

**Biogenic volatile
organic compound
emissions**

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**Biogenic volatile organic compound
emissions during BEARPEX 2009
measured by eddy covariance and
flux-gradient similarity methods**

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Abstract

The Biosphere Effects on AeRosols and Photochemistry EXperiment (BEARPEX) took place in Blodgett Forest, a Ponderosa pine forest in the Sierra Nevada Mountains of California, during summer 2009. We deployed a Proton Transfer Reaction – Mass Spectrometer (PTR-MS) to measure fluxes and concentrations of biogenic volatile organic compounds (BVOCs). Eighteen ion species including the major BVOC expected at the site were measured sequentially at 5 heights to observe their vertical gradient from the forest floor to above the canopy. Fluxes of the 3 dominant BVOCs methanol, 2-Methyl-3-butene-2-ol (MBO), and monoterpenes, were measured above the canopy by the eddy covariance method. Canopy scale fluxes were also determined by the flux-gradient similarity method (K -theory). A universal K (K_{univ}) was determined as the mean of individual K 's calculated from the measured fluxes divided by vertical gradients for methanol, MBO, and monoterpenes. This K_{univ} was then multiplied by the gradients of each observed ion species to compute their fluxes. The flux-gradient similarity method showed very good agreement with the Eddy Covariance method. Fluxes are presented for all measured species and compared to historical measurements from the same site, and used to test emission algorithms used to model fluxes at the regional scale. MBO was the dominant emission observed followed by methanol, monoterpenes, acetone, and acetaldehyde. The flux-gradient similarity method is shown to be a useful, and we recommend its use especially in experimental conditions when fast measurement of BVOC species is not available.

1 Introduction

Defining source strengths of biogenic volatile organic compounds (BVOC; e.g. isoprene and monoterpenes) and understanding their role in ozone (O_3) and secondary organic aerosol (SOA) formation are critical issues in atmospheric chemistry and climate science (Chameides et al., 1988; Andreae and Crutzen, 1997; Fuentes et al., 2000; Jang

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et al., 2002). On the global scale, VOC emissions from terrestrial vegetation are estimated to be an order of magnitude greater than those from fossil fuel combustion (Guenther et al., 1995). Additional unmeasured organics in the atmosphere are assumed to exist in both the gas and particle phases (Goldstein and Galbally, 2007). Substantial evidence has also been presented for the emissions of highly reactive BVOCs from forest ecosystems that have yet to be adequately quantified and included in BVOC emission models (Kurpius and Goldstein, 2003; De Carlo et al., 2004; Holzinger et al., 2005). More comprehensive observations are needed to better constrain the full range of BVOC emissions from ecosystems and their importance for atmospheric chemistry.

Landscape-scale BVOC flux measurement techniques (e.g. eddy covariance, relaxed eddy accumulation, flux-gradient similarity and mixed boundary layer) have been increasingly implemented in recent years (Goldstein et al., 1996; Schade and Goldstein, 2001; Karl et al., 2002; Lee et al., 2005). The eddy covariance (EC) method using proton transfer reaction – mass spectrometry (PTR-MS) is currently the most direct and reliable measurement for measuring BVOC fluxes at the canopy scale (Karl et al., 2002, Lee et al., 2005). However, this instrument allows continuous EC measurement of fluxes for only a limited number of compounds because the EC method requires fast measurement response time (e.g. > 1 Hz data) to capture concentration changes in the eddies over the forest and the quadrupole mass filter in PTR-MS requires a dwell time of order 0.2 s per mass to obtain adequate signal. The flux-gradient similarity approach, generally called *K*-theory, can be used for measuring fluxes at the canopy scale. In this method, the flux is calculated by multiplying the atmospheric eddy diffusivity (*K*) by the vertical concentration gradient of trace gases (Goldstein et al., 1996). In past applications for measuring BVOC fluxes using this approach, *K* was determined from flux and vertical gradient measurements of H₂O, CO₂, or sensible heat and multiplied by a 30 min average vertical BVOC gradient measured by in-situ GC (Goldstein et al., 1998, Schade et al., 2000). Disadvantage of this approach is that the vertical source and sink profiles of H₂O, CO₂, or sensible heat may differ significantly from those of BVOCs resulting in errors in the similarity assumption for *K* among these

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different tracers. Thus, more accurate determination of BVOC fluxes by this approach for a wide range of species should be possible using EC flux and vertical gradient measurements of a few dominant BVOCs to determine K , and then multiplying K by vertical gradients for each of the observed BVOCs.

5 The Biosphere Effects on Aerosols and Photochemistry Experiment (BEARPEX) 2009 was designed to study the emissions of BVOC and their role in SOA formation, oxidant, and ozone photochemistry. As part of BEARPEX 2009, we deployed a PTR-MS (Ionicon QMS) to measure EC fluxes and vertical gradients of BVOCs in a ponderosa pine plantation in the Sierra Nevada mountains of California with the goals to (1) determine fluxes of BVOCs (17 species) by a combination of direct EC flux measurements and the flux-gradient similarity approach, (2) compare measured fluxes with historical measurements from the same site, in order to highlight vegetation change effects on emissions, and (3) test emission algorithm for the main BVOC species emitted by the vegetation and provide basal emission factors which could be used to model fluxes at regional scale.

2 Experiment

2.1 Measurement site

20 The BEARPEX 2009 intensive field campaign took place in Blodgett Forest from 15 June to 31 July 2009. This site is a Ponderosa pine forest located in the western foothills of the Sierra Nevada Mountains of California (38.90° N, 120.63° W, 1315 m elevation), ~ 75 km downwind of Sacramento, and has been described in detail elsewhere (Goldstein et al., 2000; Lamanna and Goldstein, 1999; Schade and Goldstein, 2001; Dillon et al., 2002). Briefly, the land is owned and operated by Sierra Pacific Industries (SPI), and the plantation is dominated by ponderosa pine (*Pinus ponderosa* L.) trees with an average tree height of 8.7 m in 2009. The understory is dominated by manzanita (*Arctostaphylos* spp.) and whitethorn (*Ceanothus cordulatus*). In summertime,

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winds at this site are predominantly westerly to southwesterly (220–280°) during the day and northeasterly at night (30–60°). This daytime wind pattern transports polluted air from the Sacramento urban area to the pine forest site, along with isoprene and its oxidation products dominantly emitted from oak forests ~ 30 km southwest of the site (Dreyfus et al., 2002). The site is characterized by a Mediterranean type climate, with the majority of precipitation occurring between September and May, and almost no rain in the summer. Average daily temperature ranged from 12°C to 26°C during the campaign period, with no rain recorded.

Two towers were set up at this site. One was a 15 m tall walk-up tower which was installed in 1997. Meteorological parameters such as air temperature, humidity, wind speed and direction, and photosynthetically active radiation (PAR) were measured continuously from this tower and stored in 30 min averaged data sets. Ecosystem-scale fluxes and mixing ratios of trace gases including CO₂, O₃ and VOCs have been measured at this original tower since its construction (Goldstein et al., 2000; Kurpius et al., 2002; Holzinger et al., 2005; Lee et al., 2005; Fares et al., 2010). A new 18 m tall scaffolding tower was built in 2007 and located ~ 10 m north from the original tower. During the 2007 and 2009 summer intensive campaigns (BEARPEX 2007 and BEARPEX 2009), a full range of atmospheric trace gases and aerosols were measured by researchers from more than 10 institutions (Bouvier-Brown et al., 2009a; Wolfe et al., 2009; Smeets et al., 2009; Ren et al., 2010). Observations of vertical gradients and fluxes of BVOCs by PTR-MS were made from 17 June (25 June for fluxes) to 29 July 2009 at the new tower. Electrical power was provided by a propane generator located approximately 125 m north of the new tower. Contamination from generator exhaust was observed occasionally at night when wind was slow and variable, but BVOC measurements were rarely affected.

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2.2 PTR-MS BVOC measurement

2.2.1 Instrument setup

A PTR-MS (Ionicon Analytik, Innsbruck, Austria) was set up to measure BVOC mixing ratios at multiple heights for determination of fluxes and vertical gradients. All measured compounds are listed by mass to charge ratios (m/z) in Table 1. The principles of the PTR-MS have been described in detail elsewhere (Lindinger et al., 1998; de Gouw and Warneke, 2007). Briefly, the PTR-MS is a chemical ionization technique that uses hydronium ions (H_3O^+) to transfer a proton (H^+) from water to the VOC of interest, thus any VOC with a proton affinity higher than water is ionized in the instrument's drift tube, introduced into the quadrupole mass spectrometer and detected by the secondary electron multiplier (SEM). The same instrument has been employed for past measurements of fluxes and vertical gradients at Blodgett Forest (Holzinger et al., 2005; Lee et al., 2005), and also for a similar measurement set up in a citrus orchard (Fares et al., 2012).

