Oligomers, organosulfates, and nitrooxy organosulfates in rainwater identified by ultra-high resolution electrospray ionization FT-ICR mass spectrometry

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Abstract. Wet deposition is an important removal mechanism for atmospheric organic matter, and a potentially important input for receiving ecosystems, yet less than 50% of rainwater organic matter is considered chemically characterized. Precipitation samples collected in New Jersey, USA, were analyzed by negative ion ultra-high resolution electrospray ionization Fourier transform ion cyclotron resonance mass spectrometry (FT-ICR MS). Elemental compositions of 552 unique molecular species were determined in the mass range 50–500 Da in the rainwater. Four main groups of organic compounds were identified: compounds containing carbon, hydrogen, and oxygen (CHO) only, sulfur (S) containing CHOS compounds, nitrogen (N) containing CHON compounds, and S- and N- containing CHONS compounds. Organic acids commonly identified in precipitation were detected in the rainwater. Within the four main groups of compounds detected in the rainwater, oligomers, organosulfates, and nitrooxy-organosulfates were assigned based on elemental formula comparisons. The majority of the compounds identified are products of atmospheric reactions and are known contributors to secondary organic aerosol (SOA) formed from gas phase, aerosol phase, and in-cloud reactions in the atmosphere. It is suggested that the large uncharacterized component of SOA is the main contributor to the large uncharacterized component of rainwater organic matter.

1 Introduction

Precipitation is an efficient removal mechanism for atmospheric organic matter, which is a mixture of organic compounds that can influence climate, air quality, and ecosystem health. Dissolved organic carbon (DOC) is a ubiquitous component of rainwater and can be higher in concentration than inorganic species such as nitric and sulfuric acids (Willey et al., 2000). Organic acids contribute significantly to rainwater acidity in urban areas (Kawamura et al., 1996; Pena et al., 2002), and can be the main contributor to acidity (80–90%) in remote areas (Andreae et al., 1988). Thus, organic acids are frequently the focus of studies on rainwater DOC. However, the contribution of organic acids to total DOC ranges from only 14–36%, with other known compound classes (e.g., aldehydes, amino acids) contributing <10% to the total DOC (Avery et al., 2006). Approximately 50% of rainwater dissolved organic matter (DOM) is considered uncharacterized at both the compound class and individual compound level (Willey et al., 2000). The complexity of rainwater DOM and the large percent considered uncharacterized has made it difficult to determine the role of rainwater DOM in regional and global carbon budgets (Raymond, 2005).

A large percentage of the uncharacterized rainwater DOM has been attributed to macromolecular organic matter like that found in atmospheric aerosols, fog water and cloud water (Zappoli et al., 1999; Krivacsy et al., 2000; Feng and Moller, 2004; Likens, 1983). Much of this macromolecular material has been termed humic like substances (HULIS) because of properties similar to terrestrial and aquatic humic and fulvic acids (Graber and Rudich, 2006). It is largely unknown how the HULIS fraction of atmospheric organic matter may
Some compounds detected have distinct primary sources; however, the majority of these compounds are formed from secondary processes, and can be incorporated into rainwater either from cloud droplets directly, or from SOA formed through gas phase, aerosol phase, and in-cloud reactions.

2 Sample collection and analysis

Rainwater was collected from two sites in the northeastern USA. The urban site Camden, NJ (Latitude 39°56′57.45″ N; Longitude 75°7′16.60″ W; elevation 11 m) is a city directly across the Delaware River from Philadelphia, PA and is in a heavily urbanized region. The rural site Pinelands, NJ is located 30 miles east of Camden in the Lebanon State Forest, though located in a forest it is impacted by urban airflow (Latitude 39°56′43.61″ N; Longitude 74°37′1.52″ W; elevation 1 m). Two samples were collected from the Camden, NJ site, one collected in spring and one collected in fall. One sample was collected from the Pinelands, NJ site in summer (Table 1). The samples were collected using wet-dry deposition collectors (Aerochem Metrics Model 301, Bushnell, FL). The rain collectors were fitted with stainless steel buckets and opened only during wetfall events. The type of collector used and the placement of the collector at the site adhere to regulations outlined by the National Atmospheric Deposition Program (Bigelow et al., 2001). Samples were collected within 12 h of each rain event to minimize microbial degradation of DOM and consumption of inorganic nutrients. Sample temperature and pH were measured immediately after collection and samples were filtered through pre-combusted glass fiber filters (Whatman GFF; baked for four hours at 500°C; then rinsed with deionized water) ensuring analysis of the dissolved constituents only. Contamination due to field sampling and laboratory sample processing was minimal (field and filter blanks <5% DOC-C). Rainwater was stored at −20°C in polypropylene screw-capped tubes until analysis.

