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## Study of heterogeneous chemistry and photochemistry of single sea-spray aerosols containing Hg(II) in air using optical trapping – Raman spectroscopy†

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Sea-spray aerosols (SSAs) contribute to atmospheric loading, bringing toxic compounds like mercury (Hg) to the atmosphere, affecting the climate and human health. Despite their importance, the investigation into surface modification, including heterogeneous chemical and photochemical reactions of SSAs, is limited. In this work, we studied the heterogeneous chemistry and photochemistry of a single suspended SSA particle and a SSA containing Hg(II) in a reactive environment using optical trapping – Raman spectroscopy. The experiments are focused on the study of hygroscopicity, heterogeneous chemical reaction with ozone (O<sub>3</sub>), photochemical reaction with UVC radiation of an optically suspended single SSA particle, and photo-reduction of Hg(II) in SSAs under UVC radiation. Results show different Raman signal responses of a single SSA particle when it is optically trapped in air under varying relative humidity conditions as the aerosol particle uptakes and loses liquid water from the surrounding environment. The state and size of the aerosol are determined through the on-time images and different single-particle Raman spectral features. Results also show that the formation of chlorate (ClO<sub>3</sub><sup>-</sup>) is a reaction product of the heterogeneous reaction between the SSA particle and O<sub>3</sub>. The photochemical reaction products, as the SSA particle suspended in air under UVC radiation, are ClO<sub>3</sub><sup>-</sup> and perchlorate (ClO<sub>4</sub><sup>-</sup>). Further, we observed that these reactions occur only on the surface of the SSA particle. Based on the results, we hypothesize that Hg(II) can be photo-reduced to Hg(I) in SSAs through UVC radiation, and the amount of Hg(I) in SSAs is minor and balanced between the photo-oxidation and photo-reduction reactions.

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### Environmental significance

Sea-spray aerosols (SSAs) are an important component of atmospheric aerosols. The physical properties of SSAs, including hygroscopic properties, phase transitions, and component composition, have been well studied. However, the chemical temporal evolution, particularly surface modification, including heterogeneous chemical reactions and photochemical reactions, is less reported. This is a significant part of the aerosol's transmission in the atmosphere. Better understanding of their temporal evolution benefits actions on climate change, as well as disease control and prevention. We studied a single suspended SSA particle under a reactive environment using optical trapping, which simulates the behavior that might occur through a mitigation process of Hg(II). In particular, we studied the heterogeneous chemical reaction with O<sub>3</sub>, the photochemical reaction with UVC light, and the photo-reduction of Hg(II) in SSA. This work casts new insights into our understanding of the chemical reactions that occur on the surface of SSAs and Hg(II)-contained SSAs in air and serves as a reference for subsequent studies.

## 1. Introduction

Sea spray is a dominant source of atmospheric aerosol loading formed in the marine boundary layer (MBL) and transmitted through wind and waves, influencing the climate, air quality,

and human health.<sup>1–3</sup> Studies associated with the physical properties of a sea-spray aerosol (SSA), such as hygroscopic properties,<sup>4–10</sup> phase transition,<sup>11–13</sup> and component composition,<sup>14–16</sup> have been widely reported. However, the chemical reactions that occur on the surface of SSAs are less well studied. Sea spray is suspended in air and transmitted from the MBL to the atmosphere with gaseous reagents or air pollution surrounding it. The surface composition and properties of SSAs are subject to continuous change due to chemical reactions. For example, in the transmission process, SSAs react with air pollutants, such as SO<sub>2</sub>, NO<sub>2</sub>, and O<sub>3</sub>. These reactions can change the chemical composition of the surface of SSAs.<sup>16–18</sup>

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Furthermore, SSA's surface properties are continuously influenced by solar radiation *via* photochemistry processes.<sup>19</sup> Without knowing the chemical composition changes or reactions that occur on the surface of SSAs, understanding the atmospheric behavior and environmental impacts of SSAs remains challenging.

SSAs are composed of a variety of inorganic and organic compounds.<sup>14–16,20</sup> Despite the complex composition, the major components are water and NaCl, which play significant roles in aerosol size, state, and morphology. Hygroscopicity is an important physical property of NaCl aerosol. A suspended aerosol particle in air continues to absorb or lose water from the ambient environment.<sup>4,5</sup> The amount of water that an aerosol particle contains is mainly determined by its environmental relative humidity (RH) which influences significantly the properties of the aerosol particle such as size.<sup>8–10,21,22</sup> When the RH decreases, the suspended SSA in air shrinks due to water evaporation and eventually crystallizes at a threshold RH value. This process is efflorescence, and the threshold RH value is termed efflorescence RH (ERH). The converse process is deliquescence, in which the solid aerosol particle starts to take up water from the surrounding environment as the RH reaches a high value. This RH value is termed deliquescence RH (DRH). The reported ERH of SSA is in the range of 35–47% depending on the composition and concentration of the solutes.<sup>11,12,23–26</sup> The DRH is typically 30% higher than the ERH. The reported DRH of SSA is 70–77%.<sup>12,23–25</sup> Normally, the SSA particle in air is in the liquid state when the environmental RH is higher than the ERH. However, if the surrounding RH of the SSA particle decreases from a high value to ERH, the solute in the liquid aerosol is supersaturated and the aerosol particle may be in an amorphous glassy state, with gel-like structures.<sup>6,11</sup> The changes in the water content subsequently affect the solute concentration in the aerosol.<sup>27,28</sup> In addition, the water content also impacts the aerosol's optical properties, *e.g.* scattering coefficient,<sup>29,30</sup> and refractive index.<sup>31</sup> On the other hand, variations in the solute concentration change the pH value of the aerosol.<sup>32</sup> Furthermore, the RH affects directly the chemical reactions that happen on the aerosol's surface. Several studies have reported that the RH affects the heterogeneous chemical reactions that happen on the surface in the formation and evolution of secondary organic and inorganic aerosols in air.<sup>33–37</sup>