Air sample inlet heights on the measurement tower are depicted in Fig. 1. During each hour, six gas sampling inlets were used. One sampling inlet and a 3-D sonic anemometer (Campbell CSAT-3) were co-located at the top of the tower (17.8 m, about twice the canopy height) to measure flux by EC during the first 30 min of each hour. During second 30 min of every hour the other five inlets were sampled sequentially to measure vertical gradients above and within the forest canopy for 6 min at each height (17.8 m, 13.6 m, 9.5 m, 5.7 m, and 1.5 m). All inlets were protected by identical Teflon filters (PFA holder, PTFE membrane, pore size $2\ \mu\text{m}$) to avoid contamination by particles in the air sample. Each inlet was plumbed to the PTR-MS with identical 30 m lengths of PFA tubing (OD 6.35 mm, ID 3.96 mm). A sample flow of $10\ \text{lmin}^{-1}$ was maintained at all times through each sample tube to minimize residence time and any memory effects from previously sampled air, and to maintain a turbulent flow for the EC flux measurement. The Teflon filters were replaced every week and no signal differences for any measured compounds were observed after changing the filters. The

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PTR-MS sequentially sub-sampled $\sim 0.6 \text{ l min}^{-1}$ from each inlet, and was maintained at an E/N (electric field to buffer gas number density) ratio of 139 Td by adjusting drift tube pressure, temperature, and voltage to 200 Pa, 50 °C, and 604 V, respectively. The reaction time in the drift tube was 100 μs and the count rate of $\text{H}_3\text{O}^+\text{H}_2\text{O}$ ions (water clusters) was less than 3% of the count rate of H_3O^+ ions. The H_3O^+ reactant ion was kept in the range $8 \pm 2 \times 10^6 \text{ counts}^{-1}$.

2.2.2 Measurement calibration

Calibrations were performed by dynamic dilution of gravimetrically mixed gas-phase standards (Apel & Riemer, USA) including 2.2–5.1 ppmv of methanol, acetaldehyde, acetone, isoprene, methyl-vinyl-ketone, benzene, α -pinene, *d*-limonene, Δ -3-carene into zero air at ambient humidity levels. Zero air was created by passing sampled ambient air through a stainless steel tube filled with Platinum-coated quartz wool (Shimadzu) heated to 350 °C, catalytically removing VOCs from the sample. This zero air was directly measured to determine instrumental background counts twice daily (02:30–02:40 and 15:00–15:10 PST), and no significant difference between daytime and nighttime background was observed. Calibrations were also done twice a day for 20 min each after background measurement by diluting with purified air to concentrations of 40 ppbv–100 ppbv. In order to check linearity of calibration curves at different concentrations of standard gases and compare them to the twice daily calibrations, we also performed 5-level multipoint calibrations (e.g. 7, 12, 24, 35, 47 ppbv for *d*-limonene) before and after instrument and site maintenance activities during the campaign, such as optimizing the PTR-MS SEM voltage and shutting down the electrical generator for oil changes. Measured mixing ratios from before-and-after calibration curves were in very good agreement for all compounds, with maximum difference within 6%, and a minimum *R*-square of 0.98. To determine mixing ratios of masses for which standard gases were not available, we calculated normalized sensitivities (counts/concentration) based on calculated proton transfer reaction rate coefficients

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and the instrument-specific transmission curve (de Gouw and Warneke, 2007). This curve was determined for m/z 33 to 219 using a gas standard mixture (methanol, acetonitrile, acetaldehyde, acetone, methacrolein, benzene, toluene, xylene, triflorobenzene, bromobenzene, trichlorobenzene, and iodotoluene) at concentrations of 100 ppb (Apel & Riemer, USA). The transmission test was conducted along with each multipoint calibration. PTR-MS signal at m/z 69 is the sum of a fragment of 2-methyl-3-butene-2-ol plus the parent mass of isoprene (MBO + isoprene). Calibration of the mixing ratio for this mass was achieved by scaling the PTR-MS measurement to equal the sum of MBO + ISOP measured simultaneously by gas chromatography-flame ionization detector (GC-FID) (C. Park and G. Schade group, Texas A&M university, correlation coefficient; slope 1, R^2 0.73). For monoterpenes (MTs), the parent ion is observed at m/z 137 and the most abundant fragment occurs at m/z 81. The signal intensity was higher at m/z 81 than m/z 137 because of higher transmission efficiency. Thus, m/z 81 was used for total monoterpene flux measurements and calibrated by using a mixed monoterpene (α -pinene, 3-carene, and d -limonene) standard gas. The sum of m/z 81 and 137 was used for vertical gradient measurement. Comparison between the monoterpene concentrations from the flux inlet determined on m/z 81 with the vertical gradient inlet at the same height (17.8 m) for the adjacent 30 min period determined on the sum of m/z 81 + 137 showed good agreement (slope 0.95, R^2 0.84). Measurement accuracy for all compounds included in the calibration gas standard cylinder was better than $\pm 20\%$. For all other compounds the concentration has been estimated using the collision rate constant which should equal the reaction rate constant within $\pm 30\%$ (Holzinger et al., 2005).

2.3 Eddy covariance flux measurements

Fluxes of four BVOC masses (m/z 33, 69, 81, and 113) were measured using the eddy covariance method. Mixing ratios were measured with dwell time of 0.2 s per mass (5 Hz measurement resolution, ~ 0.8 s per a cycle), resulting in 2120 measurement cycles per 30 min flux period. Wind speed and temperature signals from 3-D sonic

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anemometer were recorded on a datalogger (Campbell Scientific CR-23x) at 10 Hz. For each 30 min flux measurement period, horizontal and vertical wind vectors were rotated according to a planar fit technique (Wilczack et al., 2001). Individual lag times for each 30-min flux period were determined by finding the time point with the maximum cross-correlation coefficient of vertical wind speed and BVOC mixing ratios in ± 60 s time windows. Lag times accounted for changes in clock synchronization between the PTR-MS computer and the data logger recording wind data, and for sampling flow and instrument response times. In cases in which a clear covariance peak was not observed, particularly at night time when vertical turbulence was low, we interpolated lags between the two nearest points having strong covariance peaks. The lag time correction has been applied to these data, and we used the same lag time for all masses. After the lag time correction, Fig. 2 shows the cross correlations between vertical wind speed and each BVOC (m/z 33, 69, and 81) by time peak at 0 s, indicating all BVOCs measured have the same lag times in one flux period.

Fluxes for 4 mass to charge ratios of interest (m/z 33, 69, 81 and 113, but we do not report m/z 113 fluxes in this paper) were determined according to the virtual (continuous flow) disjunct eddy covariance method (Rinne et al., 2001; Karl et al., 2002), which can be regarded as a variant of the EC method. The vertical fluxes of BVOCs were calculated as the mean covariance between deviations of the vertical wind speed and each BVOC mixing ratio for a flux period (30 min), thus we used a subsample of the vertical wind speed (w) data after subtracting the lag time (Δt) due to slower data acquisition frequency for each measured BVOC (5 Hz) than the wind (10 Hz). Flux was determined according to the equation:

$$F_{\text{EC}} = \frac{\sigma}{N} \sum_{i=1}^N w' \cdot c'(i) \quad (1)$$

where σ is the air density (mol m^{-3}), w' is the instantaneous deviation of the vertical wind speed from its average, c' is that of the BVOC mixing ratio (nmol mol^{-1}) (i.e.

$w' = w - \bar{w}$, $c' = c - \bar{c}$), and N is the total number of data points in the measurement period.

The EC method requires that the estimated fluxes within an analyzed period are stationary. To test for stationarity, we divided each 30 min flux period into 5 segments (6 min data for each), calculated the EC flux for each segment, and compared the average to the EC flux for the full 30 min period. We rejected fluxes for any times when the segmental flux was not within $\pm 30\%$ of the full 30 min flux (Lee et al., 2004; Foken and Wichura, 1996). More than 92 % of daytime (09:00–15:00 PST) and 86 % of nighttime (21:00–03:00 PST) data passed these filtering criteria.

We tested for potential flux errors from several known sources. Spectral attenuation by sensor separation and inlet dampening effects were estimated using transfer functions described elsewhere in detail (Moore, 1986; Massman, 1991), and these had less than 1 % affect on our measured EC fluxes. Loss of flux signal can arise from inadequate sensor response time and can be estimated as;

$$\frac{F_{\text{meas}}}{F_{\text{true}}} = \frac{1}{1 + (2\pi \cdot f_m \cdot \tau_c)^\alpha} \quad (2)$$

where F_{meas} is the measured flux, F_{true} is the true flux (non-attenuated flux), f_m is the frequency of the peak at which the frequency weighted cospectrum maximizes, τ_c is the first-order response time of the instrument, and α is 7/8 for neutral and unstable stratification within the surface layer and 1 for stable condition (Horst, 1997). Based on this consideration, if we consider only daytime (09:00 to 17:00 PST) fluxes with τ_c of 1 s and f_m of ranges between 0.01 and 0.02 for day time, fluxes could be underestimated by 8–14 %. This is a similar range to what was estimated during a previous study at this site (Lee et al., 2005), and we did not correct flux data for this effect.