Table 1. Bulk properties of rainwater samples collected in New Jersey, USA.

<table>
<thead>
<tr>
<th>Date Collected</th>
<th>Storm Trajectory</th>
<th>Rainfall Amount (cm)</th>
<th>Temp °C</th>
<th>pH</th>
<th>NO$_3^-$ (µM)</th>
<th>NH$_4^+$ (µM)</th>
<th>DON (µM)</th>
<th>DOC (µM)</th>
<th>PO$_4^{3-}$ (µM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>14 May 2002 Camden</td>
<td>SW</td>
<td>3.0</td>
<td>15</td>
<td>4.0</td>
<td>32</td>
<td>44</td>
<td>29</td>
<td>259</td>
<td>0.1</td>
</tr>
<tr>
<td>5 October 2002 Camden</td>
<td>W</td>
<td>0.4</td>
<td>21</td>
<td>4.1</td>
<td>44</td>
<td>94</td>
<td>33</td>
<td>263</td>
<td>0.5</td>
</tr>
<tr>
<td>20 July 2002 Pinelands</td>
<td>WNW</td>
<td>2.1</td>
<td>22</td>
<td>3.7</td>
<td>59</td>
<td>42</td>
<td>38</td>
<td>224</td>
<td>0.2</td>
</tr>
</tbody>
</table>
2.2 Chemical analyses

Bulk nutrients \((\text{NO}_3^- + \text{NO}_2^- + \text{NH}_4^+, \text{PO}_4^{3-})\) were measured on the three rain samples with an automated nutrient analyzer and standard colorimetric methods (Lachat, Inc). DOC was measured by high-temperature catalytic oxidation with a Shimadzu Total Organic Carbon (TOC) 5000A analyzer (Sharp et al., 1993). Dissolved organic nitrogen was determined as the difference between total dissolved nitrogen measured with an Antek 7000 TN Analyzer (Seitzinger and Sanders, 1999) and dissolved inorganic nitrogen \((\text{NO}_3^- + \text{NO}_2^- + \text{NH}_4^+)\). Calibrations were performed using potassium hydrogen phthalate standards for DOC and urea and nitrate standards for TDN.

2.3 Ultra-high resolution electrospray ionization FT-ICR MS

Analyses were performed on the three rainwater samples with a 7-T Fourier transform ion cyclotron resonance mass spectrometer equipped with an ESI source (FT-ICR MS) and operated in the negative ion mode. The sample was diluted with methanol 50:50 v/v immediately before injection to limit solvent interaction. The sample was introduced into the ESI source by direct infusion with a flow rate of 5 \(\mu\text{L min}^{-1}\). The needle voltage was 3.02 kV, the capillary voltage was \(-9.44\) V, the capillary temperature was 260 \(^\circ\)C, and the tube lens was \(-57\) V. The spectra were mass calibrated with standard ions using an external calibrant (G2421A Agilent “tuning mix”) and the residual root mean square error after calibration was 1.1–1.6 ppm. The mass lists were processed and exported using Xcalibur v2.0 SR2 (ThermoFisher Scientific).

