

Water uptake and
composition of
biogenic secondary
aerosols

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Water uptake is independent of the inferred composition of secondary aerosols derived from multiple biogenic VOCs

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Abstract

We demonstrate that the water uptake properties derived from sub- and super-saturated measurements of chamber-generated biogenic secondary organic aerosol (SOA) particles are independent of their degree of oxidation determined using both online and offline methods. SOA particles are formed from the photooxidation of five structurally different biogenic VOCs representing a broad range of emitted species and their corresponding range of chemical reactivity: α -pinene, β -caryophyllene, limonene, myrcene and linalool. The fractional contribution of mass fragment 44 to the total organic signal (f_{44}) is used to characterise the extent of oxidation of the formed SOA as measured online by an aerosol mass spectrometer. Results illustrate that the values of f_{44} are dependent on the precursor, the extent of photochemical ageing as well as on the initial experimental conditions. SOA generated from a single biogenic precursor should therefore not be used as a general proxy for biogenic SOA. Similarly, the generated SOA particles exhibit a range of hygroscopic properties depending on the precursor, its initial mixing ratio and photochemical ageing. The activation behaviour of the formed SOA particles show no temporal trends with photochemical ageing. The average κ values derived from the HTDMA and CCNc are generally found to cover the same range for each precursor under two different initial mixing ratio conditions. A positive correlation is observed between the hygroscopicity of particles of a single size and f_{44} for α -pinene, β -caryophyllene, linalool and myrcene, but not for limonene SOA. The investigation of the generality of this relationship reveal that α -pinene, limonene, linalool and myrcene are all able to generate particles with similar hygroscopicity ($\kappa_{\text{HTDMA}} \sim 0.1$) despite f_{44} exhibiting a relatively wide range of values (~ 4 to 11 %). Similarly, κ_{CCN} is found to be independent of f_{44} . The same findings are also true when sub- and super-saturated water uptake properties of SOA are compared to the averaged carbon oxidation state ($\overline{\text{OS}}_{\text{C}}$) determined using an off-line method. These findings do not necessarily suggest that water uptake and chemical composition are not related. Instead, they suggest that either f_{44} and $\overline{\text{OS}}_{\text{C}}$ do not represent the main dominant composition-related

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factors controlling water uptake of SOA particles, or they emphasise the possible impact of semi-volatile compounds on limiting the ability of current state-of-the-art techniques to determine the chemical composition and water uptake properties of aerosol particles.

1 Introduction

Atmospheric oxidation of volatile, semi-volatile and intermediate volatility organic compounds (VOCs, SVOCs, and IVOCs) produces lower volatility species that partition into the condensed phase and form secondary organic aerosol (SOA) (Seinfeld and Pankow, 2003; Hallquist et al., 2009; Robinson et al., 2007; Donahue et al., 2011).

Biogenic hydrocarbons are a dominant contributor to the global VOC budget and are known for their high reactivity towards the main atmospheric oxidants (e.g. hydroxyl radical, nitrate radical and ozone), which makes them major contributors to the total atmospheric burden of SOA (Atkinson and Arey, 2003; Kanakidou et al., 2005; Hallquist et al., 2009; Goldstein and Galbally, 2007). Understanding the formation and properties of biogenic SOA is a necessary step towards assessing the influence of atmospheric aerosols on major areas of concern such as climate change and human health.

Over the last three decades, a large body of literature has been established focusing on the formation of SOA from biogenic and anthropogenic hydrocarbons (Hallquist et al., 2009; Kroll and Seinfeld, 2008; Kanakidou et al., 2005). In situ investigation of key physical and chemical processes during biogenic SOA formation in the atmosphere is complicated by the vast number of species involved and their generally low mixing ratios. Reaction chambers have therefore often been used to study these processes. The conventional reaction chamber (also referred to as smog chamber) is a large confined volume used as a batch reactor in which mixtures of atmospherically relevant trace gases (e.g. hydrocarbons, nitrogen oxides, sulphur dioxide) are oxidised in purified air under dark conditions or with the use of sunlight or artificial illumination. Such experiments can be useful in understanding the chemical and physical parameters that

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control the formation and transformation of secondary organic aerosols, their properties and atmospheric impacts. Hallquist et al. (2009) have listed the characteristics of selected smog chambers and reviewed recent developments in smog chamber SOA studies.

5 Much of the early effort has focused on the investigation of gas-particle partitioning of VOC oxidation products through quantification of the aerosol formation potential (aerosol yield) of small aromatic and biogenic hydrocarbons (Pandis et al., 1992; Odum et al., 1996; Griffin et al., 1999). Other studies have attempted to identify the molecular composition of the oxidation products of anthropogenic and biogenic precursors, mainly aromatic and monoterpene compounds, and have offered detailed reaction mechanisms for the formation of various chemical species (Forstner et al., 1997; Glasius et al., 2000; Jang and Kamens, 2001; Jaoui and Kamens, 2003; Kleindienst et al., 2004). Over the last decade or so, a number of studies have suggested that the first reported SOA yield values for some the most commonly studied VOCs were too low. Revised, higher SOA yield values have been reported for a number of SOA systems including α -pinene (Ng et al., 2006; Chan et al., 2007; Shilling et al., 2008), isoprene (Chan et al., 2007; Kroll et al., 2006), toluene (Hildebrandt et al., 2009) and m-xylene (Ng et al., 2007). A large fraction of the studies reported in the literature have focused on the characterisation of the chemical and/or microphysical properties of SOA systems produced from the oxidation of an individual VOC precursor. Examples of these studies include the effect of NO_x concentration (Presto et al., 2005b) and UV radiation (Presto et al., 2005a) on α -pinene SOA formation; the formation of SOA from isoprene oxidation (Carlton et al., 2009; Kroll et al., 2006; Dommen et al., 2006; Dommen et al., 2009); the contribution of second generation oxidation products to β -caryophyllene SOA formation (Li et al., 2011). Other studies have investigated the SOA formation for a range of VOC compounds under comparable experimental studies. For example, Li et al. (2006a) reported SOA yields and gas-phase oxidation products from the ozonolysis of 10 terpene compounds. In a successive study, the same researchers reported SOA yield values and detailed characterisation of gas-phase oxidation products

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from a series of photooxidation experiments conducted using 16 terpene compounds (Lee et al., 2006b). A more detailed analysis of the contribution of first and second-generation oxidation products to SOA production and formation mechanisms, in the latter set of experiments, were presented by Ng et al. (2006). More recently, Chhabra et al. (2011) compared spectral and elemental ratio data from the high-resolution Aerodyne aerosol mass spectrometer to offline species identification analysis in order to investigate changes in functional and elemental composition during SOA formation and transformation using a wide range of VOC precursors.

Simultaneously, several efforts have been made over the last decade or so to characterise the cloud condensation nuclei (CCN) activity and hygroscopic properties of SOA particles produced during the ozonolysis or photooxidation of biogenic and anthropogenic VOCs. VanReken et al. (2005) investigated the potential of 5 biogenic SOA types, produced in dark ozonolysis experiments, to act as CCN. Varutbangkul et al. (2006) studied the sub-saturated water uptake properties of SOA formed by photooxidation of monoterpenes, sesquiterpenes and oxygenated terpenes. The influence of the initial precursor mixing ratio on the composition and/or hygroscopic properties have been reported for SOA produced from oxidation of α -pinene (Duplissy et al., 2008; Shilling et al., 2009; King et al., 2009) and β -caryophyllene (Alfarra et al., 2012). The influence of particle volatility on composition, hygroscopic properties and CCN activity have been studied for a number of SOA systems including α -pinene (Meyer et al., 2009; Tritscher et al., 2011) and β -caryophyllene (Asa-Awuku et al., 2009; Tang et al., 2012; Frosch et al., 2013).

Attempts to relate the chemical composition of SOA particles to their hygroscopic and CCN properties have recently been made. Massoli et al. (2010) investigated the relationship between the oxidation level (O:C ratio) and the hygroscopic and CCN properties of laboratory SOA particles generated via OH radical oxidation of α -pinene, trimethylbenzene (TMB) and *m*-xylene in separate flow reactor experiments. The study found that the hygroscopic properties at 90 % relative humidity (RH) increased linearly with O:C value for the studied systems. Similarly, an increase in the CCN activity

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was reported as a function of O : C. However, the later relationship was reported to be system-dependent and not a linear one. A semi-empirical parameterisation of hygroscopic growth at 90 % RH and O : C was proposed, whereas a general parameterisation for the CCN activity vs. O : C relationship was reported to be difficult to establish.

5 Similarly, Forsch et al. (2011) found that the level of oxygenation of α -pinene SOA increased with decreasing precursor mixing ratio and with chemical ageing, whereas CCN activity was largely independent of O : C ratio. Duplissy et al. (2011) determined a simple empirical linear relation between sub-saturated hygroscopic growth and the level of oxygenation of SOA produced from the photooxidation of α -pinene, isoprene
10 and TMB. The same analysis was also performed on ambient organic aerosol data at two contrasting locations, where similar findings were reported.

In this study, we extend the investigations of SOA composition and properties using a carefully selected set of 5 structurally different biogenic VOCs covering a range of chemical reactivity. Specifically, we will (i) illustrate the variability in the chemical
15 composition and sub- and super-saturated water uptake properties of SOA generated from the photooxidation of the selected VOCs under controlled and comparable conditions in a reaction chamber; and (ii) investigate and discuss the relationship between the sub- and super-saturated water uptake of the generated SOA particles and their chemical composition determined using on- and off-line analytical methods.

20 2 Experimental

2.1 Chamber description

Experiments were conducted in the photochemical aerosol reaction chamber at the University of Manchester (Alfarra et al., 2012; Hamilton et al., 2011). The chamber is run as a batch reactor where the composition of the gaseous precursors, oxidising
25 environment, relative humidity and temperature are controlled. It comprises an 18 m³ (3 m (H) × 3 m (L) × 2 m (W)) FEP Teflon bag mounted on three horizontal rectangular

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aluminium frames. The central rigid frame is fixed, with the upper and lower frames free to move vertically, allowing the bag to expand and collapse as sample air is introduced and extracted. Air is supplied to the chamber by a blower at a flow of $3 \text{ m}^3 \text{ min}^{-1}$. The air is dried and filtered for gaseous impurities and particles using a series of Purafil (Purafil Inc., USA), charcoal and HEPA filters (Donaldson Filtration, USA), prior to humidification with ultrapure deionised water. Halogen bulbs and a 6 kW Xenon arc lamp are mounted on the inside of the enclosure housing the bag, which is coated with reflective “space blanket”, serving to maximise the irradiance in the bag and to ensure even illumination. The combination of illumination has been tuned and evaluated to mimic the atmospheric actinic spectrum over the wavelength range 290–800 nm, and has a maximum total actinic flux of $0.7 \times 10^{18} \text{ (photon s}^{-1} \text{ m}^{-2} \text{ nm}^{-1}\text{)}$ over the region 460–500 nm. The calculated $j\text{O}(^1\text{D})$ value during the reported experiments was $3.6 \times 10^{-5} \text{ s}^{-1}$ (290–340 nm) and $j\text{NO}_2$ was $6 \times 10^{-4} \text{ s}^{-1}$ (290–422 nm).