The storage of BVOC emission between the ground and the flux measurement height may be important during times when the air is in a stable stratified condition or turbulent mixing is weak, thus plant or soil emissions in such periods may not be well represented in the above canopy flux measurements. We incorporated this into our flux calculation

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using

$$F = F_{\text{EC}} + \frac{\partial}{\partial t} \int_0^h [\text{BVOC}](z) dz \quad (3)$$

where, z and h is the height of vertical gradient and EC flux measurements, and [BVOC] is the mixing ratio of each compound. The storage term for methanol (m/z 33), MBO + isoprene (m/z 69), and monoterpenes (m/z 81) contributed 57 %, 8 %, and 65 % respectively compared to EC flux in the morning hours 06:00–08:00 PST, and were extremely small during the daytime (less than 0.7 %).

2.4 Vertical gradient measurements

Vertical gradients were measured for 18 masses as listed in Table 1. These masses were chosen to observe the most important compounds at this site based on previous BVOC observations performed by gas chromatography (GC) and PTR-MS from 1997 to 2007 (Lamanna and Goldstein, 1999; Schade et al., 2000; Holzinger et al., 2005; Bouvier-Brown et al., 2009a). Using multiple ion detection mode in the PTR-MS software (Balzer QS 422), these masses were scanned for 14 cycles over 6 min for each height with a dwell time on each mass of 1 s, except for m/z 205 for which we used a dwell of 5 s to improve detection limits for this mass. The first 4 cycles after switching inlet heights were discarded, and data for the last 10 cycles for each height were used to compute hourly averaged datasets.

3 Results

3.1 Concentration and vertical gradients

Figure 3a–c shows the full time series of volume mixing ratios for methanol (m/z 33), MBO + isoprene (m/z 69), and the sum of monoterpenes (m/z 81 + m/z 137), with

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5 mixing ratios averaged from all measurement heights. All these compounds appear dependant on temperature shown in Fig. 3d. Methanol is most abundant, followed by MBO + isoprene, acetone, acetaldehyde, MVK + MACR, and total monoterpenes as summarized in Table 1. Figure 4 presents the average diurnal profile of vertical gradients for methanol, MBO + isoprene, and monoterpenes in addition to ozone. Mixing ratios of all these BVOCs decrease with increased inlet height indicating emission from the forest, and this higher mixing ratio at lower height within the canopy is due to more biomass near the surface. In contrast, the ozone gradient is inversed, suggesting active BVOC oxidation processes and/or stomatal uptake by plants exist within the canopy (Kurpius and Goldstein, 2003; Fares et al., 2010). Diurnal patterns of these 3 BVOCs regularly peak in the morning and evening. This is mainly due to breakup of the atmospheric boundary layer (ABL) in the morning, while the ABL lowers in the evening coupled with stomatal opening/closing by plant circadian cycle, with the peaks corresponding to the transition time of ABL height observed by Choi et al. (2011). After these morning peaks, mixing ratios decrease because of dilution of emissions into a larger mixing layer and faster oxidation of BVOCs during daytime from high daytime ozone mixing ratios, which were observed up to 100 ppbv. High mixing ratios and strong gradients for methanol and monoterpenes for nighttime reflect temperature-dependent emission, but MBO + isoprene emissions are known to be temperature and light dependent (Harley et al., 1998; Schade et al., 2000; Guenther et al., 2006). Distinctively higher methanol mixing ratio near the surface (1.5 m height) suggests that soil, leaf litter, or understory plant emission may significantly contribute to methanol emission at this site (Schade and Goldstein, 2001). The same diurnal pattern has been shown previously at the site (Holzinger et al., 2005; Bouvier-Brown et al., 2009a). In addition to these compounds, Fig. 5 shows vertical gradients and diurnal cycles for MVK + MACR and sesquiterpenes. MVK and MACR are well known as major secondary products from the atmospheric oxidation of isoprene, but isoprene emission from this site is minor without strong vertical gradients observed during daytime, meaning that local isoprene emissions and MVK + MACR production are small. The maximum mixing ratio peak showed

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around hours 15:00–16:00 PST. This pattern is because the air parcel from oak forest, which is a high isoprene emitter, undergoes oxidation of isoprene and is transported to the site with isoprene oxidation products (i.e. MVK and MACR), this result being consistent with previous study (Dreyfus et al., 2002). Sesquiterpenes are not easy to measure in the ambient air because: (1) they exist at low concentration; (2) many of them have a very short lifetime; (3) their large molecule structure and low volatility makes them harder to sample through the Teflon tubing due to stickiness. However, we successfully measured sesquiterpenes by setting dwell time of mass scanning for 5 s and having a fast and continuous sample flow in the tube (residence time of ~ 2.3 s), yet the measurement of sesquiterpenes are still highly uncertain due to their low transmission efficiency passing through PTR-MS quadrupole lens and losses in the sampling inlet and instrument's internal inlet surfaces. Nevertheless, vertical gradients of sesquiterpenes showed a very similar pattern to monoterpenes, indicating emission patterns are similar.

An averaged daytime (10:00–17:00 PST) sesquiterpene mixing ratio of 84 ppt at 1.5 m above the ground was observed. This is about twice greater than GC measurement conducted in the similar season during BEARPEX 2007 (Bouvier-Brown et al., 2009a), but the GC data included only 6 speciated sesquiterpenes, suggesting that additional sesquiterpene species could be emitted in this ponderosa pine forest site.

3.2 Eddy covariance fluxes

The full time series of fluxes for methanol (m/z 33), MBO + isoprene (m/z 69) and monoterpenes (m/z 81) are depicted in Fig. 3f–h. Generally, fluxes of all 3 species show strong temperature dependent emission throughout the measurement period. For example, low emissions were observed around day 190 when temperature is relatively low, with high emissions observed during the high temperatures of day 200. Similar diurnal cycles between the three masses are also observed, with emission starting at sunrise, increasing as air temperature and light intensity increased during daytime, and stopping after sunset. Based on the vertical gradient for methanol and monoterpenes

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shown in Fig. 4a, b, nighttime emissions seem to occur, but no significant fluxes were observed due to lower vertical turbulent mixing.

The sum of MBO and isoprene is quantified as m/z 69, however MBO is the main contributor to m/z 69 fluxes since isoprene emission at this site is minor, as described above. Therefore, MBO is the predominant emission from the site with a day average $\sim 0.90 \text{ mg C m}^{-2} \text{ h}^{-1}$, and is about 5 times greater than either methanol or monoterpene emissions.

4 Analysis and discussion

4.1 Flux estimation by flux-gradient relationship (K -theory)

Fluxes of the other 14 species selected for vertical gradients were not measured, because the PTR-MS can measure EC fluxes only for a few compounds at a time due to limitations of the quadrupole mass filter. Instead, we used a flux-gradient similarity approach, also known as K -theory, for determining fluxes of all compounds not measured by the EC method (Goldstein et al., 1996, 1998). The trace gas flux (F) is assumed to be proportional to the time-averaged mixing ratio gradient (dC/dz) above the forest for intervals longer than the time scale of the slowest significant turbulence events:

$$F = K \frac{dC}{dz} \quad (4)$$

where K is the eddy diffusivity and is determined for each hour of measurements. In previous studies which used K -theory for quantifying VOC fluxes, K was computed from flux and vertical gradient measurements of carbon dioxide (CO_2), water (H_2O), or sensible heat, and then multiplied by VOC vertical gradients (Goldstein et al., 1996, 1998; Schade et al., 2000). However, this approach may have uncertainties which arise from different sources and sinks for each scalar. In order to improve this, we derived K directly from BVOC fluxes and gradients (K_{m33} for methanol, K_{m69} for MBO + isoprene,

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and K_{m81} for monoterpenes), and we further derived a universal K (K_{univ}) by averaging K computed from the three different BVOCs according to the equation:

$$K_{univ} = \frac{(K_{m33} + K_{m69} + K_{m81})}{3} \approx \frac{F_{m33}}{dC_{m33}/dz} \approx \frac{F_{m69}}{dC_{m69}/dz} \approx \frac{F_{m81}}{dC_{m81}/dz} \quad (5)$$