Midas Formula Calculator Software (v1.1) was used to calculate all mathematically possible formulas for all ions with a signal to noise ratio \(\geq 10\) using a mass tolerance of \(\pm 1\) ppm. An unlimited number of \(^{12}\)C and \(^{1}H\), up to fifteen \(^{14}\)N and \(^{16}\)O, and one of each \(^{32}\)S and \(^{31}\)P were allowed in the molecular formula calculations. The average mass error for all assignments was 0.3 ppm. Boundary values for molecular elemental ratios were applied as a filtering tool (Koch et al., 2005). The O:C, H:C, and N:C ratios were limited to \(<5\), \(\geq 0.3\), and \(<2\) respectively. Ions were also characterized by the number of rings plus double bonds (i.e., double bond equivalents (DBE)), calculated from Eq. (1) (McLafferty, 1993):

\[
\text{DBE} = c - \frac{1}{2}h + \frac{1}{2}(n + p) + 1
\]

for elemental composition, \(C_cH_hO_oN_nP_pS_s\)

3 Sample comparison

Though the three rainwater samples analyzed were from different locations and seasons, the storm trajectories and bulk properties of the samples are similar (Table 1) and fall within the ranges reported for continental precipitation (Seitzinger et al., 2003; Willey et al., 2000). The composition of the DOM in all three samples was also very similar, as discussed below. In the Camden spring sample, there were 161 masses detected in the mass range of 50–300 Da. However, only 13% of them were unique to that sample, i.e., 87% of the masses detected appeared in one or both of the other samples. Though the Pinelands sample was analyzed over a slightly smaller mass range (50–400 Da) than the fall Camden sample (50–500 Da), the samples are similar in the total number of masses detected, 388 and 383, respectively. Of the 388 masses detected in the Pinelands sample, 64% were detected in one of the other two samples (i.e., 36% are unique). For the fall Camden sample, 84% of the masses were detected in one of the other two samples (i.e., 16% are unique).

The masses detected fall into four main compound classes: carbon, hydrogen and oxygen only (CHO), sulfur (S) containing CHOS compounds, nitrogen (N) containing CHON compounds, and S- and N-containing CHONS compounds. The number of compounds in each compound class was very similar in both the Camden fall and Pinelands summer samples (Fig. 1). Approximately 50% of the compounds in the negative ion mode are composed only of CHO. Approximately 25–30% of the compounds are composed of CHOS, and \(~5\)% are composed of CHONS. The remaining 10–25%
Comparison of the elemental ratios of negative ion mode compounds in precipitation samples collected in July and October 2002 in the Pinelands and Camden, NJ (USA), respectively.

The analysis is based solely on negative ion FT-ICR MS, and as such focuses only on compounds that can be detected in the negative ion mode (e.g., carboxylic acids). It should be noted that for each elemental composition assigned based on the measured mass, multiple structural isomers are possible, and thus the reported total number of compounds is likely an underestimate. The Van Krevelen diagrams, which plot the hydrogen to carbon (H:C) ratio as a function of the oxygen to carbon (O:C) ratio for each compound, can be used to display compositional characteristics and to compare the elemental ratios of each compound in a sample (Wu et al., 2004). As evident in the Van Krevelen diagrams, the elemental ratios of the compounds in each group (i.e., CHO, CHOS, CHONS) are also similar in both the Camden and Pinelands samples.

In the analysis that follows (e.g., Fig. 3, Table 2), we have combined the elemental formulas for each of the three samples due to the high percentage of overlap in the types of elemental formulas (i.e., compound class; Fig. 1) the similarities in number and type of elemental formulas (Fig. 2), and the small number of samples. Thus we will consider this combined data representative of a precipitation sample collected from an urban impacted (i.e., downwind of a major city) location in New Jersey, USA. As an example, all of the negative ion elemental formulas determined in the 20 July 2002 sample are listed with their corresponding m/z (Supplemental Information: http://www.atmos-chem-phys.net/9/2533/2009/acp-9-2533-2009-supplement.pdf).

### 4 Organic acids and oligomers

Eight organic acids identified in the rainwater based on their elemental formulas have been previously detected in precipitation samples: glyoxylic, glycolic, pyruvic, oxalic, lactic, malonic, and succinic acids (Kieber et al., 2002). Formic acid is commonly detected in precipitation, but has a molecular weight less than the lower mass detection limit of the FT-ICR MS m/z 50; therefore, although it is likely present in the samples, it is not detectable.