Precursor VOCs are introduced into the bag via injection into a heated glass bulb, which is continually flushed with a flow of filtered, high purity nitrogen (ECD grade, 99.997%). The desired concentration of NO_x is controlled by injection of NO_2 from a cylinder into the charge line. Relative humidity (RH) and temperature are measured at the centre and the edge of the chamber (by dewpoint hygrometer and a cross-calibrated thermocouple and resistance probe), and are controlled by diverting air through the inlet humidification circuit when filling the bag and by controlling the set point of the air conditioning, which feeds the airspace between the bag and its enclosure. Cycling between experiments is facilitated by automated computer control and monitoring of key chamber conditions. Pre- and post-experiment cleaning processes consists of a series of fill/flush cycles with $3 \text{ m}^3 \text{ min}^{-1}$ flow of clean air, enabled by control of electro-pneumatic valves followed, post-experiment, by an overnight soaking at high mixing ratios of ozone (2–2.5 ppmV, supplied by a high capacity ozone generator). Each cycle takes approximately 12 min and cleaning is normally achieved after approximately 6 cycles. A particle mass background value of $0.1\text{--}0.2 \mu\text{g m}^{-3}$ is attained following these cleaning procedures.

2.2 Experimental conditions and methodology

SOA particles were formed in experiments involving the photooxidation of individual precursors in the presence of NO_x and in the absence of seed particles. Precursor VOCs limonene (R-(+)-Limonene, Fluka, $\geq 99.0\%$), myrcene ($\geq 95.0\%$, Fluka), linalool ((-)-Linalool, $\geq 98.5\%$, Fluka), α -pinene (98%, Aldich) and β -caryophyllene (Aldrich, purity not specified) were investigated, each at two different nominal initial mixing ratios of 50 and 250 ppbV, in order to study the effect of initial precursor mixing ratio on SOA properties and composition. VOC measurements (described in Sect. 2.3) were made during some of the experiments and the measured initial precursor mixing ratios during those experiments are reported in Table 1. Where not measured, the nominal initial precursor mixing ratios were estimated from the amount injected into the heated bulb and introduced into the chamber via the high flow rate inlet without accounting for losses. Experimental protocol consisted of the addition of required amounts of the precursor, NO_x and water vapour during the final filling cycle of the chamber; ensuring that all ingredients are well-mixed by the end of the cycle as a result of the high flow rate at which the clean air is introduced. Air conditioning was switched on shortly before the arc lamp and halogen bulbs were illuminated, marking the start of the photochemistry and the experiment (t_0). An overview of the conditions used in each experiment is given in Table 1.

2.3 Instrumentation

Real-time broad chemical characterisation of the SOA was made using a compact Time-of-Flight Aerosol Mass Spectrometer (cToF-AMS, Aerodyne Research Inc., USA). A detailed description of the instrument, its operation and calibrations can be found elsewhere (Drewnick et al., 2005; Canagaratna et al., 2007). The instrument was operated in the standard configuration, taking both mass spectrum (MS) and particle-time-of-flight (PToF) data and was calibrated for ionisation efficiency using 350 nm monodisperse ammonium nitrate particles. The vapouriser was set at approximately

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600 °C and data were collected at a time resolution of 2 min. A collection efficiency value of unity was applied to these data, based on evidence from a previous chamber study (Alfarra et al., 2006).

A hygroscopicity tandem differential mobility analyser (HTDMA) was used to measure on-line size resolved water uptake at 90 % RH. A technical description of the instrument was provided in Good et al. (2010a) and Cubison et al. (2006). Briefly, the HTDMA dries the aerosol sample to < 10 % RH using a Nafion[®] drier (Perma Pure, MD-110-12, Toms River, NJ, USA). A DMA (BMI, Haywood, CA, USA) selects particles of a single mobility. In this work, diameters chosen were larger than the mode of the number size distribution, thus avoiding the sampling of a significant fraction of multi-charged particles. The sample is then humidified to 90 % RH using a humidifier, comprising a Gore-Tex[®] tube running through a controlled humid counter-flow of air. The humidified size selected sample is then passed through a residence coil for 15 s. A second DMA (BMI, Haywood, CA, USA) and water-based condensation particle counter (WCPC 3782, TSI Inc., USA) are then used to measure the size distribution of the humidified sample. The operating procedure of the HTDMA was validated by sampling ammonium sulphate and sodium chloride test aerosols, following Good et al. (2010a) and inverted using TDMAInv (Gysel et al., 2009). The data from the HTDMA are reported in terms of the hygroscopic growth factor ($GF_{D_0, RH}$), the wet particle diameter at a given RH divided by the particle's dry diameter (D_0).

The cloud condensation nuclei (CCN) activity of the particles was characterised using a continuous flow CCN counter (Droplet Measurement Technologies, USA) combined with a Vienna style DMA (Winklmayr et al., 1991) and a condensation particle counter (CPC 3010, TSI Inc., USA). The DMA was used to generate monodisperse aerosol in the size range between 20 nm and 500 nm dry diameter. These particles were directed in parallel to the CPC and CCN counters. The inlet flow in the CCN counter was 0.5 Lpm and it was operated at different supersaturations in the range between 0.07 % and 1 % for each particle size. The calibration and quality assurance procedures carried out for this setup is described in Good et al. (2010a). Using the in-

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5 instrumental set-up described above CCN and CN number size distributions were measured at a temperature difference (ΔT) equivalent to super-saturations between 0.07 % and 1.0 %. The CCN and CN number size distributions were then inverted to account for charging efficiency and multiple charging (Good et al., 2010a). The inverted number size distributions were then used to calculate the activated fraction (CCN/CN) as
10 a function of dry size. Sigmoidal functions were then fitted to the activation curve, from which the dry diameter at which 50 % of the particles were activated was calculated. Given that the aerosol was internally mixed, the diameter at which 50 % of the particles activate (D_{50}) was reasonably interpreted to be the dry diameter at which the
15 particles are activated as CCN at the set-point super-saturation (Good et al., 2010b). κ values (Petters and Kreidenweis, 2007) are then iteratively derived from the CCN derived critical super-saturation and dry diameter.

Total particle number concentrations were measured using a water based condensation particle counter with a minimum particle size cut off of 2.5 nm (WCPC 3786, TSI Inc., USA). NO and NO₂ mixing ratios were measured using a chemiluminescence gas analyser (Model 42i, Thermo Scientific, MA, USA). Ozone was measured using a UV photometric gas detector (Model 49C, Thermo Scientific, MA, USA).

20 The gas phase organic compounds within the chamber were measured using Chemical Ionisation Reaction Time-of-Flight Mass Spectrometry (CIR-TOF-MS). This technique has been described in (Wyche et al., 2007; Jenkin et al., 2012). The CIR-TOF-MS instrument comprises a bespoke, temperature controlled ($40^{\circ}\text{C} \pm 1^{\circ}\text{C}$) radioactive (^{241}Am) ion source/drift tube assembly, coupled via a system of ion transfer optics to an orthogonal time-of-flight mass spectrometer (Kore Technology, UK). In this instance, hydrated hydronium ions ($\text{H}_3\text{O}^+ \cdot (\text{H}_2\text{O})$), generated from a humidified N₂ carrier gas
25 (purity = 99.9999 %), were employed as the primary chemical ionisation reagent. Under the conditions of the applied drift tube parameters, ion-molecule reaction between the analyte VOC/OVOC, M, and the hydrated hydronium ion would yield a protonated VOC/OVOC product ion (MH^+) for mass spectrometric analysis, providing the analyte has a proton affinity greater than that of the water dimer (808 kJ mol^{-1}). Depending

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on the energies involved and chemical structure of the VOC/OVOC analyte, the MH^+ product may undergo fragmentation to produce certain daughter ions. Mass spectrometric data were recorded by the CIR-TOF-MS over the range 1 and 300 Da with a time resolution of 1 min and the signals of interest were converted to ppbV via both experimental calibration and theoretical calculation using the methods described in Jenkin et al. (2012).

2.4 Offline analysis: fourier transform ion cyclotron resonance mass spectrometry

Aerosol samples were collected onto 47 mm quartz fibre filters (Whatman) at a flow rate of $3\text{ m}^3\text{ min}^{-1}$. For this purpose, filters were placed in a holder, positioned in the chamber vent line. After sampling, filters were immediately placed in pre-cleaned glass vials and stored below -18°C until analysis. Approximately an area of 7/8th of the filter sample was extracted into high purity water, filtered and reduced to 1 ml using a vacuum solvent evaporator (Biotage, Sweden) and the remaining portion retained for other analysis. In total 8 filters were analysed, corresponding to the experiments labelled in Table 1.

Samples were analysed at high mass resolution using a Bruker APEX 9.4 T Fourier Transform Ion Cyclotron Resonance Mass Spectrometer. Extracts were sprayed at a flow rate of $2\ \mu\text{L min}^{-1}$, into an Apollo II electrospray interface with ion funnelling technology. Spectra were acquired in both positive and negative ion mode over the scan range m/z 100–3000 using the following MS parameters: nebulising gas flow: 0.9 L min^{-1} , drying gas flow: 5 L min^{-1} , drying temperature: 190°C , collision cell accumulation: 0.05–0.5 s, and data acquisition size: 2 Mb (yielding a target resolution of 130 000 at m/z 400). Data were analysed using DataAnalysis 4.0 software (Bruker Daltonics, Bremen, Germany). The instrument was calibrated using protonated (positive ion mode) or deprotonated (negative ion mode) arginine clusters.

The data analysis software employed a user defined set of conditions to determine the molecular formulae. For the analysis reported in this paper, the following molecular

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configurations were allowed; unlimited numbers of C, H and O and 0–3N. Chemical formulae were assigned to the masses of singly charged ions $100 < m/z < 600$ after internal recalibration using fatty acid signals (a solvent contaminant present in all samples) and a maximum error of 5 ppm for ions with a relative intensity of greater than 0.01 %. Isotopic ratios were used for further confirmation of the formulae. Background contaminant peaks also seen in pure water and blank extracted filters were manually removed from the analysis.

