



**HAL**  
open science

## Development of a parallel sampling and analysis method for the elucidation of gas/particle partitioning of oxygenated semi-volatile organics : a limonene ozonolysis study

Stéphanie Rossignol, Laura Chiappini, Emilie Perraudin, Caroline Rio,  
Sébastien Fable, R. Valorso, Jean-François Doussin

### ► To cite this version:

Stéphanie Rossignol, Laura Chiappini, Emilie Perraudin, Caroline Rio, Sébastien Fable, et al.. Development of a parallel sampling and analysis method for the elucidation of gas/particle partitioning of oxygenated semi-volatile organics : a limonene ozonolysis study. *Atmospheric Measurement Techniques*, 2012, 5 (6), pp.1459-1489. 10.5194/amt-5-1459-2012 . ineris-00961778

**HAL Id: ineris-00961778**

**<https://ineris.hal.science/ineris-00961778>**

Submitted on 20 Mar 2014

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



# Development of a parallel sampling and analysis method for the elucidation of gas/particle partitioning of oxygenated semi-volatile organics: a limonene ozonolysis study

S. Rossignol<sup>1,2</sup>, L. Chiappini<sup>1</sup>, E. Perraudin<sup>2,\*</sup>, C. Rio<sup>1</sup>, S. Fable<sup>1</sup>, R. Valorso<sup>2</sup>, and J. F. Doussin<sup>2</sup>

<sup>1</sup>Institut National de l'Environnement Industriel et des Risques, Parc Technologique ALATA, 60550 Verneuil-en-Halatte, France

<sup>2</sup>LISA, Universités Paris-Est-Créteil et Paris Diderot, CNRS UMR7583, 61 Av. du Général de Gaulle, 94010 Créteil, France

\* now at: EPOC, University of Bordeaux, CNRS UMR5805, 351 Cours de la Libération, 33400 Talence, France

Correspondence to: L. Chiappini (laura.chiappini@ineris.fr)

Received: 9 January 2012 – Published in Atmos. Meas. Tech. Discuss.: 6 February 2012

Revised: 3 May 2012 – Accepted: 25 May 2012 – Published: 27 June 2012

**Abstract.** The gas/particle partitioning behaviour of the semi-volatile fraction of secondary organic matter and the associated multiphase chemistry are key features to accurately evaluate climate and health impacts of secondary organic aerosol (SOA). However, today, the partitioning of oxygenated secondary species is rarely assessed in experimental SOA studies and SOA modelling is still largely based on estimated partitioning data. This paper describes a new analytical approach, solvent-free and easy to use, to explore the chemical composition of the secondary organic matter at a molecular scale in both gas and particulate phases. The method is based on thermal desorption (TD) of gas and particulate samples, coupled with gas chromatography (GC) and mass spectrometry (MS), with derivatisation on sampling supports. Gaseous compounds were trapped on Tenax TA adsorbent tubes pre-coated with pentafluorobenzylhydroxylamine (PFBHA) or N-Methyl-N-(*t*-butyldimethylsilyl)trifluoroacetamide (MTBSTFA). Particulate samples were collected onto quartz or Teflon-quartz filters and subsequently subjected to derivatisation with PFBHA or MTBSTFA before TD-GC/MS analysis. Method development and validation are presented for an atmospherically relevant range of organic acids and carbonyl and hydroxyl compounds. Application of the method to a limonene ozonolysis experiment conducted in the EUPHORE simulation chamber under simulated atmospheric conditions of low concentrations of limonene precursor and relative humidity, provides an overview of the method capabilities. Twenty-

five compounds were positively or tentatively identified, nine being in both gaseous and particulate phases; and twelve, among them tricarboxylic acids, hydroxyl dicarboxylic acids and oxodicarboxylic acids, being detected for the first time.

## 1 Introduction

Atmospheric formation of particulate matter from the gas-to-particle transfer of organics during their atmospheric oxidation is now considered as one of the main phenomena involved in both air quality and climate issues (IPCC, 2007; Hallquist et al., 2009). A better understanding of the inter-related processes and an improved representation in models are still required (Kanakidou et al., 2005). Secondary organic aerosol (SOA) formation is commonly described by gas-to-particle conversion of semi- and non-volatile secondary species formed in the gas phase by atmospheric oxidation of volatile organic compound (VOC) precursors (e.g. Odum et al., 1996; Seinfeld and Pankow, 2003; Asher and Pankow, 2006). However, for SOA formation and evolution pathways, in addition to the initial gas-phase oxidation processes, a significant number of studies in the last decade have pointed out the role of multiphase chemical processes in the particle phase (e.g. Graber and Rudich, 2006; Kroll et al., 2007; Healy et al., 2008; Monks et al., 2009) or at the gas-particle interface (e.g. Rudich, 2003; Rudich et al., 2007). These processes involve continuous modification

of the chemical composition of SOA and its physicochemical properties and interdependently influence the partitioning of semi-volatile organic species (SVOCs) between gaseous and aerosol phases, thereby consuming or forming SVOCs (Healy et al., 2008).

Hence, understanding multiphase chemical processes is today a key issue to assess both aerosol climate impacts and health effects. It can improve climate impact evaluation through modelling enhancement of formation and evolution pathways and characterisation of aerosol physical properties. In terms of health issues, knowledge of SOA formation pathways and fine particle composition in different outdoor/indoor conditions is required for toxicological evaluation.

In spite of large progress made in recent years (e.g. Donahue et al., 2006; Camredon et al., 2010; Parikh et al., 2011), current SOA formation and evolution modelling often suffer from discrepancies between modelling results and observations (e.g. Volkamer et al., 2007; Chan et al., 2009; Pankow, 2011), notably revealing gaps in our knowledge of SOA formation and aging pathways. Atmospheric chemistry models integrating SOA formation are usually based on the parameterisation of SVOC<sub>*i*</sub> gas-particle partitioning proposed by Pankow (1994). This parameterisation is based on the definition of an equilibrium partitioning coefficient  $K_{p,i}$  ( $\text{m}^3 \mu\text{g}^{-1}$ ):

$$K_{p,i} = \frac{(\text{ng} \mu\text{g}^{-1})_{\text{particle phase}}}{(\text{ng} \text{m}^{-3})_{\text{gas phase}}} = \frac{P_i}{G_i M}, \quad (1)$$

where  $P_i$  is the mass concentration ( $\text{ng} \text{m}^{-3}$  air) of the SVOC<sub>*i*</sub> in the particle phase,  $G_i$  is the mass concentration ( $\text{ng} \text{m}^{-3}$  air) of the SVOC<sub>*i*</sub> in the gas phase and  $M$  is the mass concentration ( $\mu\text{g} \text{m}^{-3}$  air) of the total sorbing particle phase. While Eq. (1) is independent of the nature of the sorption process – i.e. adsorption or absorption (Lazaridis, 1999) – most of the models assume an absorptive gas/liquid partitioning of SVOCs. This mechanism is expected to be predominant in SOA formation (Pankow, 1994; Odum et al., 1996) and links  $K_{p,i}$  with the saturation vapour pressure of the SVOC<sub>*i*</sub> through the Eq. (2):

$$K_{p,i} = \frac{760 RT}{10^6 \overline{MW}_{\text{om}} \gamma_i P_{L,i}^0}, \quad (2)$$

where  $R$  ( $\text{m}^3 \text{atm} \text{mol}^{-1} \text{K}^{-1}$ ) is the ideal gas constant,  $T$  (K) is temperature,  $\overline{MW}_{\text{om}}$  ( $\text{g} \text{mol}^{-1}$ ) is the average molecular weight of the absorbing particulate material,  $\gamma_i$  is the activity coefficient of compound  $i$  in the particulate phase, and  $P_{L,i}^0$  (Torr) is the vapour pressure of compound  $i$  as a pure liquid (subcooled, if necessary).

SOA modelling based on chemical oxidation schemes (e.g. Jenkin, 2004; Camredon et al., 2007; Bessagnet et al., 2008; Chan et al., 2009; Valorso et al., 2011) generally estimate the theoretical  $P_{L,i}^0$  using group contribution methods

(e.g. Arp and Goss, 2009; Barley and McFiggans, 2010) and usually sets  $\gamma_i$  to unity, owing to experimental determination difficulties. This can lead to uncertainties that are suspected to account for observed discrepancies between models and ambient observations in SOA yields. Another likely major source of discrepancies could be the poor consideration of multiphase chemical processes as a sink or source of SVOCs (Chan et al., 2009). An alternative modelling approach to the mechanistic description of SOA is the volatility basis set (VBS) (Donahue et al., 2006, 2009; Cappa and Jimenez, 2010; Farina et al., 2010; Tsimpidi et al., 2010) which is based on distribution of the considered organic matter mass into volatility classes according to their saturation mass concentration  $C_i^*$  (corresponding to the inverse of the Pankow type partitioning coefficient). In this case, the dilution process and the evolution of particulate mass are the driving features. While the VBS scheme does not directly integrate chemical mechanisms, it takes reactivity into account by considering a redistribution of the organic mass from one volatility bin to another. VBS consequently requires data to link atmospherically relevant partitioning behaviour to the origin of the organic aerosol and its composition. This context highlights the requirement for improving our comprehension of multiphase chemical mechanisms, especially those driving the formation of SVOCs and their consumption, and providing atmospherically relevant data on the partitioning behaviour of SVOCs. There is therefore a strong need to simultaneously explore the molecular composition of both gas and particulate phases in SOA studies, and to systematically determine the observed partitioning of SVOCs. The related experimental data have to be evaluated not only with low-concentration simulation chamber experiments but also using ambient atmospheres to assess the relevance of the obtained results.

However, since atmospheric secondary organic matter results from complex and successive oxidation processes, it is highly functionalised and reactive. It contains oxygenated moieties such as aldehyde, keto, hydroxyl, and carboxylic acid groups. Exploring SOA chemical composition and the associated gas phase at a molecular scale is consequently a true analytical challenge (Prather et al., 2008). This is emphasised when the atmospherically relevant constraints of low concentration levels and relative humidity are taken into account.

Until today, only a few studies have assessed the partitioning of secondary organic compounds through determination of partitioning coefficients  $K_p$  for individual secondary species. Some papers report the use of a thermodenuder (de Gouw and Warneke, 2007; Grieshop et al., 2009; Lee et al., 2011) in order to evaluate the volatility of SOA formed from different precursors without information on the chemical composition. In these studies, SOA formed in simulated atmospheres flows through and evaporates in a temperature-controlled tube, previously calibrated with compounds with a known vapour pressure. By this method SOA mass is divided

into volatility classes with known vapour pressure ranges, providing a volatility distribution. This method is very useful to provide atmospheric relevant constraints for modelling based on the Volatility Basis Set (Grieshop et al., 2009) and evaluate the semi-volatile and non-volatile fractions of SOA. Nevertheless, it remains incomplete as it provides no insights into chemical processes.

Studies aiming at elucidating chemical processes focus on the characterisation of the chemical composition of secondary organic matter at a molecular scale. However, while the aerosol phase is most often analysed by chromatographic off-line methods to explore its highly complex molecular composition, the gas-phase products are most often monitored by an on-line method providing limited insights into the molecular composition. Today, proton transfer reaction mass spectrometry (PTR-MS) is the most widely employed on-line technique for assessing the chemical composition of the gas phase (e.g. Blake et al., 2004; Kroll et al., 2006), as it provides continuous molecular information and quantification of gas-phase compounds at very low concentration levels (around 10 ppt) without interference of water, making it usable in the ambient atmosphere. Nevertheless, PTR-MS is not specific as it does not provide detailed structural information.

Chromatographic methods using an on- or off-line approach, often coupled with thermal desorption, are sometimes applied to elucidate the molecular composition of the gas phase. However, apart from a few approaches using derivatisation strategies (e.g. Ho and Yu, 2002, 2004; Temime et al., 2007; Pacolay et al., 2008), these techniques are restricted to non-polar or mono-functionalised compounds (e.g. Wedel et al., 1998; Xu et al., 2003; Sinha et al., 2010) with limited applications to secondary chemistry studies.

Studies aiming at elucidating the underlying mechanisms of SOA formation focus on the chemical composition of the particulate phase. For this purpose, a wider range of powerful analytical methods exists, mainly because of the chemical complexity of the aerosol and its diversity, which can be divided into on-line and off-line methods (Hallquist et al., 2009). Over the past decade, on-line real-time aerosol mass spectrometry (AMS) has become a technique of choice to explore the chemical composition of the aerosol phase with a high time resolution (Sullivan and Prather, 2005; Canagaratna et al., 2007). Combining direct thermal extraction of sampled airborne aerosol (flash vaporisation upon impaction on a hot surface,  $\sim 600^\circ\text{C}$ , under high vacuum) with electron ionisation (EI) and mass spectrometry, the AMS technique provides data on the bulk chemical nature of particulate organic matter – sulphate, nitrate, ammonium and chloride contents, and the oxidation state with function type identification (alcohols, carboxylic acids and carbonyls) – and is consequently a powerful tool for the chemical and physical characterisation of SOA. It has been successfully applied to both ambient (e.g. Zhang et al., 2011) and simulated atmospheres

(e.g. Sato et al., 2010). However, complex EI mass spectra resulting from extensive fragmentation do not allow for the unambiguous identification of individual organic compounds, which limits the exploration of the composition at a molecular scale.

Another interesting in-situ method is thermal desorption aerosol gas chromatography (TAG instrument), based on coupling thermal desorption of the aerosol with gas chromatography/mass spectrometry (GC/MS) or GC-flame ionisation detection (GC-FID) (Williams et al., 2006). Recently improved by GC  $\times$  GC separation (2D-TAG, Goldstein et al., 2008; Isaacman et al., 2011), TAG provides detailed data on the volatility distribution of the particle components with a good time resolution (from 1 to 2 h). However, GC separation limits TAG detection and the identification range to non-polar and mono-functionalised compounds, so that it does not allow to obtain insights into SOA chemistry.

Off-line chromatographic methods are required to address the high complexity of SOA composition. The methods differ in a first instance by the extraction step. Hays and Lavrich (2007) provided an interesting review on thermal desorption-gas chromatography/mass spectrometry (TD/GC/MS) applied to fine aerosol. TD is a very convenient extraction method as it can be fully automated and is inexpensive, sensitive and solvent-free. However, like on-line TAG, this technique is limited by the polarity of SOA components. To overcome this limitation, an in-situ derivatisation step can be introduced. A few recent studies have integrated this principle, exposing sample filters to a derivatisation reagent for methylation or trimethylsilylation during thermal desorption for the detection and quantification of mono- and dicarboxylic acids (Beiner et al., 2009; Sheesley et al., 2010; Orasche et al., 2011). Other extraction techniques, more scarcely used, have also been reported. As an example, supercritical fluid extraction (SFE) couples the extraction step directly to the analytical process. Its efficiency for extracting organic compounds from complex matrices has been demonstrated by coupling SFE with GC/MS or liquid chromatography/GC/MS (LC/GC/MS) (Hansen et al., 1995; Forstner et al., 1997; Castells et al., 2003). The process has the advantage in that it allows derivatisation of polar compounds during the static extraction step (Shimmo et al., 2004; Chiappini et al., 2006). Nevertheless, the large majority of studies exploring the chemical composition of fine organic aerosol is based on a first solvent sample extraction step as it provides several possibilities for further analytical processes. Solvent extraction type and steps to be considered (solvents choice, ultrasonication, Soxhlet, pre-concentrations steps, derivatisation, etc.) are selected according to both the fraction of the organic mass studied (water-soluble organic compounds, polycyclic aromatic hydrocarbons, carbonyl, carboxyl and/or hydroxyl compounds, organosulphates, organonitrates, oligomers, humic-like substances, etc.) and the analytical technique employed (GC, LC, ion chromatography – IC, electrophoresis, etc.). Mass

spectrometric analysis is widely used for identification and quantification. Among the various possible combinations, those aiming at analysis of polar and functionalised compounds (compounds containing, for example, carbonyl, carboxyl and/or hydroxyl groups) are mainly divided between GC/MS analysis with prior derivatisation (e.g. Nolte et al., 2001; Claeys et al., 2004, 2007; Edney et al., 2005; Szmigielski et al., 2007; Chan et al., 2010; Vivanco et al., 2011) and LC/soft ionisation-MS analysis [electrospray ionisation (ESI-MS) or atmospheric pressure chemical ionisation (APCI-MS)] (e.g. Wan and Yu, 2007; Gómez-González et al., 2008; Kitanovski et al., 2011). These techniques are constantly being improved and made more complex in order to improve detection limits, enlarge the range of separable species (GC  $\times$  GC, LC  $\times$  LC), or increase the structural characterisation capability through higher-order MS ( $MS^n$ ) or high-resolution MS (e.g. ion trap – ITMS, Time of Flight – ToFMS, Fourier Transform Ion Cyclotron Resonance Mass Spectrometry – FTICR-MS) (e.g. Hamilton et al., 2005; Pol et al., 2006; Claeys et al., 2009; Yasmeen et al., 2011; Hamilton et al., 2011). Nevertheless, while solvent extraction has proven useful for subsequent quantification and precise molecular structure identification, related protocols are time-consuming, and are prone to contamination and losses, especially for the semi-volatile fraction, during the different extraction steps.

Global approaches considering gas and particulate phases simultaneously are required to elucidate secondary organic matter at a molecular scale for both phases and to provide relevant partitioning data for SVOCs. Most works aiming at studying the partitioning of SVOCs are based on off-line chromatographic techniques coupled to mass spectrometry, for both phases. This approach involves simultaneous sampling of both gas and particulate phases. One of the most common sampling methods is to trap gas-phase compounds on one or more annular glass denuders (XAD-4 coated) placed in series with a filter collecting the aerosol phase. This sampling technique has been applied without any derivatisation step prior to gas GC/MS analysis to investigate gas/particle partitioning of non-polar compounds – alkanes, PAHs (Volckens and Leith, 2003), or small carbonyls in diesel exhaust (Lee et al., 2004; Jakober et al., 2008). In more specific SOA studies targeting poly-functionalised and reactive compounds (carbonyls, hydroxyls and carboxyls), a derivatisation step has been introduced prior to the chromatographic separation. After solvent extraction of both denuder and filter, each sample is derivatised, in the liquid phase, by oxidation with pentafluorobenzylhydroxylamine (PFBHA). Extracts are reduced in volume and blown to dryness before reconstitution in a suitable solvent. Double derivatisation is then performed by additional trimethylsilylation with *N,O*-bis(trimethylsilyl)trifluoroacetamide (BSTFA). Subsequently, aliquots of the doubly-derivatised extracts are injected into the GC/MS instrument (e.g. Woo and Kim, 1999; Kalberer et al., 2000; Jaoui et al., 2003). Alternative derivati-

sation reagents have occasionally been employed, such as the  $BF_3$ -methanol reagent (Jaoui and Kamens, 2003). In spite of its powerful capacity to identify and quantify individual species in complex mixtures of poly-functionalised compounds, this procedure presents the disadvantage of solvent extraction. More recently, the same simultaneous sampling technique of both gas and particulate phases onto a denuder/filter pack has been improved for carbonyl measurements, where PFBHA is used prior to sampling to coat the XAD-4 denuder and impregnate the filter(s) (Temime et al., 2007; Healy et al., 2008, 2009; Ortiz et al., 2009). The sample treatment is thus reduced to a simple extraction of the sampling supports, denuders and filters, possibly followed by a solvent reduction step and/or a filtering step. However, this method is reserved to carbonyls and solvent extraction is still required.

In this study, we present a new analytical approach to explore, simultaneously and in parallel, the chemical composition of both gas and particulate phases, and to quantify the partitioning of individual and poly-functionalised SVOCs. Reducing both solvent use and time-consuming sample treatment steps, the new method is based on thermal desorption of both gas and particulate phases sampling supports, coupled to GC/MS. It involves on-sorbent derivatisation of gas-phase compounds and on-filter derivatisation of particulate-phase compounds.

The validation of the method is presented as well as its application to smog chamber experiments. As to the latter experiments, two series of simulated atmosphere experiments were performed in the EUPHORE European reactor. A first series focuses on limonene ozonolysis chemical system, whereas a second one focuses on the isoprene photo-oxidation chemical system. Results from these experiments are used here to illustrate the capability of the method to explore the chemical composition of the secondary organic matter in both gas and particulate phases.

## 2 Materials

### 2.1 Chemical

#### 2.1.1 Standards

The standards used for the method development were chosen in order to fulfil the following purposes: (i) represent a wide range of functionalised semi-volatile secondary organic compounds; and (ii) facilitate the identification and quantification of secondary products that are expected from limonene ozonolysis (as this system has been chosen as a model system). Each series of compounds used for the different method characterisation tests is chosen in order to meet both experimental requirements and objectives. All chemicals were purchased from Sigma-Aldrich Co. (St. Louis, USA) with the exception of 4-oxopentanal, available from Diverchim SA



### 2.2.1 Thermal desorption – gas chromatography/mass spectrometry system and procedures

The thermal desorption (TD) system was composed of (1) an Ultra 50:50 (Markes International, Llantrisant, UK) multi-tubes auto-sampler equipped with an injected sample split flow automated re-collection system, (2) a Unity 1 (Markes International) platform desorber for single tubes, and (3) an Air Server (Markes International) used in the mass flow controller mode. Samples were desorbed at 300 °C for 15 min at a flow rate of 50 ml min<sup>-1</sup> and were cryogenically trapped on-Tenax<sup>®</sup> TA (mesh 60/80) adsorbent at -10 °C. The trap was desorbed for GC injection at 300 °C, for 15 min in order to clean the trap, with a split ratio of 1:9. The “re-collection” option of the commercial thermal desorption system was used: the split flow from the injected sample is driven through the original, previously desorbed, Tenax<sup>®</sup> TA tube (in the case of a gas-phase sample) or through a freshly cleaned Tenax<sup>®</sup> TA tube (in the case of the aerosol-phase filter samples), allowing the collection, at ambient temperature, of compounds that otherwise would be lost through the vent exit to be subsequently analysed. The gas chromatography system was a 6890A type instrument from Agilent Technologies equipped with an Rxi<sup>®</sup>-5Sil MS column (60 m, 0.25 mm i.d., film thickness: 0.1 µm, Restek Corporation, Bellefonte, USA). Samples were chromatographically separated with the following temperature program: from 40 to 305 °C with a thermal ramp of 10 °C min<sup>-1</sup>. The mass spectrometry system was a 5973 type instrument from Agilent Technologies. It was equipped either with an electron ionisation (EI) source (operated at an energy of 70 eV), used in the case of the first injections of samples for structural identification and quantification, or a chemical ionisation source (CI in positive mode, methane as reagent gas), used in the case of the injection of the re-collected samples to provide complementary structural information.

