Chapter 12

Effects of Inorganic Seeds on Secondary Organic Aerosol (SOA) Formation

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Additional information is available at the end of the chapter

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1. Introduction

Atmospheric aerosol has significant influences on human health (Kaiser, 2005), visibility degradation (Cheng et al., 2011), and climate change (Satheesh and Moorthy, 2005). It was found that organic aerosols (OA) was the most abundant component of atmospheric aerosol (He et al., 2001) and more than 50% of the total OA are secondary organic aerosols (SOA) (Duan et al., 2005). SOA are produced from the oxidation of volatile organic compounds (VOCs) followed by gas-particle partitioning of the semivolatile organic products. Among the various VOCs, aromatic hydrocarbons are one type of SOA precursors which have drawn the most attention due to their abundance in the air and high SOA contribution to urban atmospheres (Lewandowski et al., 2008). Toluene and m-xylene are the two of the most abundant aromatic hydrocarbon species.

The detailed mechanism and controlling factors of SOA formation are not fully understood yet, which leads to the lower SOA level prediction from air quality models than the ambient measurements (Volkamer et al., 2006). Using smog chamber, SOA formation process can be investigated under controlled experimental conditions. Series of smog experiments have been conducted by different research groups to investigate the effects of background seed aerosols on SOA formation (Cao and Jang, 2007, Czoschke et al., 2003, Gao et al., 2004, Jang et al., 2002, Liggio and Li, 2008). Increased SOA formation and SOA yields were observed with the presence of acid seed aerosols. The effects of acidic seeds suggest that aerosol phase reactions may play an important role on SOA formation (Jang et al., 2002). Interactions between the organic and inorganic components of aerosols are important for further understanding the SOA formation process. Most research concludes that acid-catalyzed aerosol-phase reactions generate additional aerosol mass due to the production of oligomeric products with large molecular weight and extremely low volatility (Cao and Jang, 2007, Czoschke et al., 2003, Gao et al., 2004) and, therefore, enhance SOA formation.
Uptake of semivolatile organic products to acidic sulfate aerosols was also found contributing to enhance SOA formation (Liggio and Li, 2008). In these studies, (NH$_4$)$_2$SO$_4$ or H$_2$SO$_4$ seed aerosols were widely used to study the effect of particle acidity on SOA formation from both biogenic and aromatic hydrocarbons.

Atmospheric aerosols always have a very complex composition. Studying the effects of (NH$_4$)$_2$SO$_4$ or H$_2$SO$_4$ seed aerosols did not draw the whole picture of the role that inorganic seed aerosols play in SOA formation. Metal-containing aerosols are important components of the atmosphere. Calcium and iron are the most abundant metal species in atmospheric aerosols and the average concentration of them in Beijing could be as high as about 1.2 μg m$^{-3}$ and 1.1 μg/m$^3$ in PM$_{2.5}$ (He et al., 2001) respectively. In this study, we tested the effect of different inorganic seeds on SOA formation using a smog chamber. Two aromatic hydrocarbon precursors toluene and m-xylene are used. Effects of various inorganic seeds, including neutral inorganic seed CaSO$_4$, acidic seed (NH$_4$)$_2$SO$_4$, transition metal contained inorganic seeds FeSO$_4$ and Fe$_2$(SO$_4$)$_3$, and a mixture of (NH$_4$)$_2$SO$_4$ and FeSO$_4$, were examined during m-xylene or toluene photooxidation with the presence of nitrogen oxides (NO$_x$).

2. Experimental section

The experiments were carried out in a smog chamber which was described in detail in Wu et al. (Wu et al., 2007). The 2 m$^3$ cuboid reactor, with a surface-to-volume ratio of 5 m$^{-1}$, was constructed with 50 μm-thick FEP-Teflon film (Toray Industries, Inc. Japan). The reactor was located in a temperature controlled room (Escpec SEWT-Z-120), with a constant temperature between 10 and 60 °C (± 0.5 °C). The reactor was irradiated by 40 black lights (GE F40T12/BLB, peak intensity at 365 nm). Based on the equilibrium concentrations of NO, NO$_2$ and O$_3$ in a photo-irradiation experiment of an NO$_2$/air mixture, the NO$_2$ photolysis rate was calculated at approximately 0.21 min$^{-1}$, using a method described by Takekawa et al. (2000, 2003).

Prior to each experiment, the chamber was flushed for about 40 h with purified air at a flow rate of 15 L/min. In the first 20 hours, the chamber was exposed to UV light at 34 °C. In the last several hours of the flush, humid air was introduced to obtain the target relative humidity (RH).