For each SOA filter analysed, the molecular formulae were determined using FTI-CRMS. Compounds were screened to remove unlikely molecular formulae. Compounds were removed from the list if they contained any of the following; H:C < 0.5, O:C > 3 and DBE > 20. Using the formulae, the intensity weighted mean O:C and H:C ratios of each sample were determined (Bateman et al., 2010) and these used to calculate the average carbon oxidation state (\overline{OS}_C) according to Kroll et al. (2011).

2.5 Rationale behind the choice of precursors

The emitted speciation of biogenic VOCs is highly complex, and published studies have typically reported up to ca. 20 species making notable contributions to monoterpene, sesquiterpene and oxygenated VOC fluxes from a variety of vegetation types at a variety of locations (Owen et al., 2001; Boissard et al., 2001; He et al., 2000; Spanke et al., 2001). Because of differences in structure and reactivity, these species are oxidised in the atmosphere on timescales ranging from minutes to days and possess a range of propensities to form SOA, which may exhibit a range of physical properties reflecting its diverse chemical composition. In this study, we have identified a subset of biogenic VOCs with an aim to represent a broad range of emitted species and the corresponding range in the rates and chemical complexity of their subsequent degradation. Based on the available information on the atmospheric lifetimes of observed biogenic VOCs with respect to reaction with OH radicals and O₃, and current understanding of their known or probable degradation pathways, it is reasonable to divide the biogenic VOCs into a series of structural subsets possessing similar characteris-

tics (Atkinson and Arey, 2003). As a result, consideration of a single compound from each category provides a reasonable starting point for simplifying the emitted speciation whilst maintaining the range in oxidation timescales and propensities to generate SOA. Table 2 provides a summary of the categories, identifying possible representatives and their atmospheric lifetimes with respect to reaction with OH and O₃ at typical mixing ratios (Atkinson and Arey, 2003). The identified compounds thus provide a series of lower reactivity (α -pinene), very high reactivity (β -caryophyllene), intermediate reactivity (limonene, myrcene) and more reactive oxygenated (e.g., linalool) species.

3 Results and discussion

3.1 Gas-phase oxidation and SOA formation characteristics

Figure 1 shows typical temporal profiles of each of the studied gaseous precursors under lower mixing ratio conditions (see Table 1), along with the concomitant production profile of a selection of their primary oxidation products. Each of the different precursors, (a) α -pinene, (b) limonene, (c) myrcene, (d) linalool and (e) β -caryophyllene, exhibited similar and reproducible oxidation profiles, with mixing ratios decaying to minimal levels between approximately 100 and 200 min after the start of the photochemistry. Each of the primary oxidation products shown in Fig. 1, i.e. (a) pinonaldehyde, (b) limononaldehyde, (c) 4-vinyl-4-pentenal (d) 4-hydroxy-4-methyl-5-hexen-1-al and (e) β -caryophyllon aldehyde, appeared immediately after lights on, increasing at various rates to peak mixing ratios in the approximate range 4–15 ppbV, between 100–200 min. Following each respective product peak and depletion of the precursor, the various primary oxidation products began to decrease in concentration at varying rates, as the magnitude of loss processes to secondary products and to the aerosol phase overtook production rates. Similar gas phase results have been obtained from several other simulation chamber studies investigating biogenic oxidation (Lee et al., 2006b; Ng et al., 2006).

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The time taken to reach half of the initial mixing ratio of the individual precursors, its half-life, is as an empirical representation of the inverse of the total loss rate of the parent hydrocarbon under the specified experimental conditions. This parameter accounts for losses via reaction with oxidants (e.g. ozone, hydroxyl radical), any potential photolysis and physical losses to the chamber walls. Assuming the latter is unimportant (Matsunaga and Ziemann, 2010), the chemical half-life can be used to infer and compare the reactivity of the parent hydrocarbons under the reported experimental conditions. The derived chemical half-life values are summarised in Table 1, and are broadly consistent with the precursor reactivity data presented in Table 2 and discussed in the rationale section (Sect. 2.5). The shortest lifetimes (i.e. highest loss rates) were observed for β -caryophyllene, while the longest lifetime was found to be that of α -pinene. It has to be noted that ozone was not used as a primary oxidant during any of experiments reported in this study, and it was formed as a result of the photochemistry. There seems to be some difference between the life-times of myrcene when higher and lower initial mixing ratios were used. Linalool, on the other hand, does not exhibit the same behaviour, as its life-times are comparable under lower and higher initial mixing ratios. It is not possible to attribute these differences to a mechanistic cause given that initial VOC/NO_x ratios were different during those experiments and that no explicit degradation mechanisms exists for these compounds.

The SOA production profiles during those same high and low initial mixing ratio experiments are also broadly consistent with the chemical half-life data; i.e. the peak particle number concentration was reached in a relatively shorter time in the higher initial mixing ratio cases compared to the lower ones as illustrated in Table 1. Further inspection of the times taken to reach peak number concentration indicates that, for the lower initial mixing ratios, the SOA formation is consistent with the gas phase reactivity of the individual precursors. For example, β -caryophyllene SOA peak number was the first to be reached, while α -pinene SOA was the slowest. Owing to the limited availability of data, SOA mass yield values have been calculated and reported for a subset of experiments. The yield is defined as the percentage ratio of the amount of SOA formed

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SOA (right panels), after about four hours of the start of the photooxidation (lights on). All the spectra have been normalised to the total SOA signal; the fractional contribution of each mass fragment to the total SOA signal ($f_{m/z}$) is shown and can be used to compare the broad chemical composition of the SOA formed from the photooxidation of each precursor and also changing chemical composition of each SOA system as a function of photochemical ageing. Detailed inspection of the mass spectral patterns reveals that one of the most significant differences among the various SOA systems is the fractional contributions of mass fragments 43 and 44 to the total SOA signal (i.e. f_{43} and f_{44}). This is a recognised feature of most AMS datasets and the two mass fragments have been widely used to discuss the level of oxygenation of organic aerosols in many laboratory and field studies (Ng et al., 2010; Alfarra et al., 2004). Other differences among the mass spectral patterns of the five SOA systems appear in m/z range between 55 and 60, where linalool and myrcene SOA share similar patterns, but which are different to those observed for the other precursors. β -caryophyllene SOA has a higher contribution from relatively larger mass fragments including 77, 79, 81 and 83 as well as 91, 93 and 95. As a result of the nature of electron impact ionisation and the lack of pre-separation methods, it is not possible to associate these mass fragments with specific compounds. However, some of those fragments (e.g. m/z 77 and 91) are traditionally associated with the fragmentation of aromatic compounds. A comparison of the mass spectral patterns of the fresh and aged SOA produced by each precursor shows hardly any change with the exception of the changes in f_{43} and f_{44} , discussed below.

The fractional contributions of mass fragments 43 and 44 to the total organic signal (f_{43} and f_{44}) have been widely used as markers for “less” and “more” oxygenated organic compounds, respectively, in several laboratory and field studies using the aerosol mass spectrometer (Ng et al., 2010; Pfaffenberger et al., 2012; Alfarra et al., 2004, 2006; Zhang et al., 2005; Lanz et al., 2007). The former is mostly associated with carbonyl-based compounds, while f_{44} is a marker for compounds containing di- and poly-carboxylic acid groups (Alfarra, 2004; Takegawa et al., 2007). It has been shown

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that atmospheric ageing leads to an increase in f_{44} (Alfarra et al., 2004; Aiken et al., 2008; de Gouw et al., 2005; Jimenez et al., 2009); therefore, this value is considered as an indicator of atmospheric ageing. The temporal trends of f_{44} for all biogenic SOA systems studied here are shown in the top panel of Fig. 3, both for the low and high initial precursor mixing ratio experiments. Similarly, those of f_{43} are shown in the bottom panel of the same figure. It can quickly be noticed that the different SOA produced using the selected biogenic precursors have a wide range of chemical characteristics expressed by f_{44} and f_{43} both in terms of actual values as well as temporal behaviour as a function of photochemical ageing. For example, the β -caryophyllene and linalool SOA show the lowest values of f_{44} and these values appear to be only slightly influenced by photochemical ageing. On the other hand, photochemical ageing appears to more than double the value of f_{44} of freshly produced myrcene SOA after about 6 h of processing (from about 4 to 10 %). SOA produced from low initial mixing ratio of α -pinene and limonene have relatively high f_{44} values (11 and 7 %, respectively) compared to the other SOA systems studied here, however these values do not change much as a result of photochemical ageing. Linalool and β -caryophyllene SOA exhibit relatively low f_{44} values around 4 %, which increase slightly to about 5–6 % as a result of photochemical ageing.

The initial fractional contribution values of mass fragment 43 (f_{43}) span quite a wide range from 12–22 %; changing to 8–23 % after six hours of photochemical ageing. The trends of f_{43} appear to show, to a large extent, a reverse pattern to those of f_{44} . With the exception of limonene SOA, f_{43} values decrease with photochemical ageing. Such trends are consistent with processing of carbonyl-containing compounds to more oxidised carboxylic acid-containing compounds. An explanation of the reverse pattern observed for limonene SOA (i.e. the increase in the f_{43} fraction with photochemical ageing) is not obvious. These results show that composition of SOA generated from a single biogenic precursor cannot and should not be used as a proxy for biogenic SOA in general. Additionally, the values of f_{44} which are widely used in the literature to indicate the level of oxygenation of organic aerosol in field and laboratory studies is

very much dependent on the precursor, the extent of photochemical ageing as well as on the initial experimental conditions.

Another intriguing observation in this dataset is that the effect of initial precursor mixing ratio (in this study 50 vs. 250 ppbV) on the chemical characteristics of the produced SOA represented by f_{44} and f_{43} is not uniform and does not appear to follow a specific pattern. Previous studies reported an increase in the f_{44} value for α -pinene SOA as mass loading or initial precursor mixing ratio were decreased (Duplissy et al., 2008; Shilling et al., 2009; Pfaffenberger et al., 2012). Our work shows that although this appears to be also true for relatively aged linalool, limonene and β -caryophyllene SOA (Alfarra et al., 2012), it is not necessarily always the case for SOA generated from other biogenic precursors. For example, changing the initial myrcene mixing ratio does not appear to have a measurable effect on the f_{44} and f_{43} values of the produced SOA regardless of photochemical ageing. The results also indicate that changing the initial precursor mixing ratio does not influence the f_{44} value for freshly produced SOA in most of the cases reported here. Of course, this finding is limited to the mixing ratio ranges explored in the current study and characterisation of the variation in composition across a broader range of initial mixing ratios is required.

