



# Application of time-of-flight aerosol mass spectrometry for the real-time measurement of particle-phase organic peroxides: an online redox derivatization–aerosol mass spectrometer (ORD-AMS)

Marcel Weloe and Thorsten Hoffmann

Department of Chemistry, Johannes Gutenberg University, 55128 Mainz, Germany

**Correspondence:** Thorsten Hoffmann (t.hoffmann@uni-mainz.de)

Received: 25 October 2019 – Discussion started: 8 November 2019

Revised: 20 July 2020 – Accepted: 28 July 2020 – Published: 28 October 2020

**Abstract.** Aerosol mass spectrometers (AMS) are frequently applied in atmospheric aerosol research in connection with climate, environmental or health-related projects. This is also true for the measurement of the organic fraction of particulate matter, still the least understood group of components contributing to atmospheric aerosols. While quantification of the organic and/or inorganic aerosol fractions is feasible, more detailed information about individual organic compounds or compound classes can usually not be provided by AMS measurements. In this study, we present a new method to detect organic peroxides in the particle phase in real-time using an AMS. Peroxides (ROOR') are of high interest to the atmospheric aerosol community due to their potentially high mass contribution to SOA, their important role in new particle formation and their function as “reactive oxygen species” in aerosol–health-related topics. To do so, supersaturated gaseous triphenylphosphine (TPP) was continuously mixed with the aerosol flow of interest in a condensation/reaction volume in front of the AMS inlet. The formed triphenylphosphine oxide (TPPO) from the peroxide–TPP reaction was then detected by an aerosol mass spectrometer (AMS), enabling the semiquantitative determination of peroxide with a time resolution of 2 min. The method was tested with freshly formed and aged biogenic VOC and ozone SOA as well as in a short proof-of-principle study with ambient aerosol.

## 1 Introduction

Atmospheric aerosols attract attention in atmospheric research owing to their effects on climate and human health. The climate relevance of aerosol particles arises from their influence on the transmissivity of the atmosphere for solar radiation and on cloud properties (McNeill, 2017). Secondary organic aerosols (SOAs) are a major contributor to tropospheric particulate matter (Hallquist et al., 2009; Pöschl, 2005). One highly relevant class of compounds which is assumed to play an essential role in SOA formation, including participation in new particle formation, are organic peroxides (Bonn et al., 2004; Mochida et al., 2006; Reinnig et al., 2008, 2009; Tobias et al., 2000; Tobias and Ziemann, 2002; Zhou et al., 2018; Bianchi et al., 2016, 2017, 2019; Ziemann, 2002, 2003; Kahnt et al., 2018; Zhang et al., 2017; Lee et al., 2019). Besides their significant contribution to SOA mass (Docherty et al., 2005; Ziemann and Atkinson, 2012), particle-phase peroxides are especially interesting due to their often very low saturation vapor pressure, particularly when they contain several hydroperoxy groups induced by autoxidation (Tröstl et al., 2016; Krapf et al., 2016). Also their chemical properties such as their oxidizing character (Claffin et al., 2018) or their function as an indicator for the oxidizing capacity of the troposphere (Thompson, 1992; Lelieveld et al., 2008) create a special interest in the analysis of organic peroxides. Finally, peroxide functionalities are likely to play an important role in health-related effects of SOA, since organic peroxides are believed to contribute to the formation of reactive organic species (ROS) by transporting oxidants on and/or in particles

into the respiratory system (Verma et al., 2015; Wragg et al., 2016; Fuller et al., 2014; Steimer et al., 2017; Tong et al., 2018).

The fact that organic peroxides are oxidizing agents is often used in their analysis. One of the oldest and most frequently applied techniques to measure and quantify peroxides is iodometry (Mertes et al., 2012; Docherty et al., 2005; Mutzel et al., 2013; Zhao et al., 2018). Alternatively, triphenylphosphine (TPP) can be used as a reacting agent. TPP rapidly reacts with hydrogen peroxide, organic (hydro)peroxides and peroxy acids to triphenylphosphine oxide (TPPO) (Docherty et al., 2004; Ruiz et al., 2001), which can be detected spectrometrically (Porter et al., 1979). There are various suggestions about the mechanism of the reaction. One of the mechanisms involves a bimolecular nucleophilic displacement of TPP on the O–O bond resulting in a bipolar transition state (Eq. 1) (Jarvie et al., 1971; Chellamani and Suresh, 1988). Two other suggested mechanisms proceed via a cyclic peroxide intermediates (Eqs. 2 and 3) (Davies and Feld, 1958; Jayanthi and Koshore, 1997). The main application of the TPP–TPPO method is the determination of hydroperoxides from the autoxidation of oils and fats (Gotoh et al., 2011; Nakamura and Maeda, 1991; Wang et al., 2016). Compared to iodometric peroxide determination, the TPP–TPPO method is insensitive to moisture, and the reaction with oxygen is much slower.

Another technique that has been used for the analysis of organic peroxides is mass spectrometry (Krawczyk and Baj, 2014), for example for the measurement of SOA-related organic peroxides by direct infusion of liquid samples into an atmospheric-pressure chemical ionization tandem mass spectrometer (Zhou et al., 2018). However, all the above-mentioned techniques are offline methods relying on filter sampling techniques, which have disadvantages such as the evaporation of semivolatile species or chemical alterations of labile compounds during sampling. Obviously the temporal resolution of offline methods is also limited. Therefore, online methods for organic peroxide measurements would be beneficial.

One of the main online techniques that is used in the field of aerosol research is the aerosol mass spectrometer (AMS), which is commercially available from Aerodyne Research Inc. (Billerica, MA, USA). Aerosol particles are introduced into a vacuum chamber through an aerodynamic lens that focuses the sampled aerosol particles to a beam (Jayne et al., 2000). The particle beam is directed onto an evaporation unit, which is heated to 600 °C, ideally volatilizing the entire particle mass. The evaporated compounds are subsequently ionized by electron ionization (EI) at 70 eV and analyzed by MS (Pratt and Prather, 2012; Jayne et al., 2000). The application of EI allows the analysis of a wide range of chemical species and provides robust quantitative information on various non-refractory ambient aerosol components since the ionization efficiency of EI is much less analyte and matrix dependent than, for example, chemical ionization techniques (Allan et

al., 2003b, a; Canagaratna et al., 2007). However, EI induces a high degree of fragmentation of organic compounds due to excess internal energy imparted during the ionization processes. Usually, these fragmentation steps impede the identification and quantification of individual particle-phase organics, thus leading to a reduced level of chemical information (Zhang et al., 2011). Therefore, recent mass spectrometric instrumental developments for organic aerosol analysis focus on softer ionization techniques such as atmospheric pressure chemical ionization (APCI-MS) (Vogel et al., 2013b, a; Brüggemann et al., 2014; Warscheid and Hoffmann, 2001a, b), flowing atmospheric-pressure afterglow mass spectrometry (AeroFAPA-MS) (Brüggemann et al., 2015, 2016, 2017) or extractive electrospray ionization (EESI-MS) (Gallimore et al., 2017). However, while these mass spectrometric techniques are applicable for a wide spectrum of organic compounds and functionalities, for example, carboxylic acids (Hallquist et al., 2009; Yatavelli et al., 2015), unambiguous identification of organic peroxides using mass spectrometric techniques is still challenging, since in contrast to other functionalities the molecular ions usually fragment resulting from the cleavage of the peroxide bond and/or further fragmentation (Reinnig et al., 2009).

