

Measurement report: Effects of NO_x and seed aerosol on highly oxygenated organic molecules (HOM) from cyclohexene ozonolysis

Meri Rätty¹, Otso Peräkylä¹, Matthieu Riva^{1,2}, Lauriane Quéléver¹, Olga Garmash¹, Matti Rissanen^{1,3}, and Mikael Ehn¹

¹Institute for Atmospheric and Earth System Research / Physics, Faculty of Science, University of Helsinki, Finland

²Univ Lyon, Université Claude Bernard Lyon 1, CNRS, IRCELYON, F-69626, Villeurbanne, France

³Aerosol Physics Laboratory, Physics Unit, Tampere University, Tampere, Finland

Correspondence: Meri Rätty (meri.ratty@helsinki.fi) and Mikael Ehn (mikael.ehn@helsinki.fi)

Abstract. Cyclohexene (C₆H₁₀) is commonly used as a proxy for biogenic monoterpenes, when studying their oxidation mechanisms and secondary organic aerosol (SOA) formation. The ozonolysis of cyclohexene has been shown to be effective at producing highly oxygenated organic molecules (HOM), a group of molecules known to be important in the formation of SOA. Here, we provide an in depth look at how ~~on a molecular level, the HOM~~ the formation and fate of the broad range of observed HOM changed with perturbations from NO_x and seed particles.

HOM were produced in a chamber from cyclohexene ozonolysis, and measured with a chemical ionisation mass spectrometer (CIMS) using nitrate (NO₃⁻) as reagent ion. As high-resolution CIMS instruments provide mass spectra with numerous ion signals and a wealth of information that can be hard to manage, we employed a primarily statistical approach for the data analysis. To utilise as many individual HOM signals as possible, each compound was assigned a parameter describing the quality of the observed signal. These parameters were then used as weights or to determine the inclusion of a given signal in further analyses.

Under unperturbed ozonolysis conditions, known HOM peaks were observed in the chamber, including C₆H₈O₉ as the largest HOM signal, and C₁₂H₂₀O₉ as the largest “dimer” product. With the addition of nitric oxide (NO) into the chamber, the spectrum changed considerably, as expected. Dimer product signals decreased overall, but an increase in dimers with nitrate functionalities was seen, as a result of NO₃ radical oxidation. The response of monomer signals to NO addition varied, and while nitrate-containing monomers increased, non-nitrate signals either increased or decreased, depending on the individual molecules.

The addition of seed aerosol increased the condensation sink, which markedly decreased the signals of all low-volatility compounds. Larger molecules were seen to have a higher affinity for condensation, but a more detailed analysis showed that the uptake was controlled mainly by the number of oxygen atoms in each molecule. ~~All non-nitrate compounds with at least 7 oxygen atoms were observed to condense onto the seed aerosol at close to equal rates.~~ Nitrates required higher mass and higher oxygen content to condense at similar rates as the non-nitrate HOM. ~~A comparison to experiments with α-pinene reported earlier, showed quite a similar relationship between elemental composition and volatility, although products from α-pinene ozonolysis appeared to require slightly higher oxygen numbers for the same decrease in volatility. In addition, two models developed for predicting volatilities of volatile organic compound (VOC) oxidation products were tested on the ozonolysis~~

~~products of cyclohexene~~ We also tested two existing elemental composition based parametrisations on their ability to reproduce the condensation observed in our cyclohexene system. Both predicted higher volatilities than observed, most likely due to the number of oxygen atoms enhancing the product uptake more than the models would suggest.

1 Introduction

30 Secondary organic aerosol (SOA) plays an important role in the climate system, and its formation and growth has hence been a major focus of research in the recent years (Shrivastava et al., 2017). ~~Through oxidation, some of the initially volatile organic species can be transformed into low volatility compounds (Jimenez et al., 2009), including~~ Low volatility organic vapours, like highly oxygenated organic molecules (HOM), ~~that can then be readily taken up in aerosol formation (Ehn et al., 2014; Bianchi et al., 2019).~~ are a crucial ingredient in many of these aerosols (Ehn et al., 2014; Bianchi et al., 2019). They are naturally formed in the atmosphere in the oxidation of initially volatile organic species (Jimenez et al., 2009).

35 Many of the hydrocarbons emitted to the atmosphere have natural sources, mainly in vegetation (Laothawornkitkul et al., 2009). The oxidation of some of these biogenic volatile organic compounds (BVOCs) have been linked to high SOA yields, for example monoterpenes ($C_{10}H_{16}$) (Tunved et al., 2006; Ehn et al., 2014; Jokinen et al., 2015). Notably, the monoterpenes that have a carbon ring structure with a double bond (i.e., endocyclic alkenes) can be oxidised by ozone without fragmentation, producing a larger oxygenated molecule (Chuong et al., 2004; Jokinen et al., 2015), often resulting in a higher SOA-forming potential.

The lower the volatility of an oxidation product is, the more it has a tendency to condense onto existing aerosol particles (Kroll and Seinfeld, 2008). In addition to typically having a relatively large molecular mass, low volatility products generally have high oxygen content. More specifically, low volatility has typically been linked to the oxygen to carbon ratio O:C (Aiken et al., 2008). However, there have also been implications that the relationship between low volatility and O:C ratio might not always be as straightforward as was previously thought (Kurtén et al., 2016). Volatility depends on molecular functionality, and products with more hydrogen-bonding functional groups have been linked to lower volatilities (Donahue et al., 2013). In addition to simple condensation, reactive uptake may also be responsible for some of the growth of SOA particles (Zhang et al., 2015).

50 The most emitted monoterpene in the atmosphere is α -pinene, contributing to SOA formation and growth especially in the Boreal regions (Spanke et al., 2001; Ehn et al., 2014; Jokinen et al., 2015). As most monoterpenes, α -pinene has a relatively complex structure. Therefore, the simple symmetrical cyclohexene (C_6H_{10}) molecule has regularly been used as its surrogate in SOA studies (e.g., Chuong et al., 2004; Rissanen et al., 2014). As the carbon ring is the common site for oxidation in BVOCs with endocyclic double bonds (Rissanen et al., 2014), the simple cyclohexene can also provide insight into the oxidation pathways of larger molecules.

In this work, we studied cyclohexene ozonolysis in an environmental chamber, with the main focus of understanding HOM the formation and fates ~~under different perturbations. We added NO of HOM products under perturbations. NO was added~~ to the chamber in order to compare the change in HOM formation pathways. ~~In order to test the volatility of the HOM products,~~

we also added seed aerosol to the chamber. We compare our findings with similar studies performed on α -pinene, to test how well cyclohexene can work as a surrogate for monoterpenes with an endocyclic double bond, and how applicable earlier volatility parametrisations are on investigate the resulting changes in the formation of HOM, while introducing seed aerosol allowed us to examine the connection between volatility and HOM elemental composition. We also assessed the applicability of two existing volatility parametrisations on our cyclohexene system. This allows investigating the link between elemental composition and volatility of cyclohexene HOM in the broader context of other volatile organic compounds (VOCs), such as the atmospherically relevant BVOCs.

2 Methods

2.1 Chamber conditions and instrumentation

The measurements were conducted during a 5-day-period in the COALA chamber, which is described in detail elsewhere (Riva et al., 2019a; Peräkylä et al., 2020) by Riva et al. (2019a) and Peräkylä et al. (2020), the latter giving also a comprehensive description of chamber dynamics relevant in our experiments as well. The chamber is 2 m^3 in volume and has fluorinated ethylene propylene (FEP) as its wall material. The cyclohexene ozonolysis was performed under dark conditions. Temperature in the chamber was $27 \pm 2\text{ }^\circ\text{C}$, and relative humidity $29 \pm 2\%$. Mass flow controllers (MKS, G-Series, 0.05-50 Lpm, Andover, MA, USA) were used to control the injections of all reactants. Ozone was generated from purified air with an ozone generator (Dasibi 1008-PC), and cyclohexene was produced by flowing air through a glass bubbler with liquid cyclohexene. Both were continuously injected into the chamber, together with purified air (AADCO model 737-14, Ohio, USA). The resulting steady-state concentrations of ozone was ozone concentration without cyclohexene in the chamber, was on average 25 ppb (23-27 ppb), and when cyclohexene was added the steady-state ozone concentration was approximately 18 ppb, while a rough estimate for cyclohexene (16-19 ppb). The cyclohexene concentration was about 100 ppb. As the concentration of cyclohexene was not measured it was and therefore had to be estimated from the difference drop in ozone concentration with and without cyclohexene (with rate constant by Stewart et al., 2013) at its introduction. This produced a rough estimation of 70 ppb of cyclohexene. The average chamber residence time was approximately 50 minutes. Ozone and NO_x levels were monitored with gas analysers (UV photometric ozone monitor, Model49p, Thermo Environmental Instruments; NO - NO_2 - NO_x analyser, Model 42i, Thermo Fisher Scientific). Concentrations of all substances in the different experiments are presented in Table A1.

Compound Concentration 18 ppb * 100 ppb NO 0 ppb[†] 1 ppb[‡] 3 ppb[‡] 10 ppb[‡] 80 nm-ABS particles $0\text{ }\mu\text{g m}^{-3}$ $9\text{ }\mu\text{g m}^{-3}$ The reactant, oxidant and particle concentrations during the conducted experiments (* steady-state concentration when no NO input, † input concentrations of NO which mostly converted quickly into)

NO_x experiments were conducted by injecting NO into the chamber at concentrations shown in table A1. As NO quickly reacts with ozone to form NO_2 , the resulting NO/NO_x ratio in the chamber was fairly low, approximately $3 \pm 1\%$ during highest injection rates. For most of the time, the concentration of NO in the chamber was not high enough to exceed a few percent at most. This meant that the NO concentration remained below the detection limit of the used analyser during

our experiments. Therefore, only the total NO_x was ~~used~~ considered in the analyses. The total NO_x concentrations in the chamber during different experiments were approximately 1, 3 or 9 ppb (Table A1)

The condensation of vapours was studied with a $9 \mu\text{g m}^{-3}$ loading of ammonium bisulfate (ABS) ($(\text{NH}_4)\text{HSO}_4$) particles
95 that were produced by nebulising aqueous ammonium sulfate solution with sulfuric acid. They were dried and size selected
(80 nm) with a differential mobility analyser (DMA) before injection. The condensation sink was calculated for sulfuric acid,
from aerosol size distribution data (Maso et al., 2005) measured with a Differential Mobility Particle Sizer (DMPS, Aalto et al.,
2001). ~~Experiments with ammonium sulfate were also carried out, but in these the loading was not sufficiently high to create
a significant change that would allow for conclusions to be drawn. Therefore, these were left out of the analysis.~~ Due to the
100 acidity of the ammonium bisulfate aerosol, ~~it should be noted that~~ there is a possibility that a fraction of the overall sink might
be a result of reactive uptake (Zhang et al., 2015). However, in their similar experiments with α -pinene, Peräkylä et al. (2020)
did not observe the uptake of HOM to differ significantly between AS-ammonium sulfate (AS) and ABS seeds.

Oxygenated gas phase products were monitored by a Chemical Ionisation Atmospheric Pressure interface Time-of-flight
mass spectrometer (CI-API-TOF, TOFWERK AG, Aerodyne, Junninen et al., 2010; Jokinen et al., 2012). It consists of three
105 components, a chemical ionisation inlet (CI), an atmospheric pressure interface (APi) and the Time-Of-Flight (ToF) mass
spectrometer, which in our setup was an HTOF that has a mass resolving power of ~ 5000 Th/Th. Nitrate ions produced by
subjecting an air stream containing nitric acid (HNO_3) to soft x-rays, are used to charge the sample molecules drawn from the
chamber into the CI inlet. After charging, ions are directed into the APi. Quadruple ion guides and an ion lens stack focus and
guide the ions through the APi chambers where the air is pumped out, with the pressures ultimately reaching 10^{-6} mbar in the
110 TOF, where the ions' mass-to-charge ratio is determined. Nitrate ions have been shown to cluster readily with HOM molecules
(Ehn et al., 2014; Bianchi et al., 2019). The clustering is not ~~necessarily~~ equally efficient between all compounds however,
and signals are therefore not ~~necessarily~~ directly representative of the actual relative concentrations of the compounds. This
selectivity of the nitrate ions has been described by e.g. Hyttinen et al. (2015).