### 2.2.2 Thermal desorption – gas chromatography – flame ionisation detection

The TD-GC-FID system was composed of a thermal desorber (TurboMatrix 650, Perkin Elmer, Inc.) coupled to a gas chromatograph (Clarus 500, Perkin Elmer, Inc.) and equipped with an Rxi<sup>®</sup>-5Sil MS column (60 m, 0.25 mm i.d., film thickness: 0.1 µm, Restek Corporation, Bellefonte, USA) and a flame ionisation detector. The system was also coupled to a mass spectrometer (Clarus 500, Perkin Elmer, Inc.). The analytical method was the same as that used on the Markes-Agilent TD-GC/MS system. No re-collection was performed for analytical development. The separation of the compounds was achieved with the following temperature program: from 40 to 305 °C with a thermal ramp of 10 °C min<sup>-1</sup>.

## 2.3 Simulation chambers

### 2.3.1 INERIS dynamic simulation chamber

The INERIS dynamic simulation chamber consists of a borosilicate glass loop (200 l volume) which has been described elsewhere (Martin et al., 2003; Gonzalez-Flesca and Frezier, 2005). It has been specifically designed to generate single or multi-component gaseous species atmospheres at known and stable concentrations (from hours to weeks). Clean and dry air was generated by a zero air generator (Claind, AZ Air purifier 2010) and atmospheres were generated by continuous dilution of a concentrated mixture (typically, standard pollutants gas tanks). The dilution process was controlled and monitored by mass flow meters. If required, relative humidity was generated from a pressurised bulb containing ultrapure water (18.2 MΩ, ElgaPure-Lab Flex, Veolia water) connected to a heated liquid mass flow controller system. The temperature in the chamber was maintained at around 22 °C by circulation of a temperature-conditioned mixture of water/ethylene glycol in the chamber double walls. Temperature, pressure and relative humidity were continuously monitored.

### 2.3.2 CESAM atmospheric simulation chamber

The CESAM atmospheric simulation chamber was used to perform determination of breakthrough volumes presented in Sect. 3.2.4. It has been described in detail elsewhere (Wang et al., 2011). Briefly, this facility consists of a cylindrical 4.2 m<sup>3</sup> stainless steel chamber. The chamber is evacuable using a powerful pumping system consisting of a root pump, a rotary pump and a turbomolecular pump which allows the evacuation down to 8 × 10<sup>-4</sup> mbar between each experiment to minimise the memory effects. Both temperature and relative humidity were measured with a HMP234 Vaisala<sup>®</sup> humidity and temperature transmitter equipped with a capacitive thin-film polymer sensor Humicap<sup>®</sup>.

Injection of compounds was performed by introducing a precisely known pressure (in the range 0.1 mbar) via a vacuum ramp from a frozen pure standard solution into a bulb of known volume ( $V = 1.0832$  l). The bulb content was then immediately flushed into the chamber.

Hydrocarbon products were monitored using a Fourier Transform InfraRed spectrometry (FTIR) instrument from Bruker GmbH. The total optical path length for the in-situ FTIR measurement was set to 192 m.

### 2.3.3 EUPHORE simulation chamber

Simulation chamber experiments on limonene ozonolysis (Sect. 4.) and isoprene photo-oxidation (Sect. 3.3.4) were conducted in chamber B of the EUphore PHOtoREactor (EUPHORE) facility in Valencia (Spain), consisting of ~200 m<sup>3</sup> outdoor hemispheric fluorinated ethane/propene (FEP) bags equipped with high power fans. For limonene

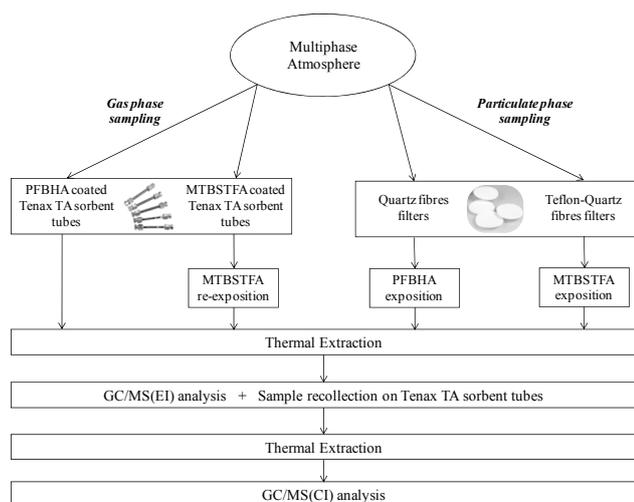


Fig. 3. Method overview.

ozonolysis experiment the chamber was protected from outdoor conditions (irradiation) by a retractable steel cover to achieve dark conditions. Before the experiment, the chamber was flushed overnight and then filled with clean air from an air purification system including an absorption dryer with molecular sieve. Air volume losses due to sampling were compensated by introduction of clean air. Prior to every experiment, an aliquot of SF<sub>6</sub> was introduced into the chamber as an inert tracer to determine the dilution rate. Limonene or isoprene was introduced using a heated clean air stream. Relative humidity was generated by water introduction through a sprayer. Humidity and temperature were monitored by a hygrometer model TS-2 (Waltz, Effeltrich, Germany). Pressure was monitored by a barometer model Air DB-VOC (Sirsa, Madrid, Spain). Ozone was produced by passing clean air (Linde, purity of 99,999 %) through a bulb equipped with a mercury UV lamp. This in-house system was previously calibrated as producing ozone with a typical rate of 20 ppb min<sup>-1</sup>, considering the volume of the chamber.

The terpene concentration was monitored by an on-line gas chromatograph coupled with a photoionisation detector (Fisons GC-8160, Thermo-Fisher, Waltham, MA, USA) and equipped with a DB-624 cyano-propylphenylpolysiloxane fused silica capillary column (J&W Scientific, 30 m, 0.32 mm id, 1.8 μm film), operated isothermally at 160 °C with a total run time of 10 min. The total aerosol mass was continuously monitored by a tapered element oscillating microbalance (TEOM, model 1400a, Ruppert and Patashnick Co. Inc, Albany, USA) with a one minute averaging time step.

### 3 Analytical development

#### 3.1 Method overview

The developed global sampling and analytical method is summarised in Fig. 3. This method is based on the simultaneous sampling of the gaseous phase on Tenax TA sorbent tubes pre-coated with one of the two derivatisation reagents (PFBHA or MTBSTFA), and the particulate phase on filters. Gaseous samples pre-coated with MTBSTFA require re-exposition to the derivatisation reagent after sampling to complete derivatisation. Particulate samples are systematically exposed to one of the two derivatisation reagents after sampling. The method involves TD-GC/MS analysis of all derivatised samples. Both EI and CI modes analyses could be performed for each sample using the sample re-collection option of the commercial thermal desorber.

For each sample type, Tenax-TA tubes and filters, PFBHA- and MTBSTFA-derivatised, the thermal desorption efficiency was evaluated for all available standard compounds for the higher mass tested (around 320 ng sample<sup>-1</sup>) by performing a second TD-GC/MS analysis just after the first one. For none of the compounds was a signal observed in the second analysis, suggesting a maximal desorption efficiency under our desorption conditions.

Laboratory blanks were also tested. Only for methylglyoxal and dimethylglyoxal on quartz filters were the observed levels significant and consequently subtracted for constructing calibration curve and estimating detection limits. However, the quartz filter blank reproducibility was not evaluated.

The developed method allows for the use of common VOCs denuders upstream of aerosol collection filters. However, in the present analytical development no denuder was tested.

#### 3.2 Carbonyl functions derivatisation in gas phase samples

For the analysis of carbonyl-bearing molecules in the gas phase, the collection of air samples collection was achieved on Tenax<sup>®</sup> TA sorbent tubes pre-coated with PFBHA. As the derivatisation reaction occurs onto the sorbent of the cartridges, which is not a classical medium for oximation with PFBHA, one of the key parts of this work was the optimisation of this process and its characterisation.

##### 3.2.1 Sorbent coating and derivatisation

Tenax<sup>®</sup> TA coating was achieved in 20 min, without using any solvent by sublimation of PFBHA: a nitrogen stream (100 ml min<sup>-1</sup> tube<sup>-1</sup>) was passed through a glass bulb that contained solid PFBHA and was connected to one or more Tenax TA sorbent tubes (max. 8 tubes, Fig. 4). A PFBHA mass of 330 μg (1.3 μmol) per tube connected was introduced. For a 12 l sample volume (corresponding to a sampling flow rate of 100 ml min<sup>-1</sup> for 2 h), this PFBHA mass

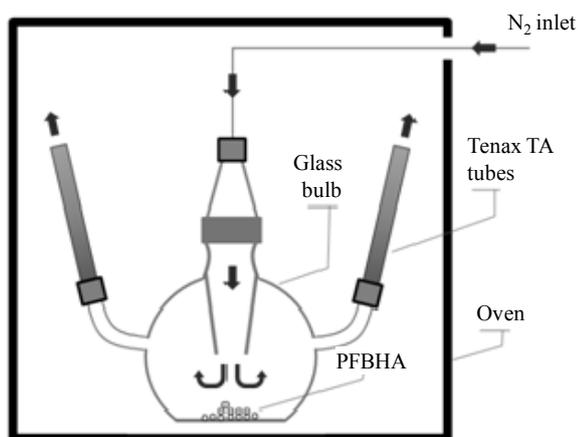


Fig. 4. The PFBHA coating process scheme for Tenax TA tubes.

is sufficient to derivatise (i.e. is 10 times higher than) a carbonyl compound at a concentration of around  $1 \text{ mg m}^{-3}$  (for an average molecular weight of 100). The glass bulb and sorbent tubes were kept at  $110^\circ\text{C}$  in an oven during the coating. Similar to Ho and Yu (2002), closed samples were stored at room temperature after sampling for a minimum of five days before TD-GC/MS analysis to allow for full derivatisation.

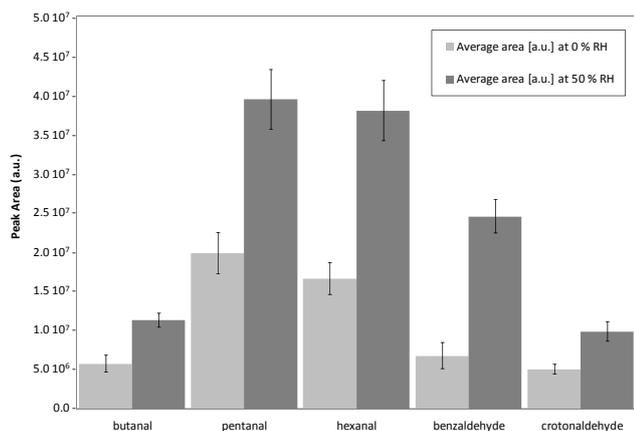
### 3.2.2 Humidity influence

For general purposes, PFBHA derivatisation is usually performed in aqueous or water containing solutions (Cancilla and Que Hee, 1992) and a molecule of water is formed for each derivatised compound (Fig. 1). A positive influence of the humidity present in the sampled air is consequently expected for the derivatisation process. On the other hand, humidity can disturb the collection of compounds by reducing the adsorption efficiency (Harper, 1993; Loedwyckx and Vansant, 2000). In order to evaluate this humidity influence a series of aldehydes was sampled from a dry (0 % RH) and wet (50 % RH) simulated atmosphere. The experiment was carried out in the INERIS dynamic simulation chamber. The system was adjusted to generate crotonaldehyde, benzaldehyde, butanal, pentanal and hexanal at concentrations of 11.7, 17.7, 12.0, 14.3, and  $16.7 \mu\text{g m}^{-3}$ , respectively. Sampling was performed during four hours at a sampling flow rate of  $150 \text{ ml min}^{-1}$ , connecting coated sorbent tubes to the chamber through a sampling cone. Five and six replicates were collected from the 0 % RH and 50 % RH atmospheres, respectively. The results are shown in Fig. 5, revealing that humidity in the sampled air promotes trapping and/or derivatisation, 1 with an average response ratio of 2.4 between wet and dry experiments (ranging from 1.9 and 3.6 depending on compounds). A complementary experiment was performed in order to evaluate bias linked with humidity variations within a realistic relative humidity range. Three sampling series of three replicates were collected at 30, 50 and 80 % RH. Sampling was performed during one hour at a sam-

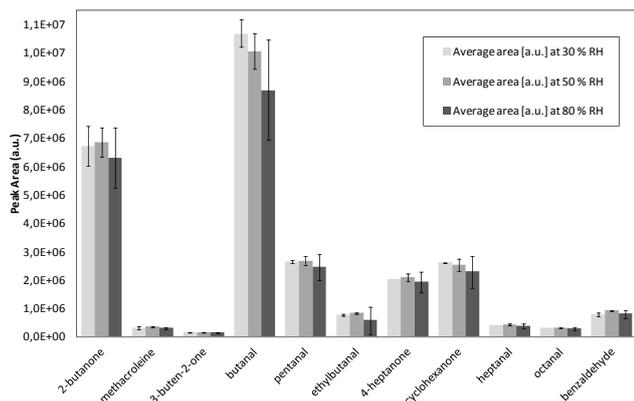
pling flow rate of  $100 \text{ ml min}^{-1}$ . The results are shown Fig. 6, revealing no significant bias linked to humidity variations. A relative humidity of 30 % appears to be sufficient to reach a maximum derivatisation yield in these conditions. However, a slight repeatability decrease is observed at 80 % RH, probably due to competition between water and adsorbed organic molecules on adsorbent surface sites. Nevertheless, this observation is consistent with the fact that high humidity levels are known to affect sampling on classic adsorbents, decreasing breakthrough volumes (Harper, 1993; Loedwyckx and Vansant, 2000). The developed method to trap and derivatise gaseous carbonyl compounds can thus be considered as suitable under realistic sampling conditions.

### 3.2.3 Derivatisation efficiency and standardisation

The derivatisation efficiency was tested by comparing in-solution derivatisation and on-Tenax<sup>®</sup> TA derivatisation. Standard solutions were prepared in a 5/95 water/acetonitrile mixture at a concentration of  $80 \text{ ng } \mu\text{l}^{-1}$  with 19 carbonyl compounds (eleven aldehydes: methacrolein, pentanal, 2-ethylbutanal, benzaldehyde, nonanal, ethyl-3-methyl-4-oxocrotonate, citronellal, citral, glyoxal, perillaldehyde and glutaraldehyde; seven ketones: butanone, 3-buten-2-one, 4-heptanone, carvone, 2,3-butadione, 3-methyl-2,4-pentadione and camphor; and one keto-aldehyde: methylglyoxal). For in-solution derivatisation, a molar excess of PFBHA corresponding to ten times the molar concentration of derivatisable functions was added to 1 ml of each solution. The solutions were kept at room temperature (around  $22^\circ\text{C}$ ) for 24 h. One  $\mu\text{l}$  of each of the five derivatised solutions was subsequently vaporised under a heated helium stream ( $270^\circ\text{C}$ ) and trapped onto five clean Tenax<sup>®</sup> TA tubes just before TD-GC-FID analysis. The repeatability of the in-solution derivatisation protocol was separately evaluated by estimating the uncertainty of the analytical response of each in-solution derivatised compound from a series of four in-solution derivatisation replicates. These uncertainties, corresponding to the standard deviation/average value ratio, were found to be satisfactory for all of the compounds, ranging from 0.3 to 29.7 %. For on-sorbent derivatisation, one  $\mu\text{l}$  of the non-derivatised solutions was injected onto PFBHA-coated Tenax<sup>®</sup> TA tubes by the same procedure. In order to expose the tubes to humidity, clean and wet (50 % RH) zero air was sampled onto the tubes during 30 min at a flow rate of  $100 \text{ ml min}^{-1}$ . Samples were kept at room temperature for 5 days before TD-GC-FID analysis. Two on-Tenax TA derivatisations per solution were performed. The comparison of the results is shown in Fig. 7. For aldehydes (Fig. 7a), on-Tenax TA derivatisation led to derivatisation efficiency close to or higher than in-solution derivatisation (ratio ranging from 102 to 205 %), except for methacrolein (9 %). For ketones (Fig. 7b), on-Tenax TA derivatisation led to a derivatisation efficiency lower than for in-solution derivatisation (ratio ranging from 12 to 41 %) (derivatised



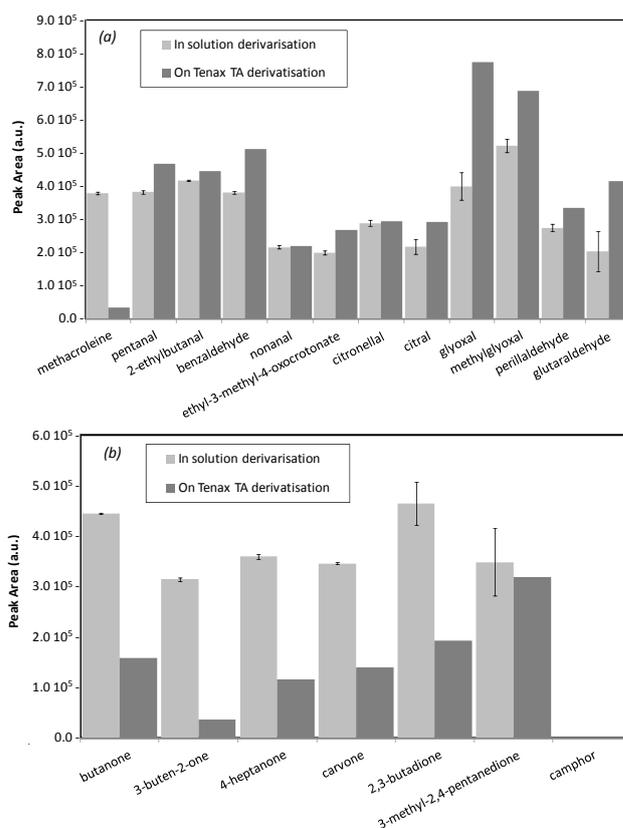
**Fig. 5.** Gaseous carbonyl compounds sampling on simulated atmosphere in dry (0 % RH) and wet (50 % RH) conditions. “Peak Area” refers to the total ion chromatogram obtained in the electron ionisation mode, in area units. Error bars represent  $\pm$  one standard deviation determined from six replicates.



**Fig. 6.** Gaseous carbonyl compounds sampling on simulated atmosphere in realistic relative humidity range (30, 50 and 80 % RH). “Peak Area” refers to the extracted ion chromatogram obtained in the electron ionisation mode, in area units (ion extraction depends on compound). Error bars represent  $\pm$  one standard deviation determined from three replicates.

camphor was not observed in both derivatisation processes). Consequently, even if this test does not allow to determine relevant on-Tenax TA derivatisation yields, it points out the fact that the two procedures (in-solution and on-Tenax TA derivatisations) are not equivalent.

Nevertheless, based on this experiment and previous ones (see Sect. 2.2.2), the repeatability of on-Tenax TA derivatisation was estimated from the standard deviation/mean ratio and determined to be equal to or lower than 10 % for compounds with more than 4 carbons and lower than 20 % for compounds with 4 carbons or less (except for the 80 % RH experiment). Thus, despite the low derivatisation yield for some compounds, a satisfactory on-Tenax TA derivatisation repeatability was observed, allowing quantification.



**Fig. 7.** Comparison of results for in-solution and on-Tenax TA carbonyl compound PFBHA derivatisation: (a) aldehydes derivatisation results, (b) ketone derivatisation results. “Peak Area” refers to FID chromatogram, in area units. Error bars represent  $\pm$  one standard deviation determined from four replicates of in-solution derivatisation. On-Tenax TA derivatisation was performed twice. Derivatised camphor was not detected by both methods.

Consequently, a calibration curve for quantification cannot be based on standards prepared from in-solution derivatisation, but has to be made by preparing standards that reproduce real sampling conditions with the method described just above and used for estimating the on-sorbent derivatisation yield estimation (by injecting non-derivatised standard solutions onto PFBHA-coated Tenax TA tubes and exposing them to a wet zero air stream to adjust humidity).

### 3.2.4 Breakthrough volumes

Experiments to determine breakthrough volumes were performed in the CESAM smog chamber. The relative humidity was maintained between 65 and 80 % as it was assumed that a high relative humidity favours breakthrough and consequently may lead to the determination of a lower limit of breakthrough volumes. A temperature of 23 °C remained constant during the experiment. Samplings were performed on five sampling trains equipped with two in series identical PFBHA-coated sorbent tubes and operated at a flow rate

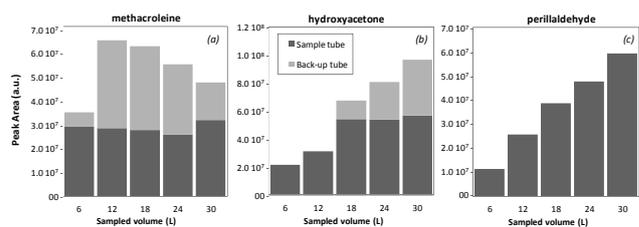
of  $100 \text{ ml min}^{-1}$ . The experiment was performed within five hours. One sampling was stopped each hour in order to test five sampling volumes.