Seed aerosols were generated by atomizing salt solutions using a constant output atomizer (TSI Model 3076). To avoid hydrolysis and precipitation in the Fe$_2$(SO$_4$)$_3$ salt solution, as little sulfuric acid as possible was added to the solution. What’s more, for generating internally mixed seed aerosols, a mixed solution of FeSO$_4$ and (NH$_4$)$_2$SO$_4$, in which the concentration ratio of FeSO$_4$ to (NH$_4$)$_2$SO$_4$ is 1:5, was used. The generated aerosols were passed through a diffusion dryer (TSI Model 3062) to remove water and a neutralizer (TSI Model 3077) to bring the aerosols to an equilibrium charge distribution. The hydrocarbon, NO and NO$_2$ were carried by purified dry air into the chamber. The concentrations were continuously monitored at a measurement point in the reactor until they were stable, ensuring the components in the reactor were well mixed. The experiment was then conducted for 6 hours with the black lights on.
A gas chromatograph (GC, Beifen SP-3420) equipped with a DB-5 column (30 m×0.53 mm×1.5 mm, Dikma) and flame ionization detector (FID) measured the concentration of the hydrocarbon every 15 min. NO$_x$ and O$_3$ were monitored with an interval of 1 min by a NO$_x$ analyzer (Thermo Environmental Instruments, Model 42C) and an O$_3$ analyzer (Thermo Environmental Instruments, Model 49C), respectively. Size distribution of particle matter (PM) was measured by a scanning mobility particle sizer (SMPS, TSI 3936) in the range of 17-1000 nm with a 6-min cycle. The volume concentration of aerosols was estimated from the measured size distribution by assuming the particles were geometrically spherical and nonporous.

3. Results and discussion

3.1. Estimating the generated SOA mass ($M_o$)

Due to deposition of particles on the Teflon film, the measured aerosol concentration had to be corrected. Takekawa et al. (2003) developed a particle size-dependent correction method, in which the aerosol deposition rate constant ($k(d_p)$, h$^{-1}$) is a four-parameter function of particle diameter ($d_p$, nm), as shown in equation (1):

$$k(d_p) = a \cdot d_p^b + c \cdot d_p^d$$  

The resulting $k(d_p)$ values for different $d_p$ (40-700 nm) were determined by monitoring the particle number decay under dark conditions at low initial concentrations (<1000 particles cm$^{-3}$) to avoid serious coagulation. Based on more than 500 sets of $k(d_p)$ values ($d_p$ ranges from 40 to 700 nm), the optimized values of parameter $a$, $b$, $c$, and $d$ were calculated to be $6.46 \times 10^{-7}$, 1.78, 13.2, and -0.957, respectively. It should be noted that the estimation of deposited aerosol concentrations using this method might introduce some error (Takekawa et al., 2003) because some scatter was recognized when fitting $k(d_p)$ values into equation (1). To reduce error due to wall deposition, SOA yields were calculated when the measured particle concentration reached its maximum in the experiments because deposited aerosols were a greater proportion of the aerosol concentration change in the reactor after that time.

Several researchers have measured SOA density, providing an estimated range of 0.6-1.5 g cm$^{-3}$ (Bahreini et al., 2005, Poulain et al., 2010, Qi et al., 2010, Song et al., 2007, Yu et al., 2008). In our study, we used a unit density (1.0 g cm$^{-3}$) to calculate SOA mass concentrations. This follows the approach used in Takekawa et al. (2003) and Verheggen et al. (2007).

3.2. Calculation of SOA yields

The fractional SOA yield ($Y$), defined as the ratio of the generated organic aerosol concentration ($M_o$) to the reacted hydrocarbon concentration ($\Delta$HC), was used to represent the aerosol formation potential of the hydrocarbon (Pandis et al., 1992). Odum et al. (1996) developed a gas/particle absorptive partitioning model to describe the phenomenon that $Y$...
largely depends on the amount of organic aerosol mass present. Equation (2) illustrates the relationship between SOA yield and organic aerosol mass concentration:

\[
Y = \frac{\Delta M_o}{\Delta HC} = M_o \sum_{i} \frac{A_i}{1 + K_{om,i}}
\]

In equation (2), \(i\) presents the serial number of the hydrocarbon reaction products, \(A_i\), \(\alpha_i\) and \(K_{om,i}\) (\(m^3 \mu g^{-1}\)) are the aerosol mass concentration, the stoichiometric coefficient based on mass and the normalized partitioning constant for product \(i\) respectively. If we assume that all semi-volatile products can be classified into one or two groups, equation (2) can be simplified to a one-product model (i.e., \(i=1\)) or two-product model (i.e., \(i=2\)). Parameters \(\alpha\) and \(K_{om}\) can be obtained by fitting the experimental SOA yield data with a least square method. Since numerous compounds are actually produced by the reaction of a hydrocarbon, parameters obtained by the simplified model only represent the overall properties of all products (Odum et al., 1996). A one-product model was proved sufficiently accurate to describe the relationship between aerosol yield and mass (Henry et al., 2008, Takekawa et al., 2003, Verheggen et al., 2007). Therefore, we used a one-product model for our experimental SOA yield data to quantify of the effects of inorganic seed aerosols on SOA formation.