In order to compare the composition of the SOA particles generated from the five biogenic precursors studied here in the context of the composition of ambient organic aerosols, a 2-dimensional representation of the relationship between f_{44} and f_{43} is shown in Fig. 4. The dotted lines in this figure mark the well-defined triangular space occupied by data points from a large number of ambient datasets collected at a large number of locations in the Northern Hemisphere and discussed in detail by Ng et al. (2010). Data of more aged, and hence more oxygenated and by inference less volatile, organic aerosols tend to show an increasing value of f_{44} and a decreasing value of f_{43} and therefore appear more towards the top of the triangle-shaped space. This space is dominated by the AMS factor defined as low volatility oxygenated organic aerosols (LV-OOA), while data in the lower half of the triangle is more representative of the AMS defined factor as semi-volatile oxygenated organic aerosols (SV-OOA) (Jimenez et al.,

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2009; Ng et al., 2010). The chamber generated SOA data from the photooxidation of myrcene and β -caryophyllene are located in the bottom right corner of the ambient triangle, indicating that the composition of those two SOA systems is consistent with ambient organic aerosols characterised by the SV-OOA factor (Ng et al., 2010). It can also be seen that photochemical ageing leads to a more significant processing of the myrcene SOA particles transforming their composition towards being more LV-OOA and less SV-OOA. On the other hand, the composition of SOA particles produced from the photooxidation of limonene, linalool and α -pinene appear to be less representative of ambient organic aerosol composition as their data points fall outside the ambient triangle featuring higher values of f_{43} . However, photochemical ageing of those three systems does, indeed, help transform their composition bringing it closer to the edge of the ambient triangle.

The representation of the chamber SOA data in Fig. 4 confirms our earlier point regarding the wide range of chemical characteristics exhibited by chamber SOA depending on the choice of precursor and the initial experimental conditions. Care should be taken not to assume that SOA produced from a specific precursor under specific conditions could be used as a proxy for biogenic SOA in general. The data points marked by open circles in Fig. 4 represent SV-OOA components from organic aerosols measured at sites likely to have strong influence from biogenic SOA sources: Duke Forest, Hyytiälä, Thompson Farm, Chebogue, and Egbert (Allan et al., 2006; Williams et al., 2007; Cottrell et al., 2008; Raatikainen et al., 2010; Slowik et al., 2010) as reported by Ng et al. (2010). Similar to our biogenic SOA data, these do not appear to be located at a specific area within the lower half of the triangle space; possibly indicating the lack of universal mass spectral signature, and therefore chemical composition, of fresh biogenic SOA. This implies that, unlike aged secondary organic aerosols, choosing or claiming a representative mass spectral signature (chemical composition) for fresh biogenic secondary organic aerosol is not likely to be a straightforward task, as it appears to be dependent on location and conditions.

3.3 Hygroscopic properties

Hygroscopicity measurements were carried out during one higher and one lower mixing ratio experiments for each SOA precursor, except for limonene experiments which were repeated three times and β -caryophyllene experiments, which were repeated four times. For all five precursors investigated, the HTDMA consistently measured a narrow particle growth factor distribution at each selected dry particle diameter, indicating that particles of the same size have similar composition at any given time (i.e. internally mixed). Being internally mixed the hygroscopicity of particles in the chamber of a given size can be well described by a single κ value (Petters and Kreidenweis, 2007) calculated from the mean of the fitted growth factor probability density function (GF-PDF) (Gysel et al., 2009). Hereafter hygroscopic growth factor refers to the mean of the GF-PDF unless otherwise stated. Figure 5 illustrates the observed range of hygroscopic properties, represented by κ_{HTDMA} , for all selected dry particle diameters (D_0) for particles generated from each initial mixing ratio of each precursor. The β -caryophyllene produced the least hygroscopic particles ($\kappa_{\text{HTDMA}} \sim 0.01$), while α -pinene, limonene, linalool and myrcene produced particles with higher hygroscopicity ($\kappa_{\text{HTDMA}} \sim 0.03$ to 0.14). The hygroscopic behaviour of each precursor is discussed in more detail below. Table 3 summarises the hygroscopic properties, represented by κ_{HTDMA} , and range of growth factors observed for each precursor.

3.3.1 α -Pinene SOA

Photooxidation of 50 ppbV α -pinene generated particles with higher hygroscopic growth factors compared to those produced using the 250 ppbV conditions. The hygroscopicity of the particles averaged over each hour of the experiment tended to increase throughout the 50 ppbV experiment, in contrast to the 250 ppbV experiment where the hourly averaged hygroscopicity increased before stabilising after approximately 4 h of photooxidation.

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3.3.2 β -Caryophyllene SOA

Photooxidation of 50 and 250 ppbV β -caryophyllene produced the least hygroscopic particles compared to all other precursors in this study. The average hygroscopic growth factors were similar for both initial mixing ratios. The κ_{HTDMA} averaged over each hour of the experiment appeared to increase slightly with time ~ 0.005 during a typical experiment, but this increase was generally within the measurement uncertainty. The β -caryophyllene hygroscopicity measurements shown here have also been presented by Hamilton et al. (2011) in the context of utilising β -caryophyllene to generate biogenic organic seed particles for chamber experiments, and by Alfarra et al., (2012) in the context of investigating the effect of initial VOC mixing ratio on hygroscopic properties.

3.3.3 Limonene SOA

Photooxidation of 50 ppbV limonene produced slightly more hygroscopic particles compared to those produced in the 250 ppbV case. The 50 ppbV experiment showed the average κ_{HTDMA} over 1 h intervals increasing slightly by 0.02 from hour 1 to hour 4. The 250 ppbV experiment showed the average κ_{HTDMA} over 1 h intervals increasing slightly by 0.02 from hour 1 to hour 5.

3.3.4 Linalool SOA

The average hygroscopic growth factors of particles produced from the photooxidation of 50 and 250 ppbV linalool were similar. However particles in the 250 ppbV case exhibited a slightly higher range. The hygroscopicity increased throughout experiments, where for the 50 ppbV case the average κ_{HTDMA} over 1 h intervals increased by 0.03 from hour 1 to hour 5. The average κ_{HTDMA} for 250 ppbV experiment, on the other hand, increased over 1 h intervals by 0.04 from hour 1 to hour 5. After ~ 1.5 h 150 nm and 200 nm diameter particle were selected alternately by the HTDMA. A consistent

difference between the hygroscopicity of the two selected particle sizes was observed of ~ 0.01 (κ_{HTDMA}), indicating that there may be difference in their composition; this though is within the measurement uncertainty of the TDMA ($\text{gf} \pm 0.02$) (Massling et al., 2011).

3.3.5 Myrcene SOA

The growth factors of particles produced from the photooxidation of 50 ppbV myrcene appeared to be slightly higher than those produced in the 250 ppbV experiment. In both cases, there was a measurable difference in the hygroscopicity of particles sampled alternately at different sizes. The difference was typically between 0.03 and 0.05 (κ_{HTDMA}) when alternating between sampling different sizes. When a larger size was selected the hygroscopicity was always lower than that of the smaller size sampled before and after. The hygroscopicity of particles at single selected size increased with time throughout the myrcene experiments. The change in particle hygroscopicity was ~ 0.1 (κ_{HTDMA}) from the start to the end of the experiments.

3.3.6 Trends in hygroscopicity

For each of the five precursors, the hygroscopicity of particles sampled throughout the experiments tended to increase, based on the hourly averaged κ_{HTDMA} values during each experiment for all dry sizes. Small particles sampled by the HTDMA at any particular time have been produced later (and exposed to shorter gaseous OVOC histories) than larger particles. Considering each selected dry size separately, there is correlation between (increasing) hygroscopicity and time (i.e. degree of oxidation). For β -caryophyllene the correlation between κ_{HTDMA} and time is sometimes large ($r^2 > 0.9$) but at other times much lower, whilst for limonene, linalool and myrcene there is generally a strong ($r^2 > 0.9$) correlation of κ_{HTDMA} and time. The increases of κ_{HTDMA} with time indicates that SOA formed from molecules which partition from the gas phase later in the experiments are more hygroscopic. Given that the observed change in

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κ_{HTDMA} is small during β -caryophyllene experiments (and of similar magnitude to the HTDMA's precision) it is only possible to assert that the hygroscopicity of molecules formed throughout the β -caryophyllene experiments is identical within measurement uncertainty.

3.4 CCN activity

Critical supersaturations derived from the D_{50} were obtained for 50 ppbV α -pinene, β -caryophyllene, limonene, linalool and myrcene experiments. D_{50} measurements were also obtained for 250 ppbV linalool and myrcene experiment. The CCNc data set is significantly limited compared to other data sets due to the fact that if a too high or too low supersaturation is set in the CCNc either all or none of the particles activate over the narrow size distribution in the chamber and a D_{50} cannot be determined. Table 4 summarises the range of κ_{CCN} values. Figure 6 illustrates the lack of temporal trends in CCN activity; this observation though is from a much more limited data set than the HTDMA. The higher mixing ratio experiments (linalool and myrcene) resulted in slightly lower κ_{CCN} values than their corresponding 50 ppbV experiments.

3.5 Reconciliation of κ_{HTDMA} and κ_{CCN}

In general, the average κ values derived from the HTDMA and CCNc are similar for each precursor. For 50 ppbV α -pinene, the HTDMA values are very slightly lower than those derived from the CCNc, the mean κ_{HTDMA} 0.12 compared to 0.13 for the CCNc. The κ_{HTDMA} values for 50 ppbV β -caryophyllene are slightly higher than those derived from the CCNc. The mean κ_{HTDMA} values are 0.014 compared to 0.009 for the CCNc. The κ_{HTDMA} values for 50 ppbV limonene SOA cover the range of κ values derived from the CCNc although it should be noted that only 3 data points were collected from the CCNc. The κ_{HTDMA} values for 50 ppbV linalool SOA are lower (mean 0.08) than those derived from the CCNc (mean 0.12). Similarly, the κ_{HTDMA} for 50 ppbV myrcene SOA (mean 0.10) are slightly lower than those derived from the CCNc (mean 0.12), but the

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range of κ values is similar. The mean κ_{HTDMA} values for the 250 ppbV myrcene and linalool experiments are close to those observed by the CCNc.

In summary κ_{HTDMA} values derived 50 ppbV α -pinene, linalool, and myrcene SOA appear slightly lower than those derived from the CCNc, this difference is largest in the first hours of the experiments, as the κ_{HTDMA} values increase towards the end of the experiments the difference between the CCNc and HTDMA is smaller. The κ values for 250 ppbV linalool and myrcene SOA are similar. The 50 ppbV β -caryophyllene κ_{HTDMA} values are slightly higher than κ_{CCN} .