The breakthrough volume is defined as the air volume required to completely elute a compound through an adsorbent tube (Baltussen et al., 2002). It defines a limit beyond which the trapping is no longer fully efficient and where losses may be encountered. It is defined for a given compound on a given mass of adsorbent and in given environmental conditions of temperature and relative humidity (CEN, 2005). Breakthrough volumes for PFBHA-coated Tenax<sup>®</sup> TA sorbent tubes were evaluated for a series of five carbonyl compounds. A mixture of methacrolein, hydroxyacetone, octanal, benzaldehyde, carvone and perillaldehyde was injected in the CESAM simulation chamber with respective concentrations of 71, 78, 121, 100, 137 and  $152 \mu\text{g m}^{-3}$ . Absolute quantification of these species was achieved by the method used for their injection: for each of them, a precisely known pressure (in the range 0.1 mbar) was applied via a vacuum ramp from a frozen pure standard solution into a bulb of known volume ( $V = 1.0832 \text{ l}$ ). The bulb content was then immediately flushed into the chamber.

For a given compound, the breakthrough volume is reached when measurable quantities of it are found in the second coated tube (back-up tube). Results are shown in Fig. 8. For most of the compounds the breakthrough was found to be satisfactory as it fell in the range needed to sample organics in air. For methacrolein, the breakthrough volume was reached within the first hour, corresponding to a sampling volume smaller than 6 l. It was reached between the first and the second hour for hydroxyacetone, corresponding to a breakthrough volume between 6 and 12 l. Nevertheless, for the other compounds – octanal, benzaldehyde, carvone and perillaldehyde – the breakthrough volumes were not reached within five hours, corresponding to a breakthrough volume higher than 30 l (results are only shown for perillaldehyde, Fig. 8c). Consequently, when the method was applied, a sampling time between one and five hours was used, depending on the atmosphere concentrations but keeping in mind that carbonyls with a number of carbon inferior to five are subjected to breakthrough, i.e. employing back-up tubes if  $\text{C}_1$ – $\text{C}_4$  carbonyl compounds require quantification.

### 3.2.5 Linearity and detection limits

Both linearity and detection limits were evaluated from calibration curves. A dilution series of five carbonyl compounds solutions (nine aldehydes and eight ketones) from 5 to  $320 \text{ ng } \mu\text{l}^{-1}$  was used. Calibration curves were performed from standards reproducing real on-Tenax TA sampling conditions (see Sect. 3.2.3). Results are given in Table 1, including calibration curve slopes, good correlation coefficients (ranging from 0.9883 to 0.9999) and satisfactory detection limits. The response linearity was evaluated from a specific  $m/z$  ion extraction for each compound and was statistically



**Fig. 8.** Evaluation of breakthrough volumes for carbonyl compounds on PFBHA-coated Tenax TA sorbent tubes (a) methacroleine, (b) hydroxyacetone (c) perillaldehyde. 3 “Peak Area” refers to the  $m/z$  181 extracted ion chromatogram obtained in the electron ionisation mode, in area units.

validated (Student’s t-test). For the negative intercepts obtained for two compounds (citronellal and citral), no explanation could be found. Uncertainties (corresponding to  $\pm 2$  standard errors) are given for a 95 % confidence level. Detection limits (corresponding to 3 times the area noise) are given in absolute mass per sample tube and in concentration units for a sampling volume of 12 l (i.e.  $100 \text{ ml min}^{-1}$  for 2 h). They range from 0.07 to 6 ng per tube, corresponding to concentrations ranging from 6 to  $500 \text{ ng m}^{-3}$  (or 1.3 to 81 ppt). These detection limits are in the same order of magnitude or better than those obtained by Ho and Yu (2004) and Li et al. (2009), who propose sampling of aldehydes onto PFPH (pentafluorophenyl hydrazine)-coated Tenax TA tubes and obtained detection limits ranging from 0.07 to 6 ng sampled per tube (corresponding to atmospheric concentrations ranging from 6 to  $500 \text{ ng m}^{-3}$  for a 12 l sample volume).

Compared to the classical 2,4-dinitrophenylhydrazine (DNPH) method involving collection of carbonyls onto DNPH-coated silica cartridges before elution and analysis by HPLC-UV, the detection limits obtained in the present study correspond to a gain of around one order of magnitude (Ho and Yu, 2004). They are in the same order of magnitude or higher than those obtained by Chi et al. (2007), which range from 2 to  $9 \text{ ng m}^{-3}$ , for a series of mono-carbonyl compounds, by DNPH derivatisation and LC-ESI-MS/MS analysis. The detection limits in the ppt range provided by the method are nevertheless more than satisfactory, considering the concentration levels of carbonyls which can be reached in the atmosphere. As an example, Müller et al. (2006) found concentrations of biogenic carbonyl compounds in a forest region to be about hundreds of ppt.

### 3.3 Derivatisation of carboxylic and hydroxyl functions in gas-phase samples

For the analysis of hydroxyl-bearing gaseous molecules, collection of the gas phase was performed on Tenax<sup>®</sup> TA sorbent tubes pre-coated with MTBSTFA reagent in order to analyse the carboxyl- and hydroxyl-containing fraction of the gaseous organic matter. This reaction requires significantly different experimental conditions compared to derivatisation

**Table 1.** Calibration curves slopes, linearities and detection limits for gaseous carbonyl compounds trapped on PFBHA-coated Tenax TA tubes.

| Standard Compound             | Extracted ion ( <i>m/z</i> ) | Calibration curve slope       | Y-intercept                  | Linearity ( <i>R</i> <sup>2</sup> ) | Detection limits      |                        |         |
|-------------------------------|------------------------------|-------------------------------|------------------------------|-------------------------------------|-----------------------|------------------------|---------|
|                               |                              |                               |                              |                                     | ng tube <sup>-1</sup> | ng m <sup>-3</sup> (*) | ppt (*) |
| <b>Aldehydes</b>              |                              |                               |                              |                                     |                       |                        |         |
| pentanal                      | 181                          | (6.8 ± 0.2) 10 <sup>5</sup>   |                              | 0.9993                              | 0.3                   | 25                     | 7.1     |
| 2-ethylbutanal                | 239                          | (1.11 ± 0.03) 10 <sup>5</sup> |                              | 0.9995                              | 0.1                   | 9                      | 2.2     |
| 3-methyl-2-butenal            | 264                          | (1.03 ± 0.01) 10 <sup>5</sup> |                              | 0.9999                              | 0.07                  | 6                      | 1.8     |
| heptanal                      | 181                          | (5.9 ± 0.1) 10 <sup>5</sup>   |                              | 0.9998                              | 0.3                   | 25                     | 5.4     |
| ethyl-3-methyl-4-oxocrotonate | 140                          | (1.97 ± 0.02) 10 <sup>4</sup> |                              | 0.9999                              | 0.3                   | 25                     | 4.3     |
| citronellal                   | 168                          | (7.01 ± 0.05) 10 <sup>4</sup> | -(1.8 ± 0.7) 10 <sup>5</sup> | 0.9997                              | 3                     | 250                    | 40      |
| citral                        | 69                           | (1.88 ± 0.03) 10 <sup>5</sup> | -(1.0 ± 0.5) 10 <sup>6</sup> | 0.9990                              | 6                     | 500                    | 80      |
| perillaldehyde                | 164                          | (6.92 ± 0.06) 10 <sup>4</sup> |                              | 0.9998                              | 0.1                   | 9                      | 1.5     |
| glutaraldehyde                | 181                          | (9.7 ± 0.6) 10 <sup>5</sup>   |                              | 0.9952                              | 0.1                   | 9                      | 2.2     |
| <b>Ketones</b>                |                              |                               |                              |                                     |                       |                        |         |
| 3-penten-2-one                | 264                          | (3.21 ± 0.07) 10 <sup>4</sup> |                              | 0.9997                              | 0.1                   | 9                      | 2.6     |
| 3-hydroxy-3-methyl-2-butanone | 181                          | (1.1 ± 0.2) 10 <sup>5</sup>   |                              | 0.9883                              | 1                     | 84                     | 20      |
| 2-hexanone                    | 181                          | (7.5 ± 0.6) 10 <sup>4</sup>   |                              | 0.9962                              | 2                     | 167                    | 41      |
| 4-heptanone                   | 128                          | (1.08 ± 0.05) 10 <sup>4</sup> |                              | 0.9988                              | 0.9                   | 75                     | 16      |
| 5-hydroxy-2-pentanone         | 279                          | (1.89 ± 0.04) 10 <sup>3</sup> |                              | 0.9992                              | 2                     | 167                    | 40      |
| (+)-dihydrocarvone            | 93                           | (3.91 ± 0.08) 10 <sup>4</sup> |                              | 0.9997                              | 1                     | 84                     | 14      |
| carvone                       | 304                          | (4.0 ± 0.4) 10 <sup>4</sup>   |                              | 0.9914                              | 0.09                  | 8                      | 1.3     |
| dimethylglyoxal               | 476                          | (6.6 ± 0.4) 10 <sup>2</sup>   |                              | 0.9970                              | 7.0                   | 584                    | 166     |
| 3-methyl-2,4-pentanedione     | 181                          | (4.2 ± 0.2) 10 <sup>5</sup>   |                              | 0.9983                              | 0.4                   | 34                     | 7.3     |
| <b>Keto-aldehydes</b>         |                              |                               |                              |                                     |                       |                        |         |
| methylglyoxal                 | 181                          | (1.7 ± 0.4) 10 <sup>6</sup>   |                              | 0.9742                              | 0.05                  | 5                      | 1.7     |
| 4-oxopentanal                 | 181                          | (6.7 ± 0.2) 10 <sup>5</sup>   |                              | 0.9997                              | 0.02                  | 2                      | 0.5     |

(\*) Given 1 for a sampling volume of 12l.

with PFBHA and is known to be sensitive to different parameters. In particular, it is sensitive to humidity and, hence, required the development of a specific protocol.

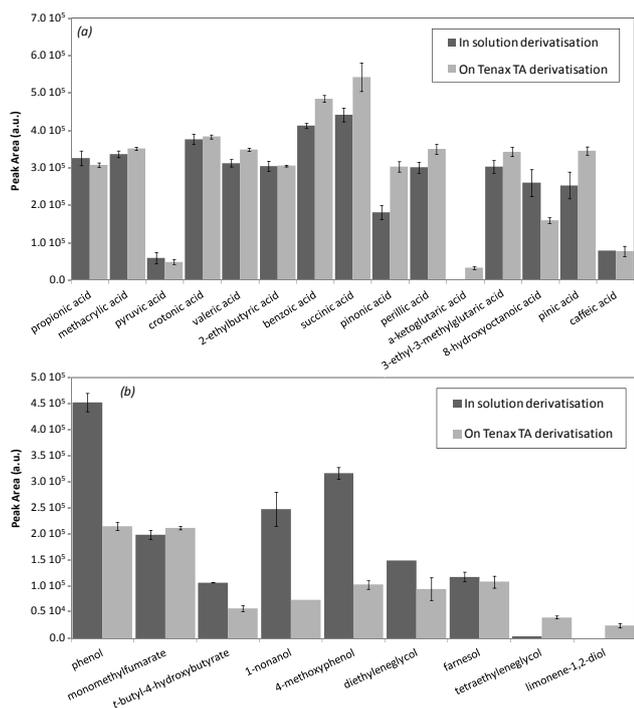
### 3.3.1 Sorbent coating and post-sampling derivatisation protocol

Similar to the PFBHA procedure, Tenax TA coating with MTBSTFA was performed without any solvent by vaporisation of 0.3 µl pure MTBSTFA (+1 % TBDMS) under a heated helium stream (270 °C, 30 ml min<sup>-1</sup>) for only 8 min. A dry purge of 2 min was applied before injection of MTBSTFA to remove traces of water in the sorbent tube. However, as previously mentioned, MTBSTFA and MTBSTFA-derivatised compounds are prone to hydrolysis, especially during sampling of humid ambient or simulated atmospheres. Consequently, in order to promote derivatisation, and considering the hydrophobic nature of Tenax TA, samples were re-exposed to MTBSTFA after sampling: samples were opened, 0.3 µl of pure MTBSTFA (+1 % TBDMS) was applied at each end of the sampling tubes and the re-closed samples were left in a furnace at 60 °C for 5 h. After

cooling to room temperature, the samples had to be rapidly analysed by TD/GC/MS within 5 h to prevent decomposition of the derivatives.

### 3.3.2 Derivatisation yield and standards preparation

Derivatisation yields were estimated comparing in-solution derivatisation and on-Tenax TA derivatisation. In-solution derivatisation was performed at room temperature within two hours by adding a molar excess of MTBSTFA (corresponding to ten times the molar concentration of derivatisable functions) to the non-derivatised solution in acetonitrile (80 ng µl<sup>-1</sup>). Derivatised solutions were then evaporated under a heated helium stream and trapped onto clean Tenax TA tubes before TD-GC-FID analysis. On-Tenax TA derivatisation was performed by preparing standard Tenax TA tubes. Clean Tenax TA tubes were spiked with the non-derivatised solution directly onto the Tenax TA adsorbent with a microsyringe. Spiked tubes were placed 3 min under a heated helium stream (270 °C, 30 ml min<sup>-1</sup>) in order to elute compounds on the Tenax surface and better represent a real sampling. Tubes, thereafter referred as carboxyls/hydroxyls

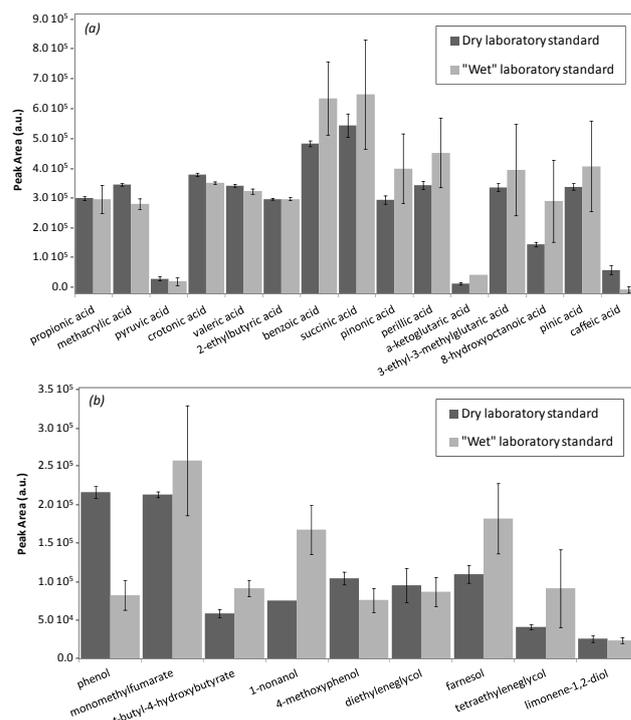


**Fig. 9.** Comparison of in-solution and on-Tenax TA carboxyl/hydroxyl compound MTBSTFA-derivatisation: **(a)** carboxylic acids derivatisation results, **(b)** hydroxyl compounds derivatisation results. “Peak Area” refers to FID chromatogram, in area units. Error bars represent  $\pm$  one standard deviation determined from three replicates for on-Tenax TA derivatisation and from two replicates for in-solution derivatisation (no error bar indicates only one replicate).

standard tubes, were then derivatised as previously presented for samples (see Sect. 3.3.1) and analysed by TD-GC-FID. The comparison of the results is shown in Fig. 9. A good agreement was found between in-solution and on-Tenax TA derivatisation yields for the acids, whereas significant discrepancies were observed for hydroxyl-bearing species. This is especially true for some alcohols such as phenol, *t*-butyl-4-hydroxy butyrate, 1-nonanol, 4-methoxyphenol – for which decreases larger than 45 % in derivatisation yield were observed.

Thus, despite the low derivatisation yields for some compounds, satisfactory on-Tenax TA derivatisation repeatability was observed, allowing quantification.

Consequently, calibration curves for quantification cannot be based on standards prepared from in-solution derivatisation, but instead have to be constructed by preparing standards reproducing real sampling conditions with the method described just above and used for estimating the on-sorbent derivatisation yield.



**Fig. 10.** Humidity influence on on-Tenax TA carboxyl/hydroxyl compounds MTBSTFA derivatisation: **(a)** carboxylic acids derivatisation results, **(b)** hydroxyl compounds derivatisation results. “Peak Area” refers to FID chromatogram, in area units. Error bars represent  $\pm$  one standard deviation determined from three replicates.

### 3.3.3 Humidity influence

The influence of humidity on the derivatisation process was evaluated by humidifying carboxyl/hydroxyl standard tubes before the post-derivatisation process. Two series of three carboxyls/hydroxyls standard tubes were prepared by spiking one  $\mu\text{l}$  of a non-derivatised solution at a concentration of  $80 \text{ ng } \mu\text{l}^{-1}$ . Wet zero air (50 % RH) was sampled for thirty minutes at a flow rate of  $100 \text{ ml min}^{-1}$  for the first series, while the second series remained closed and in dry conditions. After the post-derivatisation process, both series were analysed by TD-GC-FID. The results are shown in Fig. 10, revealing no significant discrepancies between wet and dry standards (except for nonanol and phenol) despite a lower repeatability for some compounds in wet conditions. Consequently, the post-derivatisation process is assumed to overcome the influence of humidity on the analytical protocol. Hence, the calibration can be performed using dry carboxyls/hydroxyls standard tubes.

From the humidified series of three standards, an estimation of the method repeatability can be given. The mean standard deviation/mean ratio was 25 % for all compounds, ranging from 12 (*t*-butyl-4-hydroxybutyrate) to 55 % (ethylene glycol) for alcohols and from 1 (crotonic acid) to 79 % (caffeic acid) for carboxylic acids.

**Table 2.** Breakthrough data, in percentage corresponding to the back-up tube/sample tube ratio for the quantified compound, for a series of hydroxyl 1 and carboxyl compounds identified from an isoprene photo-oxidation experiment vs. compound type, molar mass, MTBSTFA-derivatised functional groups and retention times (indicatives).

| Identified compound                               | Compound type                       | Molecular Weight (MTBSTFA-derivatised functional groups) | RT (min) | Breakthrough (%) |
|---------------------------------------------------|-------------------------------------|----------------------------------------------------------|----------|------------------|
| methacrylic acid                                  | C <sub>4</sub> mono acid            | 86 (1)                                                   | 11,87    | 6                |
| crotonic acid                                     | C <sub>4</sub> mono acid            | 86 (1)                                                   | 12,80    | 25               |
| valeric acid                                      | C <sub>5</sub> mono acid            | 102 (1)                                                  | 13,36    | 3                |
| 3-ethylbutyric acid                               | C <sub>6</sub> mono acid            | 116 (1)                                                  | 13,75    | 21               |
| 3-methylpentanoic acid                            | C <sub>6</sub> mono acid            | 116 (1)                                                  | 14,68    | 52               |
| levulinic acid                                    | C <sub>5</sub> keto mono acid       | 116 (1)                                                  | 15,58    | <1               |
| heptanoic acid                                    | C <sub>7</sub> mono acid            | 130 (1)                                                  | 15,99    | 0                |
| 1-methylcyclohexanecarboxylic acid <sup>(*)</sup> | C <sub>8</sub> mono acid            | 142 (1)                                                  | 16,10    | 0                |
| glycolic acid                                     | C <sub>2</sub> hydroxy acid         | 76 (2)                                                   | 16,37    | <1               |
| 2-hydroxypropanoic acid                           | C <sub>3</sub> hydroxy acid         | 90 (2)                                                   | 17,25    | <1               |
| oxalic acid                                       | C <sub>2</sub> di acid              | 90 (2)                                                   | 17,92    | <1               |
| maleic acid                                       | C <sub>4</sub> di acid              | 116 (2)                                                  | 20,25    | 0                |
| succinic acid                                     | C <sub>4</sub> di acid              | 118 (2)                                                  | 20,41    | <1               |
| glutaric acid                                     | C <sub>5</sub> di acid              | 132 (2)                                                  | 21,45    | 0                |
| glycerol                                          | C <sub>3</sub> tri alcohol          | 92 (3)                                                   | 21,59    | 0                |
| 2,3-dihydroxypropanoic acid                       | C <sub>3</sub> di hydroxy mono acid | 106 (3)                                                  | 22,34    | 0                |
| adipic acid                                       | C <sub>6</sub> di acid              | 146 (2)                                                  | 22,56    | 0                |
| suberic acid                                      | C <sub>8</sub> di acid              | 174 (2)                                                  | 24,48    | 0                |
| 2-hydroxypentanedioic acid                        | C <sub>5</sub> hydroxy di acid      | 148 (3)                                                  | 31,23    | 0                |

(\*) Identified from a limonene ozonolysis experiment (see Sect. 4).

### 3.3.4 Breakthrough volumes

Given the difficulty to generate stable atmospheres of hydroxyl and carboxyl compounds with low vapour pressures, precise breakthrough volumes could not be determined. An evaluation of the breakthrough of MTBSTFA-coated sorbent tubes is given here as breakthrough data for a series of carboxyl and hydroxyl compounds generated in an isoprene photo-oxidation experiment. This experiment was selected because it produced a wider range of volatile oxygenated species compared to the limonene ozonolysis experiment, and was conducted in the EUPHORE chamber. The initial isoprene concentration was 55 ppb for an initial H<sub>2</sub>O<sub>2</sub> concentration of 4 ppm. The relative humidity and temperature were 25 % and 303 K, respectively, during sampling time, after exposing the chamber to sunlight and system stabilisation. Sampling was performed onto two in series MTBSTFA pre-coated Tenax TA sorbent tubes at a flow rate of 100 ml min<sup>-1</sup> during two hours (sampled volume: 12 l). The second tube (following air flux) was employed as a “back-up tube” to collect the compounds that were not completely trapped onto the first one. The results are shown in Table 2, which gives an evaluation of the breakthrough in %, corresponding to the ratio between the amount of compound measured in the back-up tube and that measured in the sample tube for each examined compounds. No breakthrough could be observed for

most examined compounds, especially for those bearing several functionalities and/or displaying a carbon skeleton with more than five carbons.

It must be pointed out that the chamber background presented contaminations, which were responsible for the detection of compounds with more than five carbons from this isoprene photo-oxidation experiment, were not subtracted here since the purpose was to determine the breakthrough volume and not to explain isoprene oxidation pathways. All the detected compounds were consequently taken into account for the evaluation of the breakthrough, contaminations and isoprene oxidation products.