The other major component of SSA is NaCl. The ionic Cl in aqueous droplets can be oxidized to chlorate ( $\text{ClO}_3^-$ ) and perchlorate ( $\text{ClO}_4^-$ ), which are extremely toxic to humans. It was reported that  $\text{Cl}^-$  can be oxidized by  $\text{O}_3$  in the natural environment.<sup>38–40</sup> The reaction of  $\text{O}_3$  with  $\text{Cl}^-$  in aqueous solution was first reported by Yeatts *et al.*<sup>41</sup> The results showed that the reaction can be catalysed by  $\text{H}^+$  ions. Kang *et al.* reported that  $\text{ClO}_4^-$  can be formed *via* homogeneous photochemical reactions of aqueous oxychloride precursors.<sup>42</sup> Later on, they reported a number of potential reaction mechanisms of  $\text{ClO}_3^-$  and  $\text{ClO}_4^-$  production in the natural environment through the oxidation of  $\text{Cl}^-$  by  $\text{O}_3$ .<sup>38</sup> However, the reactions observed in the references mentioned above are measured from bulk solutions, where the heterogenous reaction between  $\text{Cl}^-$  and  $\text{O}_3$  can only happen on the interface between air and solution. Therefore,

the reaction is extremely slow. Different from the bulk solutions, NaCl aerosols suspended in air are on the order of micrometer in size with a high surface-to-volume ratio, which provides sufficient surface area for interactions with gaseous phase atoms and molecules, so the  $\text{O}_3$  can rapidly overwhelm the suspended droplet. In this case, the oxidation reaction that occurs on the surface of SSAs in the atmosphere would be rapid.

In addition to water and NaCl, toxic substances that SSAs carry, including mercury (Hg), are transmitted from the MBL to terrestrial environments.<sup>43,44</sup> This toxic compound can directly cause serious impacts on the environment and humans. The ocean is a significant reservoir for Hg. It is estimated that ocean waters contain  $3.5 \times 10^8$  kg of Hg.<sup>45</sup> The Hg from ocean emissions to the atmosphere is estimated at  $2.68 \times 10^6$  kg per year, which contributes approximately 40% of the total Hg input.<sup>46,47</sup> A significant amount of Hg that exists in SSAs is transmitted to terrestrial environments. Holmes *et al.* used a chemical box model to predict that about 85% of Hg is partitioned into SSAs under typical MBL conditions.<sup>48</sup> Moreover, the Hg concentration in SSAs is suggested to be higher than that found in seawater. Malcolm *et al.* reported that the Hg concentrations in SSAs can be up to 3000 times greater than in the marine system if seawater is the only source of the Hg to the aerosols.<sup>49</sup> In this case, a complete understanding of atmospheric mercury's chemical reactions and physical transformations that happened on the surface of SSAs is crucial to predicting mercury emissions.

Hg exists in the atmosphere in three different forms: elemental gaseous Hg (EGM or  $\text{Hg}(0)$ ), reactive gaseous Hg (RGM or  $\text{Hg}(II)$ ), and particulate bound Hg (PBM or  $\text{Hg}(p)$ ).<sup>50,51</sup> As most  $\text{Hg}(II)$  compounds are water-soluble, the Hg in SSAs is primarily  $\text{Hg}(II)$ , with the majority being  $\text{HgCl}_2$ . In the process of transmission of SSAs, the state of  $\text{Hg}(II)$  in the aqueous droplets will be affected by the surrounding environment. For example, SSAs contribute most RGM and PBM in the atmosphere in terms of the evaporation of  $\text{Hg}(II)$  compounds to the air and deposition of these species on the surface of aerosols.<sup>52</sup> Then, the RGM or PBM may further be reduced to  $\text{Hg}(0)$  under the specific condition.<sup>47,50,51</sup> The GEOS-Chem global model has postulated that  $\text{Hg}(II)$  photoreduction that occurs on the aqueous droplets in the atmosphere takes place in tropospheric clouds.<sup>52–54</sup> Malcolm *et al.* reported that SSAs could scavenge the RGM and significantly reduce  $\text{Hg}(II)$  levels in the air.<sup>49</sup> Tong *et al.* demonstrated that  $\text{Hg}(II)$  can be photolyzed to  $\text{Hg}(0)$  on the surface of NaCl aerosols under solar radiation.<sup>55</sup> Hedgecock *et al.* modeled the SSA effects on the photoreduction of  $\text{Hg}(II)$  and reported that the effect of the SSA is more related to the scavenging and deposition of  $\text{Hg}(II)$  than the catalytic reduction from  $\text{Hg}(II)$  to  $\text{Hg}(0)$ .<sup>56</sup> Many researchers have reported the redox reaction of  $\text{Hg}(II)$  and  $\text{Hg}(0)$  in the aquatic system.<sup>57–60</sup> However, all results from the references mentioned above are only known from computations or have been measured and tested experimentally from bulk systems (ocean and bulk aqueous solution). The reduction or photodissociation mechanisms of  $\text{Hg}(II)$  in the atmosphere remain uncertain. Due to the lack of understanding of the chemical-reaction mechanism of this process, the model



cannot provide an accurate description of the global mercury distribution.

