model to approximate leaf temperature using an energy balance for the sunlit 205 and shaded portion of each canopy layer. Emissions are estimated for sunlit and shaded fractions 206 of the canopy and summed over the two layers for total canopy emissions.

A simple two big leaf (sun and shade) temperature model was developed based on a radiation balance. The leaf radiation balance is solved for both the sun (Eq. 1) and shaded (Eq. 2) leaf sides in each layer.

sun leaf

211
$$R_{\rm sun} + IR_{\rm in} - IR_{out} - H - \lambda E_{sun} + G = 0$$
(1)

shade leaf

213
$$R_{\text{shade}} + IR_{\text{in}} - IR_{out} - H - \lambda E_{\text{shade}} + G = 0$$
(2)

214 Where IR_{in} is the incoming infrared radiation, IR_{out} is the outgoing infrared radiation, λ is the latent heat of evaporation, E_{sun} and E_{shade} are the latent heat flux from sun and shade leaves 215 respectively, *H* is the sensible heat flux, and *G* is the soil heat flux. To maintain the same energy 216 balance as WRF it was assumed that E scales linearly with sunlit and shaded fractions of the 217 218 canopy. Note, that conventionally G is positive when the soil is being heated and negative when the soil is cooling while the sign convention of the other variables are relevant to heating and 219 220 cooling of the atmosphere. R_{sun} is the total incoming solar radiation from the meteorological model and R_{shade} is modeled using the attenuation, scattering and diffuse radiation from (Weiss 221 222 and Norman, 1985).

223 The infrared budget is parameterized as

224
$$IR_{\rm in} = \varepsilon_{atm} \sigma T_{atm}^4$$
 (3)

225
$$IR_{\text{out}} = \varepsilon_{leaf} \sigma T_{leaf}^{4}$$
 (4)

226 Where ε_{atm} and ε_{leaf} are the emissivities of the atmosphere and leaf respectively, σ is the Stephan 227 Bolzman constant and T_{atm} and T_{leaf} are the atmospheric and leaf temperatures respectively.

E is parameterized as

229
$$E = \rho_{atm} \frac{e_s(T_{leaf}) - e_a}{R_{w,leaf}P_{atm}}$$
(5)

Where ρ_{atm} is the atmospheric density, $e_s(T_{leaf})$ is the saturation vapor pressure at the leaf, e_a is the atmospheric vapor pressure, $R_{w,leaf}$ is the resistance to water vapor transport from the leaf to atmosphere and P_{atm} is the atmospheric pressure at the surface.

233 The saturation vapor pressure of the leaf is defined as

234
$$e_s(T_{leaf}) = ae^{\frac{b(T_{leaf} - 273.15)}{T_{leaf} - c}}$$
 (6)

- Where the empirical coefficients are a = 611.0 Pa, b = 17.67, and c = 29.65 °C.
- *H* is parameterized following the WRF Pleim-Xiu (PX) land surface model (Skamarock et al.,
 2008) as

238
$$H = \frac{\rho_{atm} C_p \left(\frac{P_0}{P_{atm}}\right)^{R_{atm}/C_p} (T_{leaf} - T_{air})}{R_{h,leaf}}$$
(7)

Where ρ_{atm} is the atmospheric density, C_p is the specific heat of air, P_0 is the STP pressure, R_{atm} is the gas constant for dry air, and $R_{h,leaf}$ is the resistance to heat advection between the atmosphere and leaf. Note, that $R_{h,leaf}$ must consider advection from both the upper, abaxial, and lower, adaxial, surfaces of the leaf.

- The T_{leaf}^4 variable and equation 6 prevents an analytical solution. Thus the approximation from (Campbell and Norman, 1998) is used.
- 245 The T_{leaf}^4 term is simplified as follows:

246
$$\varepsilon_{leaf} \sigma T_{leaf}^{4} \approx \varepsilon \sigma T_{atm}^{4} + \frac{\rho_{atm} c_p \left(\frac{P_0}{P_{atm}}\right)^{R_{atm}/c_p} (T_{leaf} - T_{air})}{R_{r,leaf}}$$
 (8)

247 Where $R_{r, \text{leaf}}$ is the atmospheric radiative resistance ~ 230 s m⁻¹ (Monteith and Unsworth, 2013).

Equation 6 is then further simplified:

249
$$\lambda \rho_{atm} \frac{e_s(T_{leaf}) - e_a}{R_{w,leaf}P_{atm}} \approx \lambda S(T_{atm}) \frac{[T_{leaf} - T_{atm}]}{R_{w,leaf}} + \lambda \rho_{atm} \frac{e_s(T_{atm}) - e_a}{P_{atm}R_{w,leaf}}$$
(9)

250 where

$$251 \qquad S = \frac{de_s(T)}{dT} \tag{10}$$

Equations 1, 3, 5, 7, 8, and 9 are algebraically combined to estimate the sunlit leaf temperature

253 assuming that $\varepsilon_{atm} = \varepsilon_{leaf}$.

254
$$T_{sun,leaf} \approx T_{atm} + \frac{R_{sun} + G - \lambda \rho_{atm} \frac{e_s(T_{atm}) - e_a}{P_{atm}R_{w,leaf}}}{\rho_{atm} \left[\left(\frac{P_0}{P_{atm}}\right)^{R_{atm}/C_p} C_p \left(\frac{1}{R_{h,leaf}} + \frac{1}{R_{r,leaf}}\right) + \lambda S \left(\frac{1}{R_{w,leaf}}\right) \right]}$$
(11)

Equations 2, 3, 5, 7, 8, and 9 are combined to estimate the shaded leaf temperature:

256
$$T_{shade,leaf} \approx T_{atm} + \frac{\frac{R_{shade} + G - \lambda \rho_{atm} \frac{e_s(T_{atm}) - e_a}{P_{atm}R_{w,leaf}}}{\rho_{atm} \left[\left(\frac{P_0}{P_{atm}}\right)^{R_{atm}/C_p} C_p \left(\frac{1}{R_{h,leaf}} + \frac{1}{R_{r,leaf}}\right) + \lambda S \left(\frac{1}{R_{w,leaf}}\right) \right]}$$
(12)