Significant breakthrough was only observed for mono-functionalised compounds with a carbon number less than seven. If these compounds are targeted, the use of back-up tubes is thus required. Poly-functionalised compounds, even with only two carbons, do not seem to be affected by breakthrough under our experimental conditions. Their breakthrough volumes are consequently estimated to be greater than 12 l. When performing measurements in the same sampling conditions of sample volume, sampling flow rate, humidity and temperature, the use of back-up tubes is thus not required. When performing measurements in different conditions, the determination of breakthrough volumes or the use of back-up tubes is recommended.

**Table 3.** Calibration curves slopes, linearities and detection limits for gaseous carbonyl compounds trapped on PFBHA-coated Tenax TA tubes.

| Standard Compound                    | Extracted ion ( <i>m/z</i> ) | Calibration curve slope       | Linearity ( $R^2$ ) | Detection limits      |                        |         |
|--------------------------------------|------------------------------|-------------------------------|---------------------|-----------------------|------------------------|---------|
|                                      |                              |                               |                     | ng tube <sup>-1</sup> | ng m <sup>-3</sup> (*) | ppt (*) |
| Alcohols                             |                              |                               |                     |                       |                        |         |
| 5-hydroxy-2-pentanone                | 75                           | $(7 \pm 2) \times 10^4$       | 0.9769              | 0.5                   | 42                     | 10      |
| hydroxycyclohexanone                 | 171                          | $(1.7 \pm 0.4) \times 10^3$   | 0.9576              | 5                     | 417                    | 89      |
| <i>t</i> -butyl-4-hydroxybutyrate    | 161                          | $(9 \pm 2) \times 10^4$       | 0.9830              | 0.2                   | 17                     | 2.6     |
| ( <i>S</i> )-3-butene-1,2-diol       | 259                          | $(9 \pm 2) \times 10^2$       | 0.9578              | 10                    | 834                    | 232     |
| 1-nonanol                            | 201                          | $(1.1 \pm 0.3) \times 10^5$   | 0.9632              | 0.08                  | 7                      | 1.2     |
| farnesol                             | 279                          | $(3.7 \pm 0.6) \times 10^3$   | 0.9840              | 3                     | 250                    | 28      |
| Carboxylic acids                     |                              |                               |                     |                       |                        |         |
| valeric acid                         | 75                           | $(2.77 \pm 0.08) \times 10^5$ | 0.9989              | 0.09                  | 8                      | 1.9     |
| levulinic acid                       | 75                           | $(1.8 \pm 0.1) \times 10^5$   | 0.9977              | 0.08                  | 7                      | 1.5     |
| heptanoic acid                       | 75                           | $(1.83 \pm 0.05) \times 10^5$ | 0.9983              | 0.05                  | 5                      | 0.9     |
| monomethyl fumarate                  | 187                          | $(3.6 \pm 0.3) \times 10^5$   | 0.9941              | 0.07                  | 6                      | 1.1     |
| 6-oxoheptanoic acid                  | 117                          | $(2.3 \pm 0.2) \times 10^5$   | 0.9926              | 0.8                   | 67                     | 11      |
| 2,2-dimethyl-3-hydroxypropionic acid | 147                          | $(1.2 \pm 0.2) \times 10^5$   | 0.9811              | 0.5                   | 42                     | 8.7     |
| succinic acid                        | 289                          | $(3.4 \pm 0.2) \times 10^5$   | 0.9956              | 0.07                  | 6                      | 1.2     |
| cis-pinonic acid                     | 171                          | $(3.2 \pm 0.3) \times 10^5$   | 0.9941              | 0.2                   | 17                     | 2.3     |
| perillic acid                        | 223                          | $(3.9 \pm 0.3) \times 10^5$   | 0.9947              | 0.09                  | 8                      | 1.2     |
| 8-hydroxyoctanoic acid               | 331                          | $(5.6 \pm 0.7) \times 10^4$   | 0.9861              | 0.2                   | 17                     | 2.6     |
| D-malic acid                         | 419                          | $(1.1 \pm 0.1) \times 10^5$   | 0.9872              | 0.8                   | 67                     | 12      |
| pinic acid                           | 171                          | $(2.7 \pm 0.2) \times 10^5$   | 0.9913              | 0.3                   | 25                     | 3.3     |
| 4-oxoheptanedioic acid               | 345                          | $(1.9 \pm 0.3) \times 10^5$   | 0.9746              | 2                     | 167                    | 23      |
| 3-carboxyhexanedioic acid            | 475                          | $(5 \pm 2) \times 10^4$       | 0.9368              | 2                     | 167                    | 21      |

(\*) Given for a sampling volume of 121.

### 3.3.5 Linearity and detection limits

Both linearity and detection limits were evaluated from calibration curves. A dilution series of five hydroxyl compounds and organic acid solutions (six alcohols and fourteen organic acids) from 5 to 320 ng  $\mu\text{l}^{-1}$  was used. Calibration curves were performed from standards reproducing real on-Tenax TA sampling conditions (see Sect. 3.3.2). 1  $\mu\text{l}$  of carbonyl compound standard solutions were applied onto Tenax TA tubes and subsequently derivatised with MTBSTFA at 60 °C for 5 h. Results are shown in Table 3. Response linearity for all compounds was statistically validated (Student's *t*-test) and uncertainties (corresponding to  $\pm 2$  standard errors) are given for a 95 % confidence level. Detection limits (corresponding to 3 times the area noise) are given in absolute mass per sample tube and in concentration units for a sampling volume of 121 (i.e. 100 ml  $\text{min}^{-1}$  for 2 h). They range from 0.05 to 10 ng per tube and from 5 to 830 ng  $\text{m}^{-3}$ . These detection limits are lower than those obtained by Wu et al. (2008) for the analysis of volatile fatty acids without derivatisation and in the same order of magnitude or lower than those obtained by Chiappini et al. (2006) or Pietrogrande et al. (2010). The detection limits in the latter studies

ranged from 1.2 to 6.8 ng per GC/MS injection for mono- and dicarboxylic acids in the aerosol phase with prior derivatisation in CO<sub>2</sub> supercritical fluid or in solution. The detection limits obtained in the present study, ranging from 1 to 232 ppt, are more than satisfactory to measure hydroxyl and carbonyl compounds at lower ambient concentration levels. As an example, the concentration levels of dicarboxylic acids measured by Legrand et al. (2005) at altitude sites in France are a few tens of ppb.

### 3.4 Derivatisation of carbonyl functions in particulate phase samples

Collection of the particulate phase for the analysis of carbonyl compounds was carried out on 47 mm quartz fibre filters (see Sect. 2.1.4) at a typical flow rate of 1  $\text{m}^3 \text{h}^{-1}$  (16.71  $\text{min}^{-1}$ ) and was done simultaneously with sampling of the gas phase. The PFBHA post-sampling derivatisation process was performed by sample introduction into stainless-steel tubes suitable for thermal desorption and exposition of the sample to the PFBHA reagent. The humidity influence on the particulate-phase derivatisation process

**Table 4.** Calibration curves slopes, linearities and detection limits for particulate carbonyl compounds trapped on quartz fibres filters and PFBHA-derivatised.

| Standard Compound             | Extracted<br>ion ( <i>m/z</i> ) | Calibration<br>curve slope    | Y-intercept                  | Linearity<br>( <i>R</i> <sup>2</sup> ) | Detection limit         |                        |
|-------------------------------|---------------------------------|-------------------------------|------------------------------|----------------------------------------|-------------------------|------------------------|
|                               |                                 |                               |                              |                                        | ng filter <sup>-1</sup> | ng m <sup>-3</sup> (*) |
| <b>Aldehydes</b>              |                                 |                               |                              |                                        |                         |                        |
| pentanal                      | 181                             | $(4.9 \pm 0.2) \times 10^5$   |                              | 0.9986                                 | 0.2                     | 0.1                    |
| 2-ethylbutanal                | 239                             | $(7.5 \pm 0.07) \times 10^4$  |                              | 0.9999                                 | 0.04                    | 0.02                   |
| 3-methyl-2-butenal            | 264                             | $(4.3 \pm 0.2) \times 10^4$   |                              | 0.9994                                 | 0.09                    | 0.05                   |
| heptanal                      | 181                             | $(2.4 \pm 0.2) \times 10^4$   |                              | 0.9968                                 | 3                       | 2                      |
| ethyl-3-methyl-4-oxocrotonate | 140                             | $(7 \pm 3) \times 10^3$       |                              | 0.8625                                 | 0.9                     | 0.5                    |
| citronellal                   | 168                             | $(7 \pm 2) \times 10^3$       |                              | 0.9532                                 | 6                       | 3                      |
| citral                        | 69                              | $(1.2 \pm 0.4) \times 10^4$   |                              | 0.9381                                 | 2                       | 1                      |
| perillaldehyde                | 164                             | $(1.0 \pm 0.2) \times 10^3$   |                              | 0.9793                                 | 5                       | 3                      |
| glutaraldehyde                | 181                             | $(4 \pm 3) \times 10^3$       |                              | 0.7846                                 | 15                      | 8                      |
| <b>Ketones</b>                |                                 |                               |                              |                                        |                         |                        |
| 3-penten-2-one                | 264                             | $(2.1 \pm 0.8) \times 10^4$   |                              | 0.8702                                 | 0.2                     | 0.1                    |
| 3-hydroxy-3-methyl-2-butanone | 181                             | $(1.0 \pm 0.4) \times 10^4$   |                              | 0.889                                  | 4                       | 2                      |
| 2-hexanone                    | 181                             | $(1.07 \pm 0.02) \times 10^5$ |                              | 0.9998                                 | 0.4                     | 0.2                    |
| 4-heptanone                   | 128                             | $(2.72 \pm 0.06) \times 10^4$ |                              | 0.9996                                 | 0.2                     | 0.1                    |
| 5-hydroxy-2-pentanone         | 279                             | $(8.8 \pm 0.2) \times 10^1$   | $-(7.0 \pm 0.4) \times 10^3$ | 0.9999                                 | 120                     | 60                     |
| (+)-dihydrocarvone            | 93                              | $(6.3 \pm 0.4) \times 10^3$   | $-(1.2 \pm 0.7) \times 10^5$ | 0.9999                                 | 23                      | 12                     |
| carvone                       | 304                             | $(1.4 \pm 0.2) \times 10^4$   |                              | 0.9848                                 | 0.2                     | 0.1                    |
| dimethylglyoxal               | 476                             | $(2.1 \pm 0.3) \times 10^4$   |                              | 0.9931                                 | 0.2                     | 0.1                    |
| 3-methyl-2,4-pentanedione     | 181                             | $(6 \pm 2) \times 10^4$       |                              | 0.9131                                 | 2                       | 1                      |
| <b>Keto-aldehydes</b>         |                                 |                               |                              |                                        |                         |                        |
| methylglyoxal                 | 181                             | $(5 \pm 2) \times 10^5$       |                              | 0.9811                                 | 0.08                    | 0.04                   |
| 4-oxopentanal                 | 181                             | $(1.79 \pm 0.07) \times 10^5$ |                              | 0.9984                                 | 0.04                    | 0.02                   |

(\*) Given for a sampling volume of 2 m<sup>3</sup>.

was expected to be similar to that observed for gas-phase derivatisation (Sect. 3.2.2).

### 3.4.1 Post-sampling derivatisation protocol

The 47 mm sample filter was divided in four parts and was introduced into a clean stainless steel thermal desorption tube. After the introduction of each quarter, 2 µl of a saturated PFBHA solution (27 mg ml<sup>-1</sup>, 90/10 acetonitrile/ultrapure water mixture) was applied onto the filter part. A glass bulb with eight outlets was then used to expose in-tube samples to a saturated PFBHA aqueous solution for 40 h. This duration was expected to be sufficient to impregnate a rolled filter with PFBHA and was convenient for protocol organisation. From one end, the filter sample tubes were connected to the bulb outlets while the other end was kept closed. 1 ml of ultrapure water was introduced in the bulb and stirred (500 rounds per min). 90 mg of pure and solid PFBHA was then added to the water. The bulb was closed and the system was kept at room temperature. After 20 h, filter sample tubes were disconnected, reversed, reconnected by the other end and left for

20 h again. Filter sample tubes were then definitely disconnected, closed and kept at room temperature for a minimum of 5 days before analysis.

### 3.4.2 Standard preparation, linearity and detection limits

Standards used to construct calibration curves were prepared and derivatised in the same way as sample filters. Clean quartz fibre filters were introduced into empty stainless steel desorption tubes with application of a PFBHA saturated solution, spiked with 1 µl of standard carbonyl compound solutions (five points from 5 to 320 ng µl<sup>-1</sup>) and left for 40 h connected to a stirred glass bulb containing 1 ml of a PFBHA saturated aqueous solution (with tube reversion after 20 h) at room temperature. Both linearity and detection limits were evaluated. Results are shown in Table 4, giving for each compound the calibration curve slopes, the correlation coefficients (which are generally good, most are ranging from 0.9131 and 0.9999, except for three compounds displaying

an  $R^2$  value ranging from 0.7846 to 0.8890), and the detection limits, which are satisfactory.

The response linearity for all compounds was statistically validated (Student's *t*-test) and uncertainties (corresponding to  $\pm 2$  standard errors) are given for a 95 % confidence level. Detection limits (corresponding to 3 times the area noise) are given in absolute mass per filter and in concentration units for a sampling volume of 2 m<sup>3</sup> (i.e. 16.71 min<sup>-1</sup> for 2 h). These detection limits, ranging from 0.04 to 120 ng per filter (corresponding to an atmospheric concentration of 0.02 and 60 ng m<sup>-3</sup> for a 2 m<sup>3</sup> sample volume), are more than sufficient to measure carbonyl compounds in the atmosphere where their concentrations are in the same order of magnitude (Liggio and McLaren, 2003).

### 3.5 Derivatisation of carboxylic and hydroxyl functions in particulate-phase samples

Collection of the particulate phase for the analysis of carboxyl and hydroxyl compounds was performed on 47 mm Teflon-quartz fibres filters (see Sect. 2.1.4) at a typical flow rate of 1 m<sup>3</sup> h<sup>-1</sup> (16.71 min<sup>-1</sup>) simultaneously with gas-phase sampling. The MTBSTFA post-sampling derivatisation procedure was performed after sample introduction into stainless-steel tubes that are suitable for thermal desorption and exposition of samples to pure MTBSTFA (+1 % TBDMS).

#### 3.5.1 Post-sampling derivatisation protocol

The 47 mm sample filter was divided in four parts and was introduced into a clean stainless steel thermal desorption tube. MTBSTFA was not applied directly onto the filter. 10  $\mu$ l of pure MTBSTFA (+1 % TBDMS) were introduced on the bottom of one tube cap and the sample filter tube was closed in vertical position above the MTBSTFA (cf. Fig. 11). The filter was exposed to MTBSTFA at room temperature during 24 h. The filter sample tube was then directly analysed by TD-GC/MS within 5 h.

#### 3.5.2 Humidity influence and standard preparation

As for gaseous compounds derivatisation on Tenax TA tubes, humidity was expected to influence the on-filter MTBSTFA derivatisation process; therefore, Teflon-quartz filters were chosen to reduce moisture loading during wet air sampling. Nevertheless, in order to evaluate the influence of this moisture loading on the derivatisation process, response factors of a series of standard compounds were compared for dry and pre-humidified Teflon-quartz filters. Five point calibration curves (from 5 to 320 ng  $\mu$ l<sup>-1</sup>) were constructed using filters spiked with standards and prepared and derivatised in the same way as sample filters. Wet Teflon-quartz filters were humidified by sampling wet zero air (50 % RH) at a flow rate of 7 l min<sup>-1</sup> for 2 h. Clean and dry or wet filters were introduced into empty stainless-steel desorption tubes and spiked

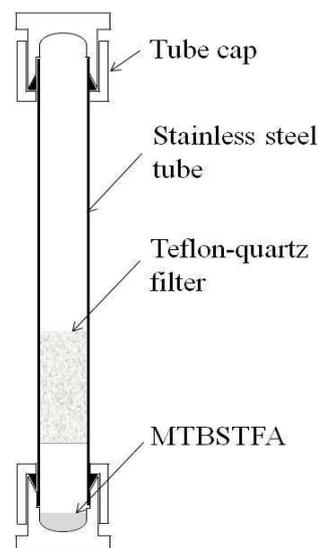


Fig. 11. Particulate-phase MTBSTFA derivatisation scheme.

with standard solutions. They were subsequently derivatised by exposition to MTBSTFA for 24 h at room temperature. The comparison of the results for compound response factors are given in Fig. 12. Surprisingly, the response factors were significantly higher for those derivatised onto humidified filters for 15 standard compounds of the 18 tested. Moreover, for keto-acids (6-oxoheptanoic acid and cis-pinonic acid), di-acids (succinic acid and pinic acid) and malic acid, the compound responses became linear compared to their responses on dry filters (not mentioned in Fig. 12). The reason why humidity positively influences on-filter derivatisation is not clear. It might be explained by competition phenomena between water and polar compounds to form bonds with the quartz surface. In this case, moisture loading may positively affect both the derivatisation reaction and the desorption capacity limiting the adsorption of polar compounds on the filter surface. Anyway, the positive influence of moisture loading onto filters has two main consequences on the method: (1) it is not necessary to dry sample filters before derivatisation with MTBSTFA, and (2) calibration curves have to be constructed using humidified Teflon-quartz filters.

#### 3.5.3 Linearity and detection limits

Both linearity and detection limits were evaluated from calibration curves that were previously used for the evaluation of the humidity influence. The results are shown in Table 5, giving for each compound the calibration curve slopes, correlation coefficients (which are satisfactory and range from 0.9433 to 0.9999), and detection limits, which are low.

Response linearity was statistically validated (Student's *t*-test) and uncertainties (corresponding to  $\pm 2$  standard errors) are given for a 95 % confidence level. Detection limits (corresponding to 3 times the area noise) are given in absolute mass

**Table 5.** Calibration curves slopes, linearities and detection limits for particulate hydroxyl compounds and organic acids trapped on Teflon-quartz 1 fibres filters and MTBSTFA-derivatised.

| Standard Compound                    | Extracted ion ( <i>m/z</i> ) | Calibration curve slope       | Linearity ( <i>R</i> <sup>2</sup> ) | Detection Limits        |                        |
|--------------------------------------|------------------------------|-------------------------------|-------------------------------------|-------------------------|------------------------|
|                                      |                              |                               |                                     | ng filter <sup>-1</sup> | ng m <sup>-3</sup> (*) |
| Alcohols                             |                              |                               |                                     |                         |                        |
| 5-hydroxy-2-pentanone                | 75                           | $(1.0 \pm 0.2) \times 10^5$   | 0.9837                              | 0.8                     | 0.4                    |
| hydroxycyclohexanone                 | 171                          | $(1.8 \pm 0.3) \times 10^4$   | 0.9791                              | 0.4                     | 0.2                    |
| <i>t</i> -butyl-4-hydroxybutyrate    | 161                          | $(2.21 \pm 0.08) \times 10^5$ | 0.9988                              | 0.04                    | 0.02                   |
| ( <i>S</i> )-3-butene-1,2-diol       | 259                          | $(8.5 \pm 0.4) \times 10^4$   | 0.9967                              | 0.04                    | 0.02                   |
| 1-nonanol <sup>a</sup>               | 201                          | $(2.9 \pm 0.5) \times 10^5$   | 0.9633                              | 0.04                    | 0.02                   |
| farnesol                             | 279                          | $(3.6 \pm 0.2) \times 10^4$   | 0.9979                              | 0.4                     | 0.2                    |
| Carboxylic acids                     |                              |                               |                                     |                         |                        |
| valeric acid                         | 75                           | $(8.6 \pm 0.3) \times 10^4$   | 0.9990                              | 0.7                     | 0.4                    |
| levulinic acid <sup>b</sup>          | 75                           | $(1.2 \pm 0.4) \times 10^5$   | 0.9433                              | 0.9                     | 0.5                    |
| heptanoic acid                       | 75                           | $(1.5 \pm 0.2) \times 10^5$   | 0.9582                              | 1.0                     | 0.5                    |
| monomethyl fumarate                  | 187                          | $(3.52 \pm 0.05) \times 10^5$ | 0.9999                              | 0.2                     | 0.1                    |
| 6-oxoheptanoic acid                  | 117                          | $(2.15 \pm 0.08) \times 10^5$ | 0.9990                              | 0.3                     | 0.2                    |
| 2,2-dimethyl-3-hydroxypropionic acid | 147                          | $(3.5 \pm 0.2) \times 10^5$   | 0.9966                              | 0.2                     | 0.1                    |
| succinic acid                        | 289                          | $(5 \pm 2) \times 10^5$       | 0.8861                              | 0.02                    | 0.01                   |
| cis-pinonic acid                     | 171                          | $(3.4 \pm 0.5) \times 10^5$   | 0.9863                              | 0.2                     | 0.1                    |
| perillic acid                        | 223                          | $(4.32 \pm 0.05) \times 10^5$ | 0.9999                              | 0.04                    | 0.02                   |
| 8-hydroxyoctanoic acid               | 331                          | $(2.33 \pm 0.07) \times 10^5$ | 0.9970                              | 0.07                    | 0.04                   |
| D-malic acid                         | 419                          | $(1.2 \pm 0.3) \times 10^5$   | 0.9623                              | 0.07                    | 0.04                   |
| pinic acid                           | 171                          | $(2.6 \pm 0.4) \times 10^5$   | 0.9887                              | 0.3                     | 0.2                    |

(\*) Given for a sampling volume of 2 m<sup>3</sup>. a: 5 points from 80 to 1280 ng. b: 4 points from 160 to 1280 ng.