3.3. Effects of CaSO\(_4\) and (NH\(_4\))\(_2\)SO\(_4\) seed aerosols on SOA formation

To investigate the effects of neutral and acid aerosols on SOA formation in \(m\)-xylene photooxidation, CaSO\(_4\) and (NH\(_4\))\(_2\)SO\(_4\) were selected as surrogates. Experimental conditions were listed in Table 1. Six seed-free experiments (Xyl-N1~6), three CaSO\(_4\)-introduced experiments (Xyl-CS1~3) and nine (NH\(_4\))\(_2\)SO\(_4\)-introduced experiments (Xyl-AS1~9) were carried out. Among these experiments, some experiments have identical initial conditions except for the seed aerosols (i.e. experiments Xyl-N5, Xyl-CS2, Xyl-AS2, Xyl-AS3, Xyl-AS9). Comparing the temporal variation of NO and O\(_3\) during these experiments with similar initial conditions (Figure 1), the results indicate that CaSO\(_4\) and (NH\(_4\))\(_2\)SO\(_4\) seed aerosols have no significant effect on gas-phase reactions. This result is consistent with the findings of Kroll et al. (2007) and Cao and Jang (2007) that (NH\(_4\))\(_2\)SO\(_4\) and (NH\(_4\))\(_2\)SO\(_4\)/H\(_2\)SO\(_4\) seed aerosols had a negligible effect on hydrocarbon oxidation.

Similarly, by comparing the temporal variation particle concentrations (Figure 2) during the experiments with identical initial conditions except for the seed aerosols, the effects of CaSO\(_4\) and (NH\(_4\))\(_2\)SO\(_4\) seed aerosols on SOA formation were identified. In Figure 2, PM\(_{corrected}\) was calculated from the measured PM concentrations plus wall deposit loss, and PM\(_0\) was the seed aerosol concentration. The results indicate that the presence of neutral aerosols CaSO\(_4\) (16-73\(\mu g\) m\(^{-3}\)) in the \(m\)-xylene/NO\(_x\) photooxidation system have no significant effect on SOA formation. Experiments with the presence of acid aerosols (NH\(_4\))\(_2\)SO\(_4\) have different particle profiles according to the concentrations of the introduced (NH\(_4\))\(_2\)SO\(_4\) seed aerosol. In Figure 2, experiment Xyl-AS2 has similar particle profile with the seed-free experiment Xyl-N5, indicating that (NH\(_4\))\(_2\)SO\(_4\) seed aerosols have little effect on SOA formation when the
initial concentration is low. However, when with high concentration of (NH₄)₂SO₄ seed aerosol introduced, SOA formation was enhanced (i.e. experiments Xyl-AS3 and Xyl-AS9) comparing with the seed-free experiment Xyl-N5. Comparing experiments Xyl-AS3 and Xyl-AS9, higher concentration of (NH₄)₂SO₄ seed aerosol resulted in higher SOA concentration. Therefore, the effects of (NH₄)₂SO₄ seed aerosol on SOA formation depend on its concentration.

| Experiment no. | HC₀ (ppm) | PM₀ (μm³ cm⁻³) | PM₀,₅ (cm² m⁻³) | NO₀ (ppb) | NOₓ₀ (ppb) | HC₀/NOₓ₀ | Mo (μg m⁻³) | ΔHC (ppm) | Y (%) |
|---------------|-----------|----------------|-----------------|------------|------------|-----------|-------------|-----------|------|
| Xyl-N1        | 0.92      | 0              | -               | 72         | 148        | 6.2       | 66          | 0.27      | 5.6  |
| Xyl-N2        | 1.26      | 0              | -               | 102        | 209        | 6.0       | 92          | 0.32      | 6.7  |
| Xyl-N3        | 1.74      | 0              | -               | 137        | 276        | 6.3       | 122         | 0.39      | 7.3  |
| Xyl-N4        | 1.68      | 0              | -               | 132        | 272        | 6.2       | 125         | 0.39      | 7.5  |
| Xyl-N5        | 2.00      | 0              | -               | 161        | 333        | 6.0       | 148         | 0.45      | 7.6  |
| Xyl-N6        | 2.51      | 0              | -               | 182        | 381        | 6.6       | 191         | 0.54      | 8.3  |
| Xyl-CS1       | 1.17      | 16             | -               | 86         | 174        | 6.7       | 67          | 0.29      | 5.4  |
| Xyl-CS2       | 2.03      | 43             | -               | 167        | 343        | 5.9       | 148         | 0.46      | 7.6  |
| Xyl-CS3       | 2.90      | 73             | -               | 232        | 471        | 6.2       | 201         | 0.59      | 8.0  |
| Xyl-AS1       | 2.85      | 11             | 3.5             | 208        | 420        | 6.8       | 208         | 0.57      | 8.6  |
| Xyl-AS2       | 2.06      | 23             | 6.7             | 166        | 337        | 6.1       | 150         | 0.45      | 7.7  |
| Xyl-AS3       | 2.15      | 47             | 11.0            | 162        | 326        | 6.6       | 169         | 0.45      | 8.8  |
| Xyl-AS4       | 0.92      | 43             | 13.2            | 70         | 137        | 6.7       | 93          | 0.27      | 8.1  |
| Xyl-AS5       | 1.73      | 45             | 12.8            | 132        | 272        | 6.4       | 165         | 0.40      | 9.7  |
| Xyl-AS6       | 2.41      | 55             | 13.0            | 178        | 365        | 6.6       | 232         | 0.53      | 10.3 |
| Xyl-AS7       | 0.92      | 63             | 16.7            | 70         | 143        | 6.4       | 110         | 0.26      | 10.1 |
| Xyl-AS8       | 1.56      | 69             | 17.1            | 132        | 269        | 5.8       | 173         | 0.35      | 11.5 |
| Xyl-AS9       | 2.07      | 74             | 17.1            | 166        | 348        | 6.0       | 249         | 0.47      | 12.3 |