257 The sunlit leaf area index, *LAI_{sun}*, is estimated following (Campbell and Norman, 1998)

258
$$LAI_{Sun} = \int_0^{LAI} e^{-k_{be}(\Psi)L} dL$$
 (13)

where *LAI* is the total canopy leaf area index, k_{be} is the extinction coefficient for direct beam incoming solar radiation as a function of the solar zenith angle, Ψ following Campbell and Norman (1998). The shaded leaf area index, *LAI*_{shade}, is then estimated as follows:

$$262 \quad LAI_{Shade} = LAI - LAI_{Sun} \tag{14}$$

BVOC emission fluxes, F_i , are estimated similar to MEGAN (Guenther et al. 2006) for sunlit and shaded fractions of the canopy

265
$$F_{i,j} = E_i \gamma_{PAR,i,j} \gamma_{T,i,j} LAI_j$$
(15)

where E_i is the emission factor or BVOC species *i*, γ_{PAR} is the emission activity factor for PAR (currently only applied to isoprene, methanol and MBO), γ_T is the emission activity factor for leaf temperature following Guenther et al. (1993), and *j* is the index for sunlit or shaded leaves. γ_{PAR} integrates the PAR emissions activity factor of Guenther et al. (1993) for sunlit and shaded layers following Niinemets et al., (2010).

271
$$\gamma_{PAR,i,Sunlit} = PAR C_L \int_0^{LAI_{Sun}} \frac{e^{-2k_{dd}L}}{\sqrt{1+\alpha^2 PAR^2 e^{-2k_{dd}L}}} dL$$
(16)

272
$$\gamma_{PAR,i,Shaded} = PAR C_L \int_{LAI_{Sun}}^{LAI} \frac{e^{-2k_{dd}L}}{\sqrt{1+\alpha^2 PAR^2 e^{-2k_{dd}L}}} dL$$
(17)

273 Where k_{dd} is the net attenuation coefficient for direct and diffuse PAR and α and C_L are empirical 274 coefficient, 0.0027 and 1.066 respectively, defined in Guenther et al. (1993).

275

276 2.4 Photochemical Model Background, Inputs, and Application

277 Chemical species are estimated using the Community Multiscale Air-Quality Model (CMAQ)

- version 5.0.2 (www.cmaq-model.org) photochemical grid model. CMAQ was applied with
- 279 SAPRC07TB gas phase chemistry (Hutzell et al., 2012), ISORROPIA II inorganic chemistry

280 (Fountoukis and Nenes, 2007), secondary organic aerosol treatment (Carlton et al., 2010) and 281 aqueous phase chemistry that oxidizes sulfur, glyoxal, and methyglyoxal (Carlton et al., 282 2008;Sarwar et al., 2013). The Weather Research and Forecasting (WRF) Advanced Research 283 WRF core (ARW) version 3.3 (Skamarock et al., 2008) was used to generate gridded meteorological inputs for CMAQ and emissions models. While not coincident with this study, 284 this WRF configuration compared well with mixing layer height and surface measurements of 285 286 temperature and winds in central California during the summer of 2010 (Baker et al., 2013). For model performance evaluation presented here, model estimates are paired with measurements 287 using the grid cell where the measurement was located. Measurements are paired in time with 288 289 hourly model estimates with the closest model hour (Simon et al., 2012).

290

The model domain covers central and northern California with 4 km square sized grid cells. The surface to 50 mb is resolved with 34 layers. Layers nearest the surface are most finely resolved with an approximate height of 38 m for layer 1. The modeling period extends from June 3 through July 31, 2009 to be coincident with the BEARPEX field campaign and minimize the influence of initial conditions on model estimates. Initial conditions and boundary inflow are from a coarser CMAQ simulation covering the continental United States. Inflow to the coarser simulation is from a global 2009 application of the GEOS-CHEM (v8-03-02) model

298 (<u>http://acmg.seas.harvard.edu/geos/</u>) (Henderson et al., 2014).

Stationary point sources are based on 2009 specific emissions where available and the 2008
National Emission Inventory (NEI) version 2 otherwise. Mobile emissions are interpolated
between 2007 and 2011 estimates provided by the California Air Resources Board (CARB) and
allocated spatially and temporally using the Spare Matrix Operator Kernel Emissions (SMOKE)
model (http://www.cmascenter.org/smoke). Other non-point and commercial marine emissions
are based on the 2008 NEI version 2 (http://www.epa.gov/ttn/chief/net/2008inventory.html).

305

306 2.5 Field Study Measurements

Between June 15 and July 31 2009, the BEARPEX study was conducted to study photochemical
reactions and products in areas downwind of urban areas with large biogenic influences. The

309 study was located at a managed ponderosa pine plantation in the foothills of the Sierra Nevada (38.90°N, 120.63°W), located near the University of California's Blodgett Research Forest 310 Station. The measurement site was near Georgetown, CA, approximately 75 km from 311 Sacramento, CA. Two research towers housed meteorology and atmospheric composition 312 measurements and inlets during BEARPEX 2009. Meteorological measurements were made on 313 the south, 12.5 m tower, including photosynthetically active radiation (PAR) measured by a LI-314 COR LI190. The second tower (17.8 m) was located approximately 10 m north of the 315 meteorological tower and housed most of the atmospheric composition measurements. The inlet 316 used to sample BVOCs was located at the top of the north tower, approximately 9 m above the 317 ponderosa pine canopy level. BVOCs including isoprene, monoterpenes, methyl vinyl ketone, 318 and methacrolein were quantified using an online gas chromatograph with a flame ionization 319 detector (GC-FID) (Park et al., 2010, 2011). BVOC samples were collected during the first 30 320 minutes of every hour, then subsequently analyzed with the GC-FID. 321

322 During June 2010, the CARES study was conducted to study the formation of organic aerosols

and the subsequent impacts on climate. The study was composed of two surface monitoring sites:

T0 and T1. The T0 was located in Sacramento, CA at the American River College campus

325 (38.65N, 121.35W), and the T1 site was in Cool, CA on the campus of Northside School

326 (38.87N, 121.02W). The T0 site was approximately 14 km northeast of downtown Sacramento,

and the T1 site was surrounded by the forested foothills of the Sierra Nevada. Isoprene and

monoterpene measurements at the Sacramento (TO) and Cool (T1) CARES ground sites were

made with GC-MS and PTRMS, respectively (Zaveri et al., 2012), and sampled via inlets at

approximately 10 m above the surface. PTRMS data were reported as 1 second measurements

approximately every 30 seconds. GC-MS data were 10 minute collections every 30 minutes. All

332 observation data was averaged to hourly concentrations before comparison with model estimates.