**Table 6.** Characteristic fragmentation of PFBHA and MTBSTFA derivatives in EI and CI modes.

| PFBHA derivatives |                                                                    | MTBSTFA derivatives |                                                                                         | CI ions | Fragments ( <i>m/z</i> )                          |
|-------------------|--------------------------------------------------------------------|---------------------|-----------------------------------------------------------------------------------------|---------|---------------------------------------------------|
| EI ions           | Fragments ( <i>m/z</i> )                                           | EI ions             | Fragments ( <i>m/z</i> )                                                                |         |                                                   |
| 181               | [C <sub>6</sub> F <sub>5</sub> CH <sub>2</sub> ] <sup>+</sup>      | 73                  | [Si(CH <sub>3</sub> ) <sub>3</sub> ] <sup>+</sup>                                       | M + 1   | [M + H] <sup>+</sup>                              |
| M – 181           | [M – C <sub>6</sub> F <sub>5</sub> CH <sub>2</sub> ] <sup>+</sup>  | 75                  | [HOSi(CH <sub>3</sub> ) <sub>2</sub> ] <sup>+</sup>                                     | M + 29  | [M + C <sub>2</sub> H <sub>5</sub> ] <sup>+</sup> |
| M – 197           | [M – C <sub>6</sub> F <sub>5</sub> CH <sub>2</sub> O] <sup>+</sup> | M – 15              | [M – CH <sub>3</sub> ] <sup>+</sup>                                                     | M + 41  | [M + C <sub>3</sub> H <sub>5</sub> ] <sup>+</sup> |
|                   |                                                                    | M – 57              | [M – C(CH <sub>3</sub> ) <sub>3</sub> ] <sup>+</sup>                                    |         |                                                   |
|                   |                                                                    | M – 115             | [M – Si(CH <sub>3</sub> ) <sub>2</sub> C(CH <sub>3</sub> ) <sub>3</sub> ] <sup>+</sup>  |         |                                                   |
|                   |                                                                    | M – 131             | [M – OSi(CH <sub>3</sub> ) <sub>2</sub> C(CH <sub>3</sub> ) <sub>3</sub> ] <sup>+</sup> |         |                                                   |

per filter and in concentration units for a sampling volume of 2 m<sup>3</sup> (i.e. 16.7 l min<sup>-1</sup> for 2 h). The linearity was not validated for two standard compounds, i.e. 4-oxoheptanedioic acid (keto-di-acid) and 3-carboxyhexanedioic acid (tri-acid), which have, besides high detection limits, close to 100 ng per filter sample. Nevertheless, the analytical responses of these two compounds were linear, as previously tested, by on-Tenax TA derivatisation with MTBSTFA. Improvement of the developed particulate-phase MTBSTFA derivatisation protocol is consequently required to specifically quantify keto-di-acids and tri-acids. Detection limits for all other compounds range from 0.04 to 1.0 ng per filter. These detection

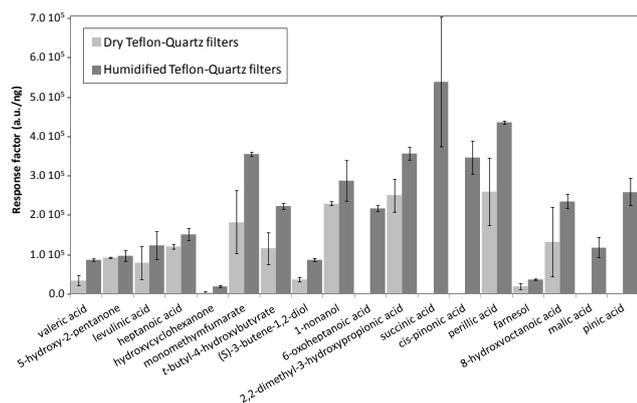
limits are significantly better than those obtained by Chiapini et al. (2006), which range from 1 to 7 ng for an analogous series of acids, keto-acids and di-acids, using derivatisation with BSTFA during supercritical fluid extraction coupled to GC/MS analysis. The detection limits are more than satisfactory to measure carboxylic acids in atmospheric particles since their concentration levels are about a few tens of ng m<sup>-3</sup> as measured by Sun and Ariya (2006).

**Table 7.** Tentative or positive identification for observed carbonyl compounds, organic acids and hydroxyls compounds from limonene ozonolysis 1 experiment. MW D.: molecular weight of the derivatised compound. MW non-D.: molecular weight of the non-derivatised compound. LOD: limit of detection. G: detected in the gas phase. P: detected in the particulate phase.

| Compound number                                       | Mass fragments                                                                                                            | Tentative assignment |                          |           |                             |                                                                                      |                            | Phase(s)                                       |     |
|-------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------|----------------------|--------------------------|-----------|-----------------------------|--------------------------------------------------------------------------------------|----------------------------|------------------------------------------------|-----|
|                                                       |                                                                                                                           | MW D.                | Functional groups number | MW non-D. | Name                        | Structure                                                                            | Assignment                 |                                                |     |
| Carbonyls compounds (PFBHA derivatives)               |                                                                                                                           |                      |                          |           |                             |                                                                                      |                            |                                                |     |
| P1                                                    | EI: 181(100), M-181=112, M=293<br>CI: M+1=294, M+29=322, M+41=334                                                         | 293                  | 1                        | 98        | 5-hexene-2-one              |    | Nist EI list               | Unique to this study                           | G-P |
| P2                                                    | EI: 42, 95(100), M-181=152, 181, M=333<br>CI: 95, 181, M+1=334, M+29=362, M+41=374                                        | 333                  | 1                        | 138       | keto-limonene               |    | Proposed from mass spectra | Forester and Wells (2009), Jaoui et al. (2006) | G   |
| P3                                                    | EI: 181(100), M-197=265, M=462<br>CI: M+1=463, M+29=491, M+41=503                                                         | 462                  | 2                        | 72        | methylglyoxal               |    | Proposed from mass spectra | Forester and Wells (2009)                      | G-P |
| P4                                                    | EI: 181(100), M-197=279, M=476<br>CI: M+1=477, M+29=505, M+41=517                                                         | 476                  | 2                        | 86        | dimethylglyoxal             |    | Standard                   | Unique to this study                           | G-P |
| P5                                                    | EI: 181(100), 279, M-197=293, M-181=309, M=490<br>CI: M+1=491, M+29=519, M+41=531                                         | 490                  | 2                        | 100       | 4-oxopentanal               |    | Standard                   | Forester and Wells (2009)                      | G-P |
| P6                                                    | EI: 181(100), 320, 558, M-197=361, M-181=377, CI: M+1=559, M+29=587, M+41=599                                             | 558                  | 2                        | 168       | limononaldehyde             |    | Proposed from mass spectra | Leungsakul et al. (2005b), Jaoui et al. (2006) | G-P |
| P7                                                    | EI: 181(100), 305, M-197=516, M=713<br>CI: <LOD                                                                           | 713                  | 3                        | 128       | dioxohexanal                |  | Proposed from mass spectra | Unique to this study                           | P   |
| P8                                                    | EI: 138, 181(100), 319<br>M-197=530, M=727<br>CI: M+1=728                                                                 | 727                  | 3                        | 142       | 3,6-dioxoheptanal           |  | Proposed from mass spectra | Leungsakul et al. (2005b)                      | P   |
| P9                                                    | EI: 94, 181(100), 347, M-197=558, M-181=574, M=755<br>CI: <LOD                                                            | 755                  | 3                        | 170       | keto-limononaldehyde        |  | Proposed from mass spectra | Leungsakul et al. (2005b), Jaoui et al. (2006) | P   |
| Hydroxyl and carboxyl compounds (MTBSTFA derivatives) |                                                                                                                           |                      |                          |           |                             |                                                                                      |                            |                                                |     |
| M1                                                    | EI: 75, 99, 145, M-15=215, M-57=173(100)<br>CI: <LOD                                                                      | 230                  | 1                        | 116       | levulinic acid              |  | Standard                   | Jaoui et al. (2006)                            | G   |
| M2                                                    | EI: 73, 116, 147, M-57=289(100), M-15=331<br>CI: <LOD                                                                     | 346                  | 2                        | 118       | succinic acid               |  | Standard                   | Jaoui et al. (2006)                            | G-P |
| M3                                                    | EI: 73(100), 123, 147, M-57=303, M-15=345<br>CI: 115, 173, 229, M-57=303, M-15=345, M+1=361, M+29=389, M+41=401           | 360                  | 2                        | 132       | 2-methylbutanedioic acid    |  | Nist EI list match 75%     | Unique to this study                           | G   |
| M4                                                    | EI: 73, 129, 147<br>M-57=303(100), M-15=345<br>CI: 115, 165, 229, M-57=303, M-15=345, M+1=361, M+29=389, M+41=401         | 360                  | 2                        | 132       | pentanedioic acid           |  | Proposed from mass spectra | Jaoui et al. (2006)                            | G-P |
| M5                                                    | EI: 73, 133, 147, 157(100), 185, 215, 289, M-57=317, M-15=359<br>CI: 215, M-57=317, M-15=359, M+1=375, M+29=403, M+41=415 | 374                  | 2                        | 146       | $\alpha$ -ketoglutaric acid |  | Standard                   | Unique to this study                           | P   |
| M6                                                    | EI: 73, 111, 141<br>M-57=317(100), M-15=359<br>CI: 257, M-57=317, M-15=359, M+1=375, M+29=403, M+41=415                   | 374                  | 2                        | 146       | hexanedioic acid            |  | Nist EI list match 92%     | Jaoui et al. (2006)                            | G-P |

Table 7. Continued.

| Compound number | Mass fragments                                                          | Tentative assignment |                          |           |                                        |                                                                                      |                            |                                                | Phase(s) |
|-----------------|-------------------------------------------------------------------------|----------------------|--------------------------|-----------|----------------------------------------|--------------------------------------------------------------------------------------|----------------------------|------------------------------------------------|----------|
|                 |                                                                         | MW D.                | Functional groups number | MW non-D. | Name                                   | Structure                                                                            | Assignment                 |                                                |          |
| M7              | EI: 73, 75, 111, 185, 317, M-57=345(100), M-15=387 CI: <LOD             | 402                  | 2                        | 174       | ketonorlimonic acid                    |    | Proposed from mass spectra | Jaoui et al. (2006)                            | P        |
| M8              | EI: 73, 75, 111, 185, 317, M-57=345(100), M-15=387 CI: <LOD             | 402                  | 2                        | 174       | 4-oxoheptanedioic acid                 |    | Standard                   | Unique to this study                           | P        |
| M9              | EI: 73, 75, 147, M-115=299, M-57=357(100), M-15=399, M=414 CI: <LOD     | 414                  | 2                        | 186       | limonic acid                           |    | Proposed from mass spectra | Jaoui et al. (2006), Leungsakul et al. (2005b) | G-P      |
| M10             | EI: 73, 75, 147, 185, 331, M-57=359(100), M-15=401 CI: <LOD             | 416                  | 2                        | 188       | ketolimonic acid                       |    | Proposed from mass spectra | Jaoui et al. (2006)                            | P        |
| M11             | EI: 73(100), 75, 99, 147, 225, 255, M-57=385, M-15=427 CI: <LOD         | 442                  | 2                        | 214       | 3-acetyl-4-methylhept-3-enedioic acid  |    | Proposed from mass spectra | Unique to this study                           | P        |
| M12             | EI: 73(100), 115, 147, 287, M-57=419, M-15=461 CI: <LOD                 | 476                  | 3                        | 134       | 2-hydroxybutanedioic acid (malic acid) |    | Standard                   | Unique to this study                           | P        |
| M13             | EI: 73(100), 133, 147, 171, 245, 273, 331, M-57=433, M-15=475 CI: <LOD  | 490                  | 3                        | 148       | 2-hydroxypentanedioic acid             |    | Nist EI list match 98%     | Unique to this study                           | P        |
| M14             | EI: 73(100), 115, 133, 147, 171, 273, M-57=447, M-15=489 CI: <LOD       | 504                  | 3                        | 162       | 3-hydroxyhexanedioic acid              |    | Proposed from mass spectra | Unique to this study                           | P        |
| M15             | EI: 73, 147, 301, M-131=387, M-57=461(100), M-15=503 CI: <LOD           | 518                  | 3                        | 176       | 3-carboxypentanedioic acid             |    | Proposed from mass spectra | Unique to this study                           | P        |
| M16             | EI: 73, 147, 269, 301, 385, M-131=401, M-57=475(100), M-15=517 CI: <LOD | 532                  | 3                        | 190       | 3-carboxyhexanedioic acid              |  | Standard                   | Unique to this study                           | P        |



**Fig. 12.** Humidity influence on on-Teflon-quartz filters organic acids and hydroxyl compounds MTBSTFA-derivatisation. Comparison is based on response factors evaluated from 5 point calibration curves (5 to 320 ng). “Response factor” refers to FID chromatogram surface peak area per ng of compound, in area units. Error bars represent  $\pm$  two standard errors given for a 95 % confidence level.

#### 4 Method application to limonene ozonolysis in a simulated atmosphere

Limonene is a significant contributor to monoterpene emissions, which accounts for around 11 % of global VOC emissions (Guenther et al., 1995). Moreover, with two double bonds, it is more reactive and has a higher SOA yield (around twice, Griffin et al., 1999) compared to mono-unsaturated monoterpenes, such as  $\alpha$ - and  $\beta$ -pinene, which are emitted in large amounts at a global scale. Limonene is consequently supposed to be a significant contributor to SOA. Furthermore, limonene is also widely present in indoor environments as it is largely employed in household cleaning products, air fresheners or essential oils (Singer et al., 2006). Limonene oxidation in indoor environments, especially through ozonolysis, has been demonstrated in many studies during the last years, pointing out strong airway irritants and ultrafine particle formation (Clausen et al., 2001; Sarwar and Corsi, 2007; Langer et al., 2008; Coleman et al., 2008; Forester and Wells, 2009; Jardine et al., 2011).

Therefore, limonene ozonolysis was chosen as a relevant system to assess the potential of the developed method to explore the chemical composition of oxygenated secondary

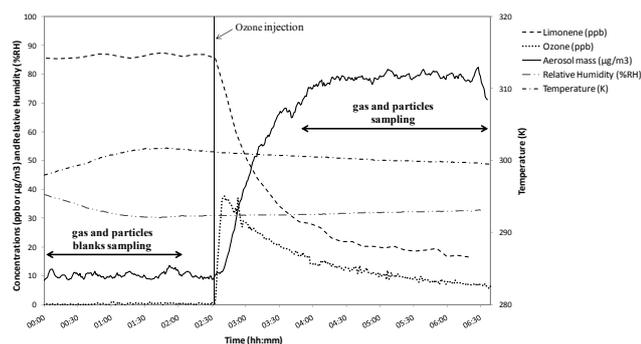


Fig. 13. Evolution of limonene ozonolysis experiment parameters.

organic matter simultaneously in gaseous and particulate phases. Moreover, this chemical system is sufficiently well characterised, allowing us to compare our results to the literature for method validation purposes (e.g. Leungsakul et al., 2005a, b; Jaoui et al., 2006; Walser et al., 2007; Forester and Wells, 2009).

#### 4.1 Experimental protocol

A limonene ozonolysis experiment was conducted in the EU-PHORE simulation chamber (cf. Sect. 2.3.3) to simulate real indoor or outdoor atmospheric conditions. Limonene was first injected into the chamber at an initial concentration level of 100 ppb. Non-acidified sulphate ammonium seeds were introduced by nebulisation of a 0.02 M solution at a concentration of  $12.5 \mu\text{g m}^{-3}$ . Water was then introduced until an initial relative humidity of 50 % was reached. Ozone was finally injected during 5 min 16 s, corresponding to an initial concentration of 105 ppb (see Sect. 2.3.3). The average temperature during the experiment was  $300 \pm 1$  K. No hydroxyl radical scavenger was introduced as this would stray the experiment from realistic indoor conditions (Docherty and Ziemann, 2003; Jonsson et al., 2008). Main parameter variations as monitored during the course of the experiment are given in Fig. 13.

In order to evaluate the fraction of the oxidative chemistry initiated by hydroxyl radicals on limonene – originating from Criegee intermediate decomposition after ozone addition onto double bonds – a simulation was run using the Master Chemical Mechanism, MCM v3.2 (Jenkin et al., 1997; Saunders et al., 2003). Initialising the MCM with our experimental conditions, it was found that around two-thirds of the initial limonene amount reacts with ozone, while one-third is expected to react with hydroxyl radicals. As a result, both ozonolysis and photo-oxidation products of limonene are likely to be detected and are thus consequently specifically investigated.

Blanks were sampled just before ozone introduction during 120 min. Gas-phase blanks were sampled on one PFBHA- and one MTBSTFA-coated Tenax TA tubes at a flow rate of  $100 \text{ ml min}^{-1}$  downstream from a Teflon fil-

ter (Zeffluor, 47 mm, Pall Life Sciences). Particulate-phase blanks were sampled in parallel on one quartz filter and one Teflon-quartz filter at a flow rate of  $16.71 \text{ min}^{-1}$ . Sampling was performed during 160 min after ozone introduction and system stabilisation (aerosol maximum mass reached within one hour). The gas phase was sampled onto three PFBHA- and three MTBSTFA-coated Tenax TA tubes at a flow rate of  $100 \text{ ml min}^{-1}$  downstream from a Teflon filter. For each derivatisation reagent, one tube was equipped with a back-up tube (see Sect. 3.2.4) which was coated in the same way to observe possible breakthrough of the sample tubes. The particulate phase was sampled onto one Teflon-quartz and two quartz filters at a flow rate of  $16.71 \text{ min}^{-1}$ . No VOC denuder was used in this method testing experiment. It is important to note that no specific set-up was used to minimise sampling artifact for these experiments, as the main objective of the experiments here was the application of the methodology presented and characterised in the previous section.

All samples were analysed by TD/GC/MS, in both EI and CI modes, as previously described (cf. Sect. 2.2.).

#### 4.2 Results

##### 4.2.1 Aerosol mass yield

The secondary aerosol growth curve for the limonene ozonolysis experiment is given in Fig. 14. The aerosol mass yield corresponds to the time-dependant aerosol mass yield  $Y$  calculated from TEOM measurements and is defined as:

$$Y = \frac{\Delta M_0}{\Delta \text{HC}}, \quad (3)$$

where  $\Delta M_0$  is the aerosol mass loading ( $\mu\text{g m}^{-3}$ ) and  $\Delta \text{HC}$  is the total reacted limonene ( $\mu\text{g m}^{-3}$ ), both corrected for dilution and wall losses. The final aerosol mass yield was around 20 %. Higher values are usually observed in the same experimental conditions ( $\sim 50$  % by Jaoui et al., 2006, Northcross and Jang, 2007, Sun et al., 2011,  $\sim 65$  % by Saathoff et al., 2009). The lower value measured here can be explained by the elevated temperature of 300 K achieved in the chamber during our experiment, as demonstrated by Saathoff et al. (2009) who examined the influence of temperature on SOA yields from limonene ozonolysis.

##### 4.2.2 Product detection and identification

Compound detection and identification were based on characteristic fragmentation patterns of PFBHA and MTBSTFA derivatives (cf. Table 6) in EI and CI modes. The mass spectra of derivatised compounds and their retention time were compared to available standard data. Compounds were positively identified in the case where standards are available. Otherwise, a tentative assignment is proposed based on the comparison of the fragmentation pattern and the retention time with surrogate standards. Compounds detected

**Table 8.** Quantification of observed carbonyl compounds, organic acids and hydroxyl compounds. LOD: limit of detection. 1 nd: not detected.

| Compound number                                            | Compound                   | Calibration standard | Gas phase concentration (ng m <sup>-3</sup> ) | Particulate phase concentration (ng m <sup>-3</sup> ) |
|------------------------------------------------------------|----------------------------|----------------------|-----------------------------------------------|-------------------------------------------------------|
| Carbonyl compounds (PFBHA derivatives)                     |                            |                      |                                               |                                                       |
| P1                                                         | 5-hexene-2-one             | 3-pentene-2-one      | 166 ±4                                        | 10 ±4                                                 |
| P2                                                         | keto-limonene              | dihydrocarvone       | 1830 ±40                                      | <LOD                                                  |
| P3                                                         | methylglyoxal              | methylglyoxal        | 2 500 ±600                                    | 70 ±30                                                |
| P4                                                         | dimethylglyoxal            | dimethylglyoxal      | 2 500 ±200                                    | 6.3 ±0.9                                              |
| P5                                                         | 4-oxopentanal              | 4-oxopentanal        | 1 760 ±60                                     | 51 ±3                                                 |
| P6                                                         | limononaldehyde            | 4-oxopentanal        | 7 800 ±300                                    | 640 ±30                                               |
| Hydroxyl compounds and organic acids (MTBSTFA derivatives) |                            |                      |                                               |                                                       |
| M1                                                         | levulinic acid             | levulinic acid       | 990 ±60                                       | nd                                                    |
| M2                                                         | succinic acid              | succinic acid        | 18 ±1                                         | 7 ±3                                                  |
| M3                                                         | 2-methylbutanedioic acid   | succinic acid        | 4.8 ±0.3                                      | nd                                                    |
| M4                                                         | pentanedioic acid          | succinic acid        | 13.9 ±0.9                                     | 4 ±1                                                  |
| M6                                                         | hexanedioic acid           | succinic acid        | 29 ±2                                         | 12 ±5                                                 |
| M9                                                         | limonic acid               | pinic acid           | 2.9 ±0.3                                      | 70 ±10                                                |
| M12                                                        | malic acid                 | malic acid           | nd                                            | 12 ±3                                                 |
| M13                                                        | 2-hydroxypentanedioic acid | malic acid           | nd                                            | 6 ±2                                                  |
| M14                                                        | 3-hydroxyhexanedioic acid  | malic acid           | nd                                            | 17 ±5                                                 |