To date, most atmospheric studies conducted in field or lab focused mainly on the composition and properties of freshly emitted SSA or SSA simulants.<sup>61</sup> However, atmospheric aging processes can transform SSA through reactions with trace gases, air pollutants, oxidants, and solar radiation.<sup>61–67</sup> Many SSA's atmospheric behaviors, such as the heterogeneous chemical reaction on aerosol modification,<sup>62–64</sup> the production of volatile organic compounds,<sup>61,65</sup> the formation of secondary organic aerosols (SOA),<sup>65</sup> and the formation of secondary marine aerosols (SMA)<sup>66,67</sup> etc., have not been well studied. This is mainly because of experimental difficulties in analyzing aerosols directly. For example, in current lab experiments, it is extremely challenging to accurately control the biological and chemical processes that lead to the formation of the SOA or SMA.<sup>65–67</sup> In the traditional method of aerosol characterization, measurements are made from the collected samples deposited on substrates, offering only information from the bulk materials. Furthermore, because of offline measurement, important information like the actual atmospheric behavior, such as changes in surface composition due to reactions and equilibrium processes, is missing. Another approach is to consider flowing aerosols in a tube or a reaction chamber. However, the aerosol particles will inevitably be affected by wall effects. To avoid these interferences and better understand the behavior of the aerosol in the atmosphere, investigation at the single-particle scale is of significance. Numerous studies that focus on the hygroscopic properties of single, suspended salt aerosols based on different techniques, including optical trapping (OT) and through an electrodynamic balance, have been reported.<sup>4–8,13,22</sup> These works only focus on physical properties, e.g., phase transition, of the aerosols. No surface-composition changes and chemical reactions of SSAs at the single-particle scale have been reported. However, this missing part is crucial for understanding SSA's influence on atmospheric processes and human health.

In this work, we study the heterogeneous chemistry and photochemistry of a single, suspended SSA particle and SSA containing Hg(II) in a reactive atmospheric environment using optical trapping – Raman spectroscopy (OT-RS). This paper consists of two parts. Part I studies the heterogeneous chemistry and photochemistry of the optically trapped single SSA particle, and Part II considers the photo-reduction reaction of Hg(II) in SSA. The major objectives of this two-part study are: (1) through the size and spectral evolution of an SSA particle trapped under different RH conditions, we estimate the size of the aerosol particle using different Raman signal features. (2) By investigating the heterogeneous chemical reaction with O<sub>3</sub> and the photochemical reaction with UVC radiation of the SSA particle, we show that the ionic Cl in the SSA can be oxidated to chlorate (ClO<sub>3</sub><sup>−</sup>) by O<sub>3</sub> and further oxidated to perchlorate (ClO<sub>4</sub><sup>−</sup>) by UVC radiation, and that these reactions occur only on the surface of SSA. (3) *Via* the study of the photochemical reaction of Hg(II) in SSA particles with UVC radiation, we shed insight on the photo-reduction and photo-oxidation reactions between Hg(II) and Hg(I) in the SSA. This study from the single particle

provides unique insight into the connections between atmospheric chemistry and aerosol properties and demonstrates how OT-RS can be used to isolate and study reactions of the single particle in a reactive environment in the laboratory under controlled conditions. This work will be a reference for subsequent research work that applies to real atmospheric environmental conditions in the future.

## 2. Methodology

### 2.1 Experimental setup

Fig. 1 is the schematic of the experimental system. The details of the experimental setup of OT-RS can be found in our previous publications<sup>68–70</sup> and are briefly described here. A 532 nm (TEM<sub>00</sub>) continuous Gaussian laser beam is used as the trapping laser beam. The laser power is set at 1740 mW in this experiment to trap all the droplet particles. A beam splitter splits the trapping laser beam into two perpendicular beams of equal intensity. These two beams pass through axicons to form hollow beams. The universal optical trap is formed in the overlapping region of these two counter-propagating hollow beams through two micro-objectives (×50, numerical aperture (NA) = 0.55) as shown in Fig. 1. The size of the optical trap can be varied from ~10 μm to 100 μm by tuning the position of the two micro-objectives. The scattering signal from a trapped droplet particle can be collected through another micro-objective (×20, NA = 0.42). The collected information passes through a dichroic beamsplitter. The short passing beam is delivered to a camera to show the trapped particle's physical information, such as size, shape, and position. The long passing beam is directed to a spectrograph (Acton 2300i, Princeton) that shows the inelastic scattering signal containing the chemical properties, e.g., the vibrational band information. The Raman information is dispersed by a 600 lines per mm grating and then projected onto an electromagnetic charge-coupled device (EMCCD, ProEM, Princeton), which has a spectral resolution of 0.05 nm per pixel. With the experimental setup in this work, the best spatial resolution that can be achieved is 600 nm × 600 nm.<sup>71</sup> The integration time of spectra is set to 60 s, and the spectrograph slit is set to 100 μm.