333 The sunlight leaf temperature in MEGAN 2.1 and the revised canopy model in BEIS 3.61 were

evaluated against observations taken in 2008 at the Blackwood Division of the Duke Forest in

Orange County, North Carolina, USA (35.97° N, 79.09° W). Details regarding the site

336 (FLUXNET, 2014), measurements, and species composition are available elsewhere (Almand-

Hunter et al., 2014). Leaf temperature measurements were taken using an infrared temperature

sensor (IRTS-P, Apogee Instruments Inc, Logan, UT) mounted on the grassland tower.

340 **3 Results**

341 3.1 Leaf temperature algorithms compared to observations

The canopy model updates for leaf temperature estimation are evaluated by comparing canopy 342 model output with infrared skin temperature measurements of a grass canopy at the Duke Forest 343 field site in central North Carolina (Figure 1). BEIS 3.61 canopy model inputs are based on field 344 measurements taken at this location coincident with the skin temperature data collection. The 345 346 infrared skin temperature measurements do not represent a mean canopy leaf temperature but rather the temperature of the portion of the canopy exposed to the atmosphere. The infrared skin 347 348 temperature measurement should be warmer than the mean leaf temperature during periods of solar irradiation and cooler during periods of radiative cooling due to the insulating effect of the 349 350 unexposed portion of the canopy. Only the estimated exposed leaf temperature (Equation 12) was used in the evaluation to account for this discrepancy between measurements and canopy model 351 352 output. Figure 1 shows observed and predicted estimates of leaf temperature and difference 353 between leaf and ambient temperature. The average temperature estimated by the BEIS 3.61 354 canopy model for the top of the canopy compares well with observations (mean bias of 0.3 K and 355 mean error 1.2 K). Top of the canopy leaf temperature estimated by MEGAN 2.1 are comparable 356 to BEIS 3.61 and the observations at the Duke Forest site.

357

358 **3.2 Evaluation of the BELD 4 land use data**

BELD 4 total forest biomass estimates were evaluated against the independent estimates of 359 360 (Blackard et al., 2008). Blackard et al. (2008) created a spatially explicit live forest biomass dataset for the United States based on FIA observations mapped to MODIS, 250 meter 361 362 aggregated NLCD, topographic and climatic data. Figure 2 shows the BELD 4 and Blackard et 363 al. (2008) estimates of forest biomass for this model domain at 4 km resolution. The Blackard et al. (2008) 250 m grid resolution data set was projected and aggregated to the CMAQ 4 km grid 364 resolution projection using rgdal and raster libraries in R (Bivand et al., 2014). The BELD 4 365 366 estimates evaluated well against those of Blackard et al. (2008) with a Pearson's correlation 367 coefficient of 0.872 (p< 0.001) and a mean and median difference in tree biomass in areas where

368 the NLCD data indicated canopy coverage was -13 kg/ha (-32%) and -0.004 kg/ha (0%) 369 respectively. BELD 4 estimates of forest biomass were greater than those of (Blackard et al., 370 2008) in the densely forested areas in the high Sierras and lower in the lower elevation areas of 371 the domain, primarily in the basin and range areas in the Sacramento valley. The prevalence of the lower elevation areas with lower biomass estimates drives the difference between the forest 372 biomass estimates. The biomass estimates of (Blackard et al., 2008) under predicted the full 373 374 range of the biomass variability with over predictions in areas with low biomass and under predictions in areas of high biomass compared to the FIA tree survey biomass observations. The 375 total biomass estimates presented here have a larger range, 0-661 kg/ha versus 0-499 kg/ha with 376 377 a median absolute deviation of 2.9 kg/ha versus 2.5 kg/ha for areas with NLCD canopy coverage. The lower biomass estimates here compared to those estimated by (Blackard et al., 2008) may be 378 due to our use of 30 m grid NLCD canopy data rather than their use of 250 m grid MODIS 379 canopy data or due to the general underestimation of 2001 NLCD canopy fraction 380

381 product(Nowak and Greenfield, 2012).

382 There are currently no continental US or global databases to quantitatively evaluate the fractional tree species data coverage developed here. However the species range maps of (Critchfield and 383 384 Little, 1966) and (Little Jr, 1971, 1976) can be used for a qualitative evaluation. The tree species that constituted the largest fraction of biomass observations in the FIA data base generally fell 385 386 within the tree species range maps (Figure 3). Note that the maps represent a binary distribution 387 of the tree species natural range and the BELD 4 estimates represent a gradient of species density. Species that did not constitute a large fraction in FIA observations typically had a much 388 smaller estimated spatial range than indicated by the range maps. This could partially be due to 389 the criteria, e.g. tree height greater than 5 m, etc., for trees carried over from the NLCD 390 classification scheme or due to sparse sampling of these tree species in the FIA data base due to 391 the species scarcity. However, these species likely represent a small fraction of the forest 392 393 coverage in the domain and a small fraction of the domain wide BVOC emissions. Also, it is possible that tree coverage has changed in California since the 1970s when the trees were 394 surveyed due to urban planning, plantations, fire, forest growth and climate change. Future 395 396 iterations of the BELD dataset and the evaluation of the BELD dataset can likely be improved by incorporating land cover data with more plant species specific information such as the California 397 398 Gap Analysis Project (David et al. 1998).

400 **3.3 Describing changes in modeled BVOC estimates in Northern California**

Biogenic VOC emissions estimated with BEIS using the new canopy model (BEIS 3.61) and 401 402 updated vegetation data (BELD 4) are shown for the northern California region in Figure 4. A 403 similar Figure of spatial biogenic emissions estimated with BEIS 3.14 and BELD 3 are shown in Figure 5. In this model domain, isoprene emissions are highest in the foothills of the Sierra 404 Nevada where high emitting isoprene vegetation (e.g. oak trees) are located. Monoterpene 405 emissions are highest in the Sierra Nevada Mountains where high emitting needle leaf trees are 406 407 located. Sesquiterpene emissions are highest in the Sacramento and San Joaquin valleys where grasses are common. Most other biogenic VOC emissions show similar spatial patterns as 408 isoprene or monoterpenes (Figure 4). 409

410 The fractional coverage of oak (high isoprene emitting species) and needle leaf trees (high monoterpene emitting species) are shown using BELD 3 and BELD 4 in Figure S2. The BELD 4 411 412 representation shows a higher intensity of fractional coverage in much of the Sierra Nevada as county level information is allocated more spatially explicitly than in BELD 3. Smearing out 413 414 vegetation coverage, as in BELD 3, will lead to lower emissions estimates where narrow features such as the band of oak trees in the western Sierra Nevada foothills exist and over predictions in 415 416 areas that get allocated vegetation that does not exist in that area. Changes in oak and needle leaf fractional coverage between BELD 3 and BELD 4 are notable for both the Cool and Blodgett 417 418 Forest sites meaning the observation data available at these locations is useful for evaluating the methodology used to generate BELD 4 (Figure S2). 419

The updated leaf canopy module increases biogenic VOC emissions throughout California (Figure 5). The changes to the vegetation input data show increases and decreases in isoprene and monoterpene emissions related to changing spatial allocation of high emitting vegetation species and changes to leaf area estimates. Sesquiterpene emissions generally decrease due to the changes in landuse and vegetation for this region (Figure 5). The new vegetation allocation approach employed here for BELD 4 provides more detailed sub-County level representation of emitting species compared to BELD 3 and those changes are reflected in biogenic VOC

427 emissions differences.