The potential sources of these eight organic acids include primary anthropogenic and biogenic emissions, as well as secondary formation from biogenic and anthropogenic precursors in the atmosphere (Kawamura et al., 2001, 2003; Sorooshian et al., 2007). Though there are primary emissions of organic acids, atmospheric aqueous phase reactions in clouds and aerosols have been hypothesized to be
the predominant formation pathway of these organic acids (Blando and Turpin, 2000). Similarly, the atmospheric concentrations of these organic acids can only be supported by a substantial secondary formation mechanism (Sorooshian et al., 2007). Thus, the organic acids detected in the rainwater are likely predominantly from secondary atmospheric processes and incorporated during in-cloud or below-cloud scavenging.

In addition to the eight identified organic acids, there were an additional 254 CHO compounds detected, representing 48% of the total compounds in the mass range 50–500 Da. The average (arithmetic mean) organic molecular weight to organic carbon weight (OM:OC) ratio for the CHO compounds in the mass range 50–500 Da was 2.1±0.7, the average H:C and O:C ratios were 1.5±0.4 and 0.7±0.5, respectively (Table 2). The elemental ratios for the CHO compounds are lower than typical organic acid ratios (OM:OC 2.5–3.8), and are more consistent with measured ratios for smog chamber generated oligomers, oligomers formed through cloud processing, aged atmospheric aerosol, and HULIS, all contributors to SOA (El-Zanan et al., 2005; Kiss et al., 2002; Kalberer et al., 2004; Altieri et al., 2008; Polidori et al., 2008).

The elemental formulas of the CHO compounds in the rainwater were compared to the elemental formulas of oligomers known to form through cloud processing reactions of methylglyoxal (Altieri et al., 2008); methylglyoxal is a water soluble compound with both biogenic and anthropogenic sources. The organic acids known to form through cloud processing of methylglyoxal were all detected in the rainwater, i.e., glyoxylic, glycolic, pyruvic, oxalic, succinic, malonic, and malic acids (Table 3). The methylglyoxal aqueous oligomerization scheme involves acid- or radical- catalyzed esterification of the organic acids with a hydroxy acid, e.g., lactic acid, leading to series of oligomers related by regular mass differences of 72.02113 Da (Altieri et al., 2008). All nine oligomer series present in the methylglyoxal cloud processing scheme were detected in the

The number of masses in each compound class and the average (arithmetic mean) elemental ratios for each compound class. The compounds contributing to the “other” category are listed below the darkened line.

<table>
<thead>
<tr>
<th>Type</th>
<th># of compounds</th>
<th>% of Total</th>
<th>H:C</th>
<th>O:C</th>
<th>N:C</th>
<th>MW</th>
<th>OM:OC</th>
</tr>
</thead>
<tbody>
<tr>
<td>CHO</td>
<td>262</td>
<td>48</td>
<td>1.5±0.4</td>
<td>0.7±0.5</td>
<td>–</td>
<td>193±58</td>
<td>2.1±0.7</td>
</tr>
<tr>
<td>CHOS</td>
<td>139</td>
<td>25</td>
<td>1.9±0.5</td>
<td>1.3±0.8</td>
<td>–</td>
<td>246±54</td>
<td>3.4±1.5</td>
</tr>
<tr>
<td>CHONS</td>
<td>28</td>
<td>5</td>
<td>1.8±0.6</td>
<td>1.7±0.9</td>
<td>2.7±1.9</td>
<td>271±60</td>
<td>4.2±1.5</td>
</tr>
<tr>
<td>CHON</td>
<td>74</td>
<td>13</td>
<td>1.9±0.8</td>
<td>1.6±1.4</td>
<td>2.6±2.4</td>
<td>192±68</td>
<td>3.7±2.2</td>
</tr>
<tr>
<td>Other</td>
<td>49</td>
<td>9</td>
<td>1.5±1</td>
<td>1.0±1.1</td>
<td>5.4±3.9</td>
<td>272±80</td>
<td>3.4±1.9</td>
</tr>
<tr>
<td>CHONP</td>
<td>29</td>
<td>5</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CHONSP</td>
<td>8</td>
<td>2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CHOP</td>
<td>7</td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CHOSP</td>
<td>5</td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 3. Organic acids and oligomers detected in the rainwater by negative ion FT-ICR MS. Compound identification based on comparison with FT-ICR MS negative ion data from methylglyoxal + -OH oligomer series as described in (Altieri et al., 2008). Compounds from all 9 methylglyoxal oligomer series are represented. n=1 and n=2 indicates the addition of one and two hydroxy acids (+C3H4O2) through esterification, respectively.