The aerosol particle is generated through a humidity diffuser and introduced directly into the chamber. A light is installed on the side of the chamber to illuminate the trapped particle. Ozone is generated through a homemade ozone generator and introduced through the same hole as the humidity diffuser after introducing the aerosol particles. At the bottom of the chamber, a hole is connected to a mechanical pump, which is controlled by a valve at a slow pumping rate of 0.01 L min<sup>−1</sup>. The ozone flows through the trapped droplet and then is pumped out through the bottom hole. The ozone flow is also set at a low rate of 0.01 to 0.2 L min<sup>−1</sup>. The pressure and RH around the trapped droplets are not affected as both the pumping and flowing rates are slow. The RH and temperature sensor are fixed inside the chamber to monitor the local conditions. In addition to the hole on the bottom, which is connected to a valve, the top hole of the chamber is open during the experiment, so the pressure and temperature (19–21 °C) inside the chamber are the same as in



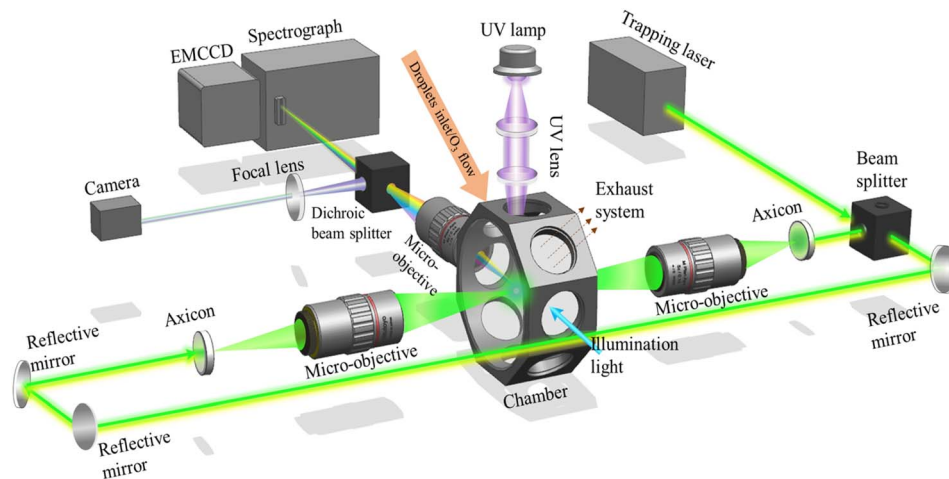


Fig. 1 Schematic of the experimental setup of the OT-RS system.

the lab environment. The humidity diffuser keeps generating droplet aerosols for 30 to 60 seconds until one droplet is stably trapped. As the chamber is not sealed, most of the excessive mist will be pumped out through the bottom hole, and the few leftovers pass out of the chamber through the open hole to the exhaust system. In the experiment, two RH sensors are installed, one installed inside of the chamber, and another installed outside of the chamber. The sensor inside the chamber shows increase of RH when introducing the droplets aerosol, and decrease of RH when the aerosol introduction is stopped. Finally, after the aerosol particle is trapped for several minutes, the RH shown by the two sensors becomes the same. In this case, after several minutes of stopping aerosol particle introduction, the RH around the trapped particle is consistent with the laboratory RH. The recorded RH in the lab for the whole year is in the range of 30–70%. For the extremely high RH cases ( $RH > 70\%$ ), an external humidifier is connected to maintain the high RH condition inside the chamber. The UVC radiation is generated from a UV lamp (90-0049-01/Photochemical Lamps, Analytik Jena US), which is fixed on the top area of the chamber. Two wavelengths can be generated, 254 nm (>95%) and 185 nm (<5%), both of which are in the UVC region, with an irradiance intensity of  $4000 \mu\text{W cm}^{-2}$ . The UVC radiation is focused into the chamber through two UV lenses. Counting the distance of 2.5 cm between the first lens and UV lamp, and the maximum energy loss of 20% at 254 nm of the two lenses, the irradiance applied on the trapping area is calculated as  $512 \mu\text{W cm}^{-2}$ .

## 2.2 Samples

The droplets are produced by ultrasonic nebulization through a humidity diffuser of aqueous solutions. SSA particles used in this work were SSA simulants generated from the 3.5% NaCl solution. The solution to generate droplets of SSA containing Hg(II) is made by adding 1%  $\text{HgCl}_2$  into 3.5% NaCl solution. The NaCl (>99.5%) was purchased from Fisher Scientific, and the  $\text{HgCl}_2$  (>99.5%) was purchased from Sigma-Aldrich.