To calculate K for each m/z , we used gradient data from 17.8 m (mixing ratio data from flux measurements were used) and 9.5 m (gradient measurements) which were above the canopy. With one instrument it is not possible to simultaneously measure mixing ratios for different heights, and one also needs to consider the time difference between flux and gradient measurements. So, all gradient data were interpolated to match the same time period as the flux measurements. In addition, data were not used if the gradient for an hour was too small to be detected reliably (less than twice standard deviation of zero air signal) since those values induce a large uncertainty in K , or if the gradient was inverted, e.g. a higher mixing ratio at 17.5 m than 9.5 m with upward flux. Based on these criteria, 75 % for methanol, 51 % for MBO + isoprene, and 77 % for monoterpene data were used, with the vast majority of unused observations occurring at night. The three K values (K_{m33} , K_{m69} , and/or K_{m81}) were calculated for every hour, and averaged to a K_{univ} of which was then applied to estimate fluxes for all compounds with measured vertical gradients. Figure 7 presents the diurnal variation of K_{m33} , K_{m69} , K_{m81} , and K_{univ} . The pattern of K is similar to the flux diurnal profile for most species with maxima during the day when vertical turbulent mixing is strongest. The three K values (K_{m33} , K_{m69} , K_{m81}) agreed well with each other (within 8 % by slope of scatter plots, not shown). This indicates that the two sampling heights have similar footprint characteristics in terms of underlying vegetation and soil, and the different time scales in photooxidation processes for these 3 compounds does not significantly affect the calculation of K at this site. To validate that K_{univ} can be properly used to calculate fluxes for the other compounds, we applied K_{univ} to vertical gradients of methanol, MBO + isoprene, and monoterpenes. Hourly fluxes computed by K_{univ} were compared

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to direct EC flux measurements, and on average these agreed within 4 % with R -square of higher than 0.83 for all three species (Fig. 8).

Fluxes of the additional 14 measured BVOC species were determined by multiplying observed gradients with K_{univ} . Figure 9 shows the resultant diurnal flux cycles for 15 BVOC species, including methanol for reference. Daily average net fluxes are presented in Table 1. Among these 14 species, acetone (m/z 59) and acetaldehyde (m/z 45) fluxes showed the most significant emissions throughout the day with maximum around noon with 0.21 and 0.12 $\text{mg C m}^{-2} \text{h}^{-1}$, respectively. After the noon peak, emission rates of both compounds decreased at hours 13:00–14:00 PST, indicating that active photochemical production/loss processes of VOC above the canopy may suppress the measurable flux strength of these compounds. This phenomenon was also observed in summer 1999 at the same site using a GC-REA (Gas Chromatography-Relaxed Eddy Accumulation) system (Schade and Goldstein, 2001). Methyl chavicol (m/z 149) and sesquiterpene (m/z 205) emissions were also apparent with daytime maxima of 0.06 and 0.08 $\text{mg C m}^{-2} \text{h}^{-1}$, respectively. Low level emissions of acetonitrile (m/z 42), hexanal and hexenols (m/z 83), hexenal (m/z 99), and m/z 111 (unidentified OVOCs) were observed with daytime maxima each below 0.05 $\text{mg C m}^{-2} \text{h}^{-1}$. Interestingly, some oxygenated BVOCs produced by photooxidation of terpenes or isoprene such as MVK + MACR (m/z 71), m/z 113 (unidentified OVOCs), m/z 151 (pinonaldehyde), and m/z 155 (linalool + unidentified OVOCs) were both emitted and deposited throughout the day, though the flux magnitudes were relatively small. This observation implies those compounds were produced by photooxidation within/above the canopy and were also deposited from the atmosphere to the ecosystem. We observed emission of nopinone (m/z 139) in the afternoon during hours 13:00–15:00 PST. Nopinone is a main β -pinene plus OH oxidation product, and β -pinene is one of the most abundant monoterpene species at this site (Lee et al., 2005; Bouvier-Brown et al., 2009a). This observation of nopinone emission from the canopy with maximum in the afternoon is consistent with the daily maximum combination of light and temperatures driving both the β -pinene emissions, and its oxidation by OH radicals. Hourly total and fractional

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BVOC fluxes for all measured species are presented in Fig. 10. A 24 h mean net total emission of $1.5 \text{ mg C m}^{-2} \text{ h}^{-1}$ was estimated with a daytime (10:00–14:00 PST) average of $4.0 \text{ mg C m}^{-2} \text{ h}^{-1}$. Emissions were dominated by MBO + isoprene (61 % to the total, and almost exclusively MBO due to minimal isoprene emission at this site by, as shown by Schade and Goldstein, 2001), followed by monoterpenes (13 %) and methanol (12 %). For the other compounds which were estimated by *K*-theory, acetone and acetaldehyde emissions were 4.9 % and 2.6 % of the total emission, followed by sesquiterpenes (1.9 %), methyl chavicol (1.0 %), with low levels of emissions for the others (less than 1 % for each). Though we did not observe substantial deposition fluxes, the maximum total deposition ($-0.015 \text{ mg C m}^{-2} \text{ h}^{-1}$) by 2 OVOCs (*m/z* 151 and 155) occurred at noontime.

4.2 BVOC emission model

To parameterize measured emissions with commonly used BVOC emission models, we categorized the observed emissions into species that are dependent on temperature versus dependent on both light and temperature. The emission of monoterpenes from this site is known from past observations to be mainly dependent on temperature (e.g. Schade and Goldstein, 2003), with a relationship that can be expressed as (Tingey 1980; Guenther et al., 1993);

$$E_T = F_{30} e^{\beta(T_{\text{air}} - 30)} \quad (6)$$

where F_{30} is the basal emission rate at 30°C , β ($^\circ\text{C}^{-1}$) is a temperature dependence coefficient, and T_{air} ($^\circ\text{C}$) represents the within-canopy air temperature. From our EC flux data, F_{30} ($0.6 \pm 0.14 \text{ mg C m}^{-2} \text{ h}^{-1}$, mean \pm standard deviation) was determined as the mean of data collected at the within-canopy air temperature range of 29°C – 31°C , and β ($0.12 \pm 0.01 \text{ }^\circ\text{C}^{-1}$) was computed by inverting Eq. (6). Canopy scale flux measurements previously reported from this site using PTR-MS EC and GC-FID REA systems had ranges of 0.47 – $1.2 \text{ mg C m}^{-2} \text{ h}^{-1}$ for F_{30} and 0.06 – $0.15 \text{ }^\circ\text{C}^{-1}$ for β (Schade and

Goldstein, 2003; Lee et al., 2005; Holzinger et al., 2006; Bouvier-Brown et al., 2012), in good agreement with the results reported here.

In contrast to monoterpenes, MBO emissions are driven by both temperature and light intensity as described here (Harley et al., 1998; Schade and Goldstein, 2001):

$$E_{T+L} = \text{BER} \cdot \gamma_P \cdot \gamma_T \quad (7)$$

where BER represents the basal emission rate at standard condition (PAR of $1000 \mu\text{mol m}^{-2} \text{s}^{-1}$ and within-canopy air temperature of 30°C), and γ_P and γ_T are temperature emission activity factor and light emission activity factor, as follows,

$$\gamma_P = \alpha \cdot c \cdot \left(\frac{\text{PAR}}{\sqrt{1 + \alpha^2 \cdot \text{PAR}^2}} \right) \quad (8)$$

$$\gamma_T = E_{\text{opt}} \left(\frac{C_{T2} \cdot e^{\frac{(1/T_{\text{opt}} - 1/T) \cdot C_{T1}}{R}}}{C_{T2} - C_{T1} \cdot \left(1 - e^{\frac{(1/T_{\text{opt}} - 1/T) \cdot C_{T2}}{R}} \right)} \right) \quad (9)$$

where α ($= 0.0011$) and c ($= 1.37$) are empirical coefficients, E_{opt} ($= 1.45$) is the maximum normalized emission capacity, T_{opt} ($= 312\text{K}$ by default) is the temperature at which E_{opt} occurs, T is air temperature (K), R is the ideal gas constant ($= 0.00831 \text{ kJ mol}^{-1} \text{ K}^{-1}$), and C_{T1} (131 kJ mol^{-1}) and C_{T2} (154 kJ mol^{-1}) are the energies of deactivation and activation, respectively. All constants used were taken from Schade and Goldstein (2001). Based on this model, BER ($3.8 \pm 0.7 \text{ mg C m}^{-2} \text{ h}^{-1}$) was derived from our MBO EC flux data. Applying this BER with PAR and temperature, the model showed good agreement with our EC flux measurement data (slope: 0.93, R^2 : 0.89, $n = 388$), with the model slightly underestimating observations on average by 7%. This basal emission rate is at least 1.2 times larger than previously reported in

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summer 1999 and the daytime emission also increased from summer 1998 and 1999 but not linearly (Baker et al., 1999; Schade and Goldstein, 2001; Bouvier-Brown et al., 2012 in press). Daytime maxima of 1.2, 1.7, and 2.6 mg C m⁻² h⁻¹ were observed for years 1998, 1999, and 2009, respectively. The shaded area below the canopy has increased as the forest matured, so lower emission capacity in lower canopy is expected while an overall increase is expected due to the increase in emitting biomass at the site. Based on tree survey, the biomass of the site in summer 2009 (1005 g m⁻²) has increased at least 5 times more from summer 1998 (184 g m⁻²).