Table 1. Initial experiment conditions and results for experiments with/without CaSO₄ or (NH₄)₂SO₄: initial m-xylene concentration (HC₀), initial seed aerosol mass concentration (PM₀), initial seed aerosol surface concentration (PM₀,₅), initial NOₓ concentrations (NO₀ and NOₓ₀-NO₀), ratio of HC₀/NOₓ₀, generated SOA mass (Mo), reacted hydrocarbon (ΔHC), and SOA yield (Y).

Further analysis found that the effects of (NH₄)₂SO₄ seed aerosol on SOA yield were positively correlated with the surface concentration of (NH₄)₂SO₄ seed aerosols. To draw the SOA yield curves shown in Figure 3, the experiments were classified into different groups (experiment Xyl-AS3 was not classified into any group since the surface concentration of (NH₄)₂SO₄ seed aerosols in this experiment was different from others) by the surface concentration of (NH₄)₂SO₄ seed aerosols. The regression lines for each group (there was no regression line for experiments XylCS1~2 and Xyl-AS1~3 since they had similar SOA yield with the seed-free experiments) were produced by fitting the data of generated SOA mass (Mo) and SOA yield (Y) into a one-product partition model. As indicated in Figure 3, experiments with higher surface concentration of (NH₄)₂SO₄ seed aerosols had higher yield curves. As proposed by most research, acid-catalyzed aerosol-phase reactions (Cao and
Jang, 2007, Czoschke et al., 2003, Gao et al., 2004) and uptake of semivolatile organic products to acidic sulfate aerosols enhance SOA formation (Liggio and Li, 2008). The observed SOA formation enhancement could be related to the acid catalytic effect of (NH$_4$)$_2$SO$_4$ seeds on particle-phase surface heterogeneous reactions and the surface uptake of semivolatile organic products.

Figure 1. Temporal evolutions of O$_3$ (a) and NO$_x$-NO (b) concentration in experiments with/without CaSO$_4$ and (NH$_4$)$_2$SO$_4$ seed aerosols.
3.4. Effects of Fe₂(SO₄)₃ and FeSO₄ seed aerosols on SOA formation

A seed-free experiment and three experiments with Fe₂(SO₄)₃ seed aerosols were carried out to investigate Fe₂(SO₄)₃ seed aerosols on photooxidation of toluene/NOₓ. The four experiments had identical initial conditions except for the concentrations of the introduced Fe₂(SO₄)₃ seed aerosol. Fe₂(SO₄)₃ seed aerosols did not have obvious effects on SOA formation as shown in the temporal variation of PM_corrected–PM₀ concentrations in Figure 4. Fe₂(SO₄)₃ seed aerosols had no obvious effect on gas phase compounds in toluene/NOₓ photooxidation either. A minimal amount of acid was added to the solution to generate Fe₂(SO₄)₃ seed aerosols. The introduced H⁺ concentration was in the range of 0.0002-0.002 µg m⁻³ in the Fe₂(SO₄)₃-
introduced experiments. This is much lower than the H⁺ concentration in the “non-acid” experiment by Cao and Jang (2007). Therefore, we presume the effect of the introduced sulfuric acid was negligible and Fe₂(SO₄)₃ seed aerosols did not have obvious effects on SOA formation in phooxidation of toluene/NOₓ.