## 3. Results and discussion

### 3.1 Part I: the study of heterogeneous chemistry and photochemistry of the optically trapped single SSA

**3.1.1 Effects of surrounding RH on size and Raman spectral features of an SSA particle.** In the experiment, the trapped droplet continues to uptake or lose water from the ambient environment during the trapping period. Fig. 2a shows the Raman spectrum of the droplet trapped in a dry environment with a RH of 37%. The Raman bands and band assignments are shown in Table S1.† Three Raman bands at  $185 \text{ cm}^{-1}$ ,  $1630 \text{ cm}^{-1}$ , and  $3430 \text{ cm}^{-1}$  are observed and assigned to O–H...O stretching mode, OH-bending mode, and OH-stretching mode of water, respectively. The spectrum of Fig. 2a is from an aerosol particle trapped under RH of 37%. Under this dry condition, the OH stretching mode is extremely weak. This is because the trapped NaCl aerosol particle experiences efflorescence, and the solute becomes supersaturated.<sup>11</sup> In this experiment, the ERH we calculated is  $39\% \pm 8\%$ , which is consistent with the reported results.<sup>11,12,23–26</sup> This value was obtained from 20 sets of data in which the Raman signals have no or very weak OH-stretching band. Therefore, the dry and wet conditions in this experiment is defined as when the RH are below and above  $39\% \pm 8\%$ , respectively. The size of the inset image is 500 pixels  $\times$  500 pixels, corresponding to approximately  $83.3 \mu\text{m} \times 83.3 \mu\text{m}$ , which was used to estimate size of each droplet shown in Fig. 2. For instance, the size of the trapped droplet in Fig. 2a is  $4.4 \mu\text{m}$ . If the RH further decreases, the SSA undergoes efflorescence and crystallization, forming a solid particle.<sup>11</sup> Fig. 2b shows the Raman spectrum of the SSA trapped in air with a RH of 55%. Different from Fig. 2a, a wide and strong Raman band of OH-stretching located around  $3430 \text{ cm}^{-1}$  is observed. This indicates that the droplet is in the aqueous state. Thus, the NaCl inside the droplet is in the ionic state. The size of the droplet shown in the on-time image in Fig. 2b was estimated to be  $9.3 \mu\text{m}$  under this RH condition. Because the amount of water a particle can absorb is affected by the surrounding RH, therefore, under different RH conditions,





Fig. 2 (a–d) Single-particle Raman spectra of the SSA particle trapped in air under different RH. The inset pictures are the on-time images corresponding to the Raman spectra, captured by the system camera. (e) A video showing the SSA particle's size evolution as the surrounding RH increases from 41% to 69%.

the intensity of the OH band is different. As shown in Fig. S1,<sup>†</sup> the intensity of the Raman signal changes as the surrounding RH changes. The on-time images, which show the size of the droplets corresponding to different RH, are shown in the insets of Fig. S1a.<sup>†</sup> The Raman signal intensity is increasing or decreasing with the changing of the size of the SSAs. Fig. S1b<sup>†</sup> shows the relationship between the intensity of Raman band at  $3430\text{ cm}^{-1}$  and the size of the trapped droplet. The result shows that the Raman band intensity is linear to the particle size, which implies that the size of the SSAs can be estimated using the Raman spectral features. Further increasing RH would increase the water content of the aqueous droplet, *i.e.*, the aqueous particle would become more diluted,<sup>11</sup> and the size of the droplets would also increase simultaneously. When the trapped droplets reach a specific size, the whisper gallery mode (WGM) can be observed from the Raman spectrum.<sup>72</sup> In the experiment, as the surrounding RH is above  $69\% \pm 5\%$  (similarly, this number was obtained from 20 sets of data), the WGM can be observed. Fig. 2c and d show the WGM in the Raman spectra of the NaCl droplet trapped in air with RH of 70% and 85%. The size of the droplets in these two cases are  $18.5\text{ }\mu\text{m}$  and  $27.8\text{ }\mu\text{m}$ , respectively. Fig. 2e is a snapshot from a video that shows the size evolution as the SSA particle trapped in air with the surrounding RH increases from 41% to 69%. The video's playing speed has been accelerated by 80 times; this whole process lasts about 22 minutes in the experiment.

**3.1.2 The heterogeneous chemical reaction between  $\text{O}_3$  and  $\text{Cl}^-$  in a single SSA particle.** Fig. 3 shows the time-resolved Raman spectra of a trapped SSA particle exposed to ozone at a RH of 62%. The ozone concentration used in this experiment is 1000 ppm. In the experiment, the  $\text{O}_3$  concentration is measured in an independent chamber on the way of  $\text{O}_3$  flowing.

So the measured  $\text{O}_3$  concentration is a direct result from the ozone generator. The  $\text{O}_3$  flows into the chamber and through the trapped aerosol particle. The  $\text{O}_3$  concentration is much diluted in the chamber as the flowing rate is very low, and the chamber is opened to air, so that the air inside and outside the chamber is constantly being exchanged. But compared with the volume of the trap, the covered area of  $\text{O}_3$  flow is much larger. In this case, 1000 ppm is an upper limit of  $\text{O}_3$  concentration involved in the reaction. In Fig. 3, one peak appears and increases slowly with time after the droplet is trapped in the ozone environment for 20 minutes. This peak is located at  $966\text{ cm}^{-1}$  and attributed to  $\text{ClO}_3^-$ . The  $\text{ClO}_3^-$  is generated through oxidation of the ionic Cl in the aqueous solution by  $\text{O}_3$ .<sup>17,38,39,73,74</sup> The reactions are as follows from (R1) to (R3.2).