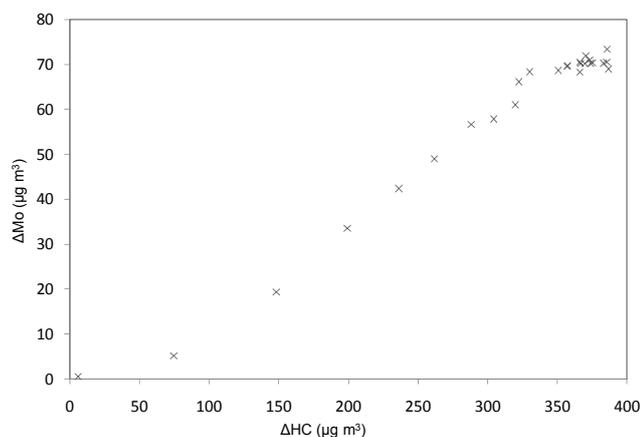
**Table 9.** Experimental partitioning coefficient  $K_i$  (m<sup>3</sup> µg<sup>-1</sup>) compared with  $K_i$  values (m<sup>3</sup> µg<sup>-1</sup>) estimated from theoretical saturation vapour pressures ( $P_{L,i}^o$ ) calculated from 3 group contribution methods at 298 K.  $K_i$  calculation assumes a temperature of 300 K, a mean MW<sub>om</sub> of 142 g mol<sup>-1</sup> and an activity coefficient  $\gamma_i$  of 1.27. MY: Myrdal and Yalkowsky, 1997. NAN: Nannoolal et al., 2008. SIM: Pankov and Asher, 2008.

| Compound number                                            | Compound          | Experimental $K_i$            | Estimated $K_i$ values |                        |                       |
|------------------------------------------------------------|-------------------|-------------------------------|------------------------|------------------------|-----------------------|
|                                                            |                   |                               | MY                     | NAN                    | SIM                   |
| Carbonyl compounds (PFBHA derivatives)                     |                   |                               |                        |                        |                       |
| P1                                                         | 5-hexene-2-one    | $1.3 \times 10^{-3} \pm 67\%$ | $4.54 \times 10^{-9}$  | $9.14 \times 10^{-9}$  | $7.26 \times 10^{-9}$ |
| P3                                                         | methylglyoxal     | $1.3 \times 10^{-3} \pm 84\%$ | $2.14 \times 10^{-9}$  | $8.77 \times 10^{-10}$ | $6.49 \times 10^{-9}$ |
| P4                                                         | dimethylglyoxal   | $7.4 \times 10^{-5} \pm 47\%$ | $7.11 \times 10^{-9}$  | $3.48 \times 10^{-9}$  | $7.00 \times 10^{-9}$ |
| P5                                                         | 4-oxopentanal     | $6.5 \times 10^{-4} \pm 34\%$ | $1.67 \times 10^{-8}$  | $1.45 \times 10^{-8}$  | $4.53 \times 10^{-8}$ |
| P6                                                         | limononaldehyde   | $1.9 \times 10^{-3} \pm 34\%$ | $4.54 \times 10^{-6}$  | $1.35 \times 10^{-6}$  | $7.50 \times 10^{-6}$ |
| Hydroxyl compounds and organic acids (MTBSTFA derivatives) |                   |                               |                        |                        |                       |
| M2                                                         | succinic acid     | $3.2 \times 10^{-2} \pm 73\%$ | $5.16 \times 10^{-4}$  | $3.00 \times 10^{-4}$  | $1.04 \times 10^{-3}$ |
| M4                                                         | pentanedioic acid | $7.8 \times 10^{-3} \pm 73\%$ | $1.90 \times 10^{-3}$  | $1.08 \times 10^{-3}$  | $2.76 \times 10^{-3}$ |
| M6                                                         | hexanedioic acid  | $6.9 \times 10^{-3} \pm 73\%$ | $6.49 \times 10^{-3}$  | $3.38 \times 10^{-3}$  | $6.50 \times 10^{-3}$ |
| M9                                                         | limonic acid      | $5.1 \times 10^{-1} \pm 50\%$ | $2.56 \times 10^{-1}$  | $2.51 \times 10^{-2}$  | $1.54 \times 10^{-1}$ |

in chamber blanks in the same order of magnitude than in samples were removed from the product list.

Around 30 compounds could be detected in the gas phase as PFBHA or MTBSTFA derivative products (20 PFBHA derivatives and 10 MTBSTFA derivatives) and around 65 in the particulate phases (30 PFBHA derivatives and 35 MTBSTFA derivatives). These products ranged from C<sub>3</sub>- to C<sub>10</sub>-

compounds and from mono- to tri-functionalised species. Compounds detected in the gas phase ranged from C<sub>3</sub> mono-carbonyl, C<sub>5</sub> mono-hydroxyl or C<sub>6</sub> mono-acid to C<sub>10</sub> di-carbonyl and C<sub>8</sub> di-hydroxyl or di-acids. In the particulate phase, tri-functionalised compounds were observed in the C<sub>8</sub>-C<sub>10</sub> range.



**Fig. 14.** Limonene ozonolysis experiment growth curve. Data corrected by dilution and wall losses.

Among all these detected products, 25 have been positively (based on standard mass spectra comparison) or tentatively identified (based on surrogate standard and/or mass spectra). An overview of the limonene oxidation products detected and tentatively or positively identified in both gas and particulate phases is presented in Table 7. Thirteen products previously identified from limonene oxidation experiments (Leungsakul et al., 2005a; Jaoui et al., 2006; Forester and Wells, 2009) or expected from theoretical oxidation schemes (Leungsakul et al., 2005b) were observed (P2, P3, P5, P6, P8, P9, M1, M2, M4, M6, M7, M9 and M10).

In addition to these species, twelve other products were, to the best of our knowledge, detected for the first time (P1, P4, P7, M3, M5, M8, M11, M12, M13, M14, M15, and M16).

This provides certainly the most comprehensive list of detected compounds in a single experiment for limonene ozonolysis in both gaseous and particulate phases.

This new information shows the potential of the method to explore the chemical composition of the secondary organic matter in a relevant range of semi-volatile compounds, and, hence, opens the way to new chemical scheme building for the production of semi-volatile organic species in the limonene ozonolysis system. An example of chromatograms is given for the analysis of carbonyl compounds in both phases (Fig. 15).

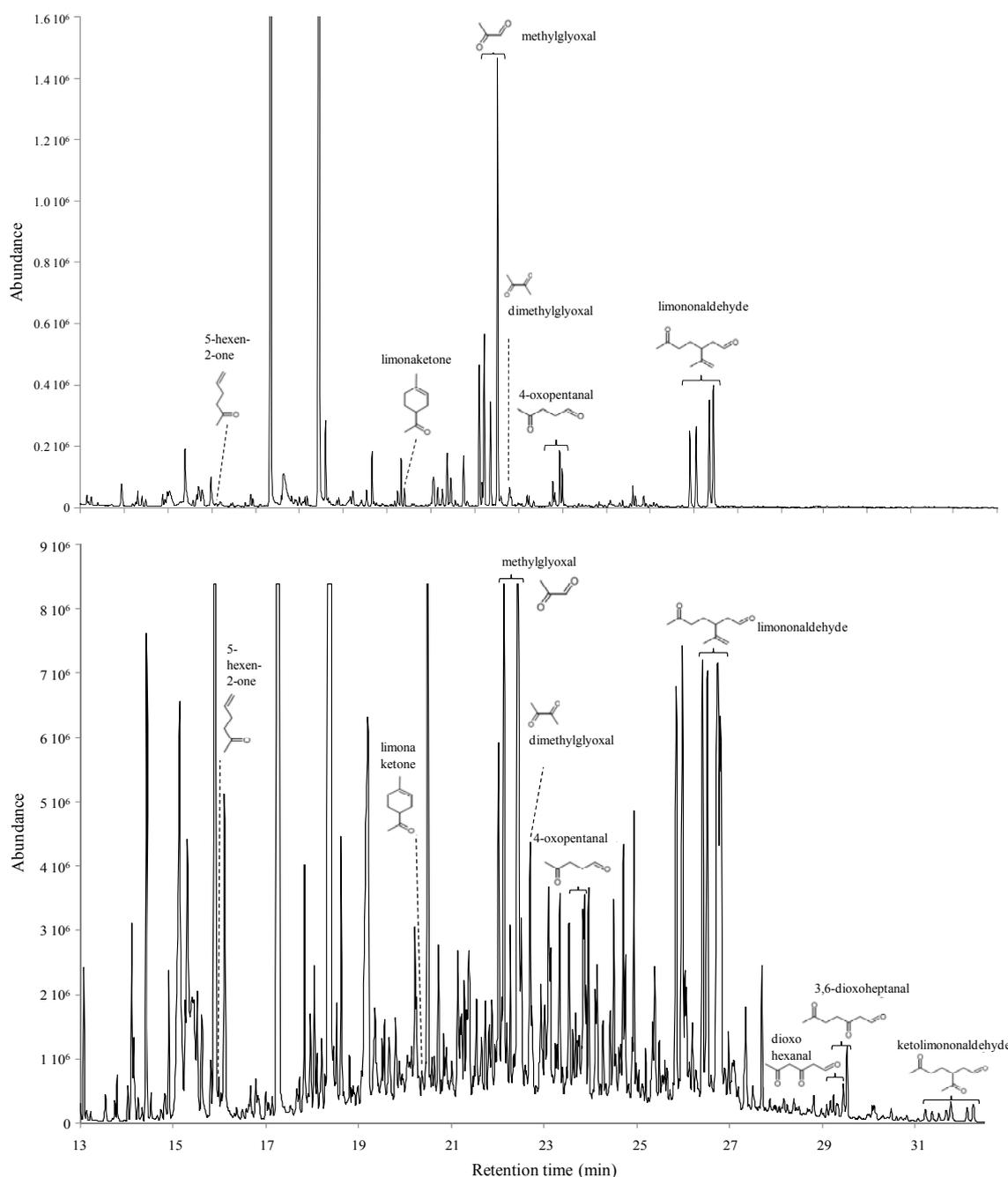
Re-collection (see Sect. 2.2.1) was used here to perform GC/MS analysis in both the EI and the CI modes successively for all gas- and particulate-phase samples, as this procedure was a convenient way to obtain complementary data for compound identification. Nevertheless, it must be indicated here that this approach is affected by an elevated detection limit obtained in CI mode for re-collected samples, especially for MTBSTFA-derivatised samples. The analysis of the re-collected samples was not validated for quantification as the re-collection efficiency was not evaluated. Furthermore, further experiments revealed that re-collected

MTBSTFA samples have to be analysed as soon as possible after first injection. Unfortunately, this was not possible in this study. Other attempts showed that a 24 h delay seems to significantly improve detection limits compared to the presented results for re-collected sample, analysed in the CI mode within 7 days.

#### 4.2.3 Quantification and estimation of partitioning coefficients

Among the 30 compounds detected in the gas phase and the 65 compounds detected in the particulate phase with the developed parallel sampling and analysis method, only those for which a relevant standard calibration in the gas and/or particulate phases (cf. Tables 1, 3, 4 and 5) could be obtained were quantified. This led to the quantification of 18 detected compounds, which are given in Table 8. Values were corrected from chamber blanks. The mentioned uncertainties correspond to the statistical uncertainties (Student's t-test) previously determined for calibration curves. The total quantified products represent around 16% of the carbon mass of the reacted limonene. The remaining non-quantified mass can be divided in three main groups: (1) non-identified compounds that were detected here but not quantified, (2) non-derivatisable species, and (3) high molecular weight compounds such as oligomers.

For compounds quantified in both the gas and particulate phases, an experimental partitioning coefficient was calculated (Table 9). However, as the whole mass of a given compound present in the chamber participates to the partitioning of this compound, quantifications values used for partitioning coefficients calculation were not corrected from chamber blank values but only from laboratory blank values. A total aerosol concentration of  $45 \pm 12 \mu\text{g m}^{-3}$  was considered, which corresponds to the average real aerosol concentration during sampling, not corrected for aerosol losses in the chamber. Uncertainties were calculated on the basis of gas-phase, particulate-phase and aerosol mass quantification uncertainties (calibration curves and TEOM uncertainty). As no denuder was used for upstream quartz and Teflon-quartz sample filters to prevent positive artifacts on the particulate-phase compounds quantification, the determined experimental partitioning coefficients represent the best estimates under the conditions applied. In order to illustrate the usefulness of determining experimental partitioning coefficients and highlight the relevance of the present method development, the determined experimental partitioning coefficients were compared to values calculated from theoretical vapour pressure data estimated for three group contribution methods (Myrdal and Yalkowsky, 1997 – MY; Nannoolal et al., 2008 – NAN; Pankow and Asher, 2008 – SIM). The following parameters were used for these calculations: an experimental temperature of 300 K, an average molecular weight in the particulate phase ( $MW_{\text{om}}$ ) of  $142 \text{ g mol}^{-1}$  and an activity coefficient ( $\gamma_i$ ) of 1.27. These two last values



**Fig. 15.** Gas chromatograms of limonene ozonolysis products derivatised by PFBHA, EI selected ion  $m/z$  181 chromatogram. Top chromatogram: gas phase. Bottom chromatogram: particulate phase.

are taken from simulation results reported by Leungskul et al. (2005a). Besides, when available, experimental partitioning coefficients were also compared with other experimental values. Limononaldehyde (P6) was the most abundant product in the gas phase, as observed by Leungskul et al. (2005a) for limonene ozonolysis experiments. However, contrary to this previous study, limononaldehyde was also the most abundant product in the particulate phase instead

of keto-limononaldehyde. The experimental partitioning coefficient for limononaldehyde obtained in the present study is  $3.5 \times 10^{-3} \text{ m}^3 \mu\text{g}^{-1}$ , a value which is three orders of magnitude higher than the estimated one by calculation and two orders of magnitude higher than the limononaldehyde experimental partitioning coefficients estimated by Leungskul et al. (2005a) (around  $6 \times 10^{-5} \text{ m}^3 \mu\text{g}^{-1}$ ).

The second most abundant products in the gas phase were methylglyoxal and dimethylglyoxal, methylglyoxal being also the second most abundant compound in the particulate phase. The experimental partitioning coefficient for methylglyoxal is also three orders of magnitude higher than estimated coefficients. However, it is similar to experimental values determined by Ortiz et al. (2010) during field ( $3.4 \times 10^{-2} \text{ m}^3 \mu\text{g}^{-1}$ ) and toluene oxidation smog chamber ( $1 \times 10^{-4} \text{ m}^3 \mu\text{g}^{-1}$ ) experiments. If this direct comparison can be affected by different aerosol properties and different sampling conditions and analytical techniques employed, our experimental value for the methylglyoxal partitioning coefficient can be considered as another clue suggesting that the eventually polymerised methylglyoxal concentration in the particulate phase, under atmospherically relevant conditions, could be higher than expected from semi-empirical group contribution methods.

Another important product detected in both gas and particulate phases was 4-oxopentanal (P5). This compound was previously detected as a limonene oxidation product only by Forester and Wells (2009). It was also observed at high levels in a forestal atmosphere, 180–1570  $\text{ng m}^3$  in the gas phase and 25–207  $\text{ng m}^3$  in the particulate phase (Matsunaga et al., 2004), indicating that 4-oxopentanal could be an important SOA contributor. However, its major source is assumed by these authors to be the oxidation of 6-methyl-5-heptene-2-one (6-MHO), which originates from squalene oxidation). However, 6-MHO was not observed in the present study, suggesting another pathway for 4-oxopentanal formation, which will not be discussed here.

Experimental partitioning coefficient estimation for 4-oxopentanal provides a value five orders of magnitude higher than estimation calculation.

Dioxohexanal (P7), 3,6-dioxoheptanal (P8) and ketolimononaldehyde (P9) were only quantified in the particulate phase, although their estimated partitioning coefficients predict a gas-phase concentration higher than  $100\,000 \mu\text{g m}^{-3}$ . More realistically, Leungsakul et al. (2005a) provide experimental partitioning coefficients of keto-limononaldehyde in the range of  $1.41\text{--}1.76 \times 10^{-3} \text{ m}^3 \mu\text{g}^{-1}$ . From these values and the keto-limononaldehyde particulate phase concentration estimated in the present study, a gas-phase concentration of around  $1400 \text{ ng m}^{-3}$  could be expected, which is still largely above detection limits achieved for di-carbonyls compounds in the gas phase. From this observation, three hypotheses could be advanced. Firstly, tri-carbonyl compounds may not exist significantly in the gas phase under the experimental conditions of this work, and the keto-limononaldehyde gas-phase concentration quantified by Leungsakul and co-workers was either mainly a positive artifact due to volatilisation from the particulate phase or linked to different experimental conditions. Secondly, these compounds are more prone to a negative artifact upon quantification in the gas phase than di-carbonyl compounds. Owing to adsorption onto the Teflon filter placed upstream from

the adsorbent tubes, this negative artifact is increasing as expected when decreasing compound volatility (Volckens and Leith, 2003). Thirdly, tri-carbonyl compounds may not be detectable when sampled onto gas collection Tenax TA tubes, owing to either derivatisation which may not be complete or to a limited desorption. Further experiments based on tri-carbonyl standards are required to support or invalidate this last hypothesis.

Levulinic acid (M1) was the most important acid observed in the gas phase. Li and Yu (2005) proposed that levulinic acid is an oxidation product of 4-oxopentanal. Limonic acid (M9) was the major acid that could be precisely quantified in the particulate phase with a concentration of  $70 \pm 10 \text{ ng m}^{-3}$ . A keto-diacid (M10: ketolimononic acid) and two tri-dicarboxylic acids (M15 and M16) were also identified in the particulate phase. However, since their standard or surrogate standards calibration curves were not linear, their concentrations were estimated on the basis of pinic acid for M10 ( $160 \mu\text{g m}^{-3}$ ) and malic acid for M15 and M16 ( $100 \mu\text{g m}^{-3}$  and  $80 \mu\text{g m}^{-3}$ ). The experimental partitioning coefficient of limonic acid was of the same order of magnitude than estimation calculation from MY and SIM. However, it was an order of magnitude higher than NAN estimation calculation and experimental values provided by Yu et al. (1999) for other biogenic di-acids: pinic acid, caric acid and sabinic acid,  $2.8\text{--}3.5 \times 10^{-2} \mu\text{g m}^{-3}$ ,  $3.3 \times 10^{-2} \mu\text{g m}^{-3}$  and  $2.7 \times 10^{-2} \mu\text{g m}^{-3}$ , respectively, at 306–308 K. Experimental partitioning coefficients of hexanedioic acid (M6) and pentanedioic acid (M4) were also consistent with theoretical calculations. The experimental partitioning coefficient of succinic acid (M2) was one or two orders of magnitude higher than theoretical calculations.

To conclude, using quantification of compounds in both gaseous and particulate phases, partitioning coefficients have been calculated. Differences can be observed in comparison with estimated and other experimental coefficients. Some hypotheses can be proposed to explain these discrepancies: (i) the estimated partitioning coefficient are calculated on the basis of estimations that may be wrong, which is one of the objectives of this study: providing partitioning coefficient obtained under real atmospheric conditions; (ii) reactions occurring in the particulate phase can disturb the equilibrium, leading to different partitioning coefficients than expected, which is another objective of our study: identifying unsuspected reaction pathways; and (iii) sampling positive artifacts upon particulate phase quantification as no denuder has been used for particulate phase sampling.

Artifacts seemed to be less important for di-acids than for carbonyl compounds. This was consistent with: (i) the Teflon-quartz nature of the filter media that is expected to reduce adsorption artifacts compared to quartz filters, even if this feature is not confirmed by Arp et al. (2007), and (ii) the artifact tendency expected to decrease compound volatility from carbonyl compounds to limonic acid (Volckens and Leith, 2003).

## 5 Conclusions and perspectives

This work has provided a useful new analytical method, based on thermal desorption coupled with GC/MS, to explore the secondary organic matter composition at a molecular scale from a global point of view in both gas and particulate phases. Using separately but in parallel two derivatisation reagents, PFBHA and MTBSTFA, to analyse carbonyl compounds and organic acids and hydroxylated compounds, respectively, the method allows one to explore the chemical composition up to tri-functionalised species. Furthermore, it combines the classical advantages of thermal desorption – solvent-free, easy to use and fast – to analyse the same sample in both EI and CI modes, providing complementary structural information through the use of the re-collection option of commercial thermal desorbers. The potential of the method to determine the partitioning behaviour of a wide range of semi-volatile compounds has been demonstrated. Quantification in both gas and particulate phases is indeed validated for mono- and difunctionalised species and for hydroxyl di-acids. This is substantial progress compared to other techniques designed for compound partitioning studies. Compared to the TAG instrument (Isaacman et al., 2011), whose main advantage is based on on-line measurements, the developed method widens the range of detectable species to polar and multi-functionalised compounds. Compared to denuder/filter techniques (e.g. Leungsakul et al., 2005a; Temime et al., 2007), it provides a simpler way to study the partitioning of the secondary organic matter, avoiding time-consuming sample work-up procedures that are prone to sample losses during solvent extraction steps, with similar detection limits. It is an easy way to systematically assess, in simulation chamber experiments and in the field, the partitioning behaviour of semi-volatile compounds, providing a set of data in a wide range of environmental conditions that allow support of a relevant integration of partitioning in models.

Application of the method to the study of limonene ozonolysis demonstrates the potential of the method to explore the chemical composition of a complex mixture in both gas and particulate phases, detecting both compounds already known as limonene oxidation products as well as a range of products detected for the first time. In the present study, 25 limonene oxidation products were positively or tentatively identified, 9 being in both gaseous and particulate phases and 12, among them tricarboxylic acids, hydroxyl dicarboxylic acids and oxodicarboxylic acids, being detected for the first time.

Quantification and partitioning coefficient estimations were provided, although as no precautions were taken to avoid or correct artifacts for the quantification of compounds in both gas and particulate phases compounds quantification, their relevance for atmospheric conclusions are limited here. Nevertheless, for decoupling gas and particulate sampling, the method allows testing a wide range of possibili-

ties to limit (e.g. denuder use) or correct (e.g. backup filter use) artifacts.

Furthermore, derivatisation protocols should be improved to extend the range of quantifiable species to tri-functionalised compounds. To improve the confidence in the experimental determination of partitioning coefficients for tricarbonyl compounds, observed in limonene ozonolysis particulate-phase samples, it appears necessary to assess detection limits in the gas phase and to provide more relevant calibration curves for the particulate phase. It is demonstrated that ketodiacids and triacids are quantifiable in the gas phase and can be detected in limonene ozonolysis particulate phase samples. An adjustment of the derivatisation protocol for the particulate phase, such as a small increase of the derivatisation temperature, could provide relevant calibration curves.