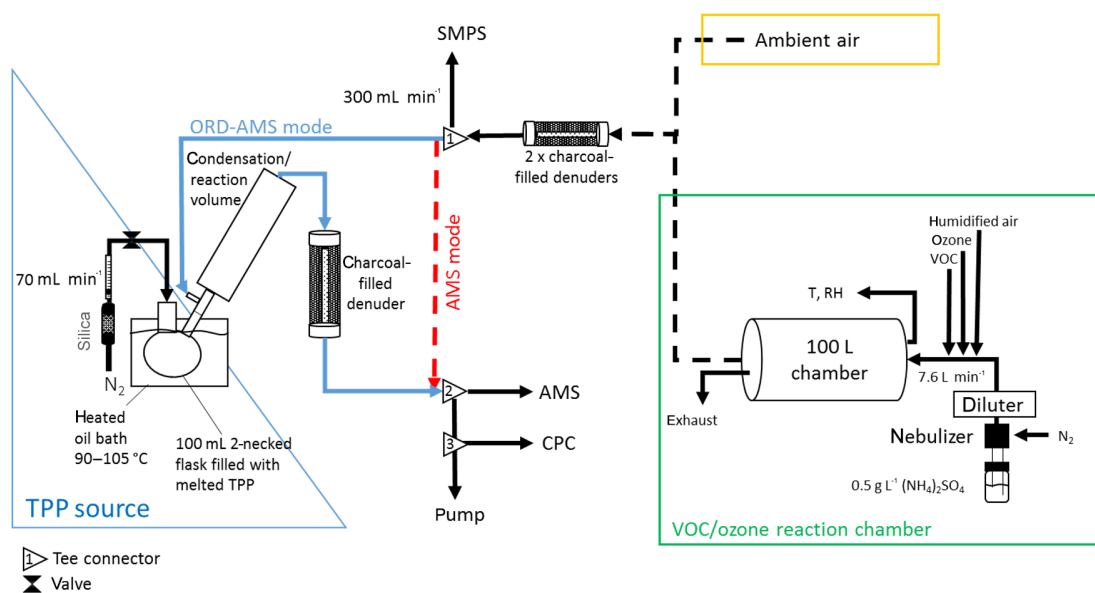
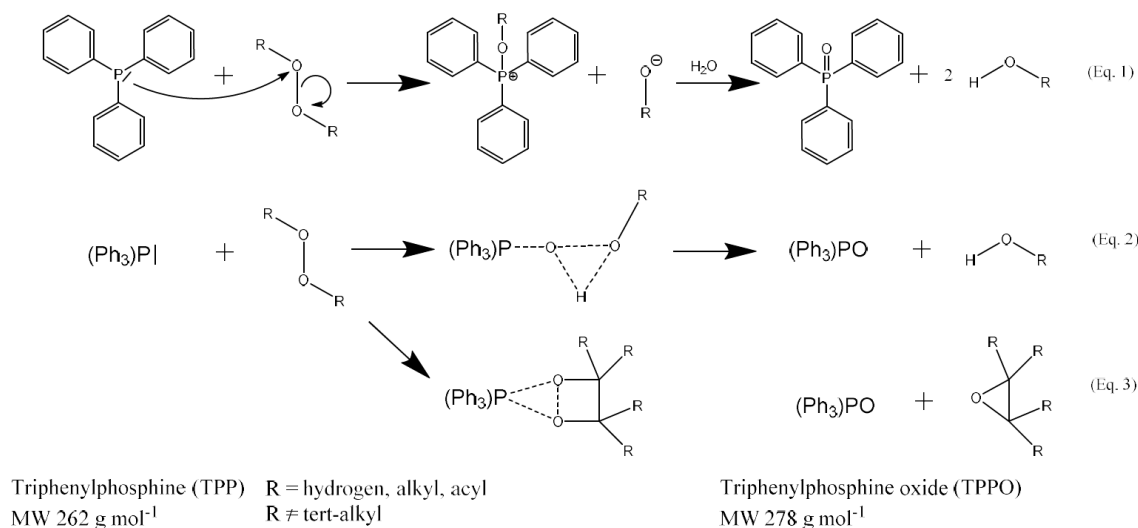
Here, we have combined the concept of selective peroxide determination by a suitable redox reaction (the TPP–TPPO system) to use this reaction as an online derivatization method to monitor the educts and products with an aerosol mass spectrometer (online redox derivatization–AMS, ORD-AMS). The development and application of the method presented in this paper is essentially a proof-of-principle study; however, the results clearly show the potential of online derivatization as an extension of regular AMS measurements, in this case for the understanding of particle-phase organic peroxides. Similar experimental setups with different derivatization agents are also conceivable in the future to further extend the usability of AMS systems.

## 2 Experimental

### 2.1 Online redox derivatization

Figure 1 shows a schematic diagram of the experimental setup for the AMS online measurements. Gas-phase organics were firstly removed by two charcoal-filled denuders (inner tube: diameter = 2 cm and length = 40 cm; outer tube: diameter = 13.5 cm, length = 40 cm), and the particle size distribution was measured using a scanning mobility particle sizer (SMPS; Grimm Aerosol Technik, Ainring, Germany).