However,  $\text{ClO}^-$  also can be reduced to  $\text{Cl}^-$  through (R4). The reaction rates  $k_1 = 3 \times 10^{-3}\text{ M}^{-1}\text{ s}^{-1}$  and  $k_2 = 30\text{ M}^{-1}\text{ s}^{-1}$  are much smaller than  $k_4 = 110\text{ M}^{-1}\text{ s}^{-1}$  at room temperature around  $20\text{ }^\circ\text{C}$ .<sup>38,75</sup> Therefore, the production of  $\text{ClO}_3^-$  is very limited. However, these reaction rates were measured from a bulk aqueous solution, in which the reaction happens only on the interface between the liquid solution and air.<sup>76</sup> Compared





Fig. 3 (a) The time-resolved Raman spectra indicate the formation of  $\text{ClO}_3^-$  as the SSA particle is exposed to  $\text{O}_3$  at 1000 ppm and RH of 62%. (b) Time variations of the intensity of the Raman band of  $\text{ClO}_3^-$  at  $966\text{ cm}^{-1}$  as the single trapped SSA particle is exposed to different  $\text{O}_3$  concentrations.

with the reaction that happens only on the interface, ozone can soon overwhelm the droplet particle as it is suspended in air because of the high surface-to-volume ratio, which greatly accelerates the reaction. Fig. 3b shows the peak intensity of  $966\text{ cm}^{-1}$  increasing as the droplets are trapped in different  $\text{O}_3$  concentrations with the RH of 54–62%. Each data point in Fig. 3b was averaged over three points and normalized to the first minute's peak intensity. In Fig. 3b, the Raman band of  $\text{ClO}_3^-$  increases slowly with the peak intensity at a rate of  $0.0015\text{ minute}^{-1}$  when the trapped droplets are exposed to  $\text{O}_3$  at 600 ppm. The reaction becomes faster with increasing ozone concentration. As shown in Fig. 3b, the intensity increases at a rate up to  $0.027\text{ minute}^{-1}$ , 18 times faster, as the  $\text{O}_3$  concentration increases to 4000 ppm. After the reaction is completed within 30 minutes, the band intensity still increases slightly. This is because that the reaction happens only on the surface of the suspended droplet (details shown in Section 3.1.3). As the ion exchanges between the surface and the interior consistently happen in the aqueous system after all the  $\text{Cl}^-$  has been oxidized on the surface, the supplied  $\text{Cl}^-$  continually reacts with ozone and produces  $\text{ClO}_3^-$ .

No change in the Raman bands is observed when the ozone concentration is below 600 ppm. This is because the concentration of  $\text{ClO}_3^-$  is too low to be observed in the first several hours. Furthermore, since (R4) is much faster than (R1), with low  $\text{O}_3$  concentrations, the  $\text{ClO}^-$  is produced at a lower rate and balanced with  $\text{Cl}^-$  in the aqueous system of the droplet, and no further reaction of (R2) and (R3) happens. Therefore, there will be less  $\text{ClO}_3^-$  produced. Nevertheless, even though the reaction is faster in the suspended droplets than in bulk solution, with the low concentration of  $\text{O}_3$  in air, this reaction is still slow in the natural environment. On the other hand, this reaction was observed only at the environmental RH of 39–69%. Experimentally, due to the loss of water, the solute in the suspended NaCl aerosol particle becomes supersaturated in the dry environment when the surrounding RH is lower than  $39\% \pm 8\%$ . Under this condition, the ionic Na and Cl are bounded to form

NaCl molecules. Therefore, the amount of  $\text{Cl}^-$  on the surface of the droplet is not sufficient to produce enough  $\text{ClO}_3^-$  that is observable.

Compared with the reported Raman band position of  $\nu_1$  ( $\text{ClO}_3^-$ ), which is located around  $940\text{ cm}^{-1}$ ,<sup>77,78</sup> the Raman band position in this study is blue-shifted to  $966\text{ cm}^{-1}$ . This may be caused by a change in the Cl–O band frequency due to the adsorbate and adsorbent interaction, which is absent in the bulk liquid state.<sup>79,80</sup> Furthermore, the concentration of NaCl also affects the force of the Cl–O band, affecting the vibrational frequency. Zachhuber *et al.* reported that the Raman band of  $\nu_1$  ( $\text{ClO}_3^-$ ) of a mixture of 50% NaCl and 50%  $\text{NaClO}_3$  shifted to  $998\text{ cm}^{-1}$ .<sup>81</sup> Additionally, changes in pH value and temperature in the aqueous environment as a result of laser heating can also slightly change the Raman band position of  $\text{ClO}_3^-$ . All the factors above are possibly attributed to the blue-shifting of the Raman band of  $\text{ClO}_3^-$  observed in this result.

**3.1.3 The photochemical reaction of single suspended SSA particle with UVC radiation.** Time-resolved Raman spectra of a SSA exposed to UVC of 185 nm and 245 nm radiation are shown in Fig. 4. The surrounding RH of the trapped droplet is



Fig. 4 Changes in the time-resolved Raman spectra indicate the formation of  $\text{ClO}_3^-$  and  $\text{ClO}_4^-$  as the SSA particle is trapped under UVC radiation with RH of 60%.