The CI-API-TOF mass spectrometry data was analysed with a Matlab (R2016a) based toolbox, tofTools (R607) (Junninen
115 et al., 2010). In further analysis the signals at different times were made comparable by normalising all time series by di-
viding them with the sum of the three reagent ion time series (NO_3^- , $\text{HNO}_3\text{NO}_3^-$, $(\text{HNO}_3)_2\text{NO}_3^-$, Jokinen et al., 2012). As
quantification of HOM remains challenging (Riva et al., 2019b), we opted for analysing only the relative changes to HOM
concentrations in this work.

2.2 Peak Fit Analysis

120 tofTools fits peaks based on a given list of compounds. Some of the compounds could potentially have very similar masses,
and fitting them correctly can sometimes prove challenging for an automatic routine (Cubison and Jimenez, 2015; Zhang et al.,
2019). Problems can also arise if the spectra are noisy, or if an observed ion is not listed in the compound list but something
with a close mass is. These types of issues may result in fitting a peak either too small or too large.

Including potentially false signals would be undesirable in the analysis. Individually estimating the quality of each peak fit
125 at each time point, is extremely time-consuming and presents an inherent challenge for high-resolution mass spectral analysis.

This motivated us to attempt to use a quantitative measure to describe the quality of the automatically generated fits. By assigning this variable for each fit, signals with poorly fitted peaks could be easily identified, and down-weighted or removed from the analysis.

130 The variable, which we will call the FitFactor, was used to quantify how well the fitted peak matched the spectra. The residual area, i.e. the difference between the spectral peak and the sum of all fitted peaks, is compared to the surface area of the fitted peak of interest. This residual-to-peak ratio is subtracted from unity, and the reached value is the FitFactor of the peak fit.

$$FitFactor = 1 - \frac{A_{residual}}{A_{peak}}$$

Thus, the better a peak fit is, the closer the value is to

135 Thus, a FitFactor of 1 (Figure A1 a.) Values would describe a perfect fit. Decreasing FitFactors indicate increasing uncertainties associated with the peak fit, and the quality of a peak fit with a FitFactor close to zero or even negative values are conversely associated with poor fits (Figure A1 d.). Something in between can be reasonable (A1 c.), but as the value decreases, uncertainty generally increases. Each compound has, is likely to be very poor. Each product was described with a single FitFactor value assigned to it. Spectra with signal below that was an average of the fits in the spectra that had reasonable signal (at least 50% of the time series maximum were ignored, because analysing for example a fit of a nitrate compound is not
140 relevant during times when no NO is being added to the chamber. The final compound FitFactor was the median of individual FitFactor calculations from the rest of the spectra).

Examples of determining the FitFactor from the residual-to-peak ratio of four different compounds. The title compounds are indicated by green lines and shading. The FitFactor value for each is shown in the upper right corner of the subplots. a) A good fit results in a FitFactor close to 1. b.) Accurate determination of the fit quality is tricky, when this many compounds and/or isotopes are fitted to the same spectral peak. Automatic fitting can artificially produce a good sum fit, which translates into a good FitFactor, but single compound fits may lack credibility. c.) The two fitted peaks match the spectra relatively well, resulting in a reasonably good FitFactor for the bigger peak. d.) A very poor fit results in low or even negative FitFactor.

By applying the FitFactor, a majority of the very poorly fitted peaks, such as the one Visual examples of the FitFactor determination from a few different peak fits are shown in Figure A1 d., can easily be identified and removed from the analysis.
150 However, some compounds may still have a relatively good FitFactor, despite the fit being somewhat questionable, like in the case of shown in Figure A1 b. These types of fits are especially tricky. The automatic peak fitting routine is capable of producing a sum peak that matches the spectra relatively well, which translates into a high FitFactor value. However, as so many peaks (3 compound peaks, 1 large isotope peak) have been closely fitted onto the same spectral peak, there is a great deal of uncertainty related to the individual peak fits. Thus, the FitFactor might not be optimal for this type of multiple fit problems,
155 but thanks to the selectivity of the nitrate CI-API-TOF, such peaks are not very prevalent in the spectra. The problem could be avoided by setting a minimum separation between fitted peaks, which we did not do in this work, alongside a discussion concerning the limitations of the method (Appendix A1). Since the peak list was generated semi-automatically by simply listing all compositions within chosen element number limits, setting a minimum FitFactor value limit, is however, a helpful

way to eliminate most, although not all, of the poorest fits. The analysis can then be focused on [the FitFactor was a helpful tool for identifying](#) the most reliable signals.

2.3 Peroxy radical chemistry and product formation

The ~~oxidation and HOM formation pathways mechanisms~~ of cyclohexene ozonolysis ~~described in detail by Rissanen et al. (2014), related autoxidation steps and other reactions leading to~~ RO₂ radical formation are detailed elsewhere (Rissanen et al., 2014). In this section, we ~~briefly outline the main steps considered when interpreting the results in this work. First, ozone oxidises cyclohexene by attaching itself on the double bond, leading to ring-opening. After rearrangement, loss of an OH radical, and an addition of an molecule, a only focus on the radical chemistry most relevant in the context of our study. The~~ primary peroxy radical ($\dot{\text{C}}_6\text{H}_9\text{O}_4$) of the chemical composition, is formed (Berndt et al., 2015). Peroxy radicals can undergo in cyclohexene ozonolysis is C₆H₉O₄ (Berndt et al., 2015), and radicals may acquire higher oxygen numbers by undergoing several steps of autoxidation (Crouse et al., 2013); ~~producing an increasingly more oxygenated radical, that may eventually terminate into a highly-oxidized end-product (i.e., HOM). However, the ultimate fate of the primary and potential subsequent radicals will depend on the exact conditions under which the oxidation takes place. This will, subsequently, greatly impact the final product molecules and their properties, such as volatility. Therefore, we outline the relevant main pathways below.~~

Before reaching termination, some radicals may be susceptible to a decrease in carbon number, following a loss of a CO group (Mereau et al., 2001; Rissanen et al., 2014):



Unimolecular termination of RO₂ typically takes place through the loss of a hydroxyl radical (OH) (Rissanen et al., 2014):



A loss of HO₂ is another previously suggested mechanism that has since been shown not to be likely under atmospheric temperatures (Hyttinen et al., 2016). Bimolecular reactions between RO₂ and HO₂, or between two RO₂ radicals also produce closed-shell species (Orlando and Tyndall, 2012; Rissanen et al., 2014). The former can typically produce ROOH molecules (Eq. (R3)), whereas the latter can lead to a production of a dimer molecule (ROOR) (Eq. (R4)) or two monomers (ROH and R[◊]CHO) (Eq. (R5)). Alternatively, bimolecular reactions can also produce reactive open-shell alkoxy radicals (RO) (Eq. R6).



Peroxy radicals are also a part of the chemistry of pollutants such as NO_x (i.e. NO and NO₂). The reactions between NO_x and RO₂ usually take one of the following forms (Orlando and Tyndall, 2012):



Typically ~~only~~ reaction (R8) is typically significant in producing closed-shell end-products, as products formed in reaction (R9) ~~are tend to be~~ thermally unstable, unless under very cold conditions (Orlando and Tyndall, 2012). ~~Under short~~
195 ~~reaction times a high concentration of has been shown to lead to dimer suppression and formation of highly oxidised nitrates in the cyclohexene ozonolysis system (Rissanen, 2018). However, here the longer reaction times and the much lower concentration are not likely to allow for a significant nitrate pool from~~ A high NO_2 reactions due to the equilibrium of reaction R9 residing
~~concentration also has the capability to suppress dimer formation, at least in shorter time scales, by terminating acylperoxy radicals ($\text{RC}(\text{O})\text{O}_2$) and producing peroxyacylnitrates that are more stable than the common RO_2NO_2 (Rissanen, 2018). Our~~
200 ~~experiments however allowed long reaction times, and the NO_2 concentration was a mere fraction of the value associated with such an effect. Therefore, the equilibrium in reaction (R9) is likely to reside~~ strongly on the reactant side, ~~and not allow a significant nitrate pool~~ under our experimental conditions.

A fraction of NO_2 radicals may ~~go on to~~ react with ozone to produce NO_3 , ~~which is also one of the~~ another main oxidising species in the atmosphere. ~~This oxidation~~ Oxidation of a BVOC compounds by NO_3 radicals produces nitrogen containing
205 peroxy radicals, ~~which can~~. Similarly to ozone derived radicals, these NO_3 -derived radicals can also form both monomer and dimer ~~species (Yan et al., 2016)~~ products (Yan et al., 2016; Zhang et al., 2020).

3 Results and Discussion

~~Figure 2~~ The left panel in figure 1 shows a HOM spectrum from cyclohexene ozonolysis (without NO addition), ~~with ten~~.
210 Ten of the biggest observed oxidation product peaks are labelled. Other large peaks shown are the nitrate dimer at 188 Th, and an unidentified peak at 234 Th. Strong signals of the HOM monomers $\text{C}_6\text{H}_8\text{O}_7$ and $\text{C}_6\text{H}_8\text{O}_9$ are observed, which ~~has been typically~~ is a typical observation reported also in ~~other earlier~~ studies (Rissanen et al., 2014; Berndt et al., 2015). A number of the largest signal peaks are from relatively little oxidised compounds, with ~~only~~ 4-5 oxygen atoms. Based on their elemental formulas, these species are most likely semi volatile organic compounds (SVOC) (~~Peräkylä et al., 2020~~), which might accumulate in the chamber ~~due to minimal wall losses (Peräkylä et al., 2020)~~, making them much more abundant than the low-
215 volatile HOM, ~~which are~~ that are much more efficiently removed by condensation. This might explain the large signals even though the charging efficiency of such compounds is generally low (Hyttinen et al., 2015). In addition to the monomers, also one dimer signal, $\text{C}_{12}\text{H}_{20}\text{O}_9$, reaches the top ten. One potential formation pathway for this dimer is a bimolecular reaction (R4) between radicals $\text{C}_6\text{H}_9\text{O}_8$ and $\text{C}_6\text{H}_{11}\text{O}_3$. The latter is the primary peroxy radical formed from OH-oxidation of cyclohexene (Berndt et al., 2015).

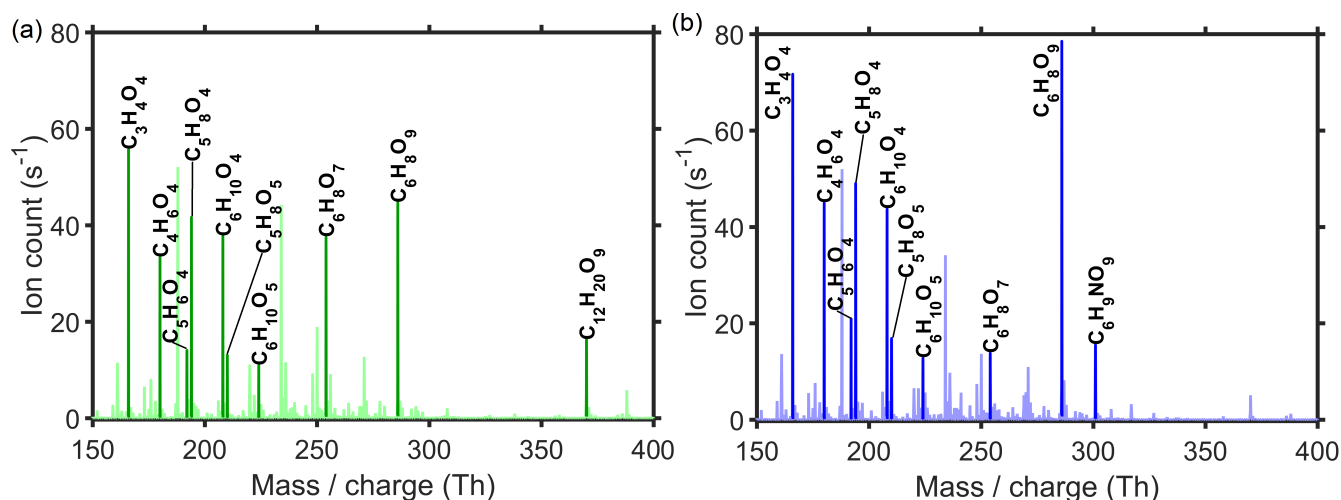


Figure 1. Two example 10-minute average spectra showing the largest product peaks that were all observed within the mass range 150-400 Th. Panel (a) shows a spectrum of cyclohexene ozonolysis without NO_x, while in panel (b) NO_x concentration was approximately 9 ppb. Ten of the biggest product peaks (darker green) are highlighted with a darker colour and labelled with their chemical composition. All products are/were detected as cluster with NO₃⁻, the mass of which is included in the masses shown in the x-axis.