<table>
<thead>
<tr>
<th>Parent</th>
<th>m/z</th>
<th>Elemental Formula</th>
<th>m/z</th>
<th>Elemental Formula</th>
<th>m/z</th>
<th>Elemental Formula</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glyoxylic acid</td>
<td>72.9931</td>
<td>C2H4O3</td>
<td>145.01426</td>
<td>C3H5O5</td>
<td>217.03546</td>
<td>C8H9O7</td>
</tr>
<tr>
<td>Glycolic acid</td>
<td>75.00875</td>
<td>C2H4O3</td>
<td>147.02991</td>
<td>C3H5O5</td>
<td>219.05107</td>
<td>C8H11O7</td>
</tr>
<tr>
<td>Pyruvic acid</td>
<td>87.0087</td>
<td>C2H4O3</td>
<td>159.0299</td>
<td>C3H5O5</td>
<td>231.05109</td>
<td>C8H11O7</td>
</tr>
<tr>
<td>Oxalic acid</td>
<td>88.98796</td>
<td>C2H2O4</td>
<td>161.00922</td>
<td>C3H5O6</td>
<td>233.03034</td>
<td>C8H9O6</td>
</tr>
<tr>
<td>Succinic acid</td>
<td>117.01935</td>
<td>C4H5O4</td>
<td>189.04032</td>
<td>C7H9O6</td>
<td>261.06157</td>
<td>C10H13O8</td>
</tr>
<tr>
<td>n/z 131</td>
<td>131.03501</td>
<td>C4H7O4</td>
<td>203.0562</td>
<td>C8H11O6</td>
<td>275.07719</td>
<td>C11H15O8</td>
</tr>
<tr>
<td>Malonic acid</td>
<td>103.00358</td>
<td>C3H4O4</td>
<td>175.02483</td>
<td>C6H8O6</td>
<td>247.04596</td>
<td>C9H13O8</td>
</tr>
<tr>
<td>Malic acid</td>
<td>133.01427</td>
<td>C4H5O5</td>
<td>205.03546</td>
<td>C7H9O7</td>
<td>277.05647</td>
<td>C10H13O9</td>
</tr>
<tr>
<td>n/z 177</td>
<td>177.04047</td>
<td>C6H9O6</td>
<td>249.0616</td>
<td>C9H13O8</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
rainwater based on elemental formula comparisons (Table 3). For each organic acid detected, compounds containing elemental formulas consistent with the addition of one and two hydroxy acids through esterification were detected. For example, moving from left to right in Table 3, glyoxylic acid ($C_2H_3O_3$), glyoxylic acid plus one addition of hydroxy acid through esterification ($C_3H_5O_5$), and glyoxylic acid plus two additions of hydroxy acid through esterification ($C_5H_9O_7$) were all detected. In addition to the elemental formulas being consistent with methylglyoxal oligomers, some of these elemental formulas are also consistent with compounds formed from photooxidation of other biogenic precursors such as α-pinene oxidation leading to hydroxy glutaric acids (Claeys et al., 2007), highlighting that multiple structural isomers of each elemental formula are possible. Without further liquid chromatography or MS-MS analysis, it is not possible to discern the exact structure of these higher molecular weight compounds. Compounds with the same elemental formulas and patterns as reported here in the rainwater were also detected in the high molecular weight fraction ($\geq m/z$ 223) of aerosol derived water soluble organic carbon (Wozniak et al., 2008).

There are CHO compounds detected in the rainwater sample that are not part of the methylglyoxal oligomer series, but also have regular mass differences indicative of oligomers and/or HULIS (Figs. 3, 4). In addition to the van Krevelen (Fig. 3), another way to visualize compounds related by regular mass differences is an island plot (Reemtsma et al., 2006a), which plots the number of carbon atoms of each molecule against the molecular weight of the molecule. When the sum of the number of carbons and oxygens for a set of compounds is the same, those compounds group as “islands” in the plot (denoted by different markers; Fig. 4). There were 19 islands detected in the mass range 50–500 Da for the compounds containing only CHO, and the sum of carbon and oxygen ranged from 4–23.