60%. The Raman band assignments corresponding to Fig. 4 are shown in Table S1.† Compared with the Raman spectra of the aerosol droplet trapped in the first minute, five more Raman bands were observed after exposure to the UVC radiation. Among them, four bands at 482 cm<sup>-1</sup>, 621 cm<sup>-1</sup>, 979 cm<sup>-1</sup>, and 1023 cm<sup>-1</sup> are attributed to ClO<sub>3</sub><sup>-</sup>, and one weak band located at 1124 cm<sup>-1</sup> is assigned to ClO<sub>4</sub><sup>-</sup>. Different from the SSAs exposed to O<sub>3</sub>, the Raman band of ν<sub>1</sub> (ClO<sub>3</sub><sup>-</sup>) at 966 cm<sup>-1</sup> shifts to 979 cm<sup>-1</sup>. As discussed in the previous section, this can be explained by the adsorbate and adsorbent interaction in the single-particle scale and the changes in concentration of the Cl<sup>-</sup> and ClO<sub>3</sub><sup>-</sup> inside the droplet. Nevertheless, the intensity of ν<sub>1</sub> (ClO<sub>3</sub><sup>-</sup>) became much stronger, and more Raman bands related to ClO<sub>3</sub><sup>-</sup> were observed, which indicates that more oxidated product of ClO<sub>3</sub><sup>-</sup> was produced due to the UVC radiation than that in the condition of the droplets exposed to ozone. Note that the vibration frequency of ν<sub>3</sub> (ClO<sub>3</sub><sup>-</sup>), which is reported to be 976 cm<sup>-1</sup> (ref. 82 and 83) blue-shifts to 1023 cm<sup>-1</sup>, but the frequency difference between ν<sub>1</sub> (ClO<sub>3</sub><sup>-</sup>) and ν<sub>3</sub> (ClO<sub>3</sub><sup>-</sup>) is the same as reported.<sup>82,83</sup> As ν<sub>1</sub> (ClO<sub>3</sub><sup>-</sup>) and ν<sub>3</sub> (ClO<sub>3</sub><sup>-</sup>) are from the stretching vibration, these two bands show consistent frequency shifts, while the bending bands of ν<sub>2</sub> (ClO<sub>3</sub><sup>-</sup>) and ν<sub>4</sub> (ClO<sub>3</sub><sup>-</sup>) retain the same frequency as reported.<sup>82,83</sup>

In this reaction, UVC radiation photolyzes oxygen to produce ozone around the trapped droplets, and O<sub>3</sub> reacts with the ionic Cl on the surface of the droplets to produce ClO<sup>-</sup>, which is the same as (R1) in Section 3.1.2. Additionally, due to the reaction product of O(<sup>3</sup>P) and O(<sup>1</sup>D) from the photolysis of O<sub>3</sub>,<sup>84</sup> more chemical reaction pathways can exist, which lead to further production of ClO<sub>3</sub><sup>-</sup>, e.g., *via* reactions (R5)–(R7) below.<sup>40,42</sup> This explains the much stronger Raman band of ClO<sub>3</sub><sup>-</sup> observed.



One Raman band that is attributed to ClO<sub>4</sub><sup>-</sup> was also observed. This indicates ClO<sub>3</sub><sup>-</sup> is further oxidated by O(<sup>1</sup>D) to ClO<sub>4</sub><sup>-</sup> *via* (R8). In addition to the Raman band related to the ClO<sub>3</sub><sup>-</sup> and ClO<sub>4</sub><sup>-</sup>, a wide Raman band located in the range 2900–2960 cm<sup>-1</sup> appears and rises slowly as the particle is trapped. This wide band indicates the formation of H<sub>2</sub>O<sub>2</sub>.<sup>85</sup> The H<sub>2</sub>O<sub>2</sub> formation is due to the reaction between O(<sup>1</sup>D) and H<sub>2</sub>O (R9). The detailed reaction pathways can be found in ref. 86 and 87, while the major reaction is (R9).

In addition to the new Raman bands of ClO<sub>3</sub><sup>-</sup> and ClO<sub>4</sub><sup>-</sup>, it is important to notice the changes in the OH-stretching band. Compared with the spectra in the first minute and after 60 minutes, the center of the band moves from 3430 cm<sup>-1</sup> to

3450 cm<sup>-1</sup>. This is mainly due to the Cl<sup>-</sup> being oxidized to ClO<sub>3</sub><sup>-</sup> and ClO<sub>4</sub><sup>-</sup>.<sup>88</sup> However, this happens only on the surface of the trapped droplets. Fig. 5 shows a contour plot of the Raman spectrum intensity distribution of the SSA trapped under UVC radiation for 60 minutes. Two places marked as S and C correspond to the surface and center of the trapped droplet, and the image shows the position of these two locations. The inset also shows the Raman spectra of the OH-stretching from these two locations. The maximum intensity of the two Raman spectra is normalized to be the same. Comparing the spectra from these two locations, the Raman band of OH-stretching is located at 3430 cm<sup>-1</sup> in position C, while it moves to 3450 cm<sup>-1</sup> in position S. This suggests that the ion distribution is also different between the surface and the center of the trapped droplet. In particular, more ClO<sub>3</sub><sup>-</sup> is located on the surface of the droplet, and more Cl<sup>-</sup> is in the center. This suggests the reaction between Cl<sup>-</sup> and O<sub>3</sub> happened only on the surface of the droplet.

Fig. 6 shows the time-resolved Raman spectra of a single SSA particle exposed to UVC radiation and trapped in a dry environment with the RH of 38%. The intensity of the OH-stretching band around 3400 cm<sup>-1</sup> becomes weaker under this condition. The Raman spectra show only one new Raman band rising as the droplets are exposed to UVC radiation under this dry condition, which is located at 2930 cm<sup>-1</sup>. This band indicates the formation of H<sub>2</sub>O<sub>2</sub>. No other band change was observed during this process. Therefore, different from the SSA trapped in a higher RH environment, no oxidation reaction of Cl<sup>-</sup> was observed. This also suggests that the reaction between Cl<sup>-</sup> and O<sub>3</sub> occurs only when the environmental RH is relatively high. As discussed in Section 3.1.2, the solute in the suspended droplet becomes saturated in a dry environment, and ionic Cl is in the state of NaCl rather than free Cl<sup>-</sup> in the aqueous system. In this case, without enough Cl<sup>-</sup> appearing on the surface of the suspended droplets, the oxidation reaction with O<sub>3</sub> becomes less likely; thus, the reaction product of ClO<sub>3</sub><sup>-</sup> is hard to observe.