It should be noted that, in both HTDMA and CCNc instruments, the RH of the SOA sample is actively controlled but not the saturation ratio of any other semi-volatile component. A smog chamber is typically an extensive continuous source of vapours of ranging volatility. It is assumed that the particles have equilibrated with the RH in the sub-saturated measurement in the HTDMA (Swietlicki et al., 2008) and supersaturated measurement in the CCNc (Roberts and Nenes, 2005). Since there is no active control of other semi-volatile components (i.e. VOCs), these molecules are free to condense or evaporate towards their equilibrium concentrations in the measurement timescales of the instruments (Topping and McFiggans, 2012). Therefore, hygroscopic growth and cloud activation of SOA particles are likely to be substantially impacted by the tendency of these semi-volatile molecules towards co-equilibration during the drying prior to initial size selection and humidification in determining their sub-saturated growth or cloud activation potential. Given the lack of active control of semi-volatile compound saturation ratios and the very different values likely to be experienced in each instrument, it may be considered surprising that the relatively large scatter in the data shown in Fig. 7 is not larger.

3.6 Relating hygroscopicity to composition

Increases in particle hygroscopicity during chamber experiments of this type (i.e. single VOC oxidation), have been shown to relate to the changing composition of the sampled particles with time (Massoli et al., 2010; Duplissy et al., 2011; Frosch et al.,

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2011). Specifically, the relative oxidation of the molecules that make up organic particles has been observed to correlate with their hygroscopicity within individual systems (α -pinene, isoprene, trimethylbenzene and *m*-xylene) reported in those studies. The extent of oxidation of organic aerosol particles is typically determined by measuring O:C ratios (Aiken et al., 2008; Chhabra et al., 2011) or averaged carbon oxidation state (Kroll et al., 2011) when a high-resolution aerosol mass spectrometer is used. Alternatively, as discussed in Sect. 3.2, f_{44} is used as a measure of oxidation of unit resolution mass spectral data. Increasing f_{44} has been used to infer the presence of more oxygenated molecules and has therefore been expected to lead to increased particle hygroscopicity.

Figure 8 shows the relationship between the hygroscopicity (κ_{HTDMA}) of particles of a single size and f_{44} measured simultaneously by the ToF-AMS for each precursor at both high and low initial mixing ratios. The linear fit parameters and coefficient of determination (r^2) for each selected particle size are listed in Table 5. As has previously been observed in various SOA systems (Massoli et al., 2010; Duplissy et al., 2011; Frosch et al., 2011), myrcene, linalool and, to a lower extent α -pinene, derived particle hygroscopicity correlate positively with f_{44} . In the cases of myrcene and linalool, the correlation sometimes weakens for the largest sizes towards the end of the experiments, when the growth factor stops increasing but f_{44} continues to rise. There is not however strong correlation between the particle hygroscopicity in the β -caryophyllene experiments and f_{44} . In the β -caryophyllene experiment the change in particle hygroscopicity is smaller than the precision of the HTDMA measurement, while the change in f_{44} is 0.3 % (over the HTDMA measurement period) so a high correlation is not expected. In the limonene experiments the growth factor at each dry size tended to increase with time. In the 250 ppbV limonene experiment κ_{HTDMA} and f_{44} correlate once f_{44} starts to increase, however overall correlation between the $\text{GF}_{D_0 = 350 \text{ nm}}$ is low, due to the fact that f_{44} drops continuously from the start of the experiment and only start to increase after ~ 2.5 h. In the 50 ppbV experiment there is a relatively large amount of scatter in consecutive HTDMA measurements which may account for the low correlation.

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Figure 9 shows mean f_{44} (± 3 s.d.) and the particle hygroscopicity derived from HTDMA (filled symbols) and CCNc (open symbols) of the largest selected particle size during the last hour of each experiment (generally within hours 4 to 6 depending upon the experiment duration) for each precursor at high and low mixing ratio. The largest selected particle size is chosen such that f_{44} (mass fraction) is the closest representation of the particles sampled by the HTDMA and CCNc (it should be remembered that the average hygroscopicity values derived from both the HTDMA and CCNc are weighted according to the number of particles sampled with a given value, but the f_{44} averages are from AMS mass-based averages). Figure 9 aims to illustrate the relationship between the absolute fraction of m/z 44 and particle hygroscopicity independent of precursor. It is apparent that α -pinene, limonene, linalool and myrcene are able to generate particle with similar hygroscopicity ($\kappa \sim 0.1$) despite f_{44} ranging from ~ 4 to 11 %. The β -caryophyllene derived particles exhibit an order of magnitude lower κ than the other systems after 4 to 6 h of photooxidation as well as the lowest f_{44} (~ 2.5 %).

A similar investigation of SOA hygroscopicity, represented by κ , and the averaged carbon oxidation state ($\overline{\text{OS}}_{\text{C}}$) determined from the off-line analysis is shown in Fig. 10. The $\overline{\text{OS}}_{\text{C}}$ values of the SOA ranged from -0.96 to -0.88 for β -caryophyllene, -0.67 to -0.47 for limonene and -0.69 for the single α -pinene experiment. Similar to the f_{44} analysis, β -caryophyllene SOA showed the lowest $\overline{\text{OS}}_{\text{C}}$ values, indicating it was the least oxidized aerosol. This is perhaps expected since the parent VOC is the least volatile and early generation oxidation products can partition to the condensed phase. Both the monoterpenes have higher $\overline{\text{OS}}_{\text{C}}$ values, with limonene higher than α -pinene. This may be a result of further oxidation of the limonene double bond, giving more oxidized products. A mean kappa value for the largest sized particles over the final hour of the experiment was used since the filter samples represent an instantaneous snapshot at the end of the experiment. From the samples available from offline analyses of limited subset of the precursors, it might be tempting to try to look for a relationship between $\overline{\text{OS}}_{\text{C}}$ and hygroscopicity. The fallibility of this approach can be seen from the

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online analysis when including the linalool and myrcene SOA data points. A direct comparison of data in Fig. 8 and 9 could only be performed using the three overlapping SOA systems of β -caryophyllene, α -pinene and limonene. Both off and online analyses methods determined lower level of oxidation for the β -caryophyllene SOA compared to that of limonene and α -pinene. It is noteworthy, however, that the \overline{OS}_C value of α -pinene SOA is lower than that of limonene, whilst the f_{44} is higher, highlighting an inconsistency in determining the degree of oxidation of particles between the employed online and offline analyses methods. Including online data for linalool and myrcene SOA systems in this analysis led to the conclusion that a relationship between f_{44} (\overline{OS}_C) and hygroscopicity could not be established and generalised for the combined SOA systems in this study.

As stated above, in the water uptake measurements, the RH of the SOA sample is controlled but the lack of control of saturation ratio of any other semi-volatile component means they are free to evaporate during drying to initial size selection and to condense on humidification in determining their sub-saturated growth or cloud activation potential. Sample treatment during online and offline chemical composition measurements are different. Particles sampled into the AMS were not actively dried. However, they were subjected to a supersonic expansion into the vacuum system, which could lead to freeze drying of water and semi-volatiles. The filter samples were collected over a relatively short period of time using a high flow rate of $3\text{ m}^3\text{ min}^{-1}$ (i.e. in approximately 6 min), and were stored at -18°C almost immediately after collection. Such a protocol should minimise sampling artefacts of condensation and evaporation. Discrepancies between online measurements of composition and hygroscopic growth when particles contain substantial amounts of semi-volatile material show that correction for this fraction are required (Gysel et al., 2007) and discrepancies between measurements of sub-saturated hygroscopic growth and droplet activation in the laboratory (Good et al., 2010b) and atmosphere (Irwin et al., 2010) are likely attributable to this effect.

4 Conclusions

Five structurally different biogenic VOCs were identified representing species emitted into the atmosphere with a range of chemical reactivity and complexity in their subsequent degradation. The studied compounds were α -pinene, β -caryophyllene, limonene, linalool and myrcene. The chemical half-life for each of the compounds derived from their decay profiles appeared to be consistent with their published reactivity data towards the main oxidants of ozone and hydroxyl radical. The shortest lifetime (i.e. highest reactivity) was observed for β -caryophyllene, while the longest lifetime was found to be that of α -pinene. The SOA particle formation profiles were found to be consistent with the derived VOC chemical lifetimes (β -caryophyllene SOA peak number was the first to be reached, while α -pinene SOA was the last). Investigations illustrated that SOA formation potential was higher when the same precursor is used at higher initial mixing ratio compared to a lower initial mixing ratio case.

The fractional contributions of mass fragments 43 and 44 to the total organic signal (f_{43} and f_{44}) were used to characterise the extent of oxidation of the SOA as measured online by an aerosol mass spectrometer. The f_{43} and f_{44} for the studied systems under lower and higher initial mixing ratios varied considerably, both in magnitude and temporal evolution. These results illustrated that SOA generated from a single biogenic precursor cannot and should not be used as a proxy for biogenic SOA in general. In addition, the values of f_{44} which are widely used in the literature to indicate the level of oxygenation of organic aerosol in field and laboratory studies are very much dependent on the type of precursor, the extent of photochemical ageing as well as on the initial experimental conditions.

Similarly, the generated SOA particles exhibited a range of hygroscopic properties, represented by κ_{HTDMA} , depending on the precursor, its initial mixing ratio and photochemical ageing. For example, the β -caryophyllene SOA particles were the least hygroscopic ($\kappa_{\text{HTDMA}} \sim 0.01$), while α -pinene, limonene, linalool and myrcene produced particles with higher hygroscopicity ($\kappa \sim 0.03$ to 0.14). Considering each selected dry size

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separately, there was a positive correlation between hygroscopicity (κ_{HTDMA}) and time (i.e. degree of oxidation). This correlation was generally strong ($r^2 > 0.9$) for limonene, linalool and myrcene SOA, whilst for the β -caryophyllene the change in κ_{HTDMA} was small and could not be resolved within the measurement uncertainty (although correlation was observed). The increases of κ_{HTDMA} with time indicated that SOA formed from molecules which partition from the gas phase later in the experiments were more hygroscopic. The CCN activity of the formed SOA particles, represented by κ_{CCN} , showed no temporal trends for the lower mixing ratio cases. The two higher mixing ratio cases of linalool and myrcene, on the other hand, showed reduced CCN activity compared to their lower mixing ratio measurements. However, these conclusions were based on a significantly limited CCNc dataset compared to other data sets. The average κ values derived from the HTDMA and CCNc were generally found to cover the same range for each precursor both for the 50 and 250 ppbV initial mixing ratios. The κ_{HTDMA} values derived for 50 ppbV α -pinene, linalool, and myrcene SOA appeared to be slightly lower than those derived from the CCNc, especially during the first hours of the experiments. Given the lack of active control of semi-volatile compound saturation ratios and the very different values likely to be experienced in each of the two instruments, the difference between the two κ values is relatively small.