220 3.1 Influence of NO on cyclohexene oxidation products

Injecting NO into the chamber has a significant impact on the chemistry within, which consequently leads to changed product signals. Although only NO mainly has an impact we expect NO to have the largest effect on the radical chemistry (see Eq. (R7) Eqs. R7-(R9)R9), due to the aforementioned technical limitations leading to a lack of separate measurements of NO, we compare signals to the total measured NO_x. NO₂ can also have some noticeable effect, by reactions with acylperoxy radicals (Eq. R9), which are expected to be abundant in this system (Rissanen et al., 2014; Rissanen, 2018), but as discussed in section 2.3, we do not expect this to be a significant process in our system. In addition, also oxidation by NO₃ radicals will take place in the chamber. As our observations are limited to the elemental composition of the products, and we have no additional information on specific structures, we do not attempt to distinguish between the effects of NO, NO₂, and NO₃.

An example spectrum during a NO_x experiment is shown in Figure ??1b. The ten biggest signals are otherwise the same as in the absence of NO_x (Figure ??1a), except for the dimer signal (C₁₂H₂₀O₉) having dropped to a fraction of the original and thus off the list, while a nitrate signal, C₆H₉NO₉ (Eq. R8), has replaced it. The monomer signal peaks are also on average larger than in Figure ?? in the presence of NO_x than without.

An example 10-minute average spectrum of cyclohexene ozonolysis in the presence of (approx. 9 ppb). Ten of the biggest product peaks (darker blue) are labelled with their chemical composition. All products are observed as clusters with, the mass of which is included in the masses shown in the x-axis.

A few examples of changes in product signals with increasing NO_x are shown in Figure 2. Species with comparable signal sizes are purposely selected (except for the nitrogen-containing dimer that has a much smaller signal). Shown are the increasing monomer $\text{C}_6\text{H}_8\text{O}_9$ signal, decreasing $\text{C}_6\text{H}_8\text{O}_7$ monomer signal, decreasing $\text{C}_{12}\text{H}_{20}\text{O}_9$ dimer signal and increasing $\text{C}_6\text{H}_9\text{NO}_9$ nitrate and $\text{C}_{12}\text{H}_{19}\text{NO}_{11}$ nitrogen-containing dimer signals. ~~When These trends are expected, as in the presence of NO_x is present in the chamber,~~ the occurrence of $\text{RO}_2 + \text{NO}_x$ reactions (Eq.(R7)-(R8)-(R9)) ~~also~~ reduce the likelihood for other bimolecular reactions (Eq. (R3)-(R6)).

~~Nitrogen-containing species are only expected to form during the~~ This figure also exemplifies the possible non-linear effects of NO_x experiments, as they are either formed in bimolecular reactions between and on different molecules. While some molecules do show a linear trend with the NO_x (Eq. Eq. R8-R9), or from radicals oxidised addition, the signal of the nitrate dimer $\text{C}_{12}\text{H}_{19}\text{NO}_{11}$ for example, initially increases with the added NO_x , as oxidation by NO_3 . ~~Nitrogen-containing dimers form in bimolecular reactions (Eq. (R4)), where one of the reacting radicals has been oxidised by a nitrate radical generates nitrogen-containing radicals, allowing for the formation of such a dimer. However, when NO_x concentration increases (>8 ppb), the suppression of $\text{RO}_2 + \text{RO}_2$ reactions by NO (and NO_2) starts to become more important and we observe a decreasing trend.~~

For a broader overview of signal changes in response to NO_x loading, Figure ~~??-3~~ shows the relative change of all non-nitrate products signals (nitrate-containing molecules are presented in Fig. 4 and discussed later) that were fitted. Signals with 9 ppb of NO_x were compared to signals without NO_x addition. All compounds with even hydrogen numbers are closed shell products, whereas ones with an odd number are radicals.

The results in Figure ~~??-3~~ paint a complicated picture on the formation of the non-nitrate compounds. While some features are expected and easy to explain, others remain less clear. Here we summarise some aspects of the findings. As adding NO into the chamber leads to peroxy radicals reacting with NO (Eq. (R7)-(R8)), other bi-molecular reactions (Eq. (R3)-R6) consequently decrease. The most clear effect is the general decrease of almost all dimers, as the RO_2 cross reactions forming them decrease. For monomers, the effects are more complicated, with some increasing and some decreasing. For example, the signals of the two largest HOM, $\text{C}_6\text{H}_8\text{O}_7$ and $\text{C}_6\text{H}_8\text{O}_9$ showed opposite behaviour, with the former decreasing and the latter increasing. ~~Speculating the reason for this difference is difficult, especially as the~~

This varied response in the monomer signals highlights how complicated the formation pathways can be, even when the initial reactant is a relatively simple molecule like cyclohexene. Speculating on the exact reasons for differences between individual signal responses is difficult. A single observed elemental composition may often contain a collection of isomers, each with distinct formation pathways. The reaction rate constants and different branching ratios of reactions between different combinations of RO_2 are also unknown, and can vary by orders of magnitude depending on the specific structures ~~(Shalleross et al., 2005; Berndt et al., 2018). Most of the~~ (Shalleross et al., 2005; Berndt et al., 2018). Therefore, any change in the reactant concentration, timescales, or other conditions, can yield very different results, with highly non-linear responses, making direct comparisons with other studies challenging.

For these reasons, we will only briefly look at certain example molecules and compound groups in more detail. The reason for the decreasing $\text{C}_6\text{H}_8\text{O}_7$ could be a decrease in bimolecular reactions of intermediate peroxyradicals. Rissanen et al. (2014) have

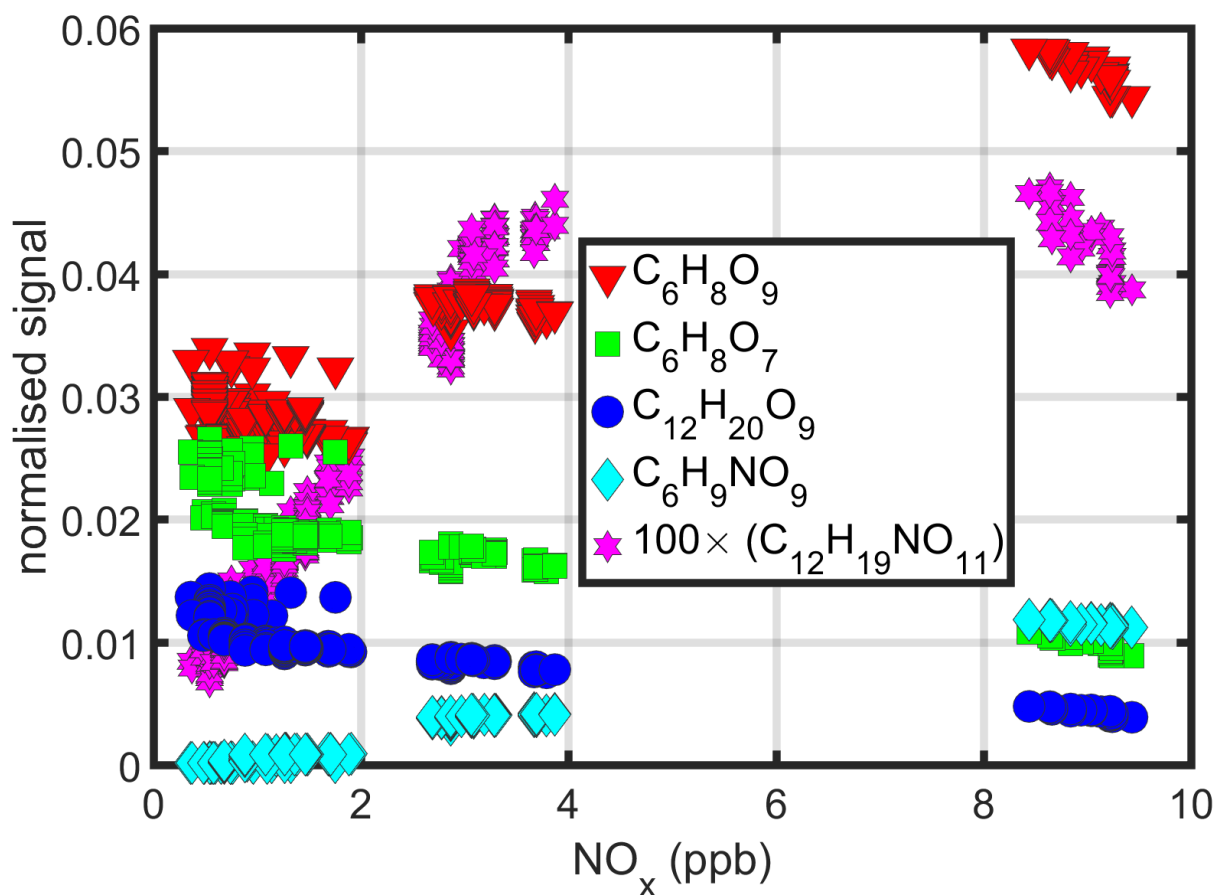


Figure 2. Examples of signal changes due to increased NO_x levels in the chamber. Shown are examples of two differently behaving monomers and a dimer without nitrogen, as well as a nitrogen-containing monomer and dimer. The last one had a signal so small that in the Figure it is multiplied by a 100. No measurements were done with NO_x concentrations between 4 and 8 ppb

275 previously suggested their likely formation pathway to involve a termination to a closed cell molecule in a $\text{RO}_2 + \text{RO}_2$ reaction. With the addition of NO , these types of reactions should decrease, and our results indeed show such a decrease (Fig. 3). The concurrent rising trend of $\text{C}_6\text{H}_8\text{O}_9$ suggests that its formation pathway may involve alkoxy radical intermediates, which can form both from $\text{RO}_2 + \text{RO}_2$ (R6) and $\text{RO}_2 + \text{NO}$ (R7) reactions. Similar to our results, (Rissanen, 2018) saw an increase of $\text{C}_6\text{H}_8\text{O}_8$ with NO , and suggested the involvement of alkoxy radicals. Overall, we noticed increasing trends for at least 20 different monomer compounds that did not contain N-atoms. As a notable exception, most of the $\text{C}_6\text{H}_{10}\text{O}_x$ compounds showed a decrease, which may be explained by a potential decrease in the bimolecular reactions R3 (R3) and (R5). The Most of the less oxidised SVOC only showed relatively small changes. Finally, NO_x will also function as a sink for OH, although