We also conducted 18 irradiated toluene/NOₓ experiments with/without FeSO₄ seed aerosols. The conditions, generated SOA mass (Mₒ), and SOA yield (Y) are shown in Table 2. FeSO₄ seed aerosols had no obvious effect on gas phase compounds either, but significantly suppressed SOA formation. Figure 5 compares the temporal variation of particle concentrations during the 4.2 ppm toluene experiments (Experiments Tol-N3, Tol-FS1, Tol-FS3, Tol-FS8 and Tol-FS12) conducted under identical initial conditions except seed aerosol concentrations. Experiments with the presence of FeSO₄ seed aerosol generated less SOA than the seed-free experiment. And experiment with a higher FeSO₄ seed aerosol concentration generated less SOA than experiment with a lower FeSO₄ concentration. So the inhibited effect of FeSO₄ aerosols on SOA yield became stronger at higher concentrations of FeSO₄ seed aerosols. At other toluene/NOₓ photooxidation concentrations, we also found similar temporal variation of particle concentrations. However, as indicated in Table 2 and Figure 5, SOA yields of experiments Tol-FS1 and Tol-FS3 are similar to corresponding seed-free experiments of Tol-N3. These two seed-introduced experiments (as well as Tol-FS2) were conducted at the lowest ratio of FeSO₄ seed aerosol mass concentration to initial toluene mass concentration (FeSO₄/toluene) and did not show obvious effect on SOA formation comparing to their corresponding seed-free experiments. In these three experiments, the mass ratios of FeSO₄/toluene (assuming particle density to be 1.898 g cm⁻³, density of FeSO₄·7H₂O because of the lack of the information the amount of hydrate water) were calculated to be lower than 4.2×10⁻⁴. It is possible that most of the ferrous iron was oxidized before significant SOA mass were generated since few FeSO₄ seed aerosols were introduced and high concentrations of oxidizing substances were generated during the

Figure 4. Variations of generated SOA mass as a function of time from toluene/NOₓ photooxidation with different concentrations of Fe₂(SO₄)₃ seed aerosols
Effects of Inorganic Seeds on Secondary Organic Aerosol (SOA) Formation

toluene/NO\textsubscript{x} photooxidation. Besides these three experiments with lowest FeSO\textsubscript{4}/toluene mass ratio, FeSO\textsubscript{4} seed aerosols suppressed SOA formation relative to the corresponding seed-free experiments. And in our experiments, the suppress ratio could be as high as 60%, as calculated from Table 2.

| Experiment No. | HC\textsubscript{0} ppm | PM\textsubscript{0} μm\textsuperscript{3} cm\textsuperscript{-3} | NO\textsubscript{0} ppb | NO\textsubscript{x,0}-NO\textsubscript{0} ppb | PM\textsubscript{0}/HC\textsubscript{0} | HC\textsubscript{0}/NO\textsubscript{x,0} ppm/ppm\textsuperscript{-1} | M\textsubscript{o} μg m\textsuperscript{-3} | ΔHC ppm | Y % |
|----------------|--------------------------|-----------------------------|-------------------|----------------|-----------------|-------------------------------|-----------------|--------|-----|
| Tol-N1         | 1.10                      | 0                           | 50                | 51            | 0               | 11.0                          | 26              | 0.20   | 3.8 |
| Tol-FS4        | 1.08                      | 1                           | 51                | 50            | 5.1×10\textsuperscript{-4} | 10.7                          | 17              | 0.22   | 2.3 |
| Tol-FS10       | 1.07                      | 4                           | 55                | 47            | 1.7×10\textsuperscript{-3} | 10.6                          | 14              | 0.20   | 2.2 |
| Tol-FS14       | 1.09                      | 10                          | 48                | 49            | 4.4×10\textsuperscript{-3} | 11.1                          | 8               | 0.19   | 1.7 |
| Tol-N2         | 3.30                      | 0                           | 165               | 160           | 0               | 10.2                          | 90              | 0.48   | 5.0 |
| Tol-FS5        | 3.21                      | 4                           | 160               | 162           | 6.1×10\textsuperscript{-4} | 10.0                          | 74              | 0.51   | 3.9 |
| Tol-FS7        | 3.31                      | 6                           | 154               | 162           | 8.4×10\textsuperscript{-4} | 10.5                          | 72              | 0.56   | 3.5 |
| Tol-FS9        | 3.19                      | 11                          | 164               | 157           | 1.5×10\textsuperscript{-3} | 9.9                           | 59              | 0.47   | 3.3 |
| Tol-FS11       | 3.28                      | 21                          | 158               | 165           | 3.0×10\textsuperscript{-3} | 10.2                          | 36              | 0.51   | 1.9 |
| Tol-N3         | 4.12                      | 0                           | 217               | 210           | 0               | 9.7                           | 123             | 0.57   | 5.8 |
| Tol-FS1        | 4.23                      | 1                           | 208               | 207           | 1.4×10\textsuperscript{-4} | 10.2                          | 105             | 0.57   | 5.0 |
| Tol-FS3        | 4.25                      | 4                           | 208               | 213           | 4.2×10\textsuperscript{-4} | 10.1                          | 115             | 0.60   | 5.2 |
| Tol-FS8        | 4.25                      | 10                          | 216               | 209           | 1.1×10\textsuperscript{-3} | 10.0                          | 81              | 0.55   | 4.0 |
| Tol-FS12       | 4.23                      | 27                          | 213               | 210           | 3.0×10\textsuperscript{-3} | 10.0                          | 47              | 0.61   | 2.1 |
| Tol-N4         | 6.10                      | 0                           | 287               | 293           | 0               | 10.5                          | 189             | 0.96   | 6.3 |
| Tol-FS2        | 6.05                      | 5                           | 295               | 306           | 3.5×10\textsuperscript{-4} | 10.1                          | 170             | 0.81   | 6.5 |
| Tol-FS6        | 6.09                      | 10                          | 299               | 306           | 7.6×10\textsuperscript{-4} | 10.1                          | 140             | 0.88   | 4.8 |
| Tol-FS13       | 6.03                      | 41                          | 296               | 310           | 3.2×10\textsuperscript{-3} | 10.0                          | 64              | 0.82   | 2.7 |