429 **3.4 CMAQ estimates compared with CARES and BEARPEX measurements**

The most recent publicly available version of BEIS (version 3.14) and BELD 3 vegetation input 430 were used to provide biogenic emissions for a 4 km CMAQ simulation covering northern and 431 432 central California for the period of time coincident with the 2009 BEARPEX field study. Additional simulations were done to illustrate the impact of updating the leaf canopy module in 433 BEIS 3.61 and also how updating vegetation input data have on biogenic VOC model 434 435 performance. Model runs were also done using satellite derived PAR as input to BEIS in addition 436 to WRF estimated solar radiation. The MEGAN 2.1 model was also run using WRF and satellite 437 estimates of PAR for the same domain and period.

438 Temperature and solar radiation used for the biogenic emissions models were compared to 439 measurements at these field sites (Sacramento, Cool, and Blodgett Forest) to determine how 440 meteorological inputs may bias model estimated BVOC. WRF model evaluation against meteorological variables is summarized in Table 3. The WRF model does well at capturing 441 442 daytime high temperatures at Blodgett Forest and slightly overestimates daily peak PAR. Daytime minimum temperatures at Blodgett Forest are largely overestimated by WRF (Figure 443 444 S3). Temperature maximums and minimums are well characterized at Sacramento and Cool 445 (Figure S4-5) and are similar at these sites during the 2009 and 2010 field study periods (Figure S3). The satellite estimated PAR underestimates the ground measurements at Blodgett Forest on 446 certain days but does better at capturing daytime peaks than WRF. In general, meteorological 447 448 model performance at Blodgett Forest and nearby areas in northern California (Figures S6) 449 should result in overestimated emissions of isoprene and monoterpenes due to model 450 overestimates in PAR and nighttime ambient temperature. While mixing layer depth has been shown to be well represented by WRF for California using the configuration used here (Baker et 451 452 al, 2013), mixing layer depth was not continuously measured at these field sites so could not be 453 directly evaluated meaning that differences between modeled and actual surface layer mixing 454 depth and also differences in local to regional scale transport could impact CMAQ estimates of biogenic VOC. 455

Field study measurements of isoprene and monoterpenes taken in 2010 at Sacramento and Cool
and 2009 at Blodgett Forest provide an opportunity to better understand if the changes to BEIS
and BELD better reflect the biogenic VOC gradient seen over these sites. Figure 6 shows the

459 observed distribution of isoprene concentrations at Sacramento and Cool from 2010, Blodgett 460 Forest in 2009, and model estimates from 2009 for the baseline CMAQ/BEIS simulation (BEIS 461 3.14 and BELD 3), canopy model updates (BEIS 3.61), vegetation data updates (BELD 4), and using satellite PAR with all formulation and other input data updates. Measured isoprene 462 concentrations are lowest in Sacramento and highest at Cool where a high density of Oak trees 463 exist. The baseline simulation predicts the highest isoprene at Blodgett Forest rather than Cool, 464 but when canopy parameterization updates and vegetation data inputs are used the modeling 465 system captures the gradient in concentration well across these three sites and also the 466 467 distribution in observations at each site (Figure 6).

Measured monoterpenes are highest at Blodgett Forest and lowest at Sacramento (Figure 7). The baseline model captured this gradient but notably overestimated monoterpenes at Cool. When BELD 4 is used as input the modeling system compares much closer to observations at Cool and begins to slightly underestimate at Blodgett Forest. The use of satellite PAR rather than solar radiation estimated by WRF does little to change model performance of isoprene. Monoterpenes are not directly sensitive to PAR input and change little due to indirect use of PAR in the canopy model.

475 The MEGAN 2.1 model generally captures the gradient in observations between sites for 476 isoprene and monoterpenes, but predicts much higher isoprene concentrations at each site 477 compared to observations (see Figure 6). This is consistent with other studies comparing 478 MEGAN 2.1 isoprene flux with measurements in the Sierra Nevada of northern California 479 (Misztal et al., 2014) and also with modeling systems using MEGAN 2.1 isoprene emissions 480 compared with ambient isoprene concentrations in Texas (Kota et al., 2015) and southern Missouri (Carlton and Baker, 2011). The airborne flux measurements of Misztal et al. (2014) are 481 lower than the MEGAN estimates for the Northern California modeling domain evaluated here 482 483 and the MEGAN canopy model behaved similarly to BEIS 3.61 (Figure 1) indicating that the 484 MEGAN over estimate in isoprene is likely due to the MEGAN 2.1 emission factors in the modeling domain. Using the MEGAN model estimates of monoterpenes resulted in 485 486 overestimates at Cool and underestimates at Blodgett Forest. Estimates of isoprene using 487 MEGAN improved when using satellite PAR as input rather than WRF solar radiation. This is 488 consistent with similar evaluation in other parts of the United States (Carlton and Baker, 2011). 489 The use of satellite PAR with MEGAN exacerbated monoterpene overestimates at Cool and

490 increased model estimates at Blodgett Forest reducing the model underestimate. First generation 491 oxidation products of isoprene (methacrolein and methyl vinyl ketones) were also measured at 492 Blodgett Forest in 2009. Model performance is similar to isoprene where BEIS estimates 493 compare favorably with measurements and MEGAN 2.1 emissions result in notable overestimates (Figure S3) similar to previous studies (Kota et al., 2015). Methacrolein can 494 further react in the atmosphere to form methacryloyl peroxynitrate (MPAN) which can form 495 methacrylic acid epoxide (MAE) and subsequently secondary organic aerosol including aerosol 496 methylglyceric acid, organic sulfates, and organic nitrates (Worton et al., 2013). CMAQ over-497 estimates MPAN at Blodgett Forest using either biogenic emisisons model, but overestimates are 498 499 greater when using MEGAN. Model performance for isoprene propagates through secondary reactions and could lead to similar over or under estimates of SOA. 500