In each island the elemental composition changes in a systematic way from one molecule to the next. Moving horizontally within an island increases the saturation of a molecule (+2 H). Moving horizontally from one island to another introduces additional oxygen (+1 O). Moving vertically within an island changes the composition by exchanging one oxygen atom for CH$_4$, hence the island number (#C+#O) remains unchanged. These clustered island patterns in the rainwater are typical of fulvic acid isolates from natural organic matter samples (Reemtsma et al., 2006a; Stenson et al., 2003), and fulvic acids and oligomers detected in water soluble aerosols (Reemtsma et al., 2006b), indicating the presence of oligomeric compounds and HULIS in the complex rainwater DOM.

5 Sulfur compounds

5.1 Organosulfates

There were 139 CHOS compounds detected (50–500 Da), representing 25% of the total compounds (Table 2). The average molecular weight and average OM:OC ratio of the CHOS compounds were higher than the CHO compounds, which is consistent with the addition of a sulfur atom onto the molecule. The number of double bond equivalents (DBEs) calculated from the CHOS elemental formulas indicates that they are likely aliphatic compounds (average DBE<3). The average H:C and O:C ratios for the CHOS compounds were also higher than the elemental ratios of the CHO compounds. The CHOS compounds (red triangles, Fig. 3) group to the right, and higher, on the van Krevelen diagram than the CHO compounds (black circles), indicating a higher degree of oxidation. This placement on the van Krevelen and the higher H:C and O:C ratios, are consistent with the addition of the sulfur as a sulfate (SO$_4^{2−}$) group. Sulfate was detected in all of the samples at $m/z$ 96,960103, and organosulfates, if present, are also expected to be seen in the negative ion mode (Surratt et al., 2008 and refs within). There were 134 CHOS compounds with oxygen to sulfur (O:S) ratios of 4–11, which allows all 134 compounds to potentially have sulfate functionalities. The high O:S ratios and the high degree of oxidation in the molecular formulas indicates that the bulk of the negative ion CHOS compounds detected in the rainwater samples are likely organosulfates.

Of the 134 CHOS compounds detected with O:S ratios >4, 28 have elemental formulas consistent with organosulfate esters detected in ambient SOA, and demonstrated to form in laboratory smog chamber experiments (Surratt et al., 2008; Romero and Oehme, 2005; Reemtsma et al., 2006b). These organosulfate esters are formed in the presence of high...
Table 4. Organosulfate compounds detected in the rainwater by negative ion FT-ICR MS. These elemental formulas were also reported in (Surratt et al., 2008) as organosulfates formed when biogenic precursor VOC’s are oxidized with high acidity sulfate seed aerosols.

<table>
<thead>
<tr>
<th>Isoprene</th>
<th>α-pinene</th>
<th>Limonene</th>
<th>α-terpinene</th>
</tr>
</thead>
<tbody>
<tr>
<td>m/z</td>
<td>Elemental Formula</td>
<td>m/z</td>
<td>Elemental Formula</td>
</tr>
<tr>
<td>138.970667</td>
<td>C$_2$H$_2$O$_5$S</td>
<td>223.028182</td>
<td>C$<em>7$H$</em>{11}$O$_6$S</td>
</tr>
<tr>
<td>152.986318</td>
<td>C$_2$H$_2$O$_5$S</td>
<td>226.987111</td>
<td>C$<em>7$H$</em>{15}$O$_6$S</td>
</tr>
<tr>
<td>154.965582</td>
<td>C$_2$H$_2$O$_5$S</td>
<td>237.043832</td>
<td>C$<em>7$H$</em>{15}$O$_6$S</td>
</tr>
<tr>
<td>168.981232</td>
<td>C$_2$H$_2$O$_5$S</td>
<td>279.054397</td>
<td>C$<em>{10}$H$</em>{15}$O$_7$S</td>
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<tr>
<td>210.991797</td>
<td>C$_2$H$_2$O$_5$S</td>
<td>281.070047</td>
<td>C$<em>{10}$H$</em>{17}$O$_8$S</td>
</tr>
<tr>
<td>213.007447</td>
<td>C$_5$H$_9$O$_3$S</td>
<td>297.064962</td>
<td>C$<em>{10}$H$</em>{17}$O$_8$S</td>
</tr>
<tr>
<td>215.023097</td>
<td>C$<em>5$H$</em>{11}$O$_2$S</td>
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</tr>
</tbody>
</table>