Table 2. Experimental conditions and results in toluene photooxidation: initial toluene concentration (HC\textsubscript{0}), initial FeSO\textsubscript{4} seed aerosol concentration (PM\textsubscript{0}), initial NO\textsubscript{x} concentrations (NO\textsubscript{0} and NO\textsubscript{x,0}-NO\textsubscript{0}), ratio of PM\textsubscript{0}/HC\textsubscript{0}, ratio of HC\textsubscript{0}/NO\textsubscript{x,0}, generated SOA mass (M\textsubscript{o}), reacted hydrocarbon (ΔHC), and SOA yield (Y)

Figure 5. Temporal evolutions of SOA generation from toluene/NO\textsubscript{x} photooxidation with different concentrations of FeSO\textsubscript{4} seed aerosols
We classified the experiments with FeSO$_4$ seed aerosols introduced into three groups by FeSO$_4$/toluene mass ratios to create SOA yield variations as a function of generated SOA mass (Figure 6). Experiments with different FeSO$_4$/toluene mass ratios seemed to fall into different yield curves. When FeSO$_4$/toluene mass ratio was lower than $4.2 \times 10^{-4}$, FeSO$_4$ seed aerosols had a negligible effect and SOA yields of these experiments with FeSO$_4$ seed aerosols coincide with the yield curve of seed-free experiments. When FeSO$_4$/toluene mass ratio was higher than $5.1 \times 10^{-4}$, the SOA yield curve indicated experiments with FeSO$_4$ seed aerosols had lower yields than seed-free experiments. Lower yield curves from the experiments with higher FeSO$_4$/toluene mass ratio were observed, indicating that a higher Fe/C ratio had a greater suppression effect on SOA formation from toluene/NO$_x$ photooxidation.

![Figure 6. SOA yield (Y) variations as a function of generated SOA mass (M_o) from toluene/NO$_x$ photooxidation with/without FeSO$_4$ seeds](image)

### 3.5. Effects of mixed (NH$_4$)$_2$SO$_4$ and FeSO$_4$ aerosols on SOA formation

Atmospheric aerosol is often a mixture of different components. We tested the effect of internal mixed (NH$_4$)$_2$SO$_4$ and FeSO$_4$ seed aerosols on SOA formation in m-xylene/NO$_x$ photooxidation. The experimental conditions, generated SOA mass (M$_o$), and SOA yield (Y) are shown in Table 3. To generate internal mixed (NH$_4$)$_2$SO$_4$ and FeSO$_4$ aerosols, a mixed solution of (NH$_4$)$_2$SO$_4$ and FeSO$_4$, in which the mass concentration ratio of (NH$_4$)$_2$SO$_4$ to FeSO$_4$ was 5:1, was used in the atomizer. So the approximately 60 $\mu$m$^3$ cm$^{-3}$ seed aerosols in the three experiments with mixed (NH$_4$)$_2$SO$_4$ and FeSO$_4$ seed aerosols (Xyl-FA1~3) contained about 10 $\mu$m$^3$ cm$^{-3}$ FeSO$_4$ seed aerosols and 50 $\mu$m$^3$ cm$^{-3}$ (NH$_4$)$_2$SO$_4$ seed aerosols.