501

502 4 Future Direction

503 The updated biomass and tree species vegetation characterization in BELD would benefit from 504 additional evaluation for other parts of the conterminous United States. It is critically important 505 to evaluate biogenic emissions models with field experiments designed for biogenic model evaluation or those that provide robust measurements of key biogenic VOC species such as those 506 507 used for this assessment. Future work is planned to evaluate BEIS against a larger field study in California designed for biogenic emissions model evaluation (2011 California Airborne BVOC 508 509 Emission Research in Natural Ecosystem Transects; CABERNET) (Karl et al., 2013; Misztal et al., 2014) and also with a field study done in the southeast United States during the summer of 510 2013 (Southern Oxidant and Aerosol Study; SOAS). Evaluation of the model in urban areas 511 would be useful although little field data exists for urban areas making this type of assessment 512 difficult. 513

514

515 **Code Availability**

516 BEIS 3.61 code is available upon request prior to the public release of CMAQ v5.1 and available now in 517 SMOKE 3.6.5 (https://www.cmascenter.org/smoke/). Please contact Jesse Bash at Bash.Jesse@epa.gov 518 for more information.

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528

529 Supporting Information

Additional model output, comparison with measurements and formulas used for data pairing areprovided in the Supporting Information.

532

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- and Physics, 12, 7647-7687, 10.5194/acp-12-7647-2012, 2012.
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741 Table 1. Species emissions estimated by BEIS and mapping to the SAPRC07T and CB6r2 gas742 phase chemical mechanism lumped species.

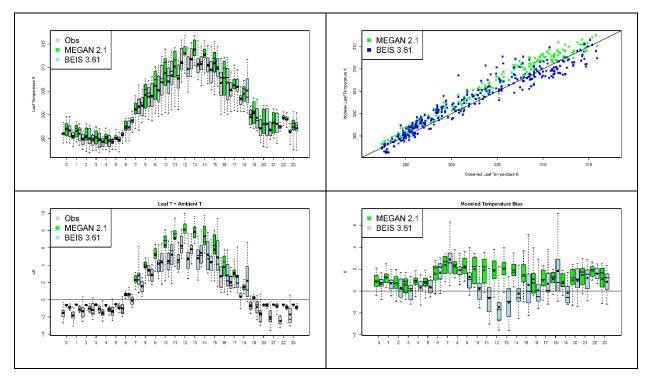
#	Emitted Specie	BEIS	SAPRC07 Species	CB6r2 Species
	•	Abbrevia	-	
1	ethene	ETHE	ETHENE	ETH
2	ethane	ETHA	ALK1	ETHA
3	methanol	METH	MEOH	MEOH
4	ethanol	ETHO	ALK3	ETOH
5	formaldehyde	FORM	HCHO	FORM
6	acetaldehyde	ACTAL	CCHO	ALD2
7	formic acid	FORAC	HCOOH	FACD
8	acetic acid	ACTAC	CCOOH	AACD
9	propene	PROPE	OLE1	33.3 % PAR + 66.7% OLE
10	hexenol	HEXE	OLE1	33.3 % PAR + 66.7 % IOLE
11	hexenylacetate	HEXY	OLE1	37.5 % PAR + 50 % IOLE + 12.5 % NR
12	butenone	BUTO	OLE1	50 % PAR + 50 % OLE
13	MBO	MBO	OLE2	60 % PAR + 40 % OLE
14	butene	BUTE	OLE2	50 % PAR + 50 % OLE
15	acetone	ACET	ACETONE	ACET
16	hexanal	HEXA	RCHO	66.7 % PAR + 33.3 % ALDX
17	Other Reactive VOCs	ORVOC	10 % OLE2 + 85 % ALK2 + 5 % NR	80 % PAR + 20 % OLE
18	Isoprene	ISOP	ISOPRENE	ISOP
19	α-pinene	APIN	TRP1	TERP
20	β-pinene	BPIN	TRP1	TERP
21	δ-3-carene	D3CAR	TRP1	TERP
22	δ-limonene	DLIM	TRP1	TERP
23	camphene	CAMPH	TRP1	TERP
24	myrcene	MYRC	TRP1	TERP
25	α-terpinene	ATERP	TRP1	TERP
26	β-phellandrene	BPHE	TRP1	TERP
27	sabinene	SABI	TRP1	TERP
28	ρ-cymene	PCYM	TRP1	TERP
29	ocimene	OCIM	TRP1	TERP
30	α-thujene	ATHU	TRP1	TERP
31	terpinolene	TRPO	TRP1	TERP
32	γ-terpinene	GTERP	TRP1	TERP
33	Sesquiterpines	SESQ	SESQ	SESQ
34	co	CO	CO	СО
35	NO	NO	NO	NO

- Table 2. Emissions (ug/m2/hr) for each specie estimated by BEIS. Median, minimum, and
- 746 maximum emission rates for each aggregated land cover/vegetation group are shown. Emission
- rates are uniform for some vegetation categories resulting in the same value for median,
- 748 minimum, and maximum.