Methanol emission is also known to be correlated with light and temperature (Nemecek-Marshall et al., 1995; Schade and Goldstein, 2001). Interestingly, our vertical gradients for methanol shown in Fig. 4 strongly indicate nighttime emission and this is similar with monoterpene gradients suggesting the main emission driver is temperature. In contrast, the daytime mixing ratio pattern is similar to that of MBO, not like monoterpenes, indicating temperature and light dependent emission is important during the day. In addition, the flux pattern is similar to that of monoterpenes in the morning when upward flux starts to increase but similar to that of MBO in the afternoon when the flux decreases. This suggests methanol emission is possibly less dependent on light, and this may be due to methanol emission from the soil. Soil methanol emission has been reported by Schade and Goldstein (2001) and Schade and Custer (2004). Based on our analysis of both emission model algorithms, the model of light + temperature for methanol (BER: 0.71 ± 0.21 mg C m⁻² h⁻¹) resulted in better correlation with EC flux measurements (slope: 0.91, R²: 0.74, n = 351). Our 2009 measurements had at least 2 times lower emission for daytime fluxes than summer 1999. This is possibly due to understory shrub removal performed in spring 1999 which may have induced an unusually large emission due to decay of plant debris during summer 1999 (Warneke et al., 1999). Methanol emission is also associated with pectin demethylation when cell walls elongate during leaf/needle expansion and plant growth is recognized as the primary global source of methanol to the atmosphere (Fall and Benson, 1996; Galbally and Kirstine, 2002). Methanol bursts from expanding needles is a phenomenon

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that may have been occurring at higher rates in summer 1999 when pine trees were younger and growing more rapidly as compared to the pine trees in 2009. Moreover, a limited understory biomass in 1999 represented a minor sink for methanol, while in 2009 dense understory vegetation may represent a sink for this OVOC.

In addition to these three dominant BVOCs, acetone, acetaldehyde, sesquiterpenes, and methyl chavicol have been reported as significant emissions from this site (Schade and Goldstein, 2001; Bouvier-Brown et al., 2009c), and our estimations of flux via K -theory are also consistent with these reports. Diurnal cycles of emissions for these compounds correlated better with that of monoterpenes than MBO, suggesting mainly temperature dependent emission. By assuming temperature as the main emission driver, we determined F_{30} and β as summarized in Table 2 for these 4 compounds. For acetone and acetaldehyde, we found slightly smaller F_{30} and β than those reported by Schade and Goldstein (2001), but within the uncertainty range. The daytime fluxes and concentrations showed similar magnitude to summer 1999, which is interesting since the biomass of the site has increased substantially over this 10 yr period.

Leaf-scale emission rates of monoterpenes, sesquiterpenes and methyl chavicol were previously reported from this site (Bouvier-Brown et al., 2009c), but canopy-scale fluxes of sesquiterpenes and methyl chavicol have not been previously reported. We estimated the canopy-scale basal emission rate by multiplying the ecosystem foliar density ($1005 \text{ g [dry weight] m}^{-2}$ for BEARPEX 2009) to leaf-scale basal emission rate and applied the emission model using the scaled F_{30} with β . Figure 11 presents temperature dependent emissions for monoterpenes and sesquiterpenes from leaf-scale emissions, canopy-scale fluxes, and temperature-algorithm basal emission model results optimized for this study. For monoterpenes, the canopy level emissions are in the middle of the range of leaf scale emissions, and this indicates little or no loss before escaping the canopy. In contrast, sesquiterpenes are known to be oxidized rapidly in the tree canopy due to their high reactivity (Ciccioli et al., 1999; Bouvier-Brown et al., 2009c), this can justify the lower emissions measured at the canopy level as compared to up-scaled emission from leaf-level measurements. For methyl chavicol, we derived

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a lower F_{30} ($0.08 \pm 0.06 \text{ mg C m}^{-2} \text{ h}^{-1}$) from the canopy scale flux data than was reported for leaf-scale measurements (range $0.16\text{--}1.1 \text{ mg C m}^{-2} \text{ h}^{-1}$, scaled for canopy-scale) by Bouvier-Brown et al. (2009c), but we observed a higher β ($0.2 \pm 0.06 \text{ }^\circ\text{C}^{-1}$ for this study, and the range of $0.12\text{--}0.2 \text{ }^\circ\text{C}^{-1}$ for leaf-scale). Bouvier-Brown et al. (2009b) reported that methyl chavicol emission accounted for 4–24 % of carbon mass emitted as MBO. Our methyl chavicol to MBO ratios were in the range 0.8–7.8 % during hours 13:00–15:00 when methyl chavicol and MBO emission were maximum, consistent with the lower end of the range previously reported.

5 Summary

Proton transfer reaction-mass spectrometry (PTR-MS) using quadrupole mass spectrometry has been applied to BVOC EC flux measurements, but only allows measurement of fluxes for a few compounds (e.g. 4 ion species applied for this study). To quantify fluxes of a broader range of BVOCs, we simultaneously measured EC fluxes and vertical gradients of BVOCs during BEARPEX 2009. Using a flux-gradient relationship (“ K -theory”), we successfully determined fluxes of 14 BVOC species in addition to MBO, monoterpenes, and methanol by EC. Comparison of fluxes measured by the eddy covariance method with fluxes estimated by K -theory for MBO, methanol, and monoterpenes showed excellent agreement. MBO (m/z 69) was the dominant BVOC emission observed ($0.90 \text{ mg C m}^{-2} \text{ h}^{-1}$ in a 24 h mean), followed by monoterpenes (m/z 81) and methanol (m/z 33) which were similar (0.20 and $0.18 \text{ mg C m}^{-2} \text{ h}^{-1}$, respectively). Fluxes of the other 14 BVOC species all showed net emission, with 24 h mean total emission of $0.2 \text{ mg C m}^{-2} \text{ h}^{-1}$, equivalent to monoterpene or methanol emissions. Of these 14, acetone (m/z 59) and acetaldehyde (m/z 45) emissions were largest at 0.07 and $0.04 \text{ mg C m}^{-2} \text{ h}^{-1}$, respectively, followed by sesquiterpene (m/z 205, $0.03 \text{ mg C m}^{-2} \text{ h}^{-1}$), methyl chavicol (m/z 149, $0.02 \text{ mg C m}^{-2} \text{ h}^{-1}$), with a lower level of emission for all the others (less than $0.01 \text{ mg C m}^{-2} \text{ h}^{-1}$ for each). By comparing with leaf-scale emission of monoterpenes and sesquiterpenes previously studied

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at the site, we found there was significant chemical loss of sesquiterpenes before they could escape the forest canopy, but not for monoterpenes, consistent with previous studies (Ciccioli et al., 1999; Bouvier-Brown et al., 2009a). Overall, estimating BVOC emissions using the flux-gradient relationship applied to direct measurements proved to be a useful method for investigation of ecosystem scale BVOC fluxes.

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**Table 1.** Mixing ratio and flux information for 18 selected ion species.

Compounds	Mass/ charge (<i>m/z</i>)	Empirical molecular for- mulae (protonated mass)	Mixing ratio 24 h-mean (day/night) ^a (nmol mol ⁻¹)	Flux 24 h-mean (day) ^a (mg C m ⁻² h ⁻¹)
Methanol	33	CH ₃ OHH ⁺	16 (13/16)	0.18 (0.48)
Acetonitrile	42	C ₂ H ₃ NH ⁺	0.15 (0.16/0.15)	0.003 (0.008)
Acetaldehyde	45	C ₂ H ₄ OH ⁺	1.9 (1.6/2.4)	0.038 (0.104)
Acetone	59	C ₃ H ₆ OH ⁺	3.6 (3.5/4)	0.073 (0.18)
MBO ^b + isoprene	69	C ₅ H ₈ H ⁺ (C ₅ H ₁₀ OH ⁺) ^c	4.2 (5.6/2.1)	0.90 (2.51)
MVK + MACR ^d	71	C ₄ H ₆ OH ⁺	1.2 (1.5/1.3)	0.013 (0.019)
Benzene	79	C ₆ H ₆ H ⁺	0.071 (0.059/0.083)	0.001 (0.003)
Monoterpenes	81, 137	C ₆ H ₈ H ⁺ , C ₁₀ H ₁₆ H ⁺	1.1 (0.6/1.3)	0.2 (0.46)
Hexanal, hexenols	83	C ₆ H ₁₀ H ⁺	0.14 (0.12/0.16)	0.010 (0.023)
Hexenals	99	C ₆ H ₁₀ OH ⁺	0.16 (0.15/0.18)	0.012 (0.031)
Unknown OVOCs	111	C ₈ H ₁₄ H ⁺ , C ₆ H ₈ O ₂ H ⁺ , C ₇ H ₁₀ OH ⁺	0.026 (0.02/0.029)	0.003 (0.006)
Unknown OVOCs	113	C ₈ H ₁₆ H ⁺ , C ₇ H ₁₂ OH ⁺ , C ₆ H ₈ O ₂ H ⁺ , C ₅ H ₄ O ₃ H ⁺	0.15 (0.18/0.15)	0.0001 (−0.0001)
Nopinone	139	C ₉ H ₁₄ OH ⁺	0.056 (0.046/0.071)	0.006 (0.016)
Methyl chavicol	149	C ₁₀ H ₁₂ OH ⁺	0.099 (0.064/0.084)	0.015 (0.038)
Pinonaldehyde	151	C ₁₀ H ₁₄ OH ⁺	0.021 (0.017/0.023)	0.002 (0.004)
Linalool, Unknown OVOCs	155	C ₁₀ H ₁₈ OH ⁺ , C ₉ H ₁₄ O ₂ H ⁺	0.015 (0.016/0.017)	0.0007 (0.003)
Sesquiterpenes	205	C ₁₅ H ₂₄ H ⁺	0.095 (0.072/0.098)	0.028 (0.06)