When the instrument was used in the ORD-AMS mode (blue path in Fig. 1) the particles to be analyzed were introduced into a 500 mL glass tube (35 cm length; 4.5 cm max. diameter), the condensation/reaction volume. Here, the aerosol was mixed with the gaseous reactant (TPP; 99 %, Acros Organics BVBA, Geel, Belgium), which was supplied



**Figure 1.** Experimental setup: the aerosol is either generated with a 100 L reaction chamber via VOC–O<sub>3</sub> reaction, usually also introducing AS seed particles, or supplied from outside of the chemistry building. The aerosol is then led to the aerosol mass spectrometer, either directly (AMS mode, red arrow) or through the online derivatization unit (ORD-AMS mode) in which the test aerosol is mixed with a nitrogen stream containing supersaturated gaseous TPP. Remaining gaseous TPP is removed by charcoal-filled denuder after the condensation/reaction volume. Most of the time the aerosol is characterized by the ToF-AMS, an SMPS system and a CPC.

from a Sinclair–La Mer setup (see Fig. 1, blue triangle). The TPP source consisted of a 100 mL two-necked, round-bottom glass filled with TPP and was heated to a temperature of 90 °C. Nitrogen was used as a carrier gas to deliver the TPP to the condensation/reaction volume, where TPP condensed on the available particle surface. After passing the condensation/reaction volume the aerosol was led through a third denuder filled with active charcoal to remove gas-phase TPP and finally introduced into the AMS and a condensation particle counter (CPC; PortaCount Plus, TSI Inc., Shoreview, MN, USA). A total flow rate of 1.74 L min<sup>-1</sup> was obtained

using an additional pump. The residence time of the aerosol in the reaction volume and the denuder to the AMS was about 35 s.

When the instrument was used in the AMS mode (red path in Fig. 1), the aerosol was led directly to the AMS and the CPC. Switching between the two modes was achieved by manually changing the PTFE connections to avoid losses or artifacts from additional valves. The data acquisition of the AMS and SMPS was interrupted during this time and continued (1–2 min) after stabilization of the particle number concentration measured by the CPC. As mentioned above, a ni-

trogen stream enriched with TPP was added to the aerosol stream in the ORD-AMS mode, resulting in a dilution of the aerosol. The correction for this dilution was done by comparing the particle number concentrations, measured in the ORD-AMS and AMS mode using the CPC. The ratio between the particle number concentrations of the two modes was used as a dilution factor.

## 2.2 ToF-AMS

The aerosol particles were introduced into the time-of-flight (ToF)-AMS (Aerodyne Research Inc., Billerica, MA, USA) at a sampling flow rate of  $1.23 \text{ mL s}^{-1}$ . An aerodynamic lens system focuses the particles in a size range of  $\sim 35\text{--}1000 \text{ nm}$  into the vacuum system. While most of the remaining gas phase is removed, the particle beam passes into the particle-sizing chamber. The ToF-AMS was either operated in the mass spectrum (MS) or the particle time-of-flight (PToF) mode. The first mode provides information about the overall composition of the total aerosol mass without any information about the particle size, while the second mode results in the size-segregated analysis of the particle composition. For the latter one, a chopper allows the transmission of only a small part of the aerosol into the PToF chamber. Due to increasing inertia with larger particle size, the particle package is separated by the particles' time of flight, and the size distribution based on the mean vacuum aerodynamic diameter of the aerosol can be determined. In this PToF mode, the EI-ToF-MS functions as a detector for the particle time of flight. To determine the concentration in the MS mode, the beam passes the chopper alternately blocked or unblocked for background corrections. After passing the particle-sizing chamber, the aerosol particles are vaporized on a heated porous tungsten surface at  $600^\circ\text{C}$ , followed by immediate ionization in electron impact (EI) mode at  $70 \text{ eV}$ . The generated ions are extracted by an orthogonal extractor into the ToF-MS, which was operated in the V mode to give higher sensitivity. The MS spectra acquisition was performed in the positive ion mode. The AMS data were processed by the Igor (Version Pro 6.37)-based programs "SQUIRREL 1.57 G" (unit mass resolution) and "PIKA 1.16H" (high resolution) (DeCarlo et al., 2006; Canagaratna et al., 2007).

## 2.3 Physical aerosol characterization and generation

A scanning mobility particle sizer was used to measure the size distributions of the particle number concentration of the incoming aerosol, which was used to obtain particle surface, volume and mass concentration. The conversion of the volume concentration into the mass concentration for the ozonolysis experiments is described in detail in the Supplement. Besides the measurement of the particle size distribution, the instrument was used to generate monodisperse particles for instrument calibration. The EI spectra of the derivatization agent TPP and its reaction product with per-

oxides TPPO ( $> 98\%$ , Merck KGaA, Darmstadt, Germany) were recorded using a nebulizer as a particle generator. Pure TPP and TPPO particles were produced from solutions in methanol (Optima<sup>®</sup> LC/MS grade, Fisher Scientific, Loughborough, UK) and *n*-heptane (HPLC grade, Carl Roth GmbH + Co. KG, Karlsruhe, Germany), respectively. For these experiments, the particles were size-selected using a differential mobility analyzer (DMA) in a setup described in the Supplement.

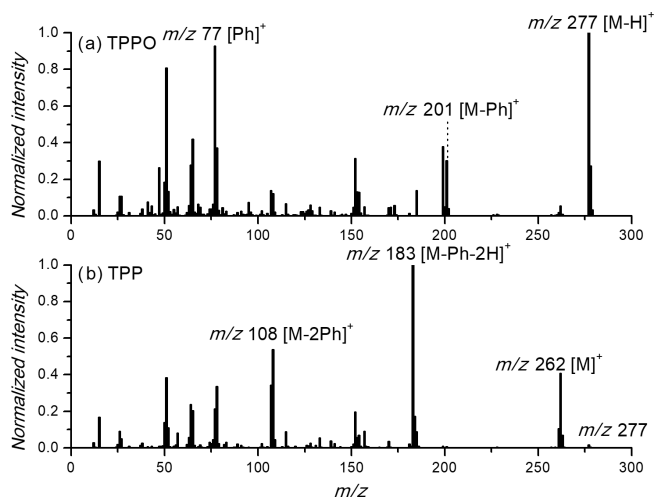
## 2.4 Reaction chamber as SOA generator