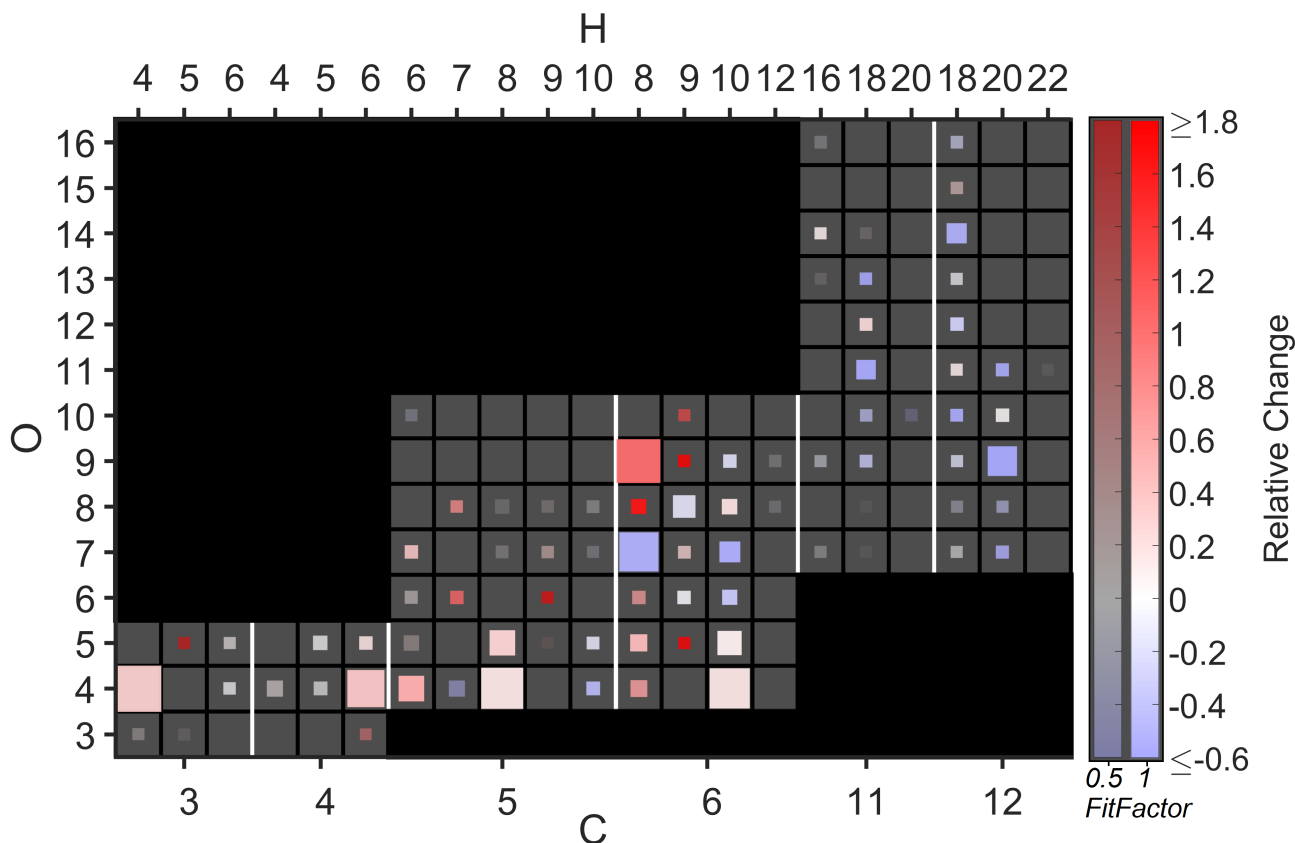


Figure 3. The effect of NO_x on observed signals of molecules that do not contain nitrogen. Signals from when NO_x concentration was 9ppb (“high”), are compared to signals during no NO injection (“zero”. Although, with small NO_x residual 1ppb of <0.9ppb). The bottom x-axis shows the carbon number of each compound, with the white vertical lines separating the different C-atom numbers. The top x-axis shows the hydrogen number, and the y-axis the oxygen number. Compounds with an odd number of H-atoms are radicals. Marker colour corresponds to the relative change in the signal size, when moving from zero NO_x to high NO_x, marker size describes the size of the signal at zero NO_x and the marker transparency describes the quality of the peak fit (if FitFactor=1, marker is fully opaque; if FitFactor<0, marker is fully transparent). A few The two colour bars show the colour scale with two different opacities (0.5 and 1). Some products had values outside the colour axis limits. (C₆H₉O₅, C₆H₉O₉ and C₁₁H₁₈O₁₁ being only such products with Fitfactors 0.7 and above, experiencing relative changes of 3.6, 4.8 and -0.7, respectively). Columns with no appreciable signals were omitted from the figure.

the-cyclohexene should still dominate the OH reactivity in the chamber even at the highest NO_x concentration. Most of the less oxidised SVOC showed relatively small changes.

The increase of the observed nitrate signals under high NO_x (9 ppb) conditions is shown in Figure 4. Many compounds were associated with an unexpectedly high signal even outside of NO_x experiments, and the size of this background signal is also shown in the figure. We opted to not simply subtract the background, i.e. the signal at zero added NO_x show the background

signal instead of simply subtracting it, because in some cases the background was much larger than the signal increase, which
285 would have caused large uncertainties when subtracting two relatively large numbers. In this way, the size of the background
is also visible, and one can identify more clearly which increases are most relevantreliable, i.e. the markers with the thickest
yellow lines. Small NO_x concentrations were measured continuously even outside the experiments, and although the NO_x
concentration during the measurement of the background signals was small in comparison to the concentration used in the
experiment (less than 10%), some of these signals could be from actual nitrate production. Alternatively, they could be residues
290 from earlier experiments, re-evaporating from the chamber walls. The signals could also be partially an artefact, as signals
assigned to organic nitrates may also have some contribution from RO₂ radicals, as the two can be difficult to separate due to
their relatively close masses (Peräkylä et al., 2020). This is perhaps the case especially with many of the nitrate monomers that
appeared to change very little. Some of them also had a relatively low FitFactor values (indicated by high transparency in the
figure) also suggesting issues with peak fitting.

295 Unsurprisingly, nitrates increased with higher Despite the difficulties with pre-existing signals, the formation of many nitrate
products was clearly observed in the experiment. C₆H₉NO₉ was clearly produced the most, and likely formed in the reaction
between NO and the radical C₆H₉O₈, which is one of the most abundant highly oxygenated RO₂ radicals in this system
(Berndt et al., 2015) (Eq. (R8)). Nitrogen-containing dimers are also detected. This implies that some of the cyclohexene had
been oxidised by NO₃, which had led to the formation of nitrogen containing radicals(Yan et al., 2016). These radicals were
300 then terminated by then likely formed dimers in reactions with ozone-derived RO₂ radicals from ozonolysis reactions, as the
observed nitrate-containing dimers only had one nitrate functionality. Similar nitrogen-containing HOM dimers have been
observed also in field measurements where monoterpenes were the dominant VOC (Zha et al., 2018; Jokinen et al., 2017), and
in studies by Yan et al. (2016) and Zhang et al. (2020) they have been directly implicated as cross reaction products of RO₂
formed in NO₃ and O₃ oxidation. The most abundant dimer was C₁₂H₁₉NO₁₁. It can could for example form in the reaction of
305 the C₆H₉O₈ radical, from ozonolysis, together with the primary RO₂ radical C₆H₁₀NO₅, from NO₃ oxidation of cyclohexene.
For the nitrate dimers to form, NO concentrations can not be so high that they completely dominate the RO₂ termination. In
Figure 2, C₁₂H₁₉NO₁₁ signal was seen to already experienced a decreasing trend when the total NO_x was above 8 ppb.

3.2 Condensation properties of cyclohexene oxidation products

The condensation sink of ABS seed aerosol was used to assess species volatilities. Compounds with low volatilities are expected
310 to rapidly condense onto the particle surfaces, whereas the more volatile compounds should remain in the gas phase. The loss
rate of the latter is mainly governed by flush out from the chamber, while the low-volatile compounds condense either onto the
walls or onto particles ($k_{\text{loss}} \approx k_{\text{CS}} + k_{\text{wall}}$). For a more detailed discussion on the dynamics of the chamber and this method,
see Peräkylä et al. (2020). Although we expect the seed aerosol to be the main driver of the changes observed during the
experiment, we cannot completely rule out the possibility for some influence from other changes in the product sources or
315 sinks. Figure 5 shows the signal response of three species against the condensation sink (CS) of the injected ABS particles.
The signals are normalised so that the median of the initial signal value before the seed experiment equals 1. As a result, the
fraction that remains following the seed addition is easy to observe. The three compounds in Figure 5 vary only by their oxygen

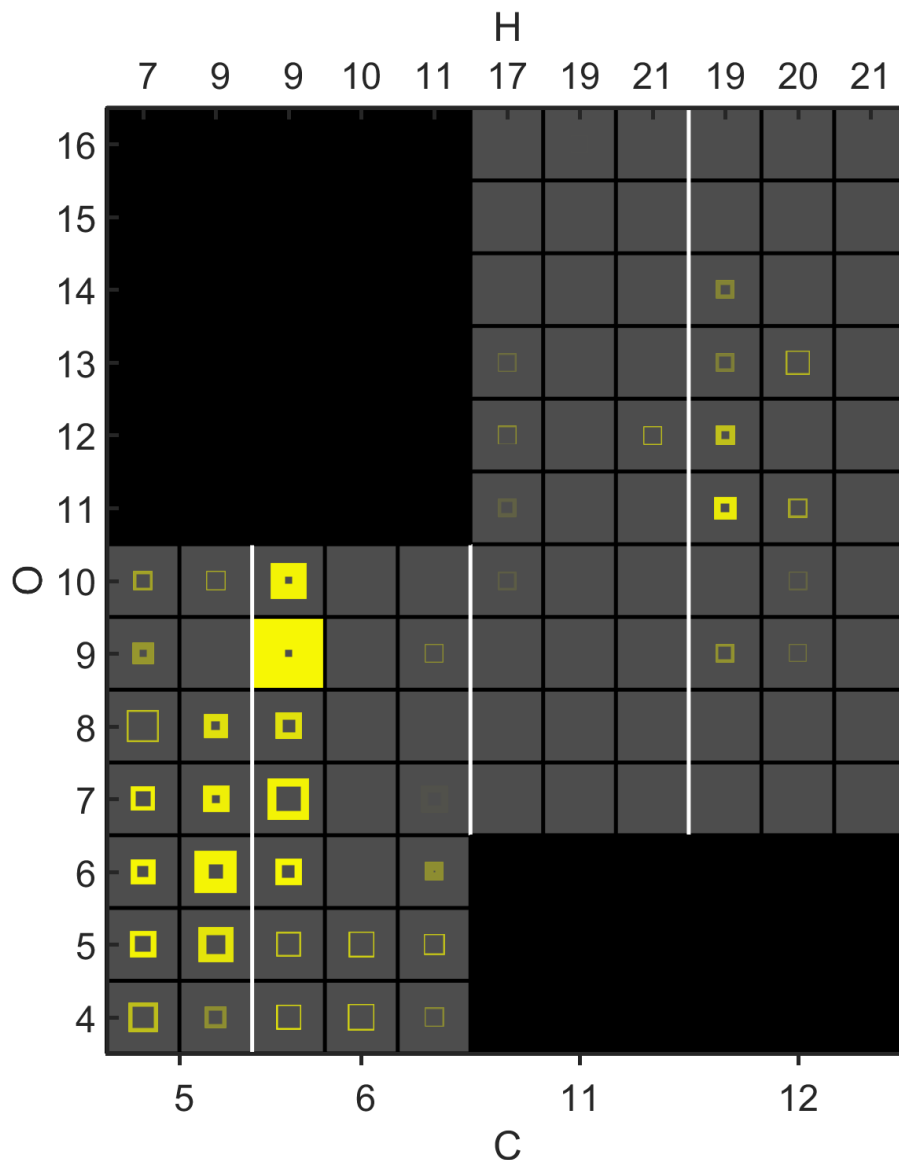


Figure 4. Signal changes in N-containing molecules upon NO_x addition during cyclohexene ozonolysis. The axes and transparency follow the same logic as in figure 5. Yellow—The area of the yellow squares show the size of the signal under high NO_x (9 ppb) conditions. The "holes" in the square correspond to the size of the background signal, i.e. the signal before NO_x addition. The large background for some peaks may be due to small amounts of being introduced to the chamber, or that these molecules were residues from chamber wall following earlier experiments.

number. The most highly oxygenated vapour, that is, $\text{C}_6\text{H}_{10}\text{O}_8$, condenses most readily and thus can be described as being the least volatile.