Table 5. Nitrooxy organosulfate compounds detected in rainwater by negative ion FT-ICR MS. These same elemental formulas were also reported by (Surratt et al., 2008) and are formed when parent biogenic VOC’s undergo reactions with high acidity sulfate seed aerosols with intermediate to high NO$_x$ levels. $^a$ The same nitrooxy organosulfates are also formed during α-terpinene, γ-terpinene, and β-pinene oxidation (Surratt et al., 2008).

<table>
<thead>
<tr>
<th>Isoprene</th>
<th>α-pinene$^a$</th>
<th>Monoterpene</th>
</tr>
</thead>
<tbody>
<tr>
<td>m/z</td>
<td>Elemental Formula</td>
<td>m/z</td>
</tr>
<tr>
<td>244.013261</td>
<td>C$<em>5$H$</em>{10}$N$_1$O$_8$S$_1$</td>
<td>294.065296</td>
</tr>
<tr>
<td>260.008175</td>
<td>C$<em>5$H$</em>{10}$N$_1$O$_8$S$_1$</td>
<td></td>
</tr>
<tr>
<td>304.992523</td>
<td>C$_5$H$_9$N$<em>2$O$</em>{11}$S$_1$</td>
<td></td>
</tr>
</tbody>
</table>

acidity sulfate seed aerosols during particle phase and aqueous phase reactions of biogenic VOCs and their oxidation products (Surratt et al., 2008). Organosulfates formed from the gas phase oxidation of isoprene, α-pinene, β-pinene, limonene, α-terpinene, and γ-terpinene were all detected in the rainwater (Table 4).

5.2 Nitrooxy-organosulfates

There were 28 CHONS compounds detected (50–500 Da), representing 5% of the total compounds. The CHONS compounds had an average H:C ratio similar to the CHOS compounds, but slightly higher O:C and OM:OC ratios (Table 2). The CHONS compounds have elemental formulas with higher oxygen content than the organosulfates and cluster even further to the right and above the CHO and CHOS compounds in the van Krevelen diagram (blue squares; Fig. 3) indicating an even higher degree of oxidation. The location of the CHONS compounds on the van Krevelen diagram, and the higher O:C ratios indicate that the nitrogen on the molecule is most likely an organic nitrate (ONO$_2$) functionality.

Five of the CHONS compounds have elemental formulas consistent with nitrooxy organosulfates detected in ambient aerosols and formed during laboratory smog chamber experiments (Table 5) conducted with high acidity sulfate seed aerosols and high NO$_x$ conditions (Surratt et al., 2007, 2008). Of the 23 remaining CHONS compounds, 18 have enough oxygen in their elemental formula for both a sulfate and nitrate functionality to be present ($\geq 7$ O) suggesting those compounds are also nitrooxy organosulfates formed from other biogenic and anthropogenic VOC precursors.

5.3 Sulfonates

Four of the CHOS compounds have an O:S ratio too low to be sulfate functionalities ($<3$), a high C content, and differ only by CH$_2$ units (C$_{16}$H$_{25}$O$_3$S, C$_{17}$H$_{27}$O$_3$S, C$_{15}$H$_{29}$O$_3$S, and C$_{19}$H$_{31}$O$_3$S). The elemental formulas of these four compounds are consistent with a class of persistent pollutants known as linear alkylbenzene sulfonates (LAS). The LAS compounds are used as synthetic surfactants and are surface active ingredients in detergents, shampoos, other cleaning compounds, and personal care products (Debelius et al., 2008; Lara-Martín et al., 2006). LAS compounds are toxic to a variety of organisms (Debelius et al., 2008) and as such are targeted in wastewater treatment plants (Reentsma, 2003). Due to their widespread use, LAS compounds have been detected at various concentrations in river water, seawater and sediments (González-Mazo and Gómez-Parra, 1996; González-Mazo et al., 2002). To our knowledge, this is the first evidence that LAS compounds can occur in atmospheric samples of any kind, and their presence in the atmosphere is difficult to explain. If the LAS compounds were able to reach
the atmosphere, the potential for negative impacts on receiving ecosystems would be extended due to the possibility of long transport times in the atmosphere before deposition.