^a Daytime and nighttime means are for hours 10:00–14:00 and 22:00–02:00 PST, respectively.^b 2-Methyl-3-butene-2-ol^c Parent MBO ion in parenthesis. MBO main fragment ion is the same as isoprene parent ion.^d Sum of methylvinylketone and methacrolein.

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Table 2. Modeled values for basal mission rate (BER and F_{30}) and temperature-dependence factor (β).

Compound	Basal emission rate at 30 °C, F_{30} (mg C m ⁻² h ⁻¹)	β (°C ⁻¹)
MBO + isoprene (m/z 69)	(3.8 ± 0.70)*	(N.A.)**
Monoterpenes (m/z 81)	0.60 ± 0.14	0.12 ± 0.01
Methanol (m/z 33)	0.54 ± 0.17 (0.71 ± 0.21)*	0.08 ± 0.01 (N.A.)**
Acetone (m/z 59)	0.24 ± 0.14	0.09 ± 0.04
Acetaldehyde (m/z 45)	0.13 ± 0.07	0.07 ± 0.05
Sesquiterpenes (m/z 205)	0.11 ± 0.10	0.07 ± 0.09
Methyl chavicol (m/z 149)	0.08 ± 0.06	0.20 ± 0.06

* Basal emission rate at PAR of 1000 $\mu\text{mol m}^{-2} \text{s}^{-1}$ and air temperature of 30 °C for light + temperature algorithm.

** Not available for light + temperature algorithm. Detailed equations are described in Eqs. (7)–(9).

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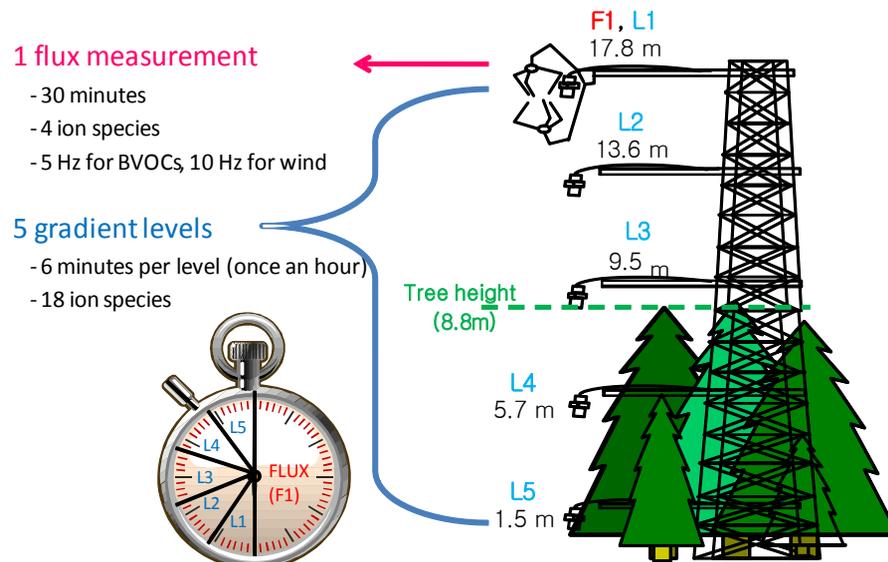


Fig. 1. Inlet configuration and sampling schedule on the 18 m tall tower during BEARPEX 2009. The PTR-MS sampled from an inlet (F1) at 17.8 m co-located with a 3-D sonic anemometer during the first 30 min of each hour for eddy covariance flux measurements, then sampled sequentially from five inlets (L1–L5) positioned at 17.8, 13.6, 9.5, 5.7, and 1.5 m above ground during the next 30 min to measure vertical gradients. Mean tree height was 8.8 m.

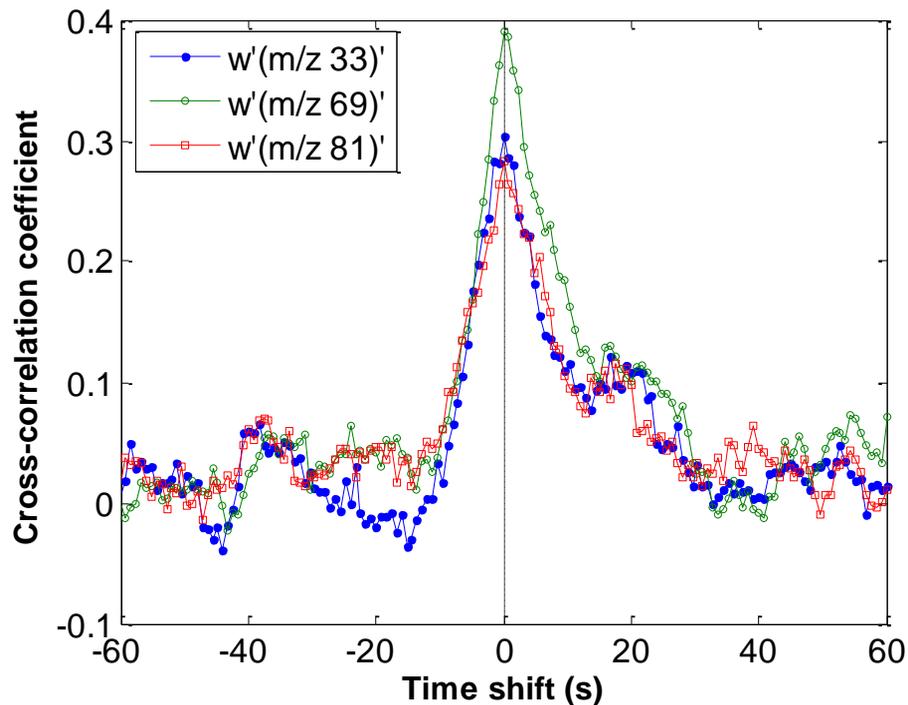


Fig. 2. Cross-correlation of vertical wind speed (w) and volume mixing ratios for 3 ion species (m/z 33, 69, and 81) at 14:00–14:30 PST on 18 July, 2009. Lag time corrections between vertical wind speed and volume mixing ratio of X seconds have been applied to these data.

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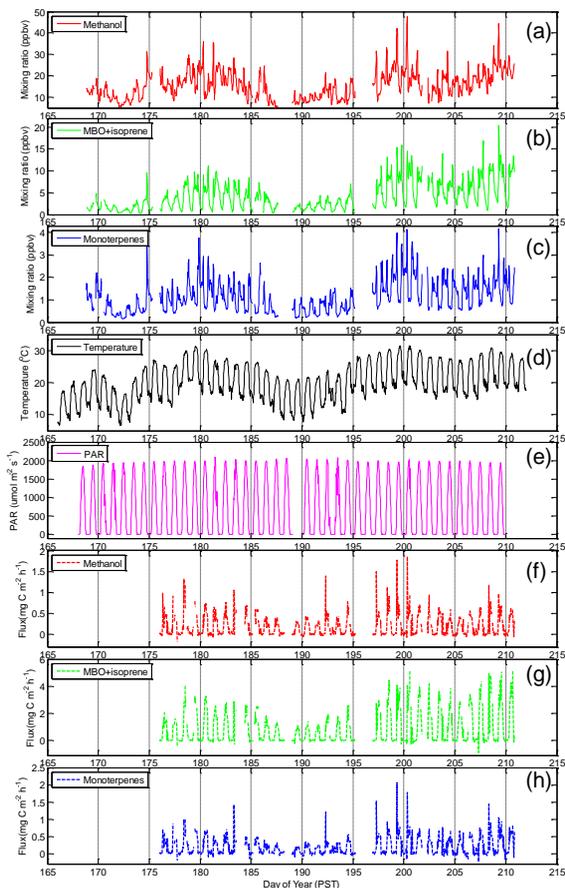


Fig. 3. Volume mixing ratio time-series of **(a)** methanol, **(b)** MBO+isoprene and **(c)** monoterpenes, **(d)** air temperature, **(e)** photosynthetically active radiation (PAR), and fluxes of **(f)** methanol, **(g)** MBO + isoprene and **(h)** monoterpenes. All data are averaged on an hourly basis.