Consistent with previous studies, a positive correlation was found between the hygroscopicity of particles (κ_{HTDMA}) of a single size and f_{44} for myrcene, linalool and α -pinene SOA (albeit the correlation for the latter was weaker). On the other hand, no such correlation was found for limonene SOA particles. Investigation of the generality of the relationship between f_{44} and κ_{HTDMA} revealed that α -pinene, limonene, linalool and myrcene were all able to generate particles with similar hygroscopicity ($\kappa_{\text{HTDMA}} \sim 0.1$) despite f_{44} exhibiting a relatively wide range of values (~ 4 to 11 %). Similarly, κ_{CCN} was found to be independent of f_{44} . The same findings were also true when sub- and super-saturated water uptake properties of SOA were compared to the averaged carbon oxidation state ($\overline{\text{OS}}_{\text{C}}$) determined off-line. These results suggest that κ values derived from sub- and super-saturated measurements of SOA particles are independent

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of their degree of oxidation determined using both online and offline methods. These findings do not necessarily suggest that water uptake and chemical composition are not related in reality. Instead, they suggest that either f_{44} and \overline{OS}_C do not represent the determinant composition-related factors controlling water uptake of SOA particles, or they may emphasise the possible impact of the lack of control of saturation ratio of semi-volatile compounds in the current state-of-the-art techniques employed to determine the chemical composition and water uptake properties of aerosol particles.

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Table 1. Overview of initial experimental conditions and SOA formation characteristics.

| Exp # | Compound | [VOC] ₀ (ppb) | [NO _x] ₀ (ppb) ² | Temp (°C) | Relative Humidity (RH%) | Chemical half-life (min) ¹ | Peak Number Conc. (p/cc) | Peak Number Time (min) | Mass Conc. (μg m ⁻³) ⁴ | Δ[VOC] (ppbV) | Δ[VOC] (μg m ⁻³) | Y _{SOA} (%) |
|-------|------------------------------|--------------------------|--|------------|-------------------------|---------------------------------------|--------------------------|------------------------|---|---------------|------------------------------|----------------------|
| 1 | Myrcene | 44.2 | 31.3 ± 0.8 | 25.5 ± 0.3 | 54.2 ± 0.9 | 49.0 | 5.47e + 4 | 29 | 34.0 | 44.2 | 246.1 | 13.8 |
| 2 | Myrcene | 254.8 | 137.2 ± 0.2 | 25.5 ± 0.1 | 52.3 ± 0.4 | 27.0 | 1.62e + 5 | 10 | 218.4 | 254.8 | 1418.8 | 15.4 |
| 3 | Linalool | 77.0 | 29.1 ± 0.7 | 25.4 ± 0.2 | 48.0 ± 0.4 | 46.7 | 6.45e + 4 | 46 | 17.4 | 77.0 | 485.5 | 3.6 |
| 4 | Linalool | 176.6 | 121.9 ± 1.2 | 24.7 ± 0.7 | 52.5 ± 1.5 | 42.0 | 1.80e + 5 | 12 | 73.3 | 176.6 | 1113.5 | 6.6 |
| 5 | α-pinene ⁸ | 45.0 | 34.2 ± 0.3 | 25.6 ± 0.5 | 71.4 ± 1.2 | 104.0 | 3.92e + 4 | 243 | 37.0 | 44.5 | 247.8 | 14.9 |
| 6 | α-pinene | 50 ⁶ | 25 ⁶ | 26.4 ± 0.3 | 51.4 ± 1.4 | - ⁷ | 3.30e + 4 ³ | 100 ³ | 51.4 ³ | - | - | - |
| 7 | β-caryophyllene ⁸ | 250 ⁶ | 128.4 ± 1.2 | 26.1 ± 0.3 | 44.4 ± 0.7 | - | 3.56e + 4 | 4 | 284.8 | - | - | - |
| 8 | β-caryophyllene ⁸ | 50 ⁶ | 28.2 ± 0.5 | 25.4 ± 0.8 | 47.9 ± 0.9 | - | 2.48e + 4 | 19 | - | - | - | - |
| 9 | β-caryophyllene | 48.8 | 34.4 ± 0.1 | 25.7 ± 0.8 | 69.7 ± 1.1 | 16.8 | 3.47e + 4 | 10 | 188.4 | 48.1 | 401.8 | 46.9 |
| 10 | β-caryophyllene ⁸ | 50.7 | 27.8 ± 0.2 | 25.9 ± 0.4 | 71.4 ± 0.9 | 32.8 | 2.76e + 4 | 11 | 220.6 | 49.9 | 416.8 | 52.9 |
| 11 | Limonene ⁸ | 250 ⁶ | 142.0 ± 0.3 | 26.3 ± 0.4 | 49.3 ± 1.0 | - | 3.26e + 4 | 17 | 861.6 | - | - | - |
| 12 | Limonene ⁸ | 50 ⁶ | 54.1 ± 0.5 | 25.5 ± 0.5 | 53.7 ± 1.3 | - | 1.51e + 4 | 48 | 79.2 ⁵ | - | - | - |
| 13 | Limonene ⁸ | 50 ⁶ | 32.1 ± 0.1 | 25.3 ± 0.2 | 53.5 ± 0.6 | - | 1.67e + 4 | 47 | 65.6 | - | - | - |
| 14 | Limonene ⁸ | 52.1 | 36.7 ± 0.3 | 25.5 ± 0.7 | 73.3 ± 2.3 | 55.1 | - | - | - | - | - | - |

¹The time taken to reach half initial mixing ratio = median life time.

²NO_x: average of 1–3 min of stable data around light on time.

³Measured by DMPS using a density of 1.3 g cm⁻¹.

⁴Wall-loss corrected mass concentration.

⁵Experiment duration 2 h.

⁶Nominal mixing ratio values estimated based on the amount introduced into the chamber without accounting for losses.

⁷- indicates that data were not available.

⁸Off-line filter analysis reported.

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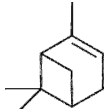
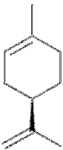
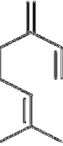
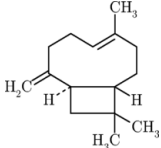
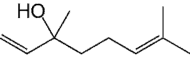
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Table 2. List of all five precursors, structures and reactivity towards OH and O₃ based on reaction lifetimes from Atkinson & Arey (2003).

| Category | Representative Compound | Structure | OH reaction lifetime | O ₃ reaction lifetime |
|------------------------------|-------------------------|---|----------------------|----------------------------------|
| Bicyclic monoterpene | <i>α</i> -pinene |  | 2.6 h | 4.6 h |
| Monocyclic diene monoterpene | limonene |  | 49 min | 2 h |
| Acyclic triene monoterpene | myrcene |  | 39 min | 50 min |
| Reactive sesquiterpene | <i>β</i> -caryophyllene |  | 42 min | 2 min |
| Reactive C10 oxygenates | linalool |  | 52 min | 55 min |

[OH] = 2.0×10^6 molecule cm⁻³, 12 h daytime average; [O₃] = 7.0×10^{11} molecule cm⁻³, 24 h average (ca. 30 ppb).

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Table 3. Mean growth factor and kappa values derived from the HTDMA measurements at 90% RH for the 50 ppbV and 250 ppbV initial precursor mixing ratios.

| Precursor | 50 ppbV | | | 250 ppbV | | |
|------------------------|---------------------|-----------|-------------------------|---------------------|-----------|-------------------------|
| | GF _{90%RH} | Range | κ_{HTDMA} | GF _{90%RH} | Range | κ_{HTDMA} |
| α -pinene | 1.27 | 1.25–1.29 | 0.121 | 1.08 | 1.07–1.13 | 0.042 |
| β -caryophyllene | 1.02 | 1.01–1.04 | 0.009 | 1.03 | 1.01–1.05 | 0.011 |
| Limonene | 1.17 | 1.11–1.23 | 0.077 | 1.15 | 1.13–1.19 | 0.069 |
| Linalool | 1.18 | 1.13–1.22 | 0.082 | 1.17 | 1.15–1.24 | 0.090 |
| Myrcene | 1.21 | 1.12–1.29 | 0.101 | 1.19 | 1.16–1.27 | 0.098 |

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Table 4. Mean and range of kappa values derived from the CCNc measurements for the 50 ppbV and 250 ppbV initial precursor mixing ratios.

| Precursor | κ_{CCN} | | 10th–90th percentile range | |
|------------------------|-----------------------|----------|----------------------------|-----------|
| | 50 ppbV | 250 ppbV | 50 ppbV | 250 ppbV |
| α -pinene | 0.13 | – | 0.10–0.16 | – |
| β -caryophyllene | 0.009 | – | 0.009–0.014 | – |
| Limonene* | 0.06 | – | – | – |
| Linalool | 0.12 | 0.10 | 0.10–0.15 | 0.06–0.14 |
| Myrcene | 0.12 | 0.10 | 0.06–0.17 | 0.06–0.13 |

* Derived from limited number of measurements.

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Table 5. Linear fit parameters and coefficient of determination (R^2) of the relationship between the hygroscopicity (represented by κ_{HTDMA}) of particles of a single size and the degree of particle oxidation (represented by f_{44}).

| Precursor | Initial mixing ratio (ppbV) | D_0 (nm) | M | C | R^2 |
|------------------------|-----------------------------|-------------------|--------------------|--------------------|-------|
| α -Pinene | 50 | 200 | -0.142 ± 0.099 | 0.024 ± 0.009 | 0.40 |
| | | 300 | -0.003 ± 0.001 | 0.004 ± 0.000 | 0.87 |
| β -Caryophyllene | 50 | 200 | 0.068 ± 0.052 | -0.001 ± 0.011 | 0.60 |
| | | 300 | -0.019 ± 0.003 | -0.067 ± 0.009 | 0.67 |
| Limonene | 250 | 250 | -0.019 ± 0.003 | -0.067 ± 0.009 | 0.67 |
| | | 300 | -0.016 ± 0.009 | 0.055 ± 0.025 | 0.10 |
| | 50 | 200 | 0.044 ± 0.034 | -0.240 ± 0.253 | 0.10 |
| | | 200 | 0.042 ± 0.037 | -0.214 ± 0.259 | 0.16 |
| Linalool | 250 | 350 | 0.168 ± 0.084 | -0.018 ± 0.016 | 0.06 |
| | | 500 | 0.055 ± 0.028 | -0.229 ± 0.149 | 0.36 |
| | 50 | 100 | 0.024 ± 0.002 | -0.032 ± 0.008 | 0.93 |
| | | 150 | 0.017 ± 0.001 | -0.008 ± 0.005 | 0.94 |
| Myrcene | 250 | 200 | -0.006 ± 0.007 | 0.119 ± 0.047 | 0.12 |
| | | 150 | 0.029 ± 0.002 | -0.019 ± 0.007 | 0.94 |
| | | 200 | 0.027 ± 0.0014 | -0.028 ± 0.006 | 0.95 |
| | 50 | 100 | 0.036 ± 0.003 | -0.188 ± 0.021 | 0.96 |
| | | 150 | 0.034 ± 0.001 | -0.200 ± 0.012 | 0.97 |
| | | 250 | 0.006 ± 0.003 | 0.025 ± 0.032 | 0.22 |
| 250 | 150 | 0.041 ± 0.011 | -0.242 ± 0.092 | 0.57 | |
| | 250 | 0.025 ± 0.001 | -0.123 ± 0.008 | 0.97 | |
| | 350 | 0.022 ± 0.003 | -0.124 ± 0.030 | 0.74 | |