		Pine			Fir		Spruce			Oak			Maple			Other Deciduous			Crops			Grass			
Number of species	40	40	40	12	12	12	9	9	9	44	44	44	13	13	13	684	684	684	42	42	42	2	2	2	
Metric	Median	Min	Max	Median	Min	Max	Median	Min	Max	Median	Min	Max	Median	Min	Max	Median	Min	Max	Median	Min	Max	Median	Min	Max	
Isoprene (ug/m2/hr)	79	79	79	170	170	170	11900	1700	11900	29750	29750	29750	43	43	43	43	43	29750	10	1	102	56	56	56	
Sesquiterpenes	70	70	210	150	150	150	150	150	150	37	37	37	37	37	37	37	37	150	29	29	29	29	29	29	
Nitric Oxide	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	4	160	0	774	58	58	58	
MBO	76	0	52675	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	11	11	11	
apinene	840	28	2100	1038	239	1472	881	449	1176	26	26	26	127	127	127	15	0	1839	8	0	102	9	9	9	
bpinene	420	0	1134	519	346	929	322	75	716	5	5	5	26	26	26	8	0	580	3	0	51	5	5	5	
d3carene	57	0	867	260	0	260	229	0	730	0	0	0	150	150	150	3	0	280	2	0	26	2	2	2	
dlimonene	48	0	1290	260	107	792	260	2	688	10	10	10	78	78	78	3	0	233	2	0	26	2	2	2	
camphene	7	0	406	260	62	260	260	57	748	6	6	6	31	31	31	3	0	210	2	0	26	2	2	2	
myrcene	37	0	611	260	39	260	218	54	1340	0	0	0	48	48	48	3	0	74	2	0	26	2	2	2	
aterpinene	0	0	96	0	0	324	0	0	78	0	0	0	3	3	3	0	0	18	0	0	0	0	0	0	
bphellandrene	0	0	221	0	0	779	78	0	488	0	0	0	0	0	0	0	0	35	0	0	0	0	0	0	
sabinene	0	0	263	0	0	260	0	0	86	0	0	0	129	129	129	0	0	61	0	0	0	0	0	0	
pcymene	0	0	462	0	0	221	2	0	173	8	8	8	0	0	0	0	0	162	0	0	0	0	0	0	
ocimene	0	0	20	0	0	0	0	0	0	10	10	10	0	0	0	0	0	248	0	0	0	0	0	0	
athujene	0	0	82	0	0	26	0	0	0	0	0	0	5	5	5	0	0	91	0	0	0	0	0	0	
terpinolene	0	0	37	0	0	75	2	0	10	9	9	9	0	0	0	0	0	34	0	0	0	0	0	0	
gterpinene	0	0	7	0	0	70	2	0	8	0	0	0	5	5	5	0	0	28	0	0	0	0	0	0	
methanol	1120	1120	1120	2400	2400	2400	2400	2400	2400	600	600	600	600	600	600	600	600	2400	480	480	480	480	480	480	
ethene	74	74	74	158	158	158	158	158	158	40	40	40	40	40	40	40	40	158	32	32	32	32	32	32	
propene	74	74	74	158	158	158	158	158	158	40	40	40	40	40	40	40	40	158	32	32	32	32	32	32	
ethanol	121	121	121	259	259	259	259	259	259	65	65	65	65	65	65	65	65	259	52	52	52	52	52	52	
acetone	102	102	102	218	218	218	218	218	218	55	55	55	55	55	55	55	55	218	44	44	44	44	44	44	
hexanal	38	38	38	82	82	82	82	82	82	20	20	20	20	20	20	20	20	82	16	16	16	16	16	16	
hexenol	156	156	156	333	333	333	333	333	333	83	83	83	83	83	83	83	83	333	67	67	67	67	67	67	
hexenylacetate	166	166	166	355	355	355	355	355	355	89	89	89	89	89	89	89	89	355	71	71	71	71	71	71	
formaldehyde	70	70	70	150	150	150	150	150	150	38	38	38	38	38	38	38	38	150	30	30	30	30	30	30	
acetaldehyde	51	51	51	110	110	110	110	110	110	28	28	28	28	28	28	28	28	110	22	22	22	22	22	22	
butene	33	33	33	70	70	70	70	70	70	18	18	18	18	18	18	18	18	70	14	14	14	14	14	14	
ethane	18	18	18	38	38	38	38	38	38	10	10	10	10	10	10	10	10	38	8	8	8	8	8	8	
formic_acid	54	54	54	115	115	115	115	115	115	31	31	31	31	31	31	31	31	115	23	23	23	23	23	23	
acetic_acid	35	35	35	75	75	75	75	75	75	20	20	20	20	20	20	20	20	75	15	15	15	15	15	15	
butenone	20	20	20	44	44	44	44	44	44	12	12	12	12	12	12	12	12	44	9	9	9	9	9	9	
Carbon monoxide	490	490	490	1050	1050	1050	1050	1050	1050	264	264	264	264	264	264	264	264	1050	210	210	210	210	210	210	
Other reactive VOC	57	0	57	122	122	122	122	122	122	31	31	31	31	31	31	31	31	122	25	25	25	25	25	25	

Table 3. Model evaluation against field campaigns and network observations.