For the proof-of-principle studies with SOA, a laboratory-scale reaction chamber in continuous-flow tank reactor mode was used. Synthetic air was supplied with a total flow of  $7.6 \text{ L min}^{-1}$  into the  $100 \text{ L}$  chamber, resulting in a residence time of  $13.5 \text{ min}$ . The total flow was composed of a gas stream through a diffusion test gas source (Thorenz et al., 2012) ( $0.600 \text{ L min}^{-1}$ ), where the VOC of interest was added to the synthetic air supply. A second gas stream ( $2.6 \text{ L min}^{-1}$ ) was humidified using a gas-washing bottle, and a third gas flow was led through an ozone generator (Dasibi Environmental Corp. Model 1008 RS O<sub>3</sub> analyzer, Glendale, CA, USA;  $2.7 \text{ L min}^{-1}$ ). To add seed particles an aqueous solution of ammonium sulfate (ca.  $0.5 \text{ g L}^{-1}$ ;  $> 99.5\%$ , Merck KGaA, Darmstadt, Germany) was nebulized using technical nitrogen and introduced into the chamber ( $1.7 \text{ L min}^{-1}$ ). Temperature and relative humidity were  $25^\circ\text{C}$  and  $61\%$  for the  $\alpha$ -pinene ozonolysis and  $22^\circ\text{C}$  and  $59\%$  for the  $\beta$ -pinene ozonolysis. They were monitored by a thermo-hygrometer (Amarell, ama-digit ad 910h, Kreuzwertheim, Germany). The chamber was set to be under slight overpressure to prevent the entry of laboratory air into the setup. The reaction chamber was wrapped with aluminum foil to avoid photochemical reactions. For the ozonolysis of  $\alpha$ -pinene ( $99\%$ , Fluka, Seelze, Germany), the chamber was first equilibrated with the terpene before the ozone generator was switched on. For the  $\beta$ -pinene ( $99\%$ , Sigma Aldrich Chemie GmbH, Steinheim, Germany) experiments, the order of the addition of the individual compounds was reversed – i.e., ozone was added before the terpene. The ozone concentration was estimated to be about  $1 \text{ ppm}$ , while the  $\alpha$ -pinene concentration was set to  $290 \text{ ppbv}$  and the  $\beta$ -pinene concentration to  $380 \text{ ppbv}$ . To remove gaseous organic compounds, the SOA formed in the reaction chamber was led through two diffusion denuders filled with activated charcoal.

## 3 Results and discussion

### 3.1 EI mass spectra

Figure 2 shows the obtained mass spectra of TPPO and TPP. The EI spectra of the two reference compounds measured by the AMS are very similar to the spectra measured by the regular MS setup (Williams et al., 1968; Stein and Slaw-



**Figure 2.** Mass spectra of (a) triphenylphosphine oxide (TPPO) and (b) triphenylphosphine (TPP).

son, 1963). The base peak for TPPO is at  $m/z$  277 resulting from a hydrogen abstraction,  $[M-1]^+$ , from the molecular ion – molar weight MW (TPPO) = 278 g mol<sup>-1</sup>). The second-largest peak is  $m/z$  77 formed from the abstraction of a phenyl cation. The base peak for TPP is located at  $m/z$  183, originating from the abstraction of two hydrogens and a phenyl group, probably a result of an aromatic stabilization of the resulting dibenzophosphole cation (Williams et al., 1968). However, the molecular ion of TPP at  $m/z$  262 is also visible. More important is the small but still observable  $[M-1]^+$  ion of TPPO in the TPP spectra. Since the peroxide determination is based on the measurement of TPPO, this background signal has to be taken into account (see below). Finally, aerosol species typically found in ambient air and measured by the AMS have mass spectra with signals mainly below  $m/z$  100 due to the fragmentation after EI ionization. As seen also in Figs. S4 and S5 in the Supplement, the signals at  $m/z$  277 and 262 are specific for TPPO and TPP, respectively. These signals can be used for the quantification of both compounds.

Experiments with peroxides (ROOR') as reference compounds were also performed, using commercially available organic peroxides, e.g. lauryl peroxide and benzoyl peroxide. While these experiments clearly showed the formation of TPPO from TPP, the use of these compounds for quantification was not possible, since the peroxide standards studied undergo a substantial change in volatility after the reaction with TPP. Therefore, lauryl peroxide and benzoyl peroxide were used to demonstrate the general proof of feasibility of the method but not, however, for calibration.

### 3.2 Quantification

The mass concentration  $c_m$  of a single compound measured by an AMS can be expressed as (Canagaratna et al., 2007)

$$c_m = \frac{10^{12} \cdot \text{MW}(\text{compound})}{\text{IE}(\text{compound}) \cdot Q \cdot N_A} \cdot \sum_{\text{all}, i} I_{\text{compound}, i}, \quad (4)$$

where MW and IE are the molar weight and the ionization efficiency of an individual compound, respectively. The ionization efficiency (IE) is a dimensionless quantity that describes the ionization and detection efficiency and is defined as the ratio of the ions detected to the parent molecules vaporized.  $Q$  is the flow rate into the AMS,  $N_A$  Avogadro's number and  $10^{12}$  a unit conversion factor.  $I_{\text{compound}, i}$  is the ion rate of an ion  $i$  formed by ionization and fragmentation of a compound. Since the fragmentation pattern can be assumed to be robust, it is possible to use the ion yield of a single ion  $I_{m/z, i}$  to calculate the concentration. Based on the stoichiometry of the reaction, the molar concentration  $c_n$  of peroxides is the same as the concentration of TPPO and can be calculated from the signal at  $m/z$  277 as

$$c_n(\text{TPPO}) = \frac{10^{12}}{\text{IE}\left(\text{TPPO}, \frac{m}{z} i\right) \cdot Q \cdot N_A} \cdot I_{\frac{m}{z} i} \quad (5)$$

$$c_n(\text{TPPO}) = \frac{10^{12}}{\text{IE}\left(\text{TPPO}, \frac{m}{z} 277\right) \cdot Q \cdot N_A} \cdot I_{\frac{m}{z} 277} \quad (6)$$

$$c_n(\text{TPP}) = \frac{10^{12}}{\text{IE}\left(\text{TPPO}, \frac{m}{z} 262\right) \cdot Q \cdot N_A} \cdot I_{\frac{m}{z} 262}. \quad (7)$$

The experimental procedure to estimate IE is discussed in more detail in the Supplement.

### 3.3 Background correction

As mentioned above (see Sect. 3.1) TPP particles, as well as pure ammonium sulfate aerosol particles passing the TPP-condensation/reaction chamber, also show a small but non-negligible TPPO signal ( $m/z$  277). This is either a consequence of TPPO impurities in the TPP itself or due to traces of peroxides in the solvents used to generate the aerosol particles when the nebulizer was used. Consequently, a background correction procedure has to be introduced to account for these effects. To do so, pure ammonium sulfate (AS) aerosol particles were investigated with different initial ammonium sulfate concentrations with the ORD-AMS setup as described above. These experiments showed a constant background contribution of 1.2 ( $\pm 0.1$ ) % TPPO resulting from the conversion of TPP [ $m/z$  262  $\Rightarrow$   $C_n$  (detected TPP)]. Therefore, the concentration of the analyte of interest [ $C_n$  (perox-

ide)] was calculated as follows:

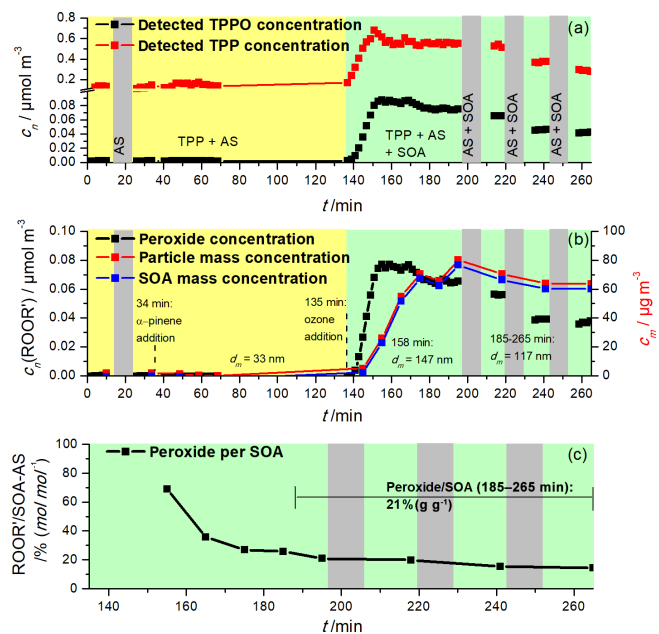
$$c_n(\text{backgr. TPPO}) = 0.012 \cdot c_n(\text{detected TPP}) \quad (8)$$

$$c_n(\text{produced TPPO}) = c_n(\text{detected TPPO}) - c_n(\text{backgr. TPPO}) \quad (9)$$

$$c_n(\text{peroxide}) = c_n(\text{produced TPPO}). \quad (10)$$

### 3.4 SOA experiments

The ozonolysis of  $\alpha$ - or  $\beta$ -pinene was performed in the presence of ammonium sulfate (AS) seed particles, and the results of a typical chamber run with  $\alpha$ -pinene are shown in Fig. 3. First, AS seed particles were injected into the chamber. After a certain period of time, when the particle concentration became constant, the biogenic VOC was introduced into the chamber (e.g., after 34 min in the run shown in Fig. 3). Following an initial period to monitor background peroxide concentrations in the AS seed aerosol–VOC system, ozone was introduced (e.g., after 135 min in the  $\alpha$ -pinene run shown in Fig. 3, green area). Periods when TPP was not added (AMS mode in Fig. 1) are marked in grey and were used to determine the dilution factor and the concentration of SOA by the AMS. As can be seen in the figure, the concentrations of both TPPO and TPP rose quickly during the first 20–25 min after the ozone addition (Fig. 3a), followed by a slow decrease towards the end of the experimental run. Why the TPP signal also rises will be explained below. However, peroxides were only detected when SOA formation began, as indicated by the time series of peroxide concentration (Fig. 3b, black) and SOA mass concentration (Fig. 3b, blue line). Figure 3c shows the temporal behavior of the relative contribution of peroxides to SOA, whereby an average molar mass of  $250 \text{ g mol}^{-1}$  for SOA was assumed. The course of the peroxide quantity per SOA quantity (Fig. 3c) shows a continuous decrease during the entire experimental run. Such concentration profiles of the organic peroxides are indicative of consecutive reactions, in which the peroxides are formed as first-generation products from the initial ozone/alkene reaction in the gas phase, followed by gas-to-particle conversion and further reactions of the peroxides to non-peroxidic products in the particle phase. Indeed, several studies suggest that organic peroxides are instantly formed in the VOC–ozone reaction (Bianchi et al., 2019; Mertes et al., 2012) and the rapid increase of the peroxide signal as shown in Fig. 3b confirms these studies. However, since the O–O bond is relatively weak, organic peroxides tend to decompose or react with other aerosol components. Possible reactions to non-peroxidic products could be reactions with alcohols or aldehydes forming aldehydes or carboxylic acids, or hydrolysis followed by the loss of  $\text{H}_2\text{O}_2$  (Zhao et al., 2018; Claffin et al., 2018; Epstein et al., 2014). In any case, the temporal evolution of the peroxide concentration (i.e., the formation/decline profile) indicates consecutive reactions and demonstrates the intermediate character of the peroxides within the VOC oxidation system.



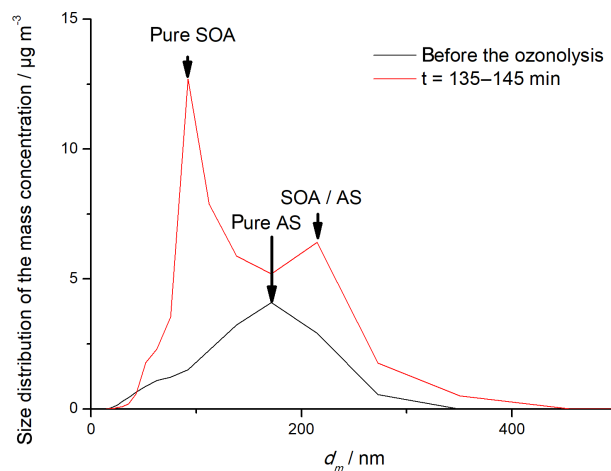
**Figure 3.** Time series of (a) the detected TPPO (black) and detected TPP (red), (b) the peroxide concentration (black) after background correction and particle mass concentration of the SOA–AS aerosol (red) and the SOA (blue), and the (c) relative contribution of peroxide to the total aerosol volume (black; time resolution 10 min). Periods without the addition of TPP (AMS mode) are marked with a grey background; the period before the addition of ozone is marked in yellow and during the ozonolysis in green (with TPP, ORD-AMS mode).  $\alpha$ -pinene was continuously added after 34 min and ozone after 135 min.

As can be seen in Fig. 3c, the contribution of peroxides to SOA was almost 70% after about 20 min. During the last 80 min of the experiment, the SOA concentration was about  $69 \mu\text{g m}^{-3}$  ( $0.28 \mu\text{mol m}^{-3}$ ), assuming again an average molar mass of the SOA components of  $250 \text{ g mol}^{-1}$ . During this time the measured peroxide concentration was about  $0.0470 \mu\text{mol m}^{-3}$ . Paul Ziemann and coworkers (Docherty et al., 2005) proposed an average molar mass of  $300 \text{ g mol}^{-1}$  for SOA peroxides. Based on this value the average peroxide mass concentration was  $14.1 \mu\text{g m}^{-3}$ , assuming the same density of peroxides as for other SOA components ( $1.2 \text{ g cm}^{-3}$ ; Kuwata et al., 2012). Thus the average peroxide/SOA yield within this period is estimated to be about 20% ( $\pm 4\%$ ). Actually, this value is very close to the value of 22% (Epstein et al., 2014) or the value of 21% (Li et al., 2016). Mertes and coworkers (Mertes et al., 2012) observed 34%, 17% and 12% for 15–35 min, 4 h and 6 h filter samples, respectively. Taking into account the influence of sampling time, duration of sampling, initial ozone and terpene concentrations as well as temperature and relative humidity, these values agree quite well.