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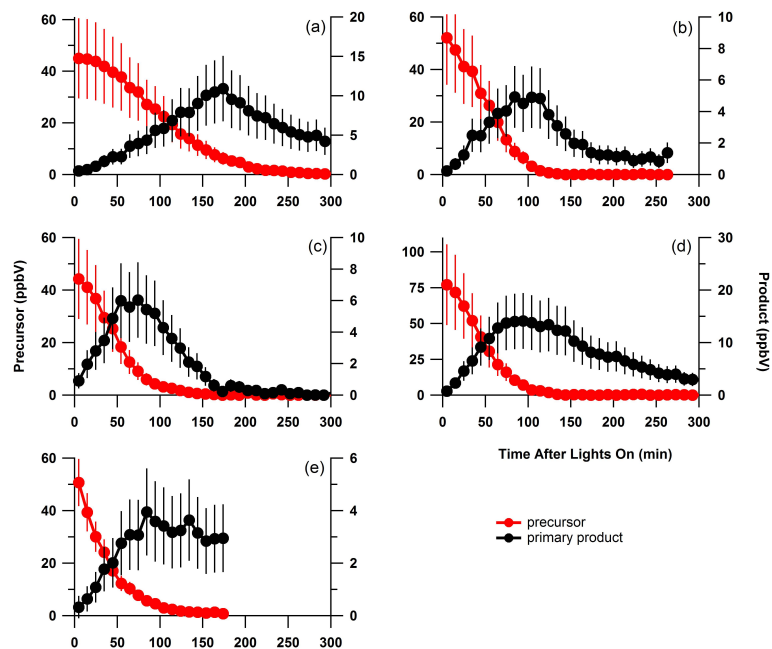


Fig. 1. Temporal profiles of each of the precursors investigated and their primary oxidation products: **(a)** α -pinene and pinonaldehyde; **(b)** limonene and limononaldehyde; **(c)** myrcene and 4-vinyl-4-pentalenal; **(d)** linalool and 4-hydroxy-4-methyl-5-hexen-1-al; **(e)** β -caryophyllene and β -caryophyllon aldehyde.

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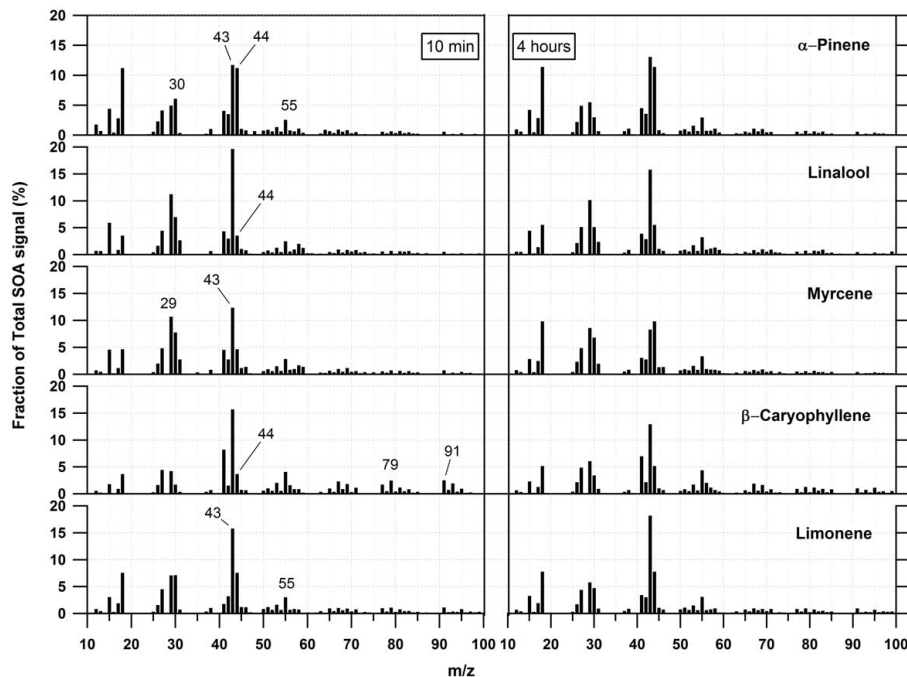


Fig. 2. Mass spectral patterns of SOA from the photooxidation of biogenic VOCs (data from the lower initial mixing ratio experiments reported in Table 1). The left hand panels include spectra measured during the first 10 min of SOA formation, while those measured after 4 h of photooxidation are displayed in the right hand panels. Each spectra has been normalised to the total SOA signal.

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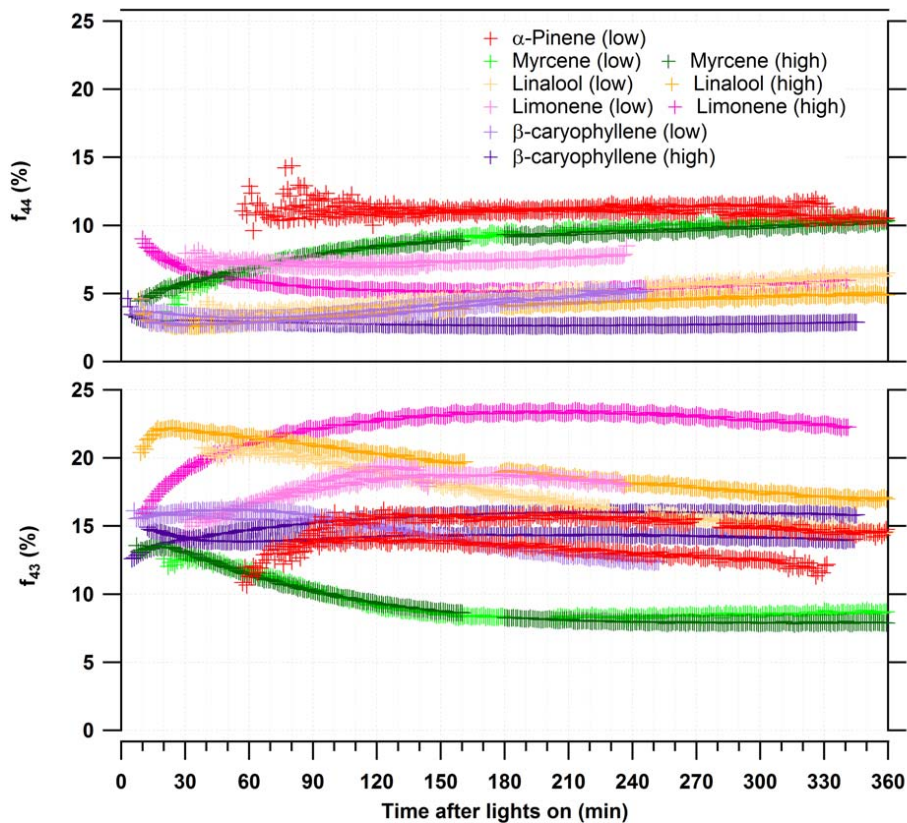


Fig. 3. Evolution of the f_{44} and f_{43} with photooxidation for low and high initial VOC mixing ratio experiments.

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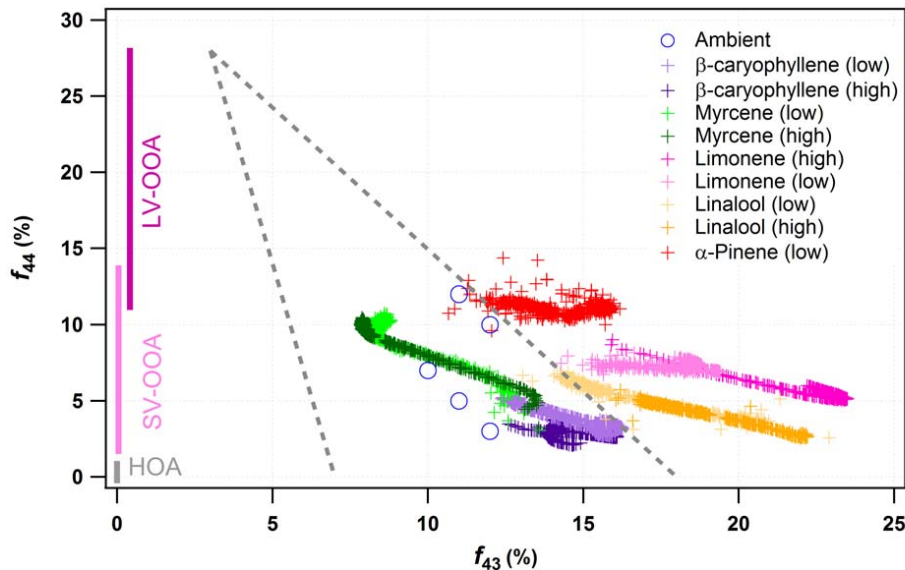


Fig. 4. A 2-dimensional representation of f_{44} and f_{43} for all SOA systems in this study. The dashed line represents the boundaries of this relationship based on previously published ambient datasets by Ng et al. (2010). The open circles are measurements conducted at biogenically-influenced locations in North America and Europe as compiled by Ng et al. (2010).

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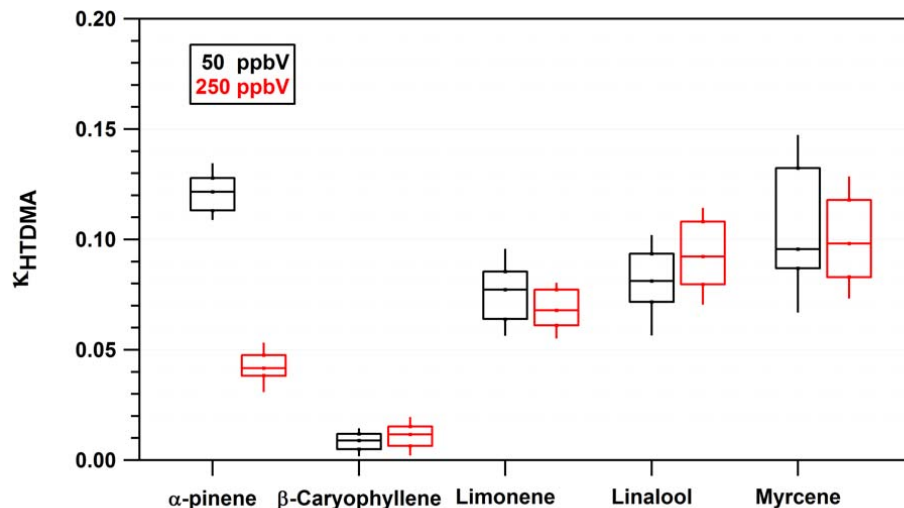


Fig. 5. A box and whisker plot illustrating the observed range of hygroscopic properties, represented by κ_{HTDMA} , for all selected dry particle diameters for particles generated from each initial mixing ratio of the studied precursors.