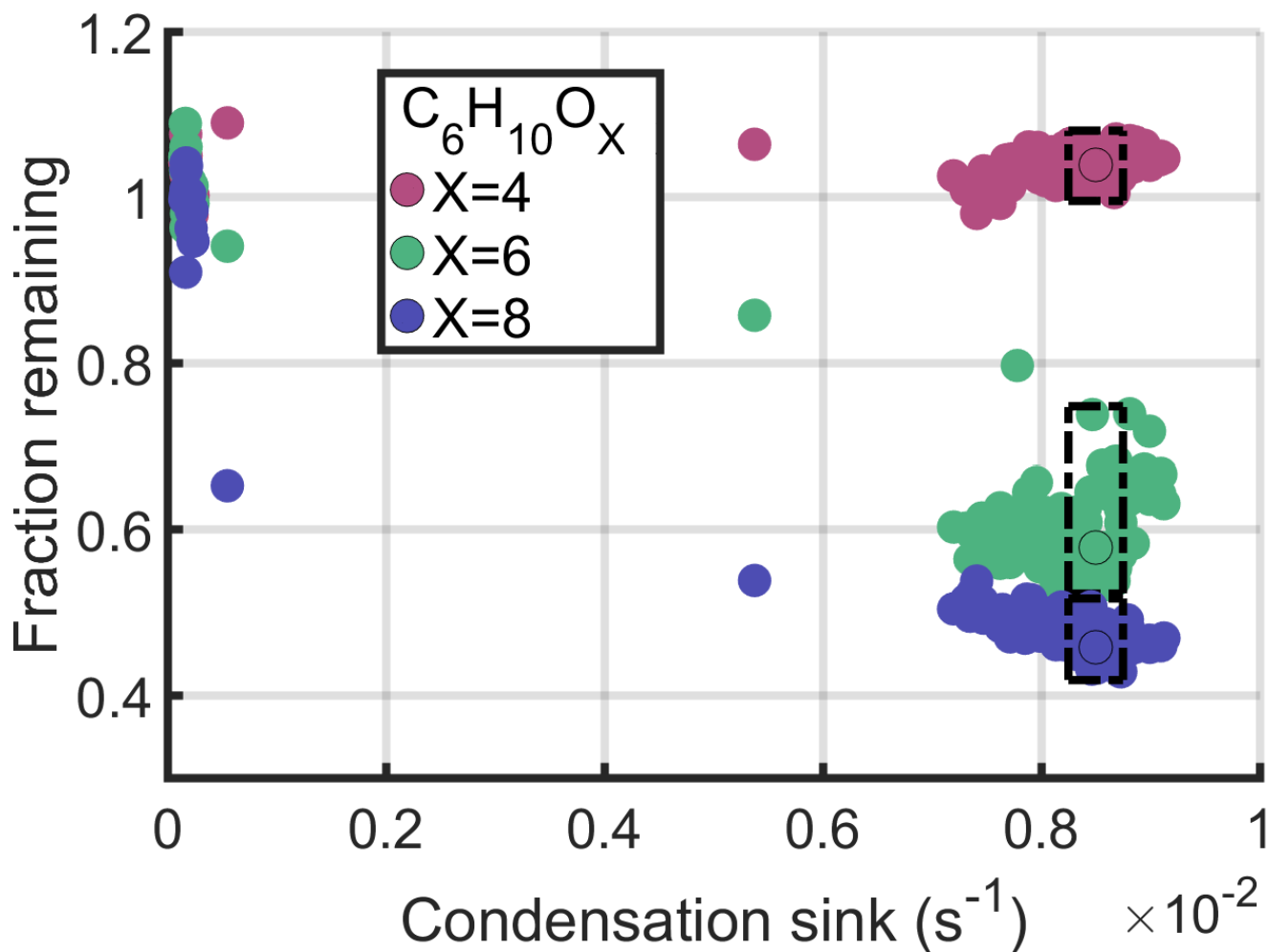


Figure 5. Condensation behaviour of three $C_6H_{10}O_x$ species. Signals are normalised so that the median of the signal before the ABS seed experiment is set to 1. Thus, shown on the y-axis is the remaining fraction of the signal of each molecules at a given condensation sink. The dashed line boxes encircle data points between the condensation sink values $0.825 \times 10^{-2} s^{-1}$ and $0.875 \times 10^{-2} s^{-1}$, and the black edged circles show the median of these boxed-in data points.

320 The fraction remaining (FR) at a condensation sink of approximately $0.85 \times 10^{-2} s^{-1}$ (see Fig. 5) was calculated for all
 compounds fitted in the mass spectra. These are plotted in Figure ??-6 against the mass of each compound. The methodology
 here closely follows that outlined by Peräkylä et al. (2020). Although a single seed experiment such as this will not provide
 us detailed information on the variation in volatility between different non-condensing or irreversibly condensing species, the
 transition between the two states can give us qualitative understanding on the main properties determining the volatility of
 cyclohexene products.
 325 Only the clearest signals are ~~shown, with the aim to bring only~~ shown in the figure, to focus on the most

meaningful ~~signal-into-focus~~ signals while minimising noise. In practice, this means compounds with FitFactor values of at least 0.7, which was deemed to strike a good balance between quality (see Figure A1) and quantity. Non-nitrate and nitrate products are both plotted, while radicals are not, since condensation is only a minor sink for these species because of their high reactivities with other radicals (Peräkylä et al., 2020).

330 In addition to observations, also predictions based on two different models are shown in Figure ??6. For both models, the modelled values were scaled to the range of observed remaining fractions, between 0.43 and 1. Estimates in the upper panel are modelled with the parametrisation by Peräkylä et al. (2020),

$$\log_{10}(C^*) = 0.18 \times n_C - 0.14 \times n_H - 0.38 \times n_O + 0.8 \times n_N + 3.1$$

that was originally based on measurements with α -pinene oxidation products. n_C , n_H , n_O , and n_N , correspond to the numbers of carbon, hydrogen, oxygen and nitrogen atoms in the molecule (charging ion excluded).

335 The model for volatility used in the lower panel figure is given by Bianchi et al. (2019), ~~although the model we used was further modified to account~~ and the modification accounting for nitrate functionalities is also included. In nitrates, 3 oxygen atoms are bonded in the nitrate group, which as a whole reduces volatility by a factor of 2.5. We assumed an activity ~~coefficient~~ coefficient (γ) of 1, so that $C^* = \gamma C^0 = C^0$. Given the small precursor molecules in our study, the formed SOA is expected to have very high O:C ratios, just like the majority of potentially condensing molecules we measure in the gas phase. Under these
340 circumstances, and considering the low RH of our experiments, an activity coefficient close to unity is likely to be a valid assumption (Donahue et al., 2011). Thereby, the equation used in this work is

$$\log_{10}(C^*) = (n_{C0} - n_C)b_C - (n_O - 3n_N)b_O - 2 \frac{(n_O - 3n_N)n_C}{n_C + (n_O - 3n_N)} b_{CO} - b_N n_N$$

Here the additional constants are $n_{C0} = 25$, $b_C = 0.475$, $b_O = 0.2$, $b_{CO} = 0.9$ and $b_N = 2.5$ (Bianchi et al., 2019). To be able to compare the modelled volatilities to our observations of FR, we converted the C^* -values to remaining fractions they would theoretically correspond to under our experimental conditions. The conversion was done similarly to Peräkylä et al. (2020),
345 who fitted a logistic curve $y = \frac{y_{max} - y_{min}}{1 + e^{-3.0(x - 0.13)}} + y_{min}$, where y_{min} and y_{max} are limits for scaling, between FR of model compounds and the common logarithm of their saturation concentration from the ADCHAM model (Roldin et al., 2014). The original conversion equation had to be modified, as the SOA concentration in our experiment was an order of magnitude lower than in the Peräkylä et al. (2020) experiment, which would lead to lesser condensation and higher FR at similar C^* . ~~To~~
By following equilibrium partitioning theory (Donahue et al., 2012), to account for the difference, we shifted the original fit
350 between FR and $\log_{10}(C^*)$ left by 1 (i.e. plugging in $x = \log_{10}(C^*) + 1$), thus adjusting the relationship between FR and C^* by a factor of 10. While certainly introducing additional uncertainty, we expect this approach to allow an adequate comparison of our findings with those of Peräkylä et al. (2020) and Bianchi et al. (2019).

Many monomers had remaining fractions above 1, indicating that the signal increased. ~~The likely explanation for this~~ A likely explanation is that these compounds are semivolatile and were still equilibrating with the chamber walls, and thus were not in
355 steady-state. ~~Another possibility~~ This type of behaviour is likely the largest source of uncertainty in our data, and the potential

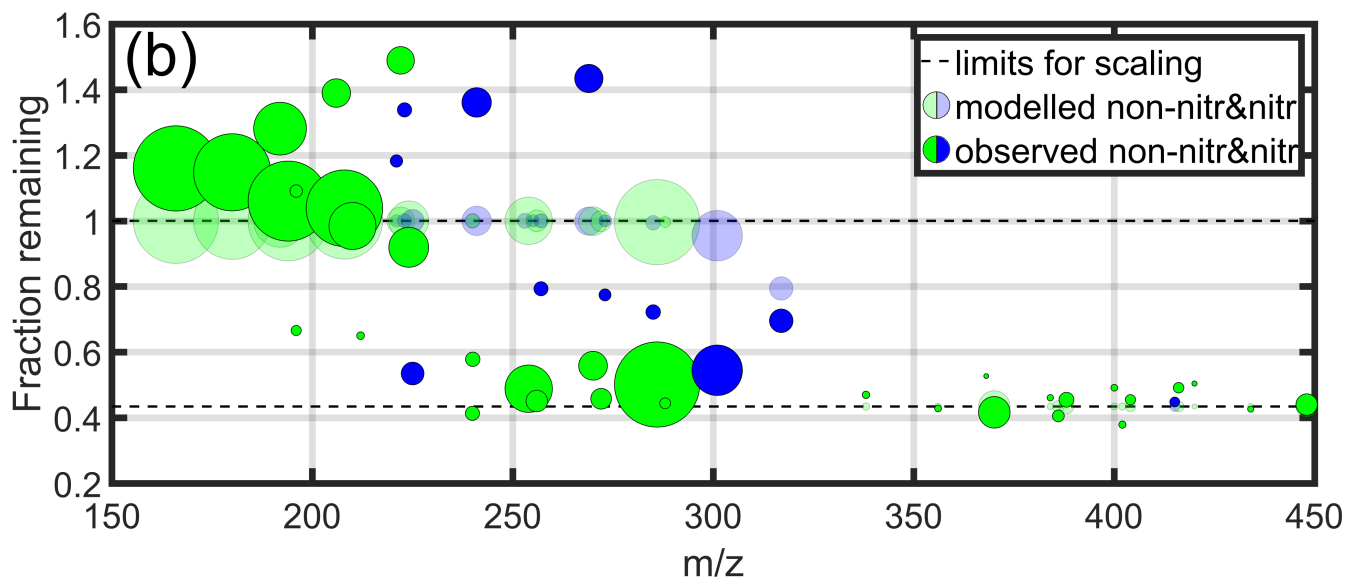
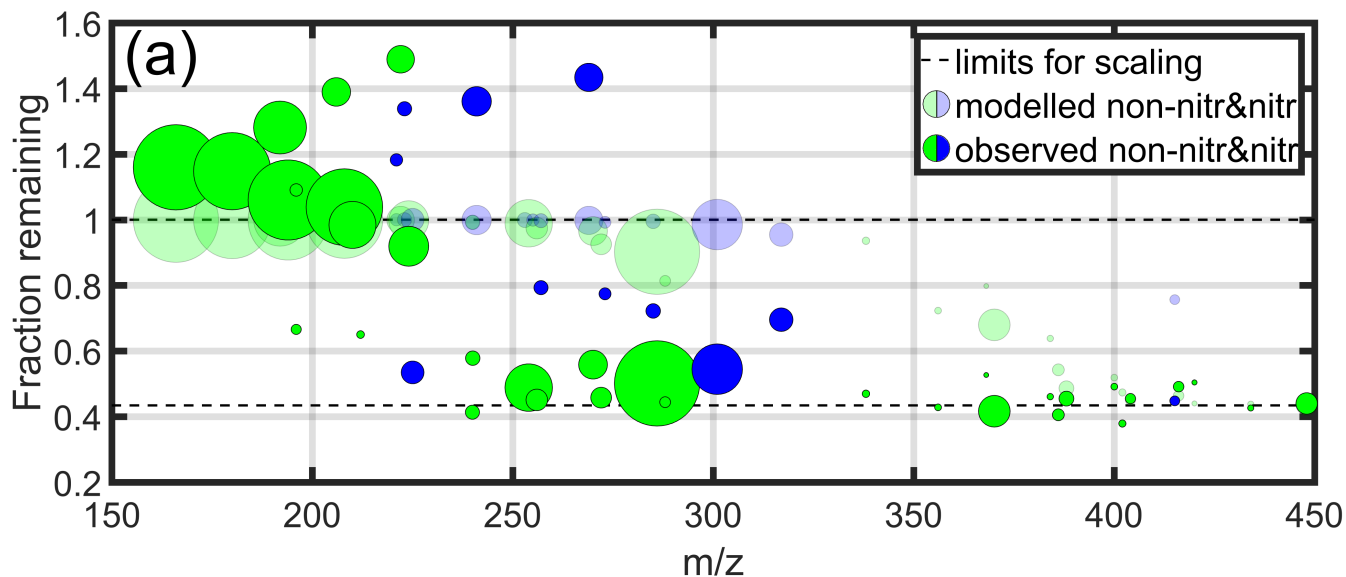