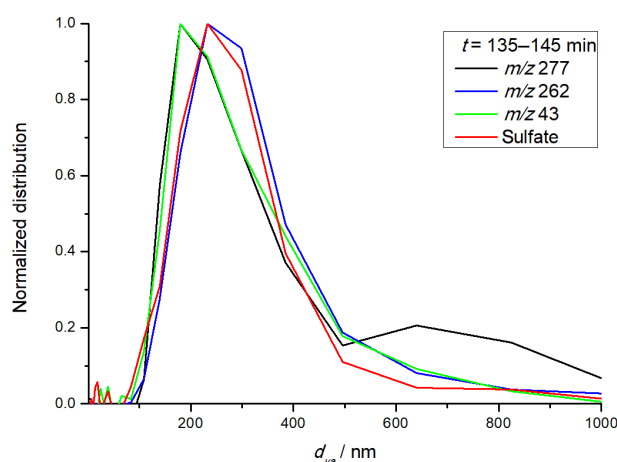
As can be seen in Fig. 3a, the TPP concentration also increased after the start of ozonolysis. This is understandable,



considering that TPP is only detected by the AMS when it condenses on particles. The more particles are introduced into the ORD setup, the higher the TPP concentration measured by the AMS. Figure 4 shows the particle size distribution measured by the SMPS system before (black) and within the first 10 min after (red) ozone was introduced into the reaction chamber. Despite the presence of AS seed particles, new particle formation from nucleation of organic compounds was observed, resulting in a bimodal size distribution at the beginning of the experiment. In principle, new particle formation is not surprising due to the high particle-formation potential of biogenic VOCs as well as the relatively high concentrations of the reactants. The same behavior was also visible in the AMS data. One of the main advantages of the AMS is the possibility to determine the size distributions of individual compounds. If the  $m/z$  is characteristic for an aerosol component, such as  $m/z$  30 and 46 for nitrate, for example, the size distribution of these components can be obtained. Figure 5 shows the  $m/z$  values as a function of particle size for TPP ( $m/z$  262; blue), TPPO ( $m/z$  277; black), SOA ( $m/z$  43; green) and sulfate (red) for the first 10 min after ozone was introduced, all normalized to the respective maximum value. Although the determined size distributions by SMPS and AMS are not identical, likely due to scanning time effects, differences in aerodynamic and mobility diameters as well as differences in the averaging procedure, the bimodal size distribution of pure organic particles and pre-existing AS-containing particles is also visible in the AMS data. While the occurrence of organic nucleation simultaneously with condensational growth resulting in two externally mixed particle populations is not surprising, a more quantitative description of the size distribution of selected compound groups is challenging. As can be seen in Fig. 5, the size profiles of SOA and TPPO are very similar, with a clear preference for the smaller particle sizes. Besides the obvious explanation of the existence of a pure organic particle population formed by homogeneous nucleation, an additional explanation is a size-dependent condensation of the organics also on the inorganic seed particles. A unique characteristic of condensational growth is that it is proportional to the Fuchs-corrected surface area of particles and not the particle volume – “surface-limited” vs. “volume-limited” growth. Because the surface area to volume ratio of particles increases as their diameter decreases, condensation tends to have a larger effect on smaller particles. Therefore, after the condensational growth of AS seed particles, the final distribution of the condensed organics can be expected to favor the smaller particles (Donahue et al., 2014; Riipinen et al., 2011). Therefore, both nucleation of organics and condensational growth will contribute to the size distribution shown in Fig. 5. Nevertheless, the results clearly show that the formation of peroxidic compounds is directly connected to SOA formation. More difficult to explain is the quite similar size distribution of TPP and ammonium sulfate seed aerosol since TPP also moves to the particle phase via condensation. One



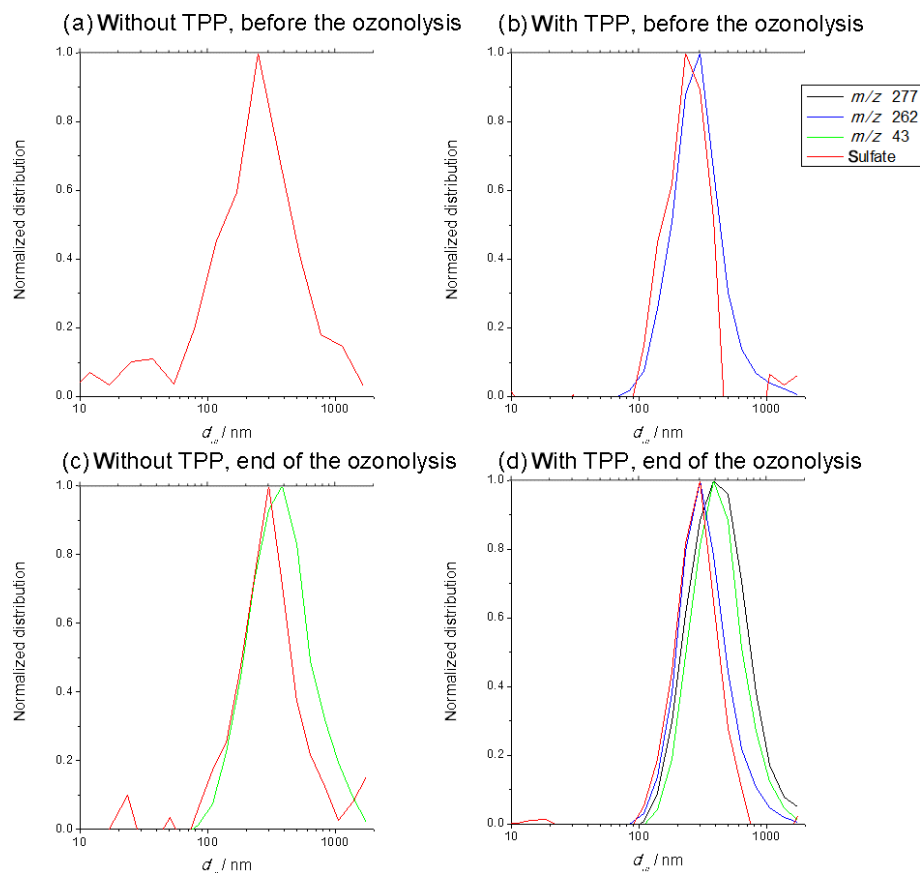
**Figure 4.** Particle mass size distribution before and 10 min after the addition of ozone.