As mentioned above, neither (NH$_4$)$_2$SO$_4$ seed aerosols nor FeSO$_4$ seed aerosols had obvious effects on gas phase compounds. And in the experiments in this section, we found that mixed (NH$_4$)$_2$SO$_4$ and FeSO$_4$ seed aerosols had no obvious effect on gas phase compounds either.
In Figure 7, after wall deposition correction and deduction of seed aerosols, temporal variation of particle concentrations in experiments conducted under identical initial conditions except seed aerosol concentrations (the initial concentration of \( m \)-xylene is 1.1ppm, 2.1ppm and 3.2 ppm in picture a, b and c, respectively) were compared.

| Experiment No. | \( \text{HC}_0 \) ppm | \( \text{PM}_0 \) \( \mu \text{m}^3 \text{ cm}^{-3} \) | \( \text{NO}_0 \) ppb | \( \text{NO}_{x,0}-\text{NO}_0 \) ppb | \( \text{HC}_0/\text{NO}_{x,0} \) ppm ppm\(^{-1} \) | \( \text{M}_0 \) \( \mu \text{g m}^{-3} \) | \( \Delta \text{HC} \) ppm | \( \text{Y} \) % |
|---------------|---------------------|------------------|----------------|-------------------|----------------|----------------|----------------|----------------|
| Xyl-N7        | 1.08                | 0                | 62            | 62                | 8.7            | 21             | 0.30           | 1.7            |
| Xyl-FS1       | 1.01                | 7                | 58            | 63                | 8.4            | 8              | 0.29           | 0.7            |
| Xyl-AS10      | 1.07                | 44               | 63            | 65                | 8.3            | 51             | 0.32           | 3.7            |
| Xyl-FA1       | 1.05                | 62               | 64            | 69                | 7.9            | 30             | 0.31           | 2.3            |
| Xyl-N8        | 2.07                | 0                | 121           | 120               | 8.6            | 57             | 0.39           | 3.4            |
| Xyl-FS2       | 2.09                | 9                | 119           | 121               | 8.7            | 29             | 0.42           | 1.6            |
| Xyl-AS11      | 2.15                | 53               | 121           | 119               | 9.2            | 119            | 0.52           | 5.4            |
| Xyl-FA2       | 2.09                | 66               | 123           | 125               | 8.5            | 56             | 0.43           | 3.1            |
| Xyl-N9        | 3.21                | 0                | 198           | 180               | 8.5            | 145            | 0.74           | 4.6            |
| Xyl-FS3       | 3.23                | 11               | 188           | 182               | 8.8            | 48             | 0.63           | 1.8            |
| Xyl-AS12      | 3.10                | 48               | 182           | 178               | 8.5            | 213            | 0.71           | 7.0            |
| Xyl-FA3       | 3.16                | 57               | 179           | 186               | 8.7            | 117            | 0.74           | 3.7            |

Table 3. Experimental conditions and results in toluene photooxidation: initial toluene concentration (\( \text{HC}_0 \)), initial FeSO\(_4\) seed aerosol concentration (\( \text{PM}_0 \)), initial NO\(_x\) concentrations (\( \text{NO}_0 \) and \( \text{NO}_{x,0}-\text{NO}_0 \)), ratio of \( \text{HC}_0/\text{NO}_{x,0} \), generated SOA mass (\( \text{M}_0 \)), reacted hydrocarbon (\( \Delta \text{HC} \)) and SOA yield (\( \text{Y} \)).

As indicated in Figure 7(a), comparing with the seed-free experiment Xyl-N7, both experiment Xyl-AS10 and experiment Xyl-FA1 had higher particle concentrations while experiment Xyl-FS1 had lower particle concentrations. So, in 1.1ppm \( m \)-xylene photooxidation, the presence of \((\text{NH}_4)_2\text{SO}_4\) aerosols and mixed aerosols (mixed \((\text{NH}_4)_2\text{SO}_4\) and FeSO\(_4\)) both increased SOA formation, while the presence of FeSO\(_4\) suppressed SOA formation. In Figure 7(b) and Figure 7(c), the effects of single \((\text{NH}_4)_2\text{SO}_4\) seed aerosols (promotion effect) and single FeSO\(_4\) seed aerosols (suppression effect) on SOA formation were consistent with Figure 7(a). However, the mixed aerosols seemed to have different effects on SOA formation in photooxidation systems with different initial concentrations of \( m \)-xylene. In Figure 7(b), experiment Xyl-FA2 had similar temporal variation of particle concentrations with its corresponding seed-free experiment Xyl-N8, and in Figure 7(c), experiment Xyl-FA3 had lower temporal variation of particle concentrations than its corresponding seed-free experiment Xyl-N9. It must be noted that the seed aerosols in experiments Xyl-FA1-3 had similar concentrations and components. So, aerosols at the same mixing ratio of \((\text{NH}_4)_2\text{SO}_4\) and FeSO\(_4\) could either enhance or suppress SOA formation depending on the experimental conditions. It seemed that the promotion effect of \((\text{NH}_4)_2\text{SO}_4\) aerosols and the suppression effect of FeSO\(_4\) aerosols competed when both of them existed. And the promotion effect of \((\text{NH}_4)_2\text{SO}_4\) aerosols was dominant with low initial hydrocarbon concentration in the competition, while the reverse was true with high initial hydrocarbon concentration. This illustrates that the interplay of different compositions of real atmosphere aerosols can lead to complex synergistic effects on SOA formation.
According to the composition of the seed aerosols, experiments with inorganic seed aerosols introduced were classified into three groups. In Figure 8, SOA yield (Y) variations as a function of generated SOA mass (M₀) from m-xylene/NOₓ photooxidation were plotted. The
Effects of Inorganic Seeds on Secondary Organic Aerosol (SOA) Formation