Figure 6. The observed (opaque) and modelled (transparent) decrease in signal during an ABS seed experiment, for non-nitrate (green) and nitrate (blue) products. The top panel compares our observations with the model presented by Peräkylä et al. (2020), while the bottom panel compares to the model by Bianchi et al. (2019). Only compounds with the best peak fits ($\text{FitFactor} \geq 0.7$) are shown. The circles sizes are based on the relative sizes of the initial signals before the ABS aerosol injection, with the exception of the two biggest signals, which were slightly scaled down. The condensation sink was approximately $0.85 \times 10^{-2} \text{ s}^{-1}$, meaning that these points correspond to the black edged circles in Figure 5. All molecules were charged with a nitrate ion, which has a mass-to-charge ratio of 62 Th.

error will be different for all molecules and therefore difficult to quantify in any detail. A rough estimate is that this would at most cause an error of 10-20 % to the relative changes reported above for the relatively fast seed addition experiments. Another possible reason for the increased signal is that these molecules have condensed-particle phase sources, causing an increased formation during the which would explain the increased formation upon seed addition. Due to these associated uncertainties, we do not try to interpret these points further, but note that although some of the. Some molecules with $FR < 1$ may also have seen have also experienced similar effects, the effects are probably smaller as these molecules but most likely to a smaller degree, as the lower volatility compounds are less likely to evaporate from particles or walls.

The higher the mass of the molecule, the more inclined it was to be taken up by the injected particles product volatilities were closely related to the molecular mass. The condensation sink caused most of the non-nitrate signals with masses above 250 Th to drop to values less than 50% of the original. Such a high net condensation is a clear indication of a low volatility, suggesting irreversible condensation and low volatilities. At masses below 225 Th, compounds were on the other hand, compounds remained fairly volatile, as all signals remained either either remained the same or increased. The exception to this are the two small green points around 200 Th, Two exceptions to this with $FR < 0.7$. These are at masses close to 200 Th were $C_5H_{10}O_4$ and $C_5H_{10}O_5$, which are less oxidised than most molecules typically observed with the nitrate adduct CI-API-TOF. This may mean that they contain multiple hydroxyl and/or carboxylic acids groups (which may improve their clustering ability with NO_3^-) that decrease their volatility more than if the O-atoms were incorporated as carbonyl and hydroperoxide functionalities, which is more typical for HOM.

Similarly to non-nitrates, the volatility of nitrates decreases with mass. However, the transition from high to low volatility appears to happen at a larger mass than for non-nitrates, which is in line with previously determined group additivity trends of volatility (Pankow and Asher, 2008; Kroll and Seinfeld, 2008). A clear outlier is $C_5H_9NO_5$ (225.04 Th), as the signal comparison suggests an unexpectedly low volatility for the compound. This might however be an artefact from issues with the fitting of multiple peaks, as shown specifically for this ion in Figure A1 b.

Both the Peräkylä et al. (2020) and Bianchi et al. (2019) models successfully predict the high volatility of the least oxidised monomers. The predicted volatilities of larger monomers are however in both cases higher than the observed. Both models predict the strongest volatility transition in, and the irreversible condensation of the largest dimers. They locate the steepest transition in volatility to the region between our observed monomers and dimers, although we observe and in disagreement with our observations, fail to capture the observed nearly irreversible condensation of many of the monomers to condense nearly irreversibly. Peräkylä et al. (2020) model predicts a more gradual more oxygenated monomers as well. The Bianchi et al. (2019) model predicts the irreversible condensation of all of the dimers, but the decrease in volatility as a in the Peräkylä et al. (2020) model is a more gradual function of mass than the Bianchi et al. (2019) model and therefore predicts a condensation weaker than observed. The discrepancy between the observations and modelled FR is also clearly visible in Figure B1.

Next, the link between elemental composition and volatility is analysed further. As the models misplace the transition between non-condensing and irreversibly condensing compounds, they also over-predict the product volatilities in general. Since we

390 cannot distinguish the differences in volatilities of compounds that condensed irreversibly in our experiments, we are not able to confirm where the observed volatilities merge with the ones predicted by the models.

In order to better understand what controls the volatility of cyclohexene products and where these models might fail, we had a more detailed look into the effect of elemental composition on FR. In Figure ??7, the fraction remaining is plotted separately against carbon, hydrogen and oxygen numbers, as well as the "effective O:C ratio" $(n_O - 2n_N)/n_C$. The incentive for choosing this ratio, rather than simply O:C, is the same as for the modification of the Bianchi et al. (2019) model, i.e. to account for nitrate functionalities. For non-nitrates this expression reduces to an O:C ratio. ~~For comparison, also α -pinene products from an experiment by Peräkylä et al. (2020) are plotted on the background. The data is from an experiment with dry ABS seed aerosol and. As the reactant concentrations and chamber conditions differed between our and their experiments, the signals are not comparable in magnitude. In Figure ??, the observations have been scaled to the same marker size and FR range. Due to the~~
400 ~~higher SOA concentration in the Peräkylä et al. (2020) experiment, the remaining fractions of α -pinene products are somewhat lower than they would be at our experiment's SOA concentration.~~

Molecules Cyclohexene oxidation products with the lowest carbon and hydrogen content hardly condensed, while the opposite was true for the molecules with the highest C and H content, mostly associated with the dimers (Fig. ??a and ??7a and 7b). However, in the regions between these extremes, e.g. around C₆ and H₈, respectively, molecules range from volatile to effectively non-volatile, and thus purely C- or H-atom content is not a good predictor for volatility. As-

However, the fraction remaining as a function of O-atom content the oxygen content of the molecule (Fig. ??e), ~~the volatility seems to be much more monotonically decreasing~~ 7c) shows the most clearly monotonically decreasing trend. The transition from non-condensing to condensing occurs at an O-atom contents of 6-7 ~~for cyclohexene HOM~~, with nitrates again being slightly more volatile than non-nitrates. Finally, ~~Figure ??~~ Figure 7d demonstrates that the effective O:C ratio ~~is~~ appears to be the
410 worst predictor for volatility out of the parameters plotted. Despite O:C ratios of organic aerosol often being used as a reference for volatility, ~~for a single molecule~~ it does not ~~have a good correlation~~ correlate well with the volatility of an individual molecule. This is ~~to be expected~~ not surprising, as high molecular weight dimers require only a relatively small number of oxygen atoms (thus having low O:C) to condense, while small molecules with 1-3 C-atoms are unlikely to condense regardless of their O:C ratios. ~~In all figures (??a-d), the~~ Therefore, the absolute oxygen number alone seems to be the best predictor for the FR.

415 Elemental composition can be a good indication of the possible functional groups in the molecule that ultimately determine the volatility of a compound (Kroll and Seinfeld, 2008; Pankow and Asher, 2008). This is essentially the basis of models, such as the two by Peräkylä et al. (2020) and Bianchi et al. (2019) that we tested (Fig. 6), that aim to estimate volatility directly from the easily measurable elemental composition. As such, they however always contain some intrinsic assumptions of the structure and the likely functional groups in the molecule. The Peräkylä et al. (2020) model is developed from an observed
420 condensation of α -pinene products (C₁₀H₁₆), while the Bianchi et al. (2019) model is also primarily constructed with larger precursors in mind. An underlying assumption of a larger carbon frame, or a lower O:C, might explain the discrepancy between our observations and the modelled FR, as an extrapolation to molecules with less C and much higher O:C, might cause the models to fail, especially in fully capturing the observed effect the O-atom content has on the uptake (Fig. 7). Incidentally, the Bianchi et al. (2019) model predicted transition in condensation (Fig. 6) appears to be misplaced by approximately the