**Figure 5.** Size distribution of individual particle-phase compounds (TPPO, TPP, SOA, sulfate) within the first 10 min after the addition of ozone.

possible explanation could be attractive forces between the nonpolar TPP molecules and AS, similar to a “salting in” effect (Hyde et al., 2017). However, possibly other effects, such as the high supersaturation of TPP in the condensation/reaction volume, also might be considered to explain the observations.

Figure 6a–d show the size distributions of  $m/z$  277 (black), 262 (blue), 43 (green) and sulfate (red) for the periods before (a, b) and during (c, d) the last 80 min of the ozonolysis, with (b, d) and without (a, c) added TPP. Again, each of the individual ratios is normalized to its respective maximum;  $m/z$  262 corresponds to the detected TPP,  $m/z$  277 to TPPO and  $m/z$  43 to SOA species (see Fig. S6 in the Supplement). Figure 6a shows the size distribution of pure AS particles. As can be seen, by comparison of Fig. 6a and b, the diameter of sulfate remained almost constant due to the condensation of TPP – if anything then the diameter



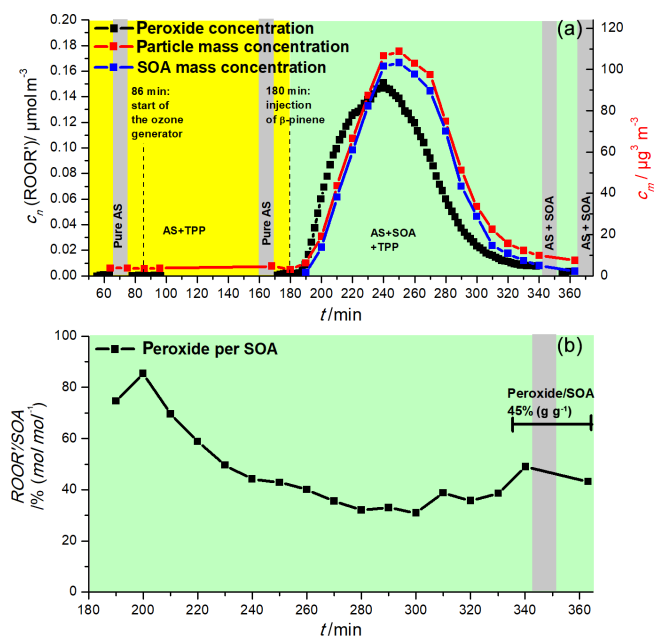
**Figure 6.** Size distribution of  $m/z$  277 (black is TPPO),  $m/z$  262 (blue is residual TPP),  $m/z$  43 (green is SOA) and sulfate (red) for following periods: before the start of the ozonolysis (a) without and (b) with adding TPP and at the end (185–265 min) of the ozonolysis (c) without and (d) with adding TPP. All distributions were normalized to their respective maximum.

slightly decreased. Generally, the vacuum-aerodynamic diameter depends on the particle density (DeCarlo et al., 2004). The larger the density, the larger  $d_{va}$  is for particles with the same physical diameter. The condensation of TPP (density:  $1.1 \text{ g cm}^{-3}$ ) will lower the particle density (density of AS:  $1.77 \text{ g cm}^{-3}$ ), which compensates for the increase in the physical diameter. Nevertheless, the condensation of SOA material finally led to an increase in the sulfate diameter (Fig. 6c). In Fig. 6d the size distributions of the four different components are shown. First to notice is again the clear overlap of  $m/z$  277 and  $m/z$  43, demonstrating the simultaneous presence of TPPO and SOA species in the particle phase. The slight shift in the size distribution to larger sizes for SOA ( $m/z$  43) and TPPO is likely a result of slightly faster growth of the pure organic particles formed by nucleation as discussed above compared to the ammonium-sulfate-containing particles.

A second biogenic VOC,  $\beta$ -pinene, was investigated with the online redox derivatization–AMS system, this time with a reverse introduction of the VOC and the oxidant to examine ozone as a direct oxidant for TPP. Again, the ozonolysis of  $\beta$ -pinene was performed in the presence of AS seed

aerosol. Figure 7 shows the temporal evolution of the peroxide concentration together with the total aerosol mass concentration determined with the SMPS. As in the case of  $\alpha$ -pinene, the peroxide concentration increased rapidly with the formation of SOA, once again demonstrating the direct connection of SOA and peroxide formation via the detection of TPPO. The comparison of the profiles of SOA (blue) and peroxide (black) demonstrates the very rapid peroxide formation followed by the decreasing contribution of peroxides to the particle phase in the later stages of the experiment as a consequence of consecutive reactions of the peroxides to non-peroxidic products. The strong decrease in total SOA after 240 min is induced by the end of  $\beta$ -pinene addition to the reaction system. The observed shift between the peroxide and SOA concentration supports the concept of organic peroxides as reaction intermediates, which contribute especially to the composition of fresh SOA and convert to non-peroxidic products in aged SOA particles. This is also clearly visible in Fig. 7b, which shows the relative contribution of peroxides to the total particle volume in the course of the experiment. However, most important for the overall method evaluation are the very low and rather constant TPPO concentrations





**Figure 7.** Ozonolysis of  $\beta$ -pinene in the presence of AS aerosol. Time series of (a) peroxide (black), particle mass concentration (red) and SOA mass concentration (blue); (b) relative contribution of peroxide to the amount of SOA. Ozone was added at 86 min,  $\beta$ -pinene at 180 min. Averages molar masses of 250 and 300  $\text{g mol}^{-1}$  for SOA and peroxides were assumed, respectively.

in the period between the ozone and  $\beta$ -pinene addition (between 86 min and 180 min) when just AS particles were introduced into the ORD-AMS system, a result that clearly excludes ozone as a direct oxidant for TPP.