regression lines for each group were produced by fitting the data of generated SOA mass ($M_o$) and SOA yield ($Y$) into a one-product partition model. As indicated in Figure 8, experiments with the presence of (NH$_4$)$_2$SO$_4$ had a higher SOA yield curve than the seed-free experiments, while experiments with the presence of FeSO$_4$ seed aerosols had a lower one, indicating the presence of (NH$_4$)$_2$SO$_4$ and FeSO$_4$ seed aerosols increased and decreased SOA yield, respectively. For the experiments with mixed seed aerosols, their SOA yield curve was similar to or a little higher than the seed-free experiments when the SOA mass load was low, but their SOA yield curve was lower than the seed-free experiments when the SOA mass load was high.

Figure 8. SOA yield ($Y$) variations as a function of generated SOA mass ($M_o$) from $m$-xylene/NO$_x$ photooxidation with/without FeSO$_4$, (NH$_4$)$_2$SO$_4$ and mixed FeSO$_4$ and (NH$_4$)$_2$SO$_4$ seed aerosols

3.6. Hypothesis for inorganic seed aerosols’ effects

In our experiment, we observed that FeSO$_4$ seed aerosols suppressed SOA formation while Fe$_2$(SO$_4$)$_3$ seed aerosols had no effect on SOA formation. It appears that the inhibiting effect of Fe(II) involves its strong reducing properties. Hydrocarbon precursors are oxidized by OH·, NO$_3$·, etc. During the gas phase reaction, the oxidized products usually have a lower saturation vapor pressure and, as a result, condense to the aerosol phase. When these oxidized condensable compounds (CCs) containing carbonyl, hydroxyl, and carboxyl groups (Gao et al., 2004, Hamilton et al., 2005) contact ferrous iron in the aerosol phase, they may react to produce ferric iron and less condensable compounds (LCCs) or incondensible compounds (ICs). The ferrous iron may stop some CCs from being further oxidized and forming low-volatility products (Hallquist et al., 2009), including oligomers (Gao et al., 2004). The experimental results also showed that the presence of neutral CaSO$_4$ seed aerosols seed aerosols have no significant effect on photooxidation of aromatic hydrocarbons, while the presence of acid (NH$_4$)$_2$SO$_4$ seed aerosols can significantly enhance SOA generation and SOA yield. A possible mechanism is shown in Figure 9. Oligomerization is one important step during SOA formation (Nguyen et al., 2011). As proposed by (Kroll et al., 2007), the effect of (NH$_4$)$_2$SO$_4$ seed aerosols may be attributed to acid catalyzed particle-phase reactions, forming high molecular weight, low-volatility products (e.g. oligomers). These processes may deplete the semivolatile CCs in the particle phase, and enhance SOA formation by shifting the gas-particle equilibrium, which is shown in
Figure 9, and, therefore force more CCs condense to aerosol phase. Since (NH₄)₂SO₄ and FeSO₄ seed aerosols may both influence the semivolatile CCs, there is a competition for CCs to form higher-volatility products (LCCs or ICs) or low-volatility products (e.g. oligomers).

Figure 9. Hypothesized mechanism for inorganic seed aerosols’ effects on SOA formation: ferrous iron Fe (II) reduces or decompose some condensable compounds (CCs), which are oligomer precursors, interrupting oligomerization and generating high volatility products (LCCs or ICs); while acid seed aerosols catalyze aerosol-phase reactions, generating oligomeric products

4. Conclusion

Effects of various inorganic seeds, including neutral inorganic seed CaSO₄, acidic seed (NH₄)₂SO₄, transition metal contained inorganic seeds FeSO₄ and Fe₂(SO₄)₃, and a mixture of (NH₄)₂SO₄ and FeSO₄ were examined during m-xylene or toluene photooxidation. Our results indicate that the presence of CaSO₄ seed aerosols and Fe₂(SO₄)₃ seed aerosols have no effect on photooxidation of aromatic hydrocarbons, while the presence of (NH₄)₂SO₄ seed aerosols and FeSO₄ seed aerosols have no effect on gas-phase reactions, but can significantly influence SOA generation and SOA yields. (NH₄)₂SO₄ seed aerosols enhance SOA formation and increase SOA yield due to acid catalytic effect of (NH₄)₂SO₄ seeds on particle-phase surface heterogeneous reactions. While FeSO₄ seed aerosols suppress SOA formation and decrease SOA yield possibly due to the reduction of some oligomer precursor CCs. These results reveal that many inorganic seeds are not inert during photooxidation process and can significantly influence SOA formation. These observed effects can be incorporated into air quality models to improve their accuracy in predicting SOA and fine particle concentrations.

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