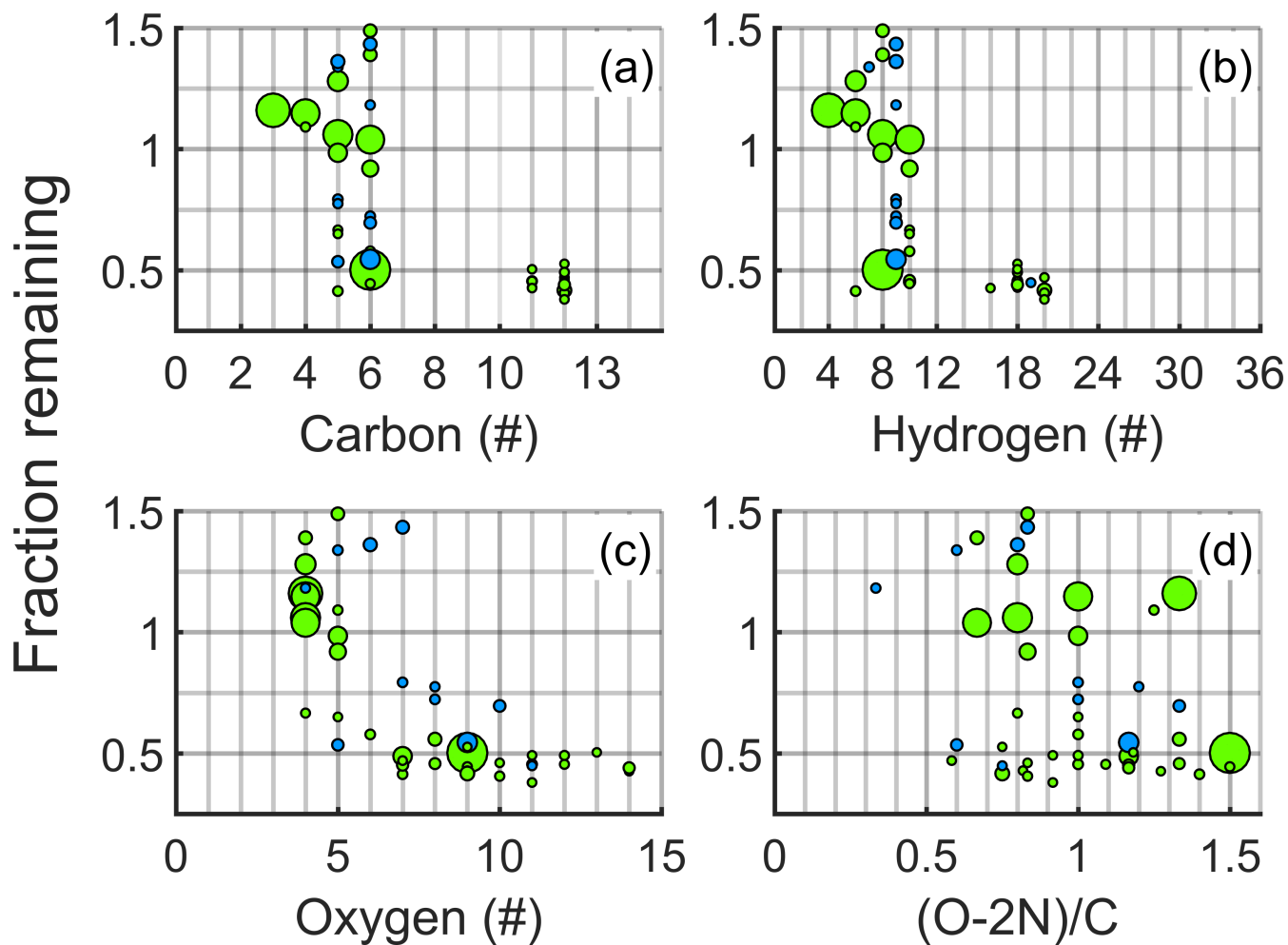


Figure 7. Fraction remaining (FR) after seed addition, as a function of elemental composition. This figure is similar to Figure 6, except that here the FR of products is plotted against the a.) carbon, b.) hydrogen and c.) oxygen content of the compounds, and against d.) the effective O:C ratio $(O-2N)/C$.

425 mass difference between monoterpenes and cyclohexene. Furthermore, if the additional 4 carbon atoms that separate the two
 compounds are imagined into the model (replacing n_C with $n_C + 4$), the modified Bianchi et al. (2019) model predicts our
 observed cyclohexene product FR remarkably well (Fig. B2). While anecdotal, it does support our conclusion that the oxygen
 content of the HOM is the primary contributor to their uptake efficiency. If the Peräkylä et al. (2020) model is adjusted similarly
 for both the number of C-atoms (+4) as well as the number of H-atoms (+6), which is also part of the parametrisation, the result
 430 is a negligible change in the modelled FR, as the added C- and cyclohexene products showed similar behaviour. The lesser sensitivity to oxygen content of the Peräkylä et al. (2020) model

could also partly explain its comparatively slow decrease in volatility as a function of mass. Peräkylä et al. (2020) themselves already contemplated that the parametrisation might not be able to predict volatilities of products with less than 10 carbon atoms, thinking that they might depend differently on the number of oxygen atoms, for example.

435 ~~Fraction remaining (FR) after seed addition, as a function of elemental composition. This figure is similar to Figure ??, except that here the FR is plotted against the a.) carbon, b.) hydrogen and c.) oxygen content of the compounds, and against d.) the effective O:C ratio (O-2N)/C. In addition to the cyclohexene (–) oxidation products, α -pinene (–) ozonolysis products from a seed experiment by Peräkylä et al. (2020) are also included. As the signals between the two studies are not directly comparable, the α -pinene products have been scaled to the same limits of FR, and to the same range of marker sizes. Another~~
440 ~~possible explanation to some of the model-observation discrepancies, is if the monomer HOM in our experiments not only condense, but are also lost by reactive uptake on the particles due to labile hydroperoxide functionalities. Separating between these effects is not possible from our data, but would be extremely important for understanding, and modelling, SOA formation. We hope our work will help spur future studies to design experiments where these open questions could be tackled. Currently, volatility estimates for HOM are highly uncertain, with different computational methods predicting vapour pressures differing~~
445 ~~by many orders of magnitude for a given HOM (Kurtén et al., 2016).~~

4 Conclusions

We have investigated the formation and fates of cyclohexene ~~oxidation-ozonolysis~~ products, with a focus on the most oxidised species. These highly oxygenated organic molecules (HOM) were measured with a CI-API-TOF mass spectrometer. A statistical approach was utilised for evaluating the ~~quality of the identification of different ions in the mass spectra. In this~~
450 ~~approach, reliability of compound signals that had been determined from the spectra by means of an automatic peak fitting procedure. By assigning each peak fit with~~ a so-called FitFactor ~~parameter was set up to compare the spectral residual and the peak area. By this simple method, evaluating thousands of peak fits was fast and feasible. Ultimately, this allowed discarding or down-weighting signals with poorly fitted peaks~~ value that compared the fitted peak and spectral residual areas, we could fast and feasibly sort through the fitted compounds and identify the most reliable signals.

455 ~~Perturbations to product signals by NO_x were analysed species-by-species, revealing that enhanced the formation of most of the observed monomer. Monomers (C₃-C₆) species, although some had a widely varied response to the addition. Although most of them increased, they did so in varying degrees, while some products also experienced an opposite effect. These effects are likely driven by increased termination of radicals by NO and decreased. This variability highlights the complexities involved in HOM formation. Even resembling compounds can have very different main formation pathways under the same~~
460 ~~conditions, leading to more alkoxy radicals, but with the current data, the detailed reasons for the individual differences can only be speculated. However, more easily interpretable effects were also observed, namely the formation of organic nitrates, and the decrease of dimer even contrasting responses to added NO_x. We expect the introduction of RO₂ + NO reactions to have caused most of the changes, and depending on whether the product benefited from resulting RO formation, or suffered from the reduced bimolecular cross-reactions, the signal either increased or decreased. Interpreting individual signal changes~~

465 any further is challenging, as any signal peak assigned to a given chemical formula is likely to contain many different isomers
with possibly distinct formation pathways, which may have different responses to changes in the prevailing conditions.

The lower rate of RO₂ cross-reactions during the NO_x experiment also resulted in a decline in the formation of dimer
products (C₁₁-C₁₂) species, due to enhanced reactions. This reaction, taking place at the expense of cross reactions, caused a
decrease in dimers without a nitrate functionality. Dimers containing a nitrate group conversely experienced a small increase.
470 This is likely due to that had no nitrate functionalities. The addition of NO_x also led to nitrate formation, and both nitrate
monomers and dimers were observed. The latter suggested multi-oxidant chemistry, as nitrate dimers are likely to form in a
bimolecular reaction involving a NO₃ radical (formed from) oxidation forming with nitrate functionalities, which can then react
with other to form dimers-derived radical. This can lead to the formation of nitrate dimers, as long as the NO concentration
is not high enough to completely dominate the RO₂ termination in the system. Monomer nitrates, on the other hand, see a
475 monotonically increasing trend as more NO is added to the chamber, as expected.

The volatility condensation of the oxidation products was probed in experiments where ammonium bisulfate seed aerosol
was added to the chamber. This resulted in a clear decrease in most HOM signals, with The concentration of lighter products
either remained the same or even increased, while after a narrow transition zone, all non-nitrate products heavier than 250 Th
condensing to condensed onto the particles at nearly equal rates. Nitrates were slightly more volatile, dropping to at least half
480 of the initial signal. Nitrates condensed less than non-nitrates with similar masses. Besides mass, a break-down A breakdown
of the elemental composition of the different products indicated that the most significant main factor determining the volatility of
the observed cyclohexene oxidation products is the oxygen number. Products having an oxygen atom content of cyclohexene
HOM is the number of oxygen atoms in the molecule. Products that had 7 or more, were all at the lowest end of the observed
volatility range oxygen atoms, all condensed irreversibly in our experiment. Carbon and hydrogen atom content also correlated
485 with volatility, but were clearly more ambiguous indicators for volatility than the oxygen content this was mostly limited to the
different condensational behaviour of monomers and dimers, while the effect of C and H content specifically was somewhat
ambiguous. For example, molecules with 5 or 6 C-atoms can range anywhere from volatile to effectively non-volatile were
observed to ranged anywhere from non-condensing to irreversibly condensing. We also found that the effective O:C ratio is by
itself not a good measure for volatility, as molecules like small monomers can have a much higher volatility than e.g., despite
490 having many dimers, even with an O:C ratio twice as high.

There have been a few models presented recently for predicting the volatility of a compound based on its elemental
composition. We tested our observed volatilities against two Several parametrisations with the aim of predicting product
volatilities based on their elemental composition have been developed. Being typically applied to larger reactants, such as
monoterpenes (C₁₀H₁₆), we investigated if two such models, by Bianchi et al. (2019) and Peräkylä et al. (2020), the model by
495 Bianchi et al. (2019) with a modification for nitrates, and another by Peräkylä et al. (2020). They are based on larger initial
reactants than cyclohexene, such as α-pinene would also be able to reproduce the condensation of cyclohexene (C₆H₁₀)
products. While both could predict the generally higher volatility of monomers compared to were able to capture the lack
of condensation in lighter monomers and the irreversible condensation of heavier dimers, they were not able to produce over
predicted the volatilities of many compounds and misplaced the transition between these two extremes to occur at a larger mass

500 than observed. This was likely primarily due to the models over estimating the impact of C, and under estimating the impact of O, on the uptake efficiency. Although misplacing the transition from non-condensing to condensing, the parametrisation by Bianchi et al. (2019) was able to reproduce the shape of the transition, suggesting that it was still able to capture the number of oxygen atoms in the cyclohexene products as the main cause for increased condensation. The Peräkylä et al. (2020) model on the other hand, is less sensitive to oxygen, and produced a volatility transition too gradual to fit our observations, and while it
505 unlike the Bianchi et al. (2019) model predicted some condensation for the largest monomers as well, it did not reproduce the observed irreversible condensation of larger monomers. This highlights the difficulty in estimating smaller dimers. This analysis demonstrates the challenges and limitations in estimating product volatilities with simple parametrisations. As cyclohexene is used as a surrogate for monoterpenes like α -pinene, we also compared the volatilities of HOM from α -pinene and cyclohexene against the different compositions. The results for both VOC imply that a high oxygen number is the most essential parameter
510 for lowered volatility. Thus, the ability to undergo efficient autoxidation may composition based parametrisations. Any parametrisation contains some intrinsic assumptions on the functional groups possibly present in the molecule. These assumptions may no longer hold true if the model is extrapolated outside its limits, which seems to be the case with the models we tested and the cyclohexene products.

The highly species-specific effects of NO_x , especially in monomers, highlights the complex formation pathways involved in
515 HOM formation, even when the reactant is the relatively simple cyclohexene. Therefore, any effects interfering with formation pathways cannot necessarily be generalised across different species. The mismatch between the parametrisations and the observed condensation of cyclohexene products highlights the role of oxygen as a primary driver for increased condensation, an effect which may be especially pronounced in the oxidation products of the smaller VOCs. As the number of oxygen atoms alone enhanced the condensation of cyclohexene oxidation products so effectively, an efficient progression of autoxidation
520 could potentially be more important for SOA formation than the size of the initial VOC reactant. Finally, the relative roles of volatility-driven condensation and potential reactive uptake remains an open question, and further studies will be needed to address this important issue.

Data availability. Data is available at <https://doi.org/10.5281/zenodo.4001576>

Appendix A: [Supporting material for Methods](#)

<u>Compound</u>	<u>concentration</u>
O ₃	<u>18 ppb</u>
C ₆ H ₁₀	<u>~70 ppb</u>
NO _x	<u>0 ppb</u>
	<u>1 ppb</u>
	<u>3 ppb</u>
	<u>9 ppb</u>
<u>80 nm ABS particles</u>	<u>0 μgm^{-3}</u>
	<u>9 μgm^{-3}</u>

Table A1. The approximate reactant, oxidant and particle concentrations during the conducted experiments. Cyclohexene concentration is a very rough estimate based on changes in ozone concentration.