In conclusion, the developed method allows to explore the chemical composition of the semi-volatile fraction of the oxygenated secondary organic matter and to assess its partitioning behaviour from mono to difunctionalised carbonyl, hydroxyl and organic acids species. Further improvement could enlarge the quantifiable species in both gas and particulate phases to trifunctionalised compounds. The method is applicable to both simulation chamber experiments under realistic humidity conditions and field campaigns.

**Supplementary material related to this article is available online at: <http://www.atmos-meas-tech.net/5/1459/2012/amt-5-1459-2012-supplement.pdf>.**

*Acknowledgements.* The authors wish to thank the EuPhoRe group at CEAM, Valencia for their contribution to simulation chamber experiments. This work was supported by the EC within the I3 project “Integrating of European Simulation Chambers for Investigating Atmospheric Processes – II” (EUROCHAMP-2, contract no. 228335).

Edited by: M. Sipilä



The publication of this article is financed by CNRS-INSU.

## References

- Arp, H. P. H. and Goss, K. U.: Ambient Gas/Particle Partitioning. 3. Estimating Partition Coefficients of Apolar, Polar, and Ionizable Organic Compounds by Their Molecular Structure, *Environ. Sci. Technol.*, 43, 1923–1929 doi:10.1021/es8025165, 2009.
- Arp, H. P. H., Schwarzenbach, R. P., and Goss, K. U.: Equilibrium sorption of gaseous organic chemicals to fiber filters used for aerosol studies, *Atmos. Environ.*, 41, 8241–8252, doi:10.1016/j.atmosenv.2007.06.026, 2007.
- Asher, W. E. and Pankow, J. F.: Vapor pressure prediction for alkenoic and aromatic organic compounds by a UNIFAC-based group contribution method, *Atmos. Environ.*, 40, 3588–3600, doi:10.1016/j.atmosenv.2005.12.004, 2006.
- Baltussen, E. B., Cramers, C. C., and Sandra, P. S.: Sorptive sample preparation – a review, *Anal. Bioanal. Chem.*, 373, 3–22, 2002.
- Barley, M. H. and McFiggans, G.: The critical assessment of vapour pressure estimation methods for use in modelling the formation of atmospheric organic aerosol, *Atmos. Chem. Phys.*, 10, 749–767, doi:10.5194/acp-10-749-2010, 2010.
- Beiner, K., Plewka, A., Haferkorn, S., Iinuma, Y., Engewald, W., and Herrmann, H.: Quantification of organic acids in particulate matter by coupling of thermally assisted hydrolysis and methylation with thermodesorption-gas chromatography-mass spectrometry, *J. Chromatogr. A*, 1216, 6642–6650, doi:10.1016/j.chroma.2009.07.054, 2009.
- Bessagnet, B., Menut, L., Curci, G., Hodzic, A., Guillaume, B., Liousse, C., Moukhtar, S., Pun, B., Seigneur, C., and Schulz, M.: Regional modeling of carbonaceous aerosols over Europe-focus on secondary organic aerosols, *J. Atmos. Chem.*, 61, 175–202, doi:10.1007/s10874-009-9129-2, 2008.
- Blake, R. S., Whyte, C., Hughes, C. O., Ellis, A. M., and Monks, P. S.: Demonstration of proton-transfer reaction time-of-flight mass spectrometry for real-time analysis of trace volatile organic compounds, *Anal. Chem.*, 76, 3841–3845, doi:10.1021/ac0498260, 2004.
- Camredon, M., Aumont, B., Lee-Taylor, J., and Madronich, S.: The SOA/VOC/NO<sub>x</sub> system: an explicit model of secondary organic aerosol formation, *Atmos. Chem. Phys.*, 7, 5599–5610, doi:10.5194/acp-7-5599-2007, 2007.
- Camredon, M., Hamilton, J. F., Alam, M. S., Wyche, K. P., Carr, T., White, I. R., Monks, P. S., Rickard, A. R., and Bloss, W. J.: Distribution of gaseous and particulate organic composition during dark  $\alpha$ -pinene ozonolysis, *Atmos. Chem. Phys.*, 10, 2893–2917, doi:10.5194/acp-10-2893-2010, 2010.
- Canagaratna, M. R., Jayne, J. T., Jimenez, J. L., Allan, J. D., Alfarra, M. R., Zhang, Q., Onasch, T. B., Drewnick, F., Coe, H., Middlebrook, A., Delia, A., Williams, L. R., Trimborn, A. M., Northway, M. J., DeCarlo, P. F., Kolb, C. E., Davidovits, P., and Worsnop, D. R.: Chemical and microphysical characterization of ambient aerosols with the aerodyne aerosol mass spectrometer, *Mass Spectrom. Rev.*, 26, 185–222, 2007.
- Cancilla, D. A. and Que Hee, S. S.: O-(2,3,4,5,6-Pentafluorophenyl)methylhydroxylamine hydrochloride: a versatile reagent for the determination of carbonyl-containing compounds, *J. Chromatogr. A*, 627, 1–16, doi:10.1016/0021-9673(92)87181-7, 1992.
- Cappa, C. D. and Jimenez, J. L.: Quantitative estimates of the volatility of ambient organic aerosol, *Atmos. Chem. Phys.*, 10, 5409–5424, doi:10.5194/acp-10-5409-2010, 2010.
- Carrasco, N., Doussin, J. F., O'Connor, M., Wenger, J. C., Picquet-Varrault, B., Durand-Jolibois, R., and Carlier, P.: Simulation chamber studies of the atmospheric oxidation of 2-methyl-3-buten-2-ol: Reaction with hydroxyl radicals and ozone under a variety of conditions, *J. Atmos. Chem.*, 56, 33–55, doi:10.1007/s10874-006-9041-y, 2007.
- Castells, P., Santos, F. J., and Galceran, M. T.: Development of a sequential supercritical fluid extraction method for the analysis of nitrated and oxygenated derivatives of polycyclic aromatic hydrocarbons in urban aerosols, *J. Chromatogr. A*, 1010, 141–151, doi:10.1016/S0021-9673(03)01121-X, 2003.
- Chan, M. N., Chan, A. W. H., Chhabra, P. S., Surratt, J. D., and Seinfeld, J. H.: Modeling of secondary organic aerosol yields from laboratory chamber data, *Atmos. Chem. Phys.*, 9, 5669–5680, doi:10.5194/acp-9-5669-2009, 2009.
- Chan, M. N., Surratt, J. D., Claeys, M., Edgerton, E. S., Tanner, R. L., Shaw, S. L., Zheng, M., Knipping, E. M., Eddingsaas, N. C., Wennberg, P. O., and Seinfeld, J. H.: Characterization and quantification of isoprene-derived epoxydiols in ambient aerosol in the Southeastern United States, *Environ. Sci. Technol.*, 44, 4590–4596, doi:10.1021/es100596b, 2010.
- Chi, Y., Feng, Y., Wen, S., Lü, H., Yu, Z., Zhang, W., Sheng, G., and Fu, J.: Determination of carbonyl compounds in the atmosphere by DNPH derivatisation and LC-ESI-MS/MS detection, *Talanta*, 72, 539–545, 2007.
- Chiappini, L., Perraudin, E., Durand-Jolibois, R., and Doussin, J. F.: Development of a supercritical fluid extraction-gas chromatography-mass spectrometry method for the identification of highly polar compounds in secondary organic aerosols formed from biogenic hydrocarbons in smog chamber experiments, *Anal. Bioanal. Chem.*, 386, 1749–1759, doi:10.1007/s00216-006-0744-3, 2006.
- Claeys, M., Graham, B., Vas, G., Wang, W., Vermeylen, R., Pashynska, V., Cafmeyer, J., Guyon, P., Andreae, M. O., Artaxo, P., and Maenhaut, W.: Formation of secondary organic aerosols through photooxidation of isoprene, *Science*, 303, 1173–1176, doi:10.1126/science.1092805, 2004.
- Claeys, M., Szmigielski, R., Kourtchev, I., Van der Veken, P., Vermeylen, R., Maenhaut, W., Jaoui, M., Kleindienst, T. E., Lewandowski, M., Offenberg, J. H., and Edney, E. O.: Hydroxycarboxylic acids: markers for secondary organic aerosol from the photooxidation of  $\alpha$ -pinene, *Environ. Sci. Technol.*, 41, 1628–1634, doi:10.1021/es0620181, 2007.
- Claeys, M., Iinuma, Y., Szmigielski, R., Surratt, J. D., Blockhuys, F., Van Alsenoy, C., Boge, O., Sierau, B., Gomez-Gonzalez, Y., Vermeylen, R., Van der Veken, P., Shahgholi, M., Chan, A. W. H., Herrmann, H., Seinfeld, J. H., and Maenhaut, W.: Terpenylic acid and related compounds from the oxidation of  $\alpha$ -pinene: implications for new particle formation and growth above forests, *Environ. Sci. Technol.*, 43, 6976–6982, doi:10.1021/es9007596, 2009.
- Clausen, P. A., Wilkins, C. K., Wolkoff, P., and Nielsen, G. D.: Chemical and biological evaluation of a reaction mixture of R-(+)-limonene/ozone: Formation of strong airway irritants, *Environ. Int.*, 26, 511–522, doi:10.1016/S0160-4120(01)00035-6, 2001.
- Coleman, B. K., Lunden, M. M., Destailats, H., and Nazaroff, W. W.: Secondary organic aerosol from ozone-initiated reactions with terpene-rich household products, *Atmos. Environ.*, 42,

- 8234–8245, doi:10.1016/j.atmosenv.2008.07.031, 2008.
- de Gouw, J. and Warneke, C.: Measurements of volatile organic compounds in the earth's atmosphere using proton-transfer-reaction mass spectrometry, *Mass Spectrom. Rev.*, 26, 223–257, 2007.
- Docherty, K. S. and Ziemann, P. J.: Effects of stabilized Criegee intermediate and OH radical scavengers on aerosol formation from reactions of  $\alpha$ -pinene with O<sub>3</sub>, *Aerosol Sci. Tech.*, 37, 877–891, doi:10.1080/02786820300930, 2003.
- Donahue, N. M., Robinson, A. L., Stanier, C. O., and Pandis, S. N.: Coupled partitioning, dilution, and chemical aging of semivolatile organics, *Environ. Sci. Technol.*, 40, 02635–02643, doi:10.1021/es052297c, 2006.
- Donahue, N. M., Robinson, A. L., and Pandis, S. N.: Atmospheric organic particulate matter: From smoke to secondary organic aerosol, *Atmos. Environ.*, 43, 94–106, doi:10.1016/j.atmosenv.2008.09.055, 2009.
- Edney, E. O., Kleindienst, T. E., Jaoui, M., Lewandowski, M., Offenberg, J. H., Wang, W., and Claeys, M.: Formation of 2-methyl tetrols and 2-methylglyceric acid in secondary organic aerosol from laboratory irradiated isoprene/NOX/SO<sub>2</sub>/air mixtures and their detection in ambient PM<sub>2.5</sub> samples collected in the eastern United States, *Atmos. Environ.*, 39, 5281–5289, doi:10.1016/j.atmosenv.2005.05.031, 2005.
- Farina, S. C., Adams, P. J., and Pandis, S. N.: Modeling global secondary organic aerosol formation and processing with the volatility basis set: Implications for anthropogenic secondary organic aerosol, *J. Geophys. Res.-Atmos.*, 115, 17, doi:10.1029/2009jd013046, 2010.
- Forester, C. D. and Wells, J. R.: Yields of carbonyl products from gas-phase reactions of fragrance compounds with OH radical and ozone, *Environ. Sci. Technol.*, 43, 3561–3568, doi:10.1021/es803465v, 2009.
- Forstner, H. J. L., Flagan, R. C., and Seinfeld, J. H.: Molecular speciation of secondary organic aerosol from photooxidation of the higher alkenes: 1-octene and 1-decene, *Atmos. Environ.*, 31, 1953–1964, doi:10.1016/s1352-2310(96)00356-1, 1997.
- Goldstein, A. H., Worton, D. R., Williams, B. J., Hering, S. V., Kreisberg, N. M., Panic, O., and Górecki, T.: Thermal desorption comprehensive two-dimensional gas chromatography for in-situ measurements of organic aerosols, *J. Chromatogr. A*, 1186, 340–347, doi:10.1016/j.chroma.2007.09.094, 2008.
- Gómez-González, Y., Surratt, J. D., Cuyckens, F., Szmigielski, R., Vermeylen, R., Jaoui, M., Lewandowski, M., Offenberg, J. H., Kleindienst, T. E., Edney, E. O., Blockhuys, F., Van Alsenoy, C., Maenhaut, W., and Claeys, M.: Characterisation of organosulfates from the photooxidation of isoprene and unsaturated fatty acids in ambient aerosol using liquid chromatography/(-) electrospray ionization mass spectrometry, *J. Mass Spectrom.*, 43, 371–382, 2008.
- Gonzalez-Flesca, N. and Frezier, A.: A new laboratory test chamber for the determination of diffusive sampler uptake rates, *Atmos. Environ.*, 39, 4049–4056, doi:10.1016/j.atmosenv.2005.03.025, 2005.
- Graber, E. R. and Rudich, Y.: Atmospheric HULIS: How humic-like are they? A comprehensive and critical review, *Atmos. Chem. Phys.*, 6, 729–753, doi:10.5194/acp-6-729-2006, 2006.
- Grieshop, A. P., Miracolo, M. A., Donahue, N. M., and Robinson, A. L.: Constraining the volatility distribution and gas-particle partitioning of combustion aerosols using isothermal dilution and thermodenuder measurements, *Environ. Sci. Technol.*, 43, 4750–4756, doi:10.1021/es8032378, 2009.
- Griffin, R. J., Cocker, D. R., III, Seinfeld, J. H., and Dabdub, D.: Estimate of global atmospheric organic aerosol from oxidation of biogenic hydrocarbons, *Geophys. Res. Lett.*, 26, 2721–2724, doi:10.1029/1999GL900476, 1999.
- Guenther, A., Hewitt, C. N., Erickson, D., Fall, R., Geron, C., Graedel, T., Harley, P., Klinger, L., Lerdau, M., McKay, W. A., Pierce, T., Scholes, B., Steinbrecher, R., Tallamraju, R., Taylor, J., and Zimmerman, P.: A global-model of natural volatile organic-compound emissions, *J. Geophys. Res.-Atmos.*, 100, 8873–8892, doi:10.1029/94jd02950, 1995.
- Hallquist, M., Wenger, J. C., Baltensperger, U., Rudich, Y., Simpson, D., Claeys, M., Dommen, J., Donahue, N. M., George, C., Goldstein, A. H., Hamilton, J. F., Herrmann, H., Hoffmann, T., Iinuma, Y., Jang, M., Jenkin, M. E., Jimenez, J. L., Kiendler-Scharr, A., Maenhaut, W., McFiggans, G., Mentel, Th. F., Monod, A., Prévôt, A. S. H., Seinfeld, J. H., Surratt, J. D., Szmigielski, R., and Wildt, J.: The formation, properties and impact of secondary organic aerosol: current and emerging issues, *Atmos. Chem. Phys.*, 9, 5155–5236, doi:10.5194/acp-9-5155-2009, 2009.
- Hamilton, J. F., Webb, P. J., Lewis, A. C., and Reviejo, M. M.: Quantifying small molecules in secondary organic aerosol formed during the photo-oxidation of toluene with hydroxyl radicals, *Atmos. Environ.*, 39, 7263–7275, doi:10.1016/j.atmosenv.2005.09.006, 2005.
- Hamilton, J. F., Rami Alfarra, M., Wyche, K. P., Ward, M. W., Lewis, A. C., McFiggans, G. B., Good, N., Monks, P. S., Carr, T., White, I. R., and Purvis, R. M.: Investigating the use of secondary organic aerosol as seed particles in simulation chamber experiments, *Atmos. Chem. Phys.*, 11, 5917–5929, doi:10.5194/acp-11-5917-2011, 2011.
- Hansen, K. J., Nansen, B. N., Cravens, E., and Sievers, R. E.: Supercritical-fluid extraction gas-chromatographic analysis of organic-compounds in atmospheric aerosols, *Anal. Chem.*, 67, 3541–3549, doi:10.1021/ac00115a025, 1995.
- Hays, M. D. and Lavrich, R. J.: Developments in direct thermal extraction gas chromatography-mass spectrometry of fine aerosols, *TrAC Trend Anal. Chem.*, 26, 88–102, 2007.
- Healy, R. M., Wenger, J. C., Metzger, A., Duplissy, J., Kalberer, M., and Dommen, J.: Gas/particle partitioning of carbonyls in the photooxidation of isoprene and 1,3,5-trimethylbenzene, *Atmos. Chem. Phys.*, 8, 3215–3230, doi:10.5194/acp-8-3215-2008, 2008.
- Healy, R. M., Temime, B., Kuprovskite, K., and Wenger, J. C.: Effect of relative humidity on gas/particle partitioning and aerosol mass yield in the photooxidation of p-xylene, *Environ. Sci. Technol.*, 43, 1884–1889, doi:10.1021/es802404z, 2009.
- Ho, S. S. H. and Yu, J. Z.: Feasibility of collection and analysis of airborne carbonyls by on-sorbent derivatization and thermal desorption, *Anal. Chem.*, 74, 1232–1240, doi:10.1021/ac015708q, 2002.
- Ho, S. S. H. and Yu, J. Z.: Determination of airborne carbonyls: Comparison of a thermal desorption/GC method with the standard DNPH/HPLC method, *Environ. Sci. Technol.*, 38, 862–870, doi:10.1021/es034795w, 2004.

- IPCC: Fourth Assessment Report: Climate Change 2007, The Physical Science Basis, Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA., 2007.
- Isaacman, G., Worton, D. R., Kreisberg, N. M., Hennigan, C. J., Teng, A. P., Hering, S. V., Robinson, A. L., Donahue, N. M., and Goldstein, A. H.: Understanding evolution of product composition and volatility distribution through in-situ GC  $\times$  GC analysis: a case study of longifolene ozonolysis, *Atmos. Chem. Phys.*, 11, 5335–5346, doi:10.5194/acp-11-5335-2011, 2011.
- Jakober, C. A., Robert, M. A., Riddle, S. G., Destailhats, H., Charles, M. J., Green, P. G., and Kleeman, M. J.: Carbonyl emissions from gasoline and diesel motor vehicles, *Environ. Sci. Technol.*, 42, 4697–4703, doi:10.1021/es7029174, 2008.
- Jaoui, M. and Kamens, R. M.: Gaseous and particulate oxidation products analysis of a mixture of  $\alpha$ -pinene +  $\beta$ -pinene/O<sub>3</sub>/Air in the absence of light and  $\alpha$ -pinene +  $\beta$ -pinene/NO<sub>x</sub>/Air in the presence of natural sunlight, *J. Atmos. Chem.*, 44, 259–297, 2003.
- Jaoui, M., Leungsakul, S., and Kamens, R. M.: Gas and particle products distribution from the reaction of  $\beta$ -caryophyllene with ozone, *J. Atmos. Chem.*, 45, 261–287, 2003.
- Jaoui, M., Corse, E., Kleindienst, T. E., Offenberg, J. H., Lewandowski, M., and Edney, E. O.: Analysis of secondary organic aerosol compounds from the photooxidation of d-limonene in the presence of NO<sub>x</sub> and their detection in ambient PM<sub>2.5</sub>, *Environ. Sci. Technol.*, 40, 3819–3828, doi:10.1021/es052566z, 2006.
- Jardine, K., Serrano, A. Y., Arneth, A., Abrell, L., Jardine, A., van Haren, J., Artaxo, P., Rizzo, L. V., Ishida, F. Y., Karl, T., Kesselmeier, J., Saleska, S., and Huxman, T.: Within-canopy sesquiterpene ozonolysis in Amazonia, *J. Geophys. Res.-Atmos.*, 116, 10, doi:10.1029/2011jd016243, 2011.
- Jenkin, M. E.: Modelling the formation and composition of secondary organic aerosol from  $\alpha$ - and  $\beta$ -pinene ozonolysis using MCM v3, *Atmos. Chem. Phys.*, 4, 1741–1757, doi:10.5194/acp-4-1741-2004, 2004.
- Jenkin, M. E., Saunders, S. M., and Pilling, M. J.: The tropospheric degradation of volatile organic compounds: a protocol for mechanism development, *Atmos. Environ.*, 31, 81–104, doi:10.1016/S1352-2310(96)00105-7, 1997.
- Jonsson, A. M., Hallquist, M., and Ljungstrom, E.: Influence of OH scavenger on the water effect on secondary organic aerosol formation from ozonolysis of limonene, delta(3)-carene, and alpha-pinene, *Environ. Sci. Technol.*, 42, 5938–5944, doi:10.1021/es702508y, 2008.
- Kalberer, M., Yu, J., Cocker, D. R., Flagan, R. C., and Seinfeld, J. H.: Aerosol formation in the cyclohexene-ozone system, *Environ. Sci. Technol.*, 34, 4894–4901, doi:10.1021/es001180f, 2000.
- Kanakidou, M., Seinfeld, J. H., Pandis, S. N., Barnes, I., Dentener, F. J., Facchini, M. C., Van Dingenen, R., Ervens, B., Nenes, A., Nielsen, C. J., Swietlicki, E., Putaud, J. P., Balkanski, Y., Fuzzi, S., Horth, J., Moortgat, G. K., Winterhalter, R., Myhre, C. E. L., Tsigaridis, K., Vignati, E., Stephanou, E. G., and Wilson, J.: Organic aerosol and global climate modelling: a review, *Atmos. Chem. Phys.*, 5, 1053–1123, doi:10.5194/acp-5-1053-2005, 2005.
- Kitanovski, Z., Grgic, I., and Veber, M.: Characterization of carboxylic acids in atmospheric aerosols using hydrophilic interaction liquid chromatography tandem mass spectrometry, *J. Chromatogr. A*, 1218, 4417–4425, doi:10.1016/j.chroma.2011.05.020, 2011.
- Kleindienst, T. E., Conner, T. S., McIver, C. D., and Edney, E. O.: Determination of secondary organic aerosol products from the photooxidation of toluene and their implications in ambient PM<sub>2.5</sub>, *J. Atmos. Chem.*, 47, 79–100, 2004.
- Kroll, J. H., Ng, N. L., Murphy, S. M., Flagan, R. C., and Seinfeld, J. H.: Secondary organic aerosol formation from isoprene photooxidation, *Environ. Sci. Technol.*, 40, 1869–1877, doi:10.1021/es0524301, 2006.
- Kroll, J. H., Chan, A. W. H., Ng, N. L., Flagan, R. C., and Seinfeld, J. H.: Reactions of semivolatile organics and their effects on secondary organic aerosol formation, *Environ. Sci. Technol.*, 41, 3545–3550, doi:10.1021/es062059x, 2007.
- Langer, S., Moldanová, J., Arrhenius, K., Ljungström, E., and Ekberg, L.: Ultrafine particles produced by ozone/limonene reactions in indoor air under low/closed ventilation conditions, *Atmos. Environ.*, 42, 4149–4159, doi:10.1016/j.atmosenv.2008.01.034, 2008.
- Lazaridis, M.: Gas-particle partitioning of organic compounds in the atmosphere, *J. Aerosol Sci.*, 30, 1165–1170, doi:10.1016/S0021-8502(98)00788-5, 1999.
- Lee, B.-H., Pierce, J. R., Engelhart, G. J., and Pandis, S. N.: Volatility of secondary organic aerosol from the ozonolysis of monoterpenes, *Atmos. Environ.*, 45, 2443–2452, doi:10.1016/j.atmosenv.2011.02.004, 2011.
- Lee, S., Jang, M., and Kamens, R. M.: SOA formation from the photooxidation of  $\alpha$ -pinene in the presence of freshly emitted diesel soot exhaust, *Atmos. Environ.*, 38, 2597–2605, doi:10.1016/j.atmosenv.2003.12.041, 2004.
- Legrand, M., Preunkert, S., Galy-Lacaux, C., Liousse, C., and Wagenbach, D.: Atmospheric year-round records of dicarboxylic acids and sulfate at three French sites located between 630 and 4360 m elevation, *J. Geophys. Res.-Atmos.*, 110, 11, doi:10.1029/2004jd005515, 2005.
- Leungsakul, S., Jaoui, M., and Kamens, R. M.: Kinetic mechanism for predicting secondary organic aerosol formation from the reaction of d-limonene with ozone, *Environ. Sci. Technol.*, 39, 9583–9594, doi:10.1021/es0492687, 2005a.
- Leungsakul, S., Jeffries, H. E., and Kamens, R. M.: A kinetic mechanism for predicting secondary aerosol formation from the reactions of d-limonene in the presence of oxides of nitrogen and natural sunlight, *Atmos. Environ.*, 39, 7063–7082, doi:10.1016/j.atmosenv.2005.08.024, 2005b.
- Li, J., Feng, Y. L., Xie, C. J., Huang, J., Yu, J. Z., Feng, J. L., Sheng, G. Y., Fu, J. M., and Wu, M. H.: Determination of gaseous carbonyl compounds by their pentafluorophenyl hydrazones with gas chromatography/mass spectrometry, *Anal. Chim. Acta*, 635, 84–93, doi:10.1016/j.aca.2008.12.041, 2009.
- Li, Y.-c. and Yu, J. Z.: Simultaneous determination of mono- and dicarboxylic acids,  $\omega$ -oxo-carboxylic acids, midchain ketocarboxylic acids, and aldehydes in atmospheric aerosol samples, *Environ. Sci. Technol.*, 39, 7616–7624, doi:10.1021/es050896d, 2005.
- Liggio, J. and McLaren, R.: An optimized method for the determination of volatile and semi-volatile aldehydes and ketones in ambient particulate matter, *Int. J. Environ. Anal. Chem.*, 83, 819–835, doi:10.1080/03067310310001597653, 2003.