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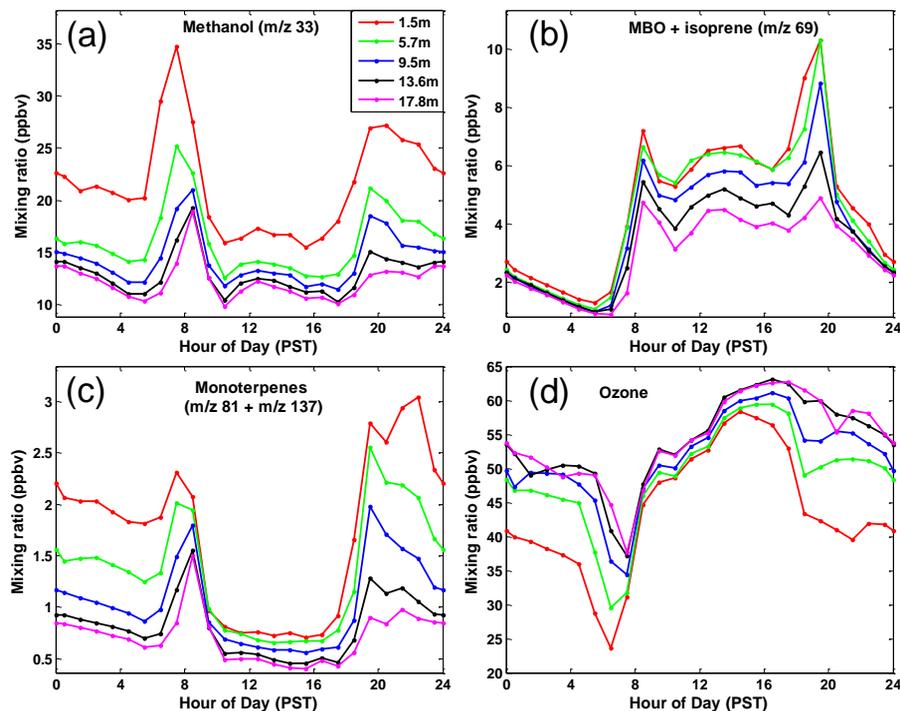


Fig. 4. Averaged diurnal cycles of vertical gradients for **(a)** methanol, **(b)** MBO + isoprene, **(c)** monoterpenes, and **(d)** ozone. Measurement heights are indicated in the legend **(a)**.

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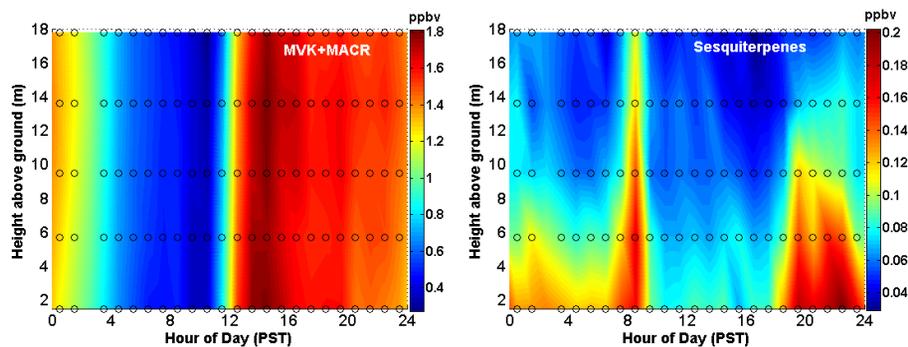


Fig. 5. Mean vertical gradient diurnal patterns for MVK + MACR (left) and sesquiterpenes (right). Color represents interpolated concentration gradients with actual measurement timing and vertical positions indicated by open circles.

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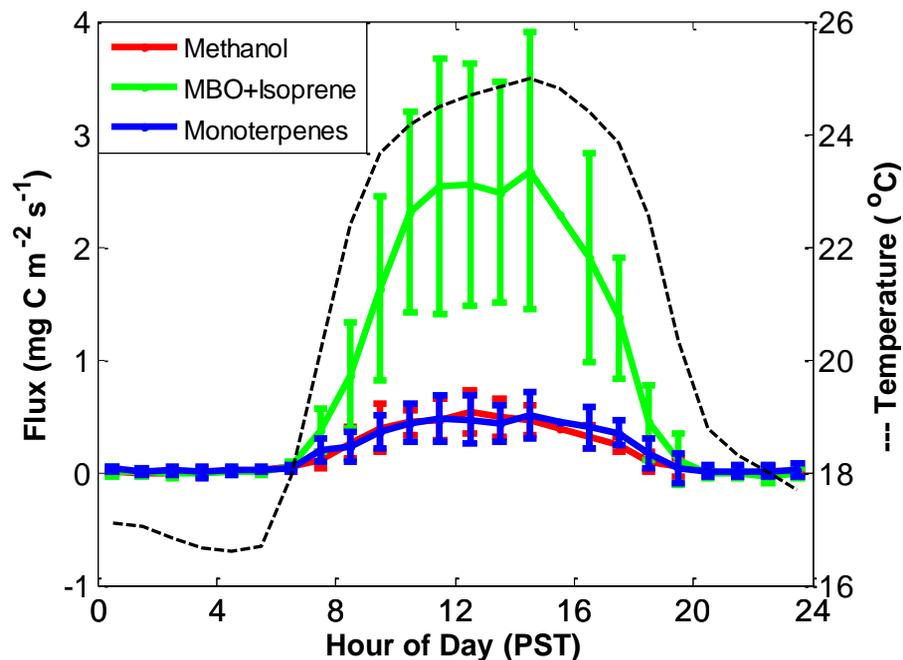


Fig. 6. Mean diurnal variation of air temperature (dashed black line) and fluxes for methanol (red line), MBO + isoprene (green line), and monoterpenes (blue line). Error bars denote standard deviations of all measurements at the respective hour of the day.

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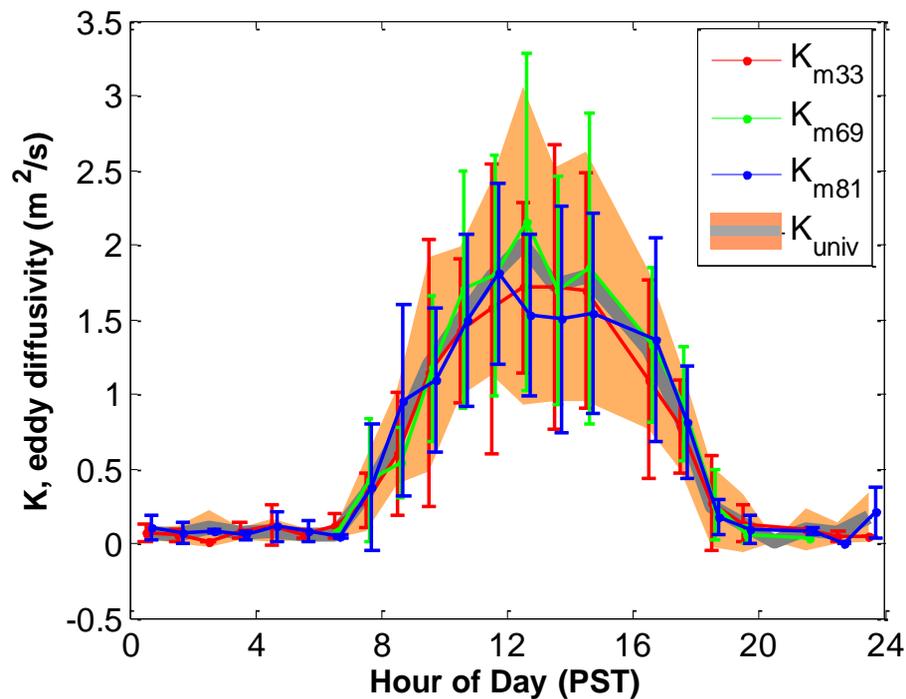


Fig. 7. Mean eddy diffusivity diurnal patterns calculated from measured fluxes and vertical gradients of methanol (K_{m33}), MBO + isoprene (K_{m69}), monoterpene (K_{m81}) and the universal K (K_{univ}) calculated according to Eq. (5). Error bars and shaded area denote standard deviations from the mean.