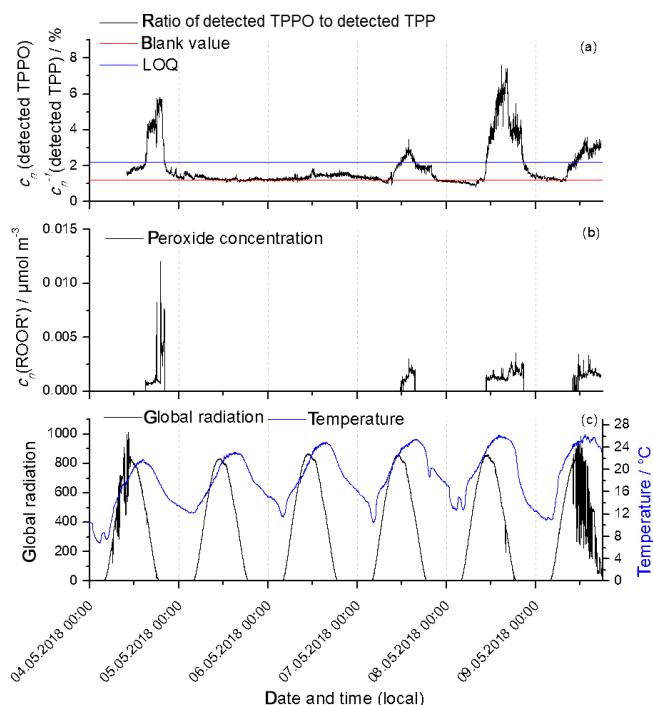
### 3.5 Discussion

As discussed above, the validation of the quantitative performance of the presented measurement method was not performed with peroxide reference compounds. Instead, the ratio between the concentration of peroxides and total SOA at the end of the ozonolysis of  $\alpha$ -pinene was compared with the results of Epstein et al. (2014), Li et al. (2016) and Mertes et al. (2012). In these studies the peroxide concentration formed from the ozonolysis of  $\alpha$ -pinene using filter or impactor sampling, followed by the extraction and iodometric analysis of the peroxide content, was measured. Several experimental parameters, such as the terpene and ozone concentration, their ratio to each other, the temperature, and the relative humidity, were slightly different. However, the comparison show very good agreement with values close to 20%. Another aspect of all SOA experiments which include chemistry, as the peroxide–TPP reaction here, that has to be considered is the question of the phase state of SOA. According to Koop et al. (2011) SOA is expected to be a liquid under the conditions used for our experiments (22–25 °C; RH 60%). Under these conditions also the AS seed particles can be ex-

pected to have liquid-like behavior (Matthew et al., 2008). As a consequence, the diffusion time of TPP within the particles should be approximately 1 s (Koop et al., 2011), which is much shorter than the residence time of about 35 s of the aerosol in the ORD setup (Fig. 1). In conclusion, based on the agreement of the peroxide/SOA ratio at a later stage of the experiments measured in this work and the results of former studies, the estimated diffusion times and the excess of the reactant TPP to SOA, a quantitative reaction of TPP with the particle-phase peroxides to TPPO can be assumed. However, in future investigations, the method should be revalidated using as aerosol particles composed of pure single-component peroxides.

### 3.6 Ambient air measurements

Finally, the ORD-AMS method was tested with ambient air directly outside the chemistry building located at the campus of the university in Mainz (49.9915928° N, 8.2312864° E). The measurements were performed between 4 May (starting time 09:30 CEST) and 9 May 2018 (17:30). Meteorological data were measured and provided by the Institute of Physics, also located on the campus of the university. During the measurement period, the air masses came from north, northeast or east, and air mass composition can be expected to be especially influenced by the Frankfurt Rhine-Main Metropolitan Region with approximately 5.8 million inhabitants. Figure S9 in the Supplement shows 72 h backward trajectories of the air masses (every 24 h) from 4 to 10 May 2018. The blank value for the seeded SOA experiments was used for the background correction, and values below the quantification limit were set to zero. Figure 8a shows the concentration ratios of TPPO to total TPP during the measurement period and Fig. 8b the course of the particle-phase peroxide concentration. Actually, the smallest values of detected TPPO to total TPP shown in Fig. 8a lie very close to the estimated blank value level (red line). This indicates that blank values of ambient air measurements and the blank value from the seeded SOA experiments are almost identical. Values of the peroxide concentration above the limit of quantification (LOQ) (blue line) were only obtained during weekdays (5 May Saturday; 6 May Sunday) and in the daytime. The enhanced concentration during weekdays in daylight conditions indicates local sources of peroxides in combination with photochemistry, an observation that is in agreement with the result from previous peroxide measurements (Hua et al., 2008; Guo et al., 2014). However, due to highly variable emission sources directly outside the chemistry building, the selected measurement site was certainly not ideal for general studies about particle-phase peroxide chemistry. Nevertheless, this short proof-of-principle study demonstrates that peroxides in atmospheric aerosols can be detected in real time with the new ORD-AMS system.



**Figure 8.** (a) Time series of the relative amount of detected TPPO to detected TPP, LOQ, and blank value, (b) diurnal variation in the estimated peroxide concentration, and (c) time series of the global radiation and temperature.

## 4 Conclusions

The real-time detection of peroxides by adding gaseous TPP to aerosol particles was successfully realized using an aerosol mass spectrometer in combination with a specifically designed online derivatization unit. The ozonolysis of  $\alpha$ - and  $\beta$ -pinene in the presence of ammonium sulfate seed aerosol was used to demonstrate the potential of the analytical procedure. The results clearly show the conversion of TPP to TPPO and hence the presence of organic peroxides in biogenic SOA. Especially in the early phase of SOA formation the contribution of peroxides is exceptionally high (up to ca. 80%  $v/v$ ). In addition, the time-resolved measurements show a rapid further reaction of the peroxides to non-peroxidic compounds indicative of consecutive reactions of the peroxides within the particle phase. The developed instrumental setup can also be used to receive particle size-resolved information about particle-phase peroxides. Again, the important role of organic peroxides in the early stages of particle formation, which was probably induced by very low-volatility multifunctional organic peroxides (HOMs), became clear. Finally, the new online redox derivatization method was successfully tested with ambient air outside the laboratory. The results presented here represent the first attempt to better characterize the role of organic peroxides in atmospheric aerosols. Future laboratory and field studies are certainly necessary

to better understand the chemistry of peroxides in organic aerosols.

**Data availability.** The data obtained by ORD-AMS measurements from the SOA experiments are available on request from Marcel Weloe (maweloe@uni-mainz.de).

**Supplement.** The supplement related to this article is available online at: <https://doi.org/10.5194/amt-13-5725-2020-supplement>.

**Author contributions.** MW and TH designed the study. MW performed the experiments to develop ORD-AMS and performed the laboratory work, calibration and the data analysis. MW and TH examined the results, provided input on the data analysis and interpretation, and prepared the paper.

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

**Financial support.** This research has been supported by the Deutsche Forschungsgemeinschaft (grant no. HO 1748/16-1, “Development and application of a real-time method for the determination of selected compound classes (especially organic peroxides) in atmospheric aerosol particles using an aerosol mass spectrometer (AMS)”).

This open-access publication was funded by Johannes Gutenberg University Mainz.

**Review statement.** This paper was edited by Charles Brock and reviewed by two anonymous referees.

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