- Martin, N. A., Marlow, D. J., Henderson, M. H., Goody, B. A., and Quincey, P. G.: Studies using the sorbent Carbopack X for measuring environmental benzene with PerkinElmer-type pumped and diffusive samplers, *Atmos. Environ.*, 37, 871–879, doi:10.1016/S1352-2310(02)01000-2, 2003.
- Matsunaga, S., Mochida, M., and Kawamura, K.: High abundance of gaseous and particulate 4-oxopentanal in the forestal atmosphere, *Chemosphere*, 55, 1143–1147, doi:10.1016/j.chemosphere.2003.10.004, 2004.
- Monks, P. S., Granier, C., Fuzzi, S., Stohl, A., Williams, M. L., Aki-moto, H., Amann, M., Baklanov, A., Baltensperger, U., Bey, I., Blake, N., Blake, R. S., Carslaw, K., Cooper, O. R., Dentener, F., Fowler, D., Fragkou, E., Frost, G. J., Generoso, S., Ginoux, P., Grewe, V., Guenther, A., Hansson, H. C., Henne, S., Hjorth, J., Hofzumahaus, A., Huntrieser, H., Isaksen, I. S. A., Jenkin, M. E., Kaiser, J., Kanakidou, M., Klimont, Z., Kulmala, M., Laj, P., Lawrence, M. G., Lee, J. D., Liousse, C., Maione, M., McFiggans, G., Metzger, A., Mieville, A., Moussiopoulos, N., Orlando, J. J., O'Dowd, C. D., Palmer, P. I., Parrish, D. D., Petzold, A., Platt, U., Pöschl, U., Prévôt, A. S. H., Reeves, C. E., Reimann, S., Rudich, Y., Sellegri, K., Steinbrecher, R., Simpson, D., ten Brink, H., Theloke, J., van der Werf, G. R., Vautard, R., Vestreng, V., Vlachokostas, C., and von Glasow, R.: Atmospheric composition change - global and regional air quality, *Atmos. Environ.*, 43, 5268–5350, doi:10.1016/j.atmosenv.2009.08.021, 2009.
- Müller, K., Haferkorn, S., Grabmer, W., Wisthaler, A., Hansel, A., Kreuzwieser, J., Cojocariu, C., Rennenberg, H., and Herrmann, H.: Biogenic carbonyl compounds within and above a coniferous forest in Germany, *Atmos. Environ.*, 40, Supplement 1, 81–91, doi:10.1016/j.atmosenv.2005.10.070, 2006.
- Myrdal, P. B. and Yalkowsky, S. H.: Estimating pure component vapor pressures of complex organic molecules, *Ind. Eng. Chem. Res.*, 36, 2494–2499, doi:10.1021/ie950242l, 1997.
- Nannoolal, Y., Rarey J., and Ramjugernath, D.: Estimation of pure component properties: Part 3. Estimation of the vapor pressure of non-electrolyte organic compounds via group contributions and group interactions, *Fluid Phase Equilib.*, 269, 117–133, doi:10.1016/j.fluid.2008.04.020, 2008.
- Nolte, C. G., Schauer, J. J., Cass, G. R., and Simoneit, B. R. T.: Highly polar organic compounds present in wood smoke and in the ambient atmosphere, *Environ. Sci. Technol.*, 35, 1912–1919, doi:10.1021/es001420r, 2001.
- Northcross, A. L. and Jang, M.: Heterogeneous SOA yield from ozonolysis of monoterpenes in the presence of inorganic acid, *Atmos. Environ.*, 41, 1483–1493, doi:10.1016/j.atmosenv.2006.10.009, 2007.
- Odum, J. R., Hoffmann, T., Bowman, F., Collins, D., Flagan, R. C., and Seinfeld, J. H.: Gas/particle partitioning and secondary organic aerosol yields, *Environ. Sci. Technol.*, 30, 2580–2585, 1996.
- Orasche, J., Schnelle-Kreis, J., Abbaszade, G., and Zimmermann, R.: Technical Note: In-situ derivatization thermal desorption GC-TOFMS for direct analysis of particle-bound non-polar and polar organic species, *Atmos. Chem. Phys.*, 11, 8977–8993, doi:10.5194/acp-11-8977-2011, 2011.
- Ortiz, R., Enya, K., Sekiguchi, K., and Sakamoto, K.: Experimental testing of an annular denuder and filter system to measure gas-particle partitioning of semivolatile bifunctional carbonyls in the atmosphere, *Atmos. Environ.*, 43, 382–388, doi:10.1016/j.atmosenv.2008.09.074, 2009.
- Ortiz, R., Enya, K., and Sakamoto, K.: Experimental Determination of the Partition Coefficient for Bifunctional Carbonyls in the Atmosphere and in Smog Chamber, in: *NATO Science for Peace and Security Series B-Physics and Biophysics*, edited by: Noto-Chal. M., 265–269, 2010.
- Pacolay, B. D., Ham, J. E., Slaven, J. E., and Wells, J. R.: Feasibility of detection and quantification of gas-phase carbonyls in indoor environments using PFBHA derivatization and solid-phase microextraction (SPME), *J. Environ. Monit.*, 10, 853–860, doi:10.1039/b801926f, 2008.
- Pankow, J. F.: An absorption model of gas/particle partitioning of organic compounds in the atmosphere, *Atmos. Environ.*, 28, 185–188, doi:10.1016/1352-2310(94)90093-0, 1994.
- Pankow, J. F.: On the ability of the gas/particle partitioning constant  $K_p$  to consider the effects of mean MW and the presence of high MW compounds, *Atmos. Environ.*, 45, 1213–1216, doi:10.1016/j.atmosenv.2010.11.041, 2011.
- Pankow, J. F. and Asher, W. E.: SIMPOL.1: a simple group contribution method for predicting vapor pressures and enthalpies of vaporization of multifunctional organic compounds, *Atmos. Chem. Phys.*, 8, 2773–2796, doi:10.5194/acp-8-2773-2008, 2008.
- Parikh, H. M., Carlton, A. G., Vizuete, W., and Kamens, R. M.: Modeling secondary organic aerosol using a dynamic partitioning approach incorporating particle aqueous-phase chemistry, *Atmos. Environ.*, 45, 1126–1137, doi:10.1016/j.atmosenv.2010.11.027, 2011.
- Pietrogrande, M. C. and Bacco, D.: GC-MS analysis of water-soluble organics in atmospheric aerosol: Response surface methodology for optimizing silyl-derivatization for simultaneous analysis of carboxylic acids and sugars, *Anal. Chim. Acta*, 689, 257–264, doi:10.1016/j.aca.2011.01.047, 2011.
- Pietrogrande, M. C., Bacco, D., and Mercuriali, M.: GC-MS analysis of low-molecular-weight dicarboxylic acids in atmospheric aerosol: comparison between silylation and esterification derivatization procedures, *Anal. Bioanal. Chem.*, 396, 877–885, 2010.
- Pol, J., Hohnova, B., Jussila, M., and Hyotylainen, T.: Comprehensive two-dimensional liquid chromatography-time-of-flight mass spectrometry in the analysis of acidic compounds in atmospheric aerosols, *J. Chromatogr. A*, 1130, 64–71, doi:10.1016/j.chroma.2006.04.050, 2006.
- Prather, K. A., Hatch, C. D., and Grassian, V. H.: Analysis of Atmospheric Aerosols, in: *Annu. Rev. Anal. Chem.*, Annual Reviews, Palo Alto, 485–514, 2008.
- Rudich, Y.: Laboratory Perspectives on the Chemical Transformations of Organic Matter in Atmospheric Particles, *Chem. Rev.*, 103, 5097–5124, doi:10.1021/cr020508f, 2003.
- Rudich, Y., Donahue, N. M., and Mentel, T. F.: Aging of organic aerosol: Bridging the gap between laboratory and field studies, *Annu. Rev. Phys. Chem.*, 58, 321–352, doi:10.1146/annurev.physchem.58.032806.104432, 2007.
- Saathoff, H., Naumann, K.-H., Möhler, O., Jonsson, Å. M., Hallquist, M., Kiendler-Scharr, A., Mentel, Th. F., Tillmann, R., and Schurath, U.: Temperature dependence of yields of secondary organic aerosols from the ozonolysis of  $\alpha$ -pinene and limonene, *Atmos. Chem. Phys.*, 9, 1551–1577, doi:10.5194/acp-9-1551-2009, 2009.

- Sarwar, G. and Corsi, R.: The effects of ozone/limonene reactions on indoor secondary organic aerosols, *Atmos. Environ.*, 41, 959–973, doi:10.1016/j.atmosenv.2006.09.032, 2007.
- Sato, K., Takami, A., Iozaki, T., Hikida, T., Shimono, A., and Imamura, T.: Mass spectrometric study of secondary organic aerosol formed from the photo-oxidation of aromatic hydrocarbons, *Atmos. Environ.*, 44, 1080–1087, doi:10.1016/j.atmosenv.2009.12.013, 2010.
- Saunders, S. M., Jenkin, M. E., Derwent, R. G., and Pilling, M. J.: Protocol for the development of the Master Chemical Mechanism, MCM v3 (Part A): tropospheric degradation of non-aromatic volatile organic compounds, *Atmos. Chem. Phys.*, 3, 161–180, doi:10.5194/acp-3-161-2003, 2003.
- Seinfeld, J. H. and Pankow, J. F.: Organic atmospheric particulate material, *Annu. Rev. Phys. Chem.*, 54, 121–140, doi:10.1146/annurev.physchem.54.011002.103756, 2003.
- Sheesley, R. J., DeMinter, J. T., Meiritz, M., Snyder, D. C., and Schauer, J. J.: Temporal trends in motor vehicle and secondary organic tracers using in situ methylation thermal desorption GC-MS, *Environ. Sci. Technol.*, 44, 9398–9404, doi:10.1021/es102301t, 2010.
- Shimmo, M., Jantti, J., Aalto, P., Hartonen, K., Hyötyläinen, T., Kulmala, M., and Riekkola, M.-L.: Characterisation of organic compounds in aerosol particles from a Finnish forest by on-line coupled supercritical fluid extraction-liquid chromatography-gas chromatography-mass spectrometry, *Anal. Bioanal. Chem.*, 378, 1982–1990, 2004.
- Singer, B. C., Destailhats, H., Hodgson, A. T., and Nazaroff, W. W.: Cleaning products and air fresheners: emissions and resulting concentrations of glycol ethers and terpenoids, *Indoor Air*, 16, 179–191, 2006.
- Sinha, V., Williams, J., Lelieveld, J., Ruuskanen, T. M., Kajos, M. K., Patokoski, J., Hellen, H., Hakola, H., Mogensen, D., Boy, M., Rinne, J., and Kulmala, M.: OH reactivity measurements within a boreal forest: evidence for unknown reactive emissions, *Environ. Sci. Technol.*, 44, 6614–6620, doi:10.1021/es101780b, 2010.
- Sullivan, R. C. and Prather, K. A.: Recent advances in our understanding of atmospheric chemistry and climate made possible by on-line aerosol analysis instrumentation, *Anal. Chem.*, 77, 3861–3886, doi:10.1021/ac050716i, 2005.
- Sun, J. and Ariya, P. A.: Atmospheric organic and bio-aerosols as cloud condensation nuclei (CCN): A review, *Atmos. Environ.*, 40, 795–820, doi:10.1016/j.atmosenv.2005.05.052, 2006.
- Sun, T., Wang, Y., Zhang, C., Sun, X., and Wang, W.: The chemical mechanism of the limonene ozonolysis reaction in the SOA formation: A quantum chemistry and direct dynamic study, *Atmos. Environ.*, 45, 1725–1731, doi:10.1016/j.atmosenv.2010.12.054, 2011.
- Szmigielski, R., Surratt, J. D., Vermeylen, R., Szmigielska, K., Kroll, J. H., Ng, N. L., Murphy, S. M., Sorooshian, A., Seinfeld, J. H., and Claeys, M.: Characterization of 2-methylglyceric acid oligomers in secondary organic aerosol formed from the photooxidation of isoprene using trimethylsilylation and gas chromatography/ion trap mass spectrometry, *J. Mass Spectrom.*, 42, 101–116, 2007.
- Temime, B., Healy, R. M., and Wenger, J. C.: A denuder-filter sampling technique for the detection of gas and particle phase carbonyl compounds, *Environ. Sci. Technol.*, 41, 6514–6520, doi:10.1021/es070802v, 2007.
- Tsimpidi, A. P., Karydis, V. A., Zavala, M., Lei, W., Molina, L., Ulbrich, I. M., Jimenez, J. L., and Pandis, S. N.: Evaluation of the volatility basis-set approach for the simulation of organic aerosol formation in the Mexico City metropolitan area, *Atmos. Chem. Phys.*, 10, 525–546, doi:10.5194/acp-10-525-2010, 2010.
- Valorso, R., Aumont, B., Camredon, M., Raventos-Duran, T., Mouchel-Vallon, C., Ng, N. L., Seinfeld, J. H., Lee-Taylor, J., and Madronich, S.: Explicit modelling of SOA formation from  $\alpha$ -pinene photooxidation: sensitivity to vapour pressure estimation, *Atmos. Chem. Phys.*, 11, 6895–6910, doi:10.5194/acp-11-6895-2011, 2011.
- Vivanco, M. G., Santiago, M., Martinez-Tarifa, A., Borrás, E., Rodenas, M., Garcia-Diego, C., and Sanchez, M.: SOA formation in a photoreactor from a mixture of organic gases and HONO for different experimental conditions, *Atmos. Environ.*, 45, 708–715, doi:10.1016/j.atmosenv.2010.09.059, 2011.
- Volckens, J. and Leith, D.: Effects of sampling bias on gas-particle partitioning of semi-volatile compounds, *Atmos. Environ.*, 37, 3385–3393, doi:10.1016/S1352-2310(03)00356-X, 2003.
- Volkamer, R., Martini, F. S., Molina, L. T., Salcedo, D., Jimenez, J. L., and Molina, M. J.: A missing sink for gas-phase glyoxal in Mexico City: Formation of secondary organic aerosol, *Geophys. Res. Lett.*, 34, 5, doi:10.1029/2007gl030752, 2007.
- Walser, M. L., Park, J., Gomez, A. L., Russell, A. R., and Nizkorodov, S. A.: Photochemical aging of secondary organic aerosol particles generated from the oxidation of d-limonene, *J. Phys. Chem. A*, 111, 1907–1913, doi:10.1021/jp066293l, 2007.
- Wan, E. C. H. and Yu, J. Z.: Analysis of sugars and sugar polyols in atmospheric aerosols by chloride attachment in liquid chromatography/negative ion electrospray mass spectrometry, *Environ. Sci. Technol.*, 41, 2459–2466, doi:10.1021/es062390g, 2007.
- Wang, J., Doussin, J. F., Perrier, S., Perraudin, E., Katrib, Y., Pangu, E., and Picquet-Varrault, B.: Design of a new multi-phase experimental simulation chamber for atmospheric photochemistry, aerosol and cloud chemistry research, *Atmos. Meas. Tech.*, 4, 2465–2494, doi:10.5194/amt-4-2465-2011, 2011.
- Wedel, A., Müller, K.-P., Ratte, M., and Rudolph, J.: Measurements of Volatile Organic Compounds (VOC) during POPCORN 1994: Applying a new on-line GC-MS technique, *J. Atmos. Chem.*, 31, 73–103, 1998.
- Williams, B. J., Goldstein, A. H., Kreisberg, N. M., and Herling, S. V.: An In-Situ Instrument for Speciated Organic Composition of Atmospheric Aerosols: Thermal Desorption Aerosol GC/MS-FID (TAG), *Aerosol Sci. Tech.*, 40, 627–638, doi:10.1080/02786820600754631, 2006.
- Woo, K.-L. and Kim, J.-I.: New hydrolysis method for extremely small amount of lipids and capillary gas chromatographic analysis as N(O)-tert.-butyldimethylsilyl fatty acid derivatives compared with methyl ester derivatives, *J. Chromatogr. A*, 862, 199–208, doi:10.1016/S0021-9673(99)00934-6, 1999.
- Wu, L. J. and Hee, S. S. Q.: A solid sorbent personal air sampling method for aldehydes, *Am. Ind. Hyg. Assoc. J.*, 56, 362–367, doi:10.1080/15428119591016980, 1995.
- Wu, T., Wang, X., Li, D., Sheng, G., and Fu, J.: Determination of trace volatile fatty acids in ambient air by capillary gas chromatography-mass spectrometry in SIM mode, *Int. J. Environ. Anal. Chem.*, 88, 1107–1115, doi:10.1080/03067310802447034, 2008.

- Xu, X., Stee, L. L. P., Williams, J., Beens, J., Adahchour, M., Vreuls, R. J. J., Brinkman, U. A., and Lelieveld, J.: Comprehensive two-dimensional gas chromatography (GC  $\times$  GC) measurements of volatile organic compounds in the atmosphere, *Atmos. Chem. Phys.*, 3, 665–682, doi:10.5194/acp-3-665-2003, 2003.
- Yasmeen, F., Szmigielski, R., Vermeylen, R., Gómez-González, Y., Surratt, J. D., Chan, A. W. H., Seinfeld, J. H., Maenhaut, W., and Claeys, M.: Mass spectrometric characterization of isomeric terpenoic acids from the oxidation of  $\alpha$ -pinene,  $\beta$ -pinene, d-limonene, and  $\Delta^3$ -carene in fine forest aerosol, *J. Mass Spectrom.*, 46, 425–442, 2011.
- Yu, J., Cocker, D. R., Griffin, R. J., Flagan, R. C., and Seinfeld, J. H.: Gas-phase ozone oxidation of monoterpenes: gaseous and particulate products, *J. Atmos. Chem.*, 34, 207–258, 1999.
- Zhang, Q., Jimenez, J. L., Canagaratna, M. R., Ulbrich, I. M., Ng, N. L., Worsnop, D. R., and Sun, Y.: Understanding atmospheric organic aerosols via factor analysis of aerosol mass spectrometry: a review, *Anal. Bioanal. Chem.*, 401, 3045–3067, 2011.