					Average	Average	Meadian	Median	Average	Average	Fractional	Fractiona
	Scenario	Location	Units	N	Observation	Prediction	Bias (%)	Error (%)	Bias	Error	Bias (%)	Error (%
soprene	BEIS v3.14	Blodgett Forest	ppb	155	1.4	2	26.0	56.0	0.5	1.1	-0.4	73.9
soprene	BEIS v3.6 WRF par	Blodgett Forest	ppb	155	1.4	1.5	-6.0	49.0	0.1	0.8	-22.3	70.3
soprene	BEIS v3.61 SAT par	Blodgett Forest	ppb	155	1.4	1.4	-18.0	49.0	0.0	0.9	-34.4	76.3
soprene	MEGAN v2.1 WRF par	Blodgett Forest	ppb	153	1.4	4.6	203.0	203.0	3.2	3.5	60.3	108.6
soprene	MEGAN v2.1 SAT par	Blodgett Forest	ppb	153	1.4	3.4	97.0	110.0	2.0	2.5	26.3	101.5
Vonoterpenes	BEIS v3.14	Blodgett Forest	ppb	855	0.7	0.8	-10.0	43.0	0.1	0.4	-13.8	58.0
Monoterpenes	BEIS v3.61 WRF par	Blodgett Forest	ppb	855	0.7	0.6	-20.0	40.0	-0.1	0.3	-31.2	57.2
Vonoterpenes	BEIS v3.61 SAT par	Blodgett Forest	ppb	855	0.7	0.6	-21.0	41.0	-0.1	0.3	-33.2	58.6
Monoterpenes	MEGAN v2.1 WRF par	Blodgett Forest	ppb	855	0.7	0.4	-42.0	44.0	-0.3	0.4	-64.1	69.2
Monoterpenes	MEGAN v2.1 SAT par	Blodgett Forest	ppb	855	0.7	0.5	-32.0	39.0	-0.2	0.3	-45.8	58.5
VVK+MACR	BEIS v3.14	Blodgett Forest	ppb	157	1.3	0.9	-29.0	33.0	-0.4	0.5	-44.5	60.8
VVK+MACR	BEIS v3.61 WRF par	Blodgett Forest	ppb	157	1.3	1.4	-4.0	43.0	0.1	0.7	-21.9	65.2
VVK+MACR	BEIS v3.61 SAT par	Blodgett Forest	ppb	157	1.3	1.3	-9.0	47.0	0.0	0.7	-31.8	69.3
VVK+MACR	MEGAN v2.1 WRF par	Blodgett Forest	ppb	155	1.3	2.5	69.0	83.0	1.2	1.6	28.3	82.7
VVK+MACR	MEGAN v2.1 SAT par	Blodgett Forest	ppb	155	1.3	1.6	12.0	61.0	0.4	1.0	-11.4	77.7
Wind Speed	WRF	Cool	m/s	920	2.1	2.8	37.0	40.0	0.7	0.9	30.4	39.3
Vind Speed	WRF	Sacramento	m/s	1266	2.1	2.8	38.0	41.0	0.8	0.9	34.0	41.8
Vind Speed	WRF	Blodgett Forest	m/s	1035	1.5	2.9	104.0	104.0	1.3	1.4	63.9	66.9
emperature	WRF	Cool	С	1786	22.2	23.1	5.0	7.0	0.9	1.6	5.3	8.1
emperature	WRF	Sacramento	С	1721	22.2	22.5	2.0	5.0	0.2	1.4	1.6	6.4
Temperature	WRF	Blodgett Forest	С	1035	18.4	22.6	28.0	29.0	4.2	5.6	28.4	34.1
PAR	WRF	Blodgett Forest	watts/m2	1056	148.3	167.6	0.0	47.0	19.2	45.5	-11.3	52.3
PAR	Satellite estimate	Blodgett Forest	watts/m2	1056	148.3	131.5	0.0	30.0	-16.8	44.3	-39.5	58.0
M2.5 organic carbon	BEIS v3.14	IMPROVE sites	ug/m3	141	1.7	1.1	-34.0	49.0	-0.6	1.0	-43.2	69.6
M2.5 organic carbon	BEIS v3.61 WRF par	IMPROVE sites	ug/m3	141	1.7	1.1	-35.0	50.0	-0.6	1.0	-44.9	70.9
M2.5 organic carbon	BEIS v3.61 SAT par	IMPROVE sites	ug/m3	141	1.7	1.1	-35.0	50.0	-0.6	1.0	-45.6	71.5
M2.5 organic carbon	MEGAN v2.1 WRF par	IMPROVE sites	ug/m3	141	1.7	1.8	8.0	43.0	0.1	1.2	-0.8	57.9
M2.5 organic carbon	MEGAN v2.1 SAT par	IMPROVE sites	ug/m3	141	1.7	2.2	11.0	47.0	0.5	1.4	9.1	62.5
03 greater than 60	BEIS v3.14	AQS sites	ppb	7125	70.9	64.8	-8.0	13.0	-6.1	11.2	-10.1	16.9
D3 greater than 60	BEIS v3.61 WRF par	AQS sites	ppb	7125	70.9	64.7	-8.0	13.0	-6.2	11.0	-10.1	16.7
D3 greater than 60	BEIS v3.61 SAT par	AQS sites	ppb	7125	70.9	64.3	-9.0	13.0	-6.6	11.0	-10.8	16.8
03 greater than 60	MEGAN v2.1 WRF par	AQS sites	ppb	7125	70.9	65.4	-9.0	14.0	-5.5	12.0	-9.5	17.8
D3 greater than 60	MEGAN v2.1 SAT par	AQS sites	ppb	7125	70.9	62.1	-12.0	14.0	-8.8	11.9	-14.1	18.3
D3 less than 60	BEIS v3.14	AQS sites	ppb	48939	32.0	41.0	29.0	33.0	8.9	11.2	30.2	36.6
D3 less than 60	BEIS v3.61 WRF par	AQS sites	ppb	48939	32.0	40.8	29.0	32.0	8.8	11.1	29.8	36.4
D3 less than 60	BEIS v3.61 SAT par	AQS sites	ppb	48939	32.0	40.7	29.0	32.0	8.7	11.0	29.4	36.2
D3 less than 60	MEGAN v2.1 WRF par	AQS sites	ppb	48939	32.0	41.7	32.0	34.0	9.7	11.8	31.9	37.9
O3 less than 60	MEGAN v2.1 SAT par	AQS sites	ppb	48939	32.0	40.7	29.0	32.0	8.7	11.0	30.0	36.4



- Figure 1. Diurnal observed, MEGAN 2.1 and BEIS 3.61 estimated leaf temperatures (top left);
- 755 MEGAN 2.1 and BEIS 3.61 leaf temperature estimates plotted against skin temperature
- observations (top right); observed, MEGAN 2.1, and BEIS 3.61 estimated gradient between leaf
- and ambient temperatures (bottom left); MEGAN 2.1 and BEIS 3.61 estimated leaf temperature
- 758 biases (model-observed) (bottom right).

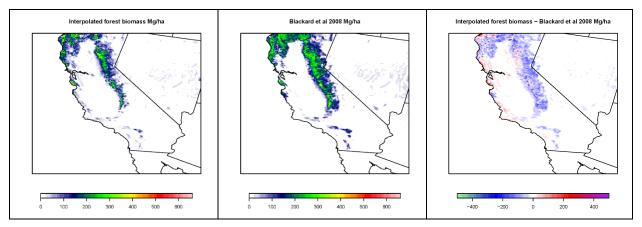


Figure 2. Total above ground forest biomass (Mg/ha) estimates for BELD 4 (left) and Blackard

- et al. 2008 (center) projected onto the 4km California model domain, and BELD 4 the 4km
- 762 projected Blackard et al. 2008 (right).

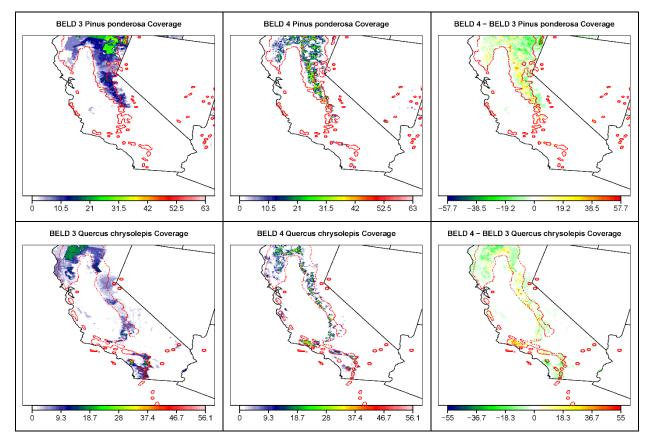


Figure 3. BELD 3 spatial allocation of Ponderosa Pine (Pinus ponderosa, top left), BELD 4

spatial allocation, (top center), and the absolute difference between the BELD 4 and BELD 3

- spatial allocation (top right). BELD 3 spatial allocation of Canyon Live Oaks (Quercus
- chrysolepis, top left), BELD 4 spatial allocation, (top center), and the absolute difference
- between the BELD 4 and BELD 3 spatial allocation (top right). The natural range maps of
- 770 Critchfield and Little (1966) and Little (1971; 1976) are represented by the dashed red lines.