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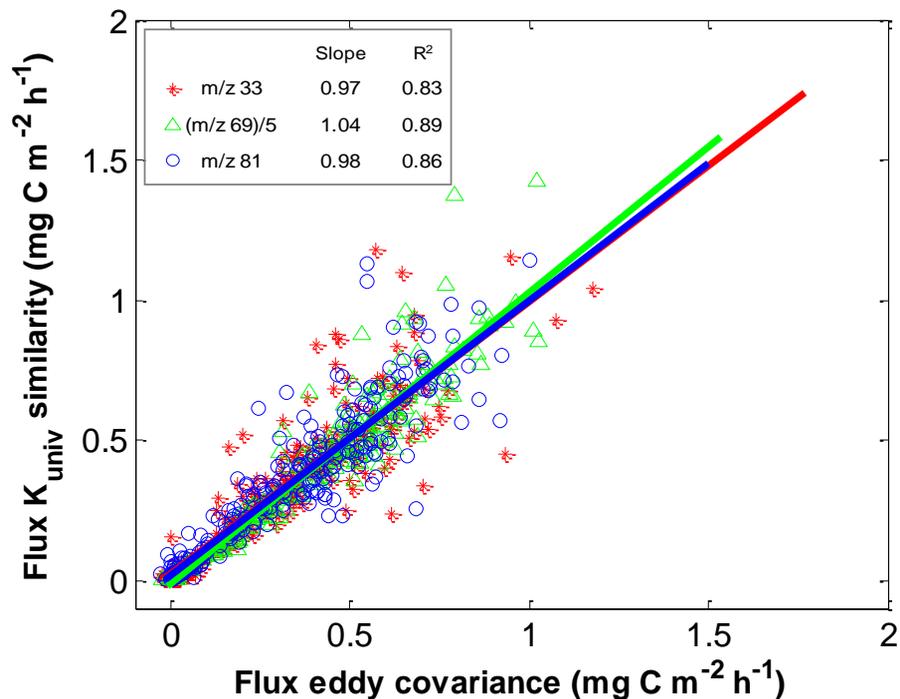


Fig. 8. Comparison between flux determined by the eddy covariance method and flux determined by the flux-gradient similarity method using the universal K (K_{univ}) for methanol (m/z 33, red star), MBO+isoprene (m/z 69, green triangle), and monoterpenes (m/z 81, blue circle). The red, green, and blue lines are the best linear fit for methanol, MBO + isoprene, and monoterpenes, respectively.

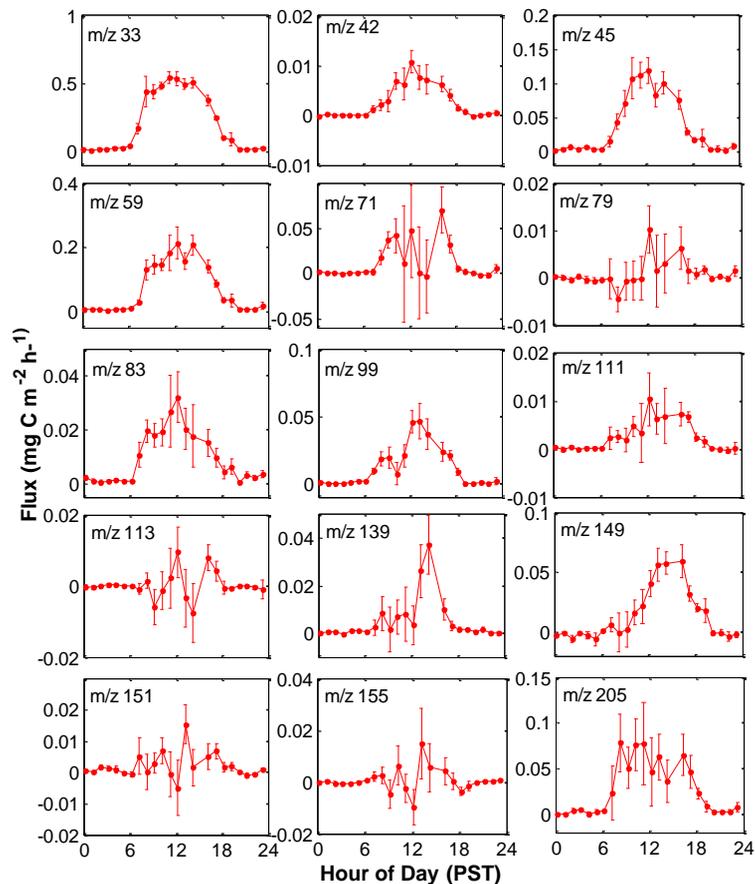


Fig. 9. Mean diurnal cycles of fluxes determined by the similarity method using the universal K for 15 ion species observed. Vertical bars denote standard errors from the mean.

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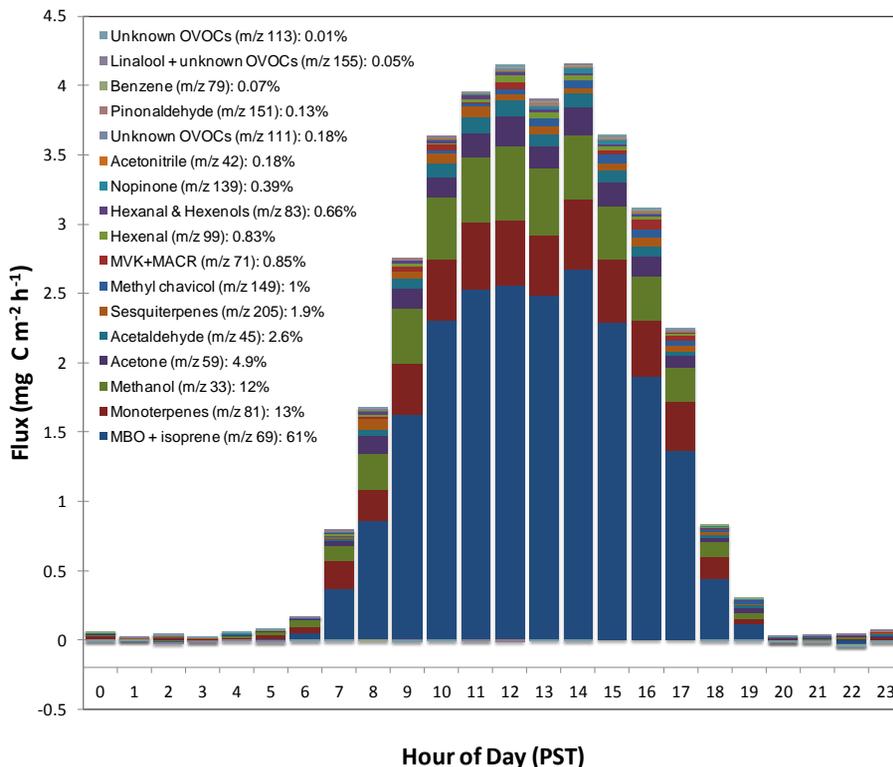


Fig. 10. Total hourly mean BVOC diurnal flux on a carbon mass basis shown as staged bar plots with contributions indicated for each of the 17 masses measured. Largest fluxes are at the bottom. The percentages in the legend indicate contributions of individual compounds to the daily total emission.

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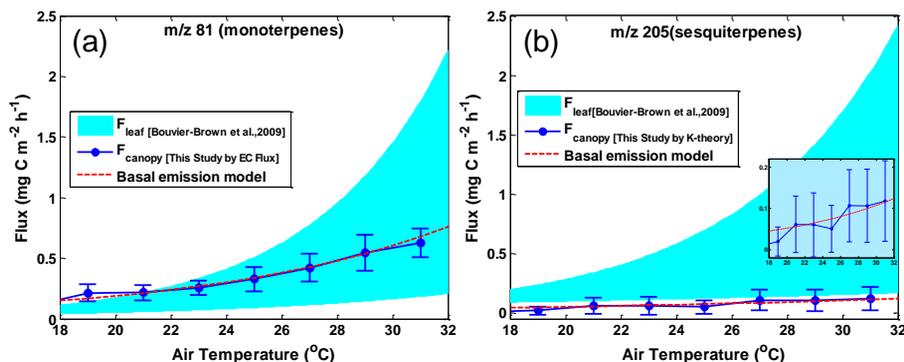


Fig. 11. Comparisons between leaf scale emissions and canopy scale emissions of **(a)** monoterpenes and **(b)** sesquiterpenes as a function of temperature. Shaded area indicates the range of expected canopy level emissions from leaf scale emissions reported previously by Bouvier-Brown et al. (2009). Blue dots represent canopy scale **(a)** eddy covariance flux for monoterpenes and **(b)** flux determined by the similarity method using the universal *K* for sesquiterpenes. All fluxes were aggregated into 2 °C temperature bins, and error bars denote standard deviations. Red dotted line represents basal emission model optimized for this study based on the canopy scale fluxes, and the inset in **(b)** enlarges the vertical scale, showing the exponential relationship between temperature and sesquiterpene flux.

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