6 Atmospheric implications

A number of lines of evidence suggest that the majority of compounds detected in these rainwater samples are from secondary processing in the atmosphere, and not primary emissions. First, the similarity in the number and types of compounds detected in the three rainwater samples from different seasons and locations in New Jersey is consistent with the organic matter being primarily from secondary processing in the atmosphere, and not local primary emissions. Second, the organic acids detected in the rainwater are likely predominantly formed through secondary processing reactions in the atmosphere, including in-cloud processing. In contrast, the sulfonates are likely primary anthropogenic compounds, and are pollutants known to be toxic to aquatic ecosystems; though the sulfonates are only four compounds out of the 552 total compounds detected. The majority of the compounds detected by negative ion FT-ICR MS in the rainwater have secondary sources both anthropogenic (e.g., organosulfates and nitrooxy organosulfates) and biogenic (e.g., oligomers from cloud processing of methylglyoxal).

Organosulfates likely contribute to the large percentage of uncharacterized water soluble organic carbon in atmospheric organic matter and could have been missed until recently because single derivatization protocols are likely to cause degradation or misinterpretation of these compounds (Surratt et al., 2007). Several organosulfates detected in the rainwater could be used as tracers for SOA that was formed under highly acidic conditions. Interestingly, in a study of DOM recovered from ice cores collected in Russia, 21% of the ions detected had organic S present in a sample from 1950, compared to 12% in a sample from 1300 AD (Grannas et al., 2006). This is consistent with an anthropogenic source of organic S capable of being transported long distances in the atmosphere before deposition, supporting the use of organosulfates as anthropogenic tracer compounds.

The nitroxy organosulfates detected in the rainwater have been seen in nighttime ambient aerosol samples, highlighting the importance of NO3 radical oxidation chemistry (Inuma et al., 2007a, b) in their formation. In the Surratt et al. (2008) study, nitroxy organosulfates were detected during both the nighttime oxidation and photooxidation experiments with acidic sulfate seed aerosol, and intermediate or high NOx conditions (~1 ppm). The nitroxy organosulfates detected in the precipitation sample could also be used as tracers for SOA that was formed under high acidity and high NOx conditions.

The oligomers, organosulfates, and nitroxy organosulfates detected in the rainwater could all contribute to the HULIS fraction of atmospheric organic matter (Graber and Rudich, 2006; Romero and Oehme, 2005). If the aqueous oligomerization of one water soluble organic compound (i.e., methylglyoxal) can account for 26 previously unidentified CHO compounds in the rainwater, it is possible that oligomerization of other known biogenic and anthropogenic precursor organic compounds through in-cloud or aerosol-phase reactions may contribute substantially to the complex HULIS fraction measured in aqueous atmospheric organic matter, including rainwater. Previous studies have suggested that no one compound comprises more than a small percentage of the total DOC in rainwater. The large number of compounds detected (552 unique masses from m/z 50 to 500), and the large contribution of multifunctional compounds (e.g., oligoesters, organosulfates, nitrooxy organosulfates) detected in the rainwater supports this idea. Unfortunately, quantification is not possible using ESI FT-ICR MS unless an authentic standard is used for each identified compound. Due to the large number of masses detected, and the lack of commercial availability of standards for these compounds, we were not able to quantify the mass contribution of each identified compound to total DOC concentration. However, as is typical for HULIS, the majority of these multifunctional compounds would not be separated and detected by traditional analytical techniques, highlighting the important advances made capable by ultra-high resolution mass spectrometry. This work provides motivation to expand studies of complex atmospheric organic matter to include quantitation, chromatographic separation and tandem MS-MS.

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