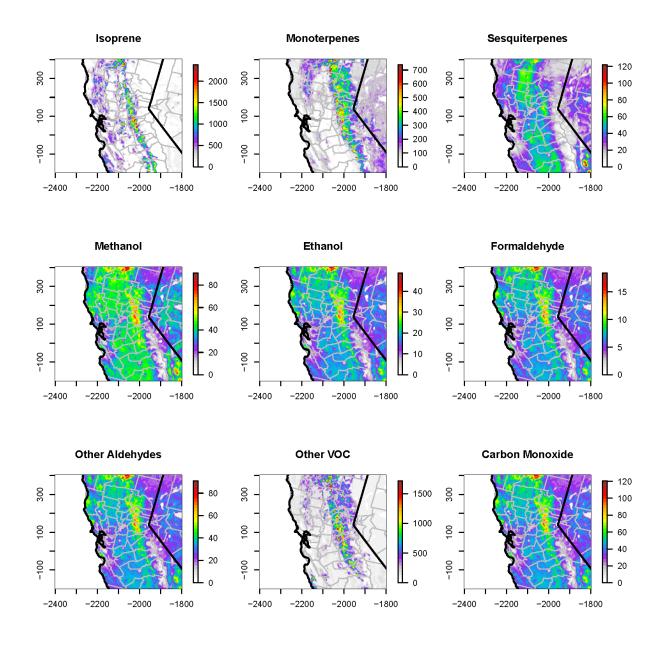
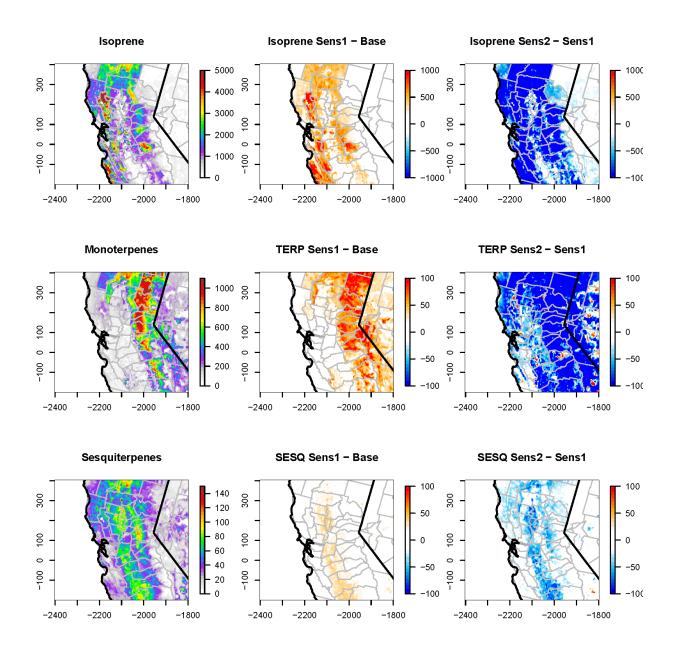




Figure 4. BEIS 3.61 /BELD 4 estimated total emissions (tons) for the modeling period.



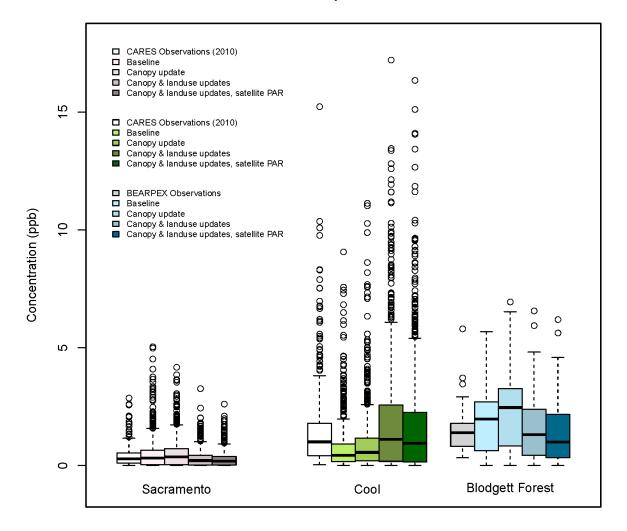
774

Figure 5. Baseline BEIS 3.14 /BELD 3 emissions (tons; left column) and difference between

canopy update and baseline BEIS 3.61 /BELD 3 (center column) and between the canopy update

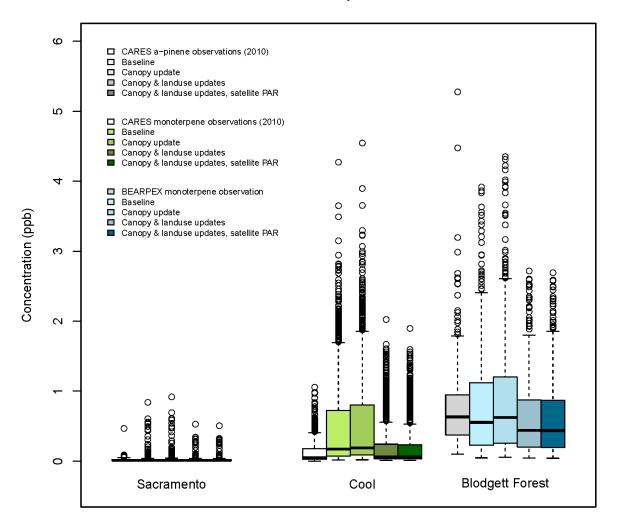
and landuse/vegetation species updates BEIS 3.61 /BELD 4 (right column).

Isoprene



- Figure 6. Distribution of observed and modeled isoprene. Observations at Sacramento and Cool
- represent June 2010. Observations at Blodgett Forest match the modeled period.

Monoterpenes



781

Figure 7. Distribution of observed and modeled monoterpenes. Observations at Sacramento and

783 Cool represent June 2010. Observations at Blodgett Forest match the modeled period.

784