525 **A1 Peak fit Analysis with the FitFactor parameter**

By applying the FitFactor, a majority of the very poorly fitted peaks, such as the one shown in Figure A1d, could easily be identified and removed from the analysis. However, some compounds could have a relatively good FitFactor, despite the fit being somewhat questionable, like in the case of $C_5H_9NO_5NO_3^-$ shown in Figure A1b. These types of fits are especially difficult. The automatic peak fitting routine is capable of producing a sum peak that matches the spectra relatively well, which translates into a high FitFactor value. However, as so many peaks (3 compound peaks, 1 large isotope peak) have been closely fitted onto the same spectral peak, there is still a great deal of uncertainty related to the individual peak fits. Thus, the FitFactor might not be optimal for this type of multiple fit problems, but thanks to the selectivity of the nitrate CI-APi-TOF, such peaks are not very prevalent in the spectra. The problem could be avoided by setting a minimum separation between fitted peaks, which we did not do in this work.

530

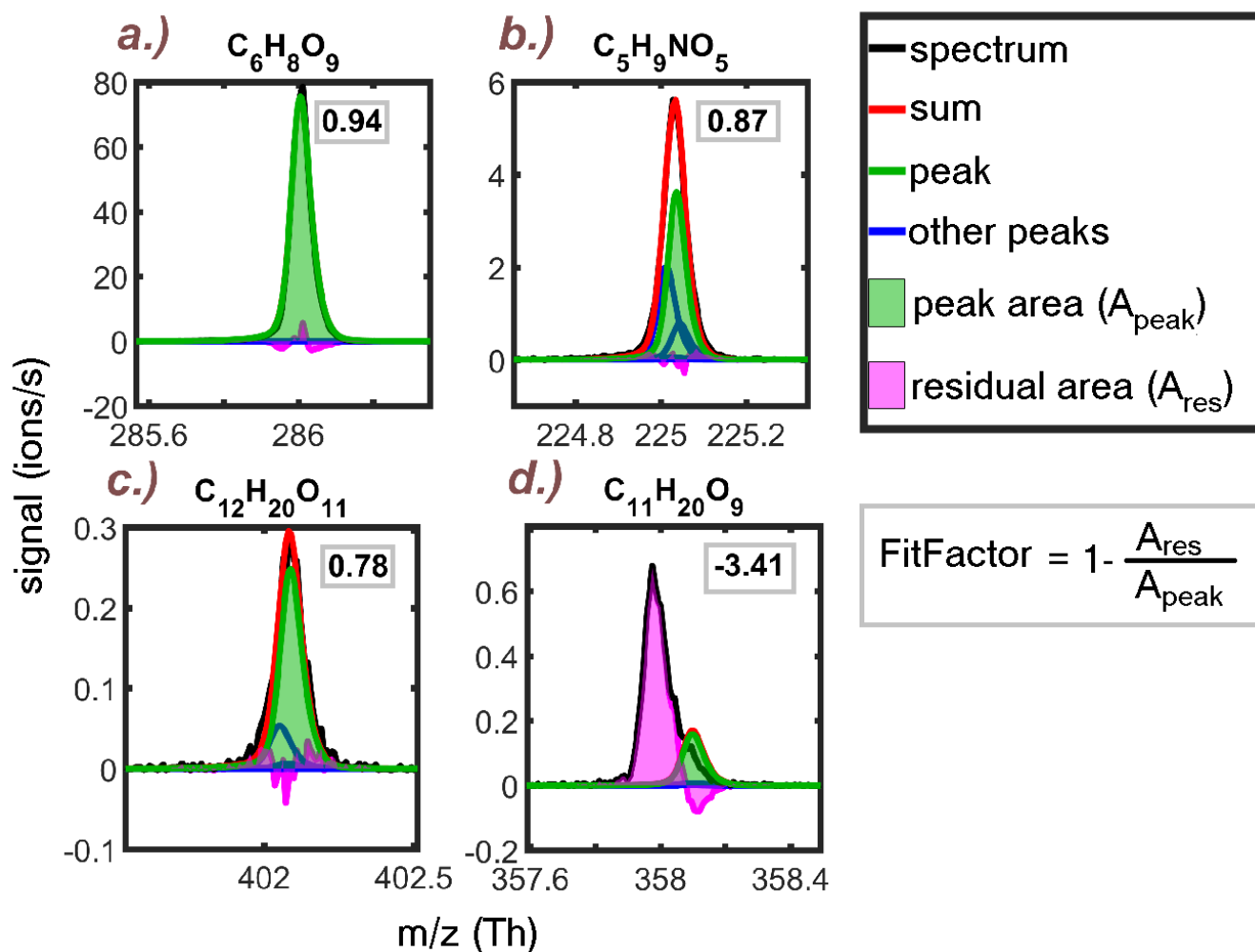


Figure A1. Examples of determining the FitFactor from the residual-to-peak ratio of four different compounds. The title compounds are indicated by green lines and shading. The FitFactor value for each is shown in the upper right corner of the subplots. a.) A good fit results in a FitFactor close to 1. b.) Accurate determination of the fit quality is tricky, when this many compounds and/or isotopes are fitted to the same spectral peak. Automatic fitting can artificially produce a good sum fit, which translates into a good FitFactor, but single compound fits may lack credibility. c.) The two fitted peaks match the spectra relatively well, resulting in a reasonably good FitFactor for the bigger peak. d.) A very poor fit results in low or even negative FitFactor.

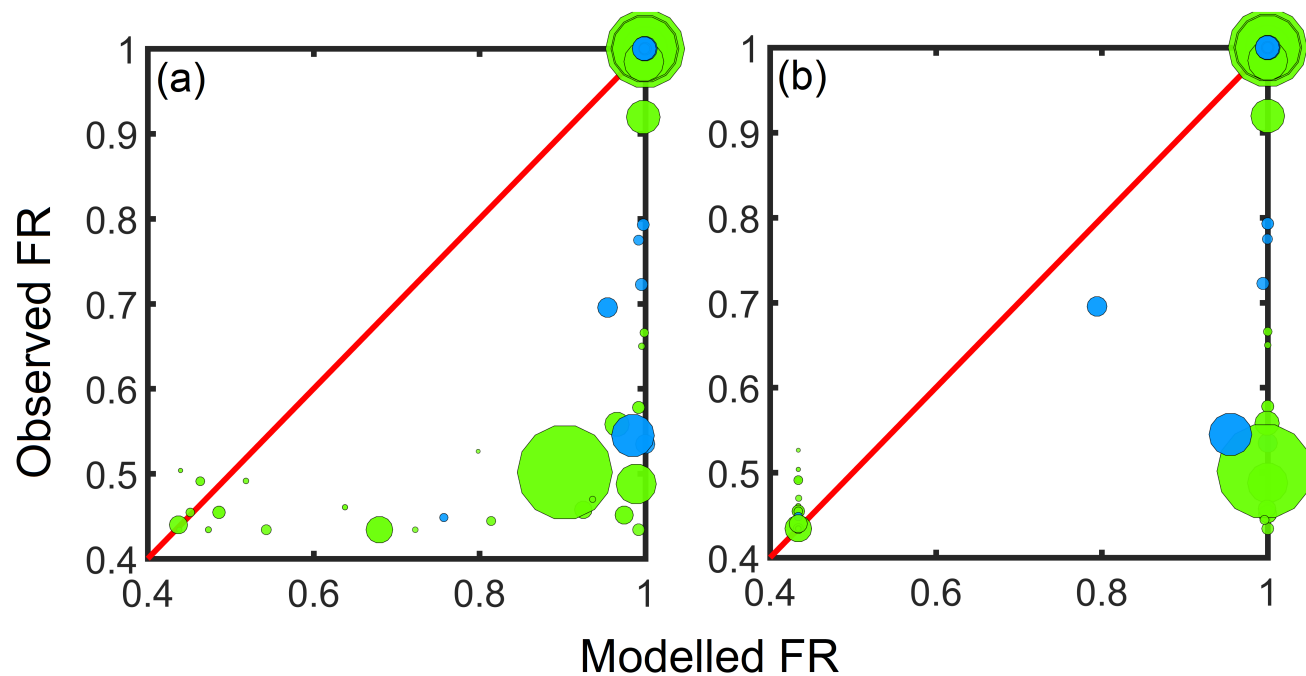


Figure B1. [Similar to Figure 6, except here the observed fraction remaining \(FR\) is plotted against the FR modelled by \(a\) the Peräkylä et al. \(2020\) parametrisation and \(b\) the Bianchi et al. \(2019\) parametrisation.](#)

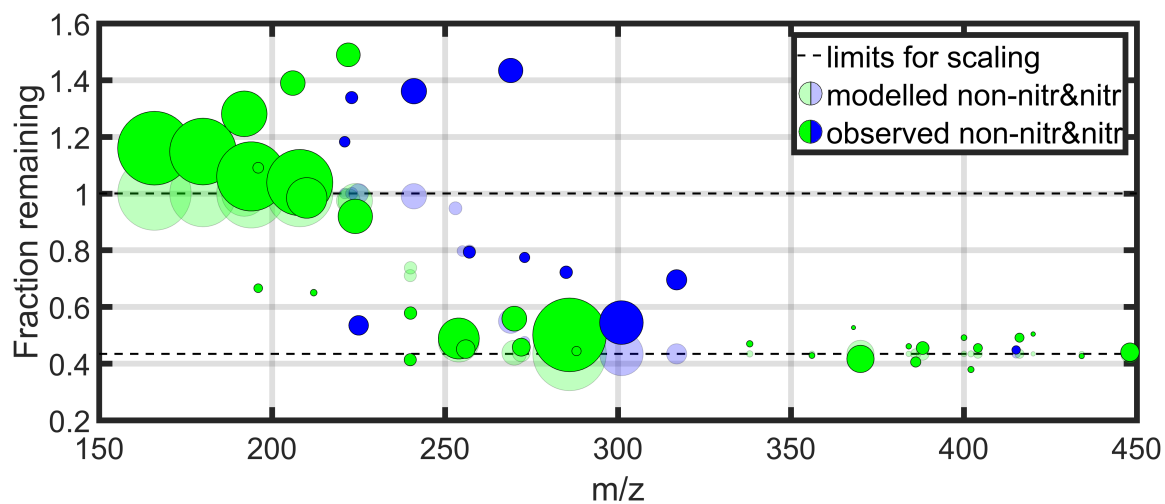


Figure B2. Similar to Figure 6, except here, the Bianchi et al. (2019) model is modified to give a volatility of a hypothetical product with 4 more carbon atoms than in our observed products:

$$\log_{10}(C^*) = (n_{C0} - (n_C + 4))b_C - (n_O - 3n_N)b_O - 2 \frac{(n_O - 3n_N)(n_C + 4)}{(n_C + 4) + (n_O - 3n_N)} b_{CO} - b_{NN}n_N$$

Author contributions. The study was conceived by ME, MRiv and MRis. The measurements were performed by MRiv, OG and LQ. MRä did the main data analysis and visualization under the supervision of ME and with additional guidance from OP and MRis. MRä wrote the manuscript, and it was reviewed and commented by the other authors.

Competing interests. The authors declare that they have no conflict of interest.

540 *Acknowledgements.* This work was supported by King Abdullah University of Science and Technology (Award OSR-2016-CRG5-3022), the European Research Council (Grant 638703-COALA), and the Academy of Finland Centre of Excellence (Grants 307331, 317380, 320094 and 326948). We thank Liine Heikkinen for assistance during the experiments, Pontus Roldin for helpful discussion, and the tofTools team for providing tools for mass spectrometry data analysis.

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