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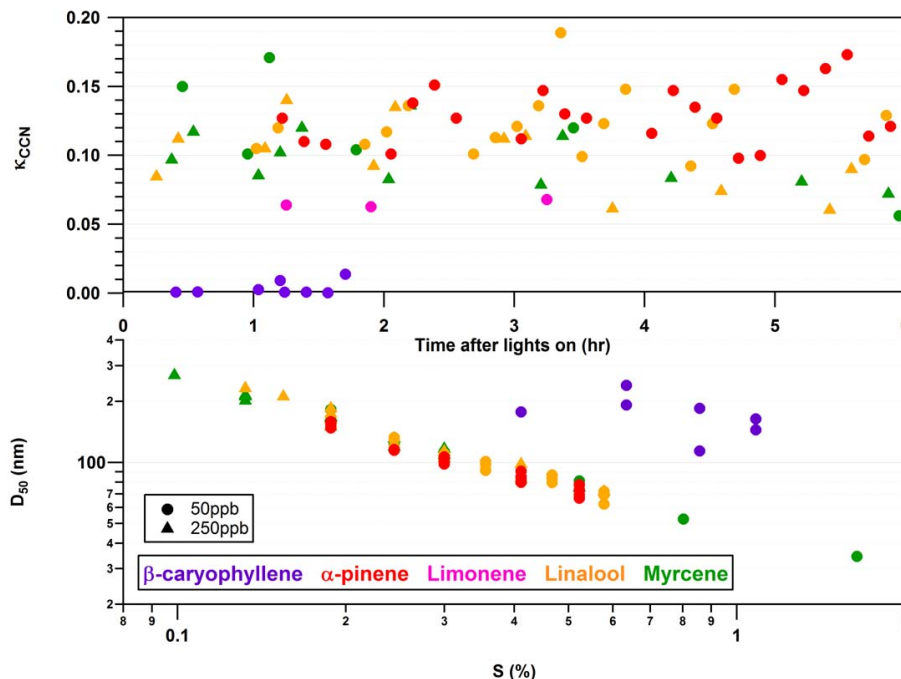


Fig. 6. Temporal profiles of CCN activity (κ_{CCN}) of the SOA particles from the five studied biogenic precursors (top panel). The relationship between the size at which 50% of the particles activated (D_{50}) and super-saturation for all of the SOA systems is shown in the bottom panel.

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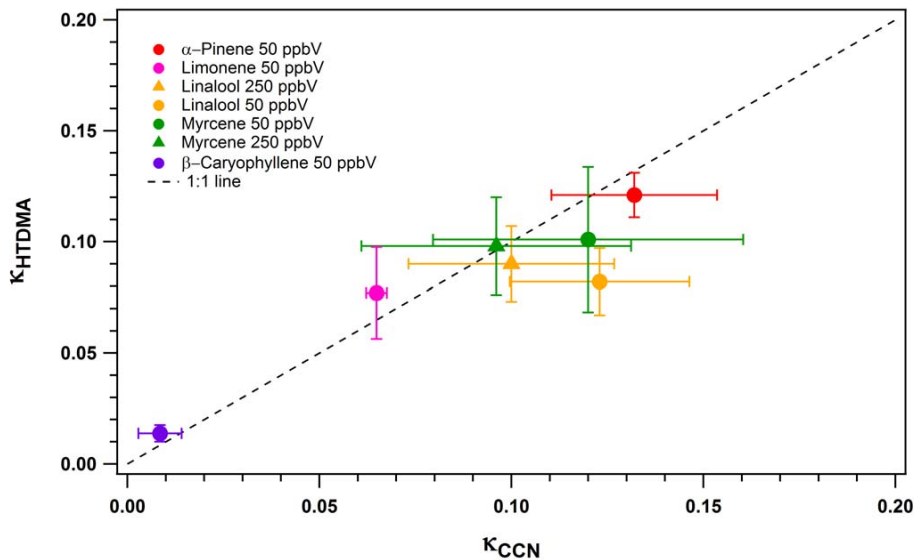


Fig. 7. Reconciliation of mean kappa values from HTDMA and CCN measurements. The vertical and horizontal variability bars represent one standard deviation for each measurement.

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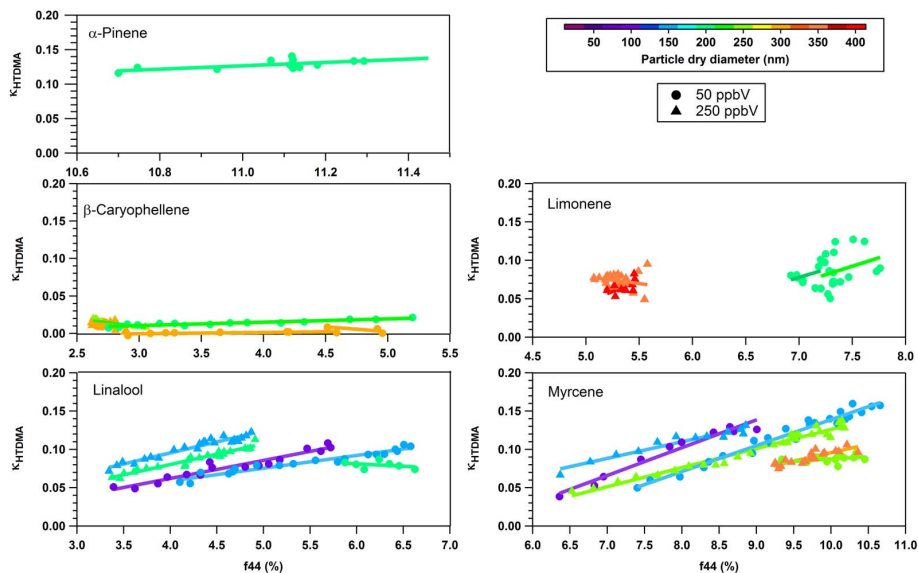


Fig. 8. The relationship between hygroscopicity of particles of a single size (represented by κ_{HTDMA}) and f_{44} measured simultaneously by the ToF-AMS for each precursor at both high and low initial mixing ratios. The linear fit parameters and coefficient of determination (r^2) for each selected particle size are listed in Table 5.

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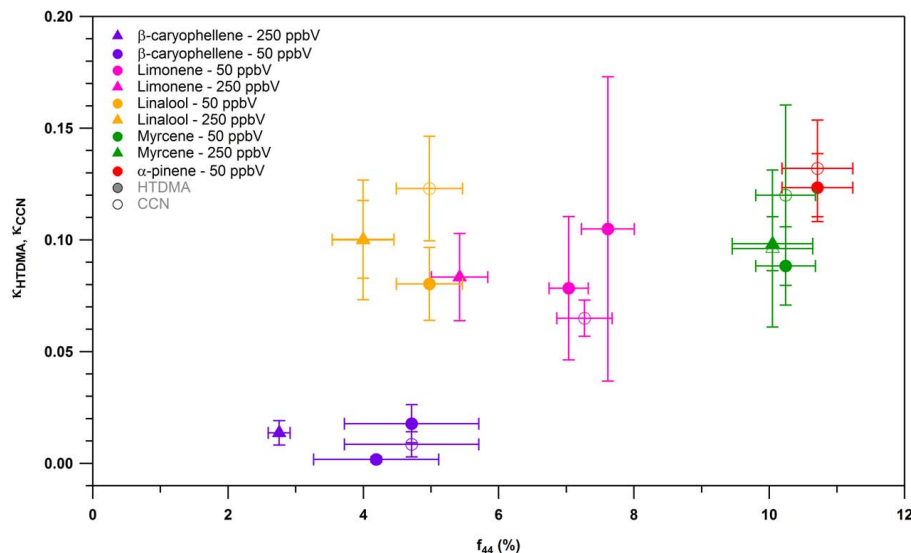


Fig. 9. The relationship between sub- and super-saturated water uptake of SOA particles and their level of oxidation. The κ_{HTDMA} (open symbols) and κ_{CCN} (filled symbols) are mean values measured for the largest size during the final hour of each experiment. The level of oxidation is inferred from the mean values of f_{44} during the corresponding period. The variability bars are (± 3 s.d.) of the measurements.

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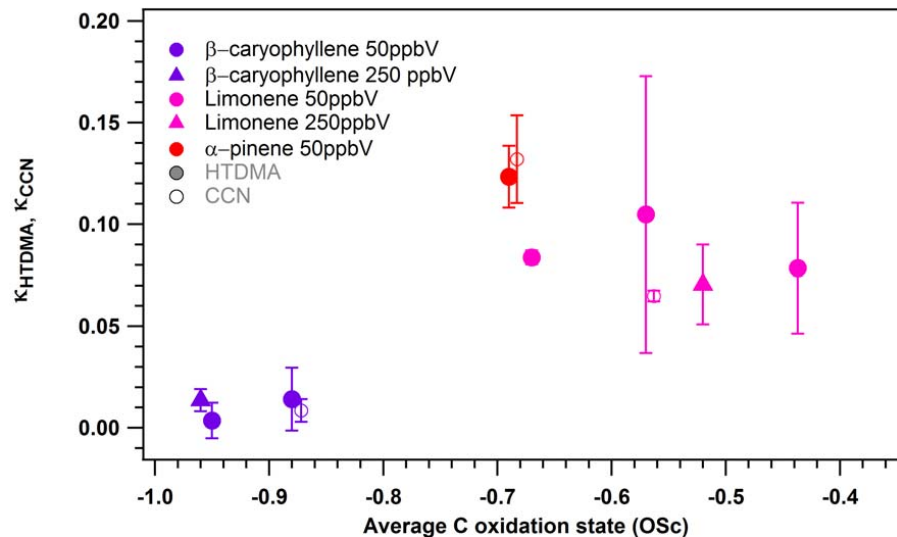


Fig. 10. κ_{HTDMA} values average over the final hour for the largest measured size and κ_{CCN} of the appropriate experiments plotted against average C oxidation state calculated from SOA filter samples collected at the end of same experiments. The filled and open symbols are for HTDMA and CCN data, respectively. The κ_{CCN} data points have been offset very slightly to the right for display purpose.

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