Under the UVC radiation, the Cl<sup>-</sup> on the surface of the droplets is oxidized to ClO<sub>3</sub><sup>-</sup> when the single SSA particle is trapped in air with a high RH condition. The reaction is much faster than that of a single SSA particle exposed to ozone at



Fig. 5 Contour plot of the spectral intensity distribution of the OH-stretching mode for Raman spectra of SSA. Two places are marked where S is the surface and C is the center. The inset picture shows the on-time image captured by the system camera; the inset figure shows the position-resolved Raman spectra from the two marked locations.





Fig. 6 Time-resolved Raman spectra of a single, suspended SSA particle is exposed to UVC radiation with RH of 38%. No Raman band changes occur except for the peak at  $2930\text{ cm}^{-1}$  appearing.

4000 ppm, even though (R1) is the same for both oxidation pathways. This is mainly because the atomic O that is photolyzed from  $\text{O}_3$  enhances this process *via* additional reactions of (R5)–(R7). The same oxidation reaction also can occur on the surface of a SSA in the atmosphere. While the high concentration of  $\text{O}_3$  and UVC radiation are not reflective of the natural atmosphere, the experiment does confirm oxidation pathways suggested previously.<sup>38–41</sup>

### 3.2 Part II: the study of heterogeneous chemical reactions and photochemistry of a single SSA containing Hg(II)

**3.2.1 Effects of surrounding RH on size and Raman spectral features of a SSA containing Hg(II).** Fig. 7 shows the Raman spectra of a SSA particle containing Hg(II) trapped in air under different RH conditions. Fig. 7a shows the Raman spectra of the droplet trapped in a dry environment with a RH of 39%. One more Raman band, compared with pure SSA, located at  $314\text{ cm}^{-1}$ , is observed, which is attributed to  $\text{HgCl}_2$ .<sup>89</sup> As discussed in Section 3.1.1, the droplets do not crystallize, but the solute in the aqueous system becomes supersaturated when the surrounding RH is low. In this case, part of the free  $\text{Hg}^{2+}$  inside

the aqueous droplets is bound to  $\text{Cl}^-$  to form  $\text{HgCl}_2$  molecules in an aqueous system. So, the Raman signal of the  $\text{HgCl}_2$  band can be observed. With different RH, the band of  $\text{HgCl}_2$  shows different intensities, indicating an increase or decrease in the amount of  $\text{HgCl}_2$  molecules. The related figures are shown in Fig. S2.† In the experiment, the ERH for SSA containing Hg(II) is calculated to be  $41\% \pm 11\%$ . This value is calculated through 20 data sets in which the Raman signals show the Raman band of  $\text{HgCl}_2$ . Due to the  $\text{HgCl}_2$  added, the ERH of SSA containing Hg(II) is slightly higher than the pure SSA. When the RH increases above the ERH of SSA containing Hg(II), as shown in Fig. 7b, the  $\text{HgCl}_2$  vibration band at  $314\text{ cm}^{-1}$  disappears, and the Raman signal of the OH-stretching band around  $3400\text{ cm}^{-1}$  is observed. This indicates that Hg(II) turns to the ionic state in the aqueous system under the higher RH condition. The changes in size are similar to the pure SSA as discussed in Section 3.1.1, which indicates that the hygroscopicity of SSA particles is not affected by the Hg(II) contained in the droplet.

**3.2.2 The heterogeneous chemical reaction between SSA containing Hg(II) and  $\text{O}_3$ .** The time-resolved Raman spectra of the droplets of SSA containing Hg(II) exposed to  $\text{O}_3$  are shown in Fig. S3.† The  $\text{O}_3$  concentration is 1000 ppm. Fig. S3a† shows the single droplet trapped in an  $\text{O}_3$  environment with a surrounding RH of 58%. This figure is similar to the pure SSA in Fig. 3 with one peak located at  $966\text{ cm}^{-1}$ , which is attributed to  $\text{ClO}_3^-$  that increases slowly with time. The peak intensity increases with the increase of the  $\text{O}_3$  concentration, and the rate is also similar to the results in Fig. 3b of pure SSA. Fig. S3b† shows the SSA containing Hg(II) exposed to  $\text{O}_3$  of 1000 ppm in a RH of 35%. As mentioned in Section 3.2.1, when the RH is below the ERH of  $41\% \pm 11\%$ , the solute in the droplets becomes saturated, and the peak at  $314\text{ cm}^{-1}$  of  $\text{HgCl}_2$  is observed. In Fig. S3b,† the peak at  $314\text{ cm}^{-1}$  is observed in the first minute and kept steady for the rest of the trapping time. In this case, the reaction mechanism of the SSA containing Hg(II) is the same as the pure SSA with the reaction product  $\text{ClO}_3^-$  when the RH is above the ERH of SSA containing Hg(II). No reaction between  $\text{O}_3$  and  $\text{HgCl}_2$  was



Fig. 7 Single-particle Raman spectra of the SSA containing Hg(II) trapped in air under different RH. The inset figure shows the Raman spectra with high resolution in the region of  $150\text{--}400\text{ cm}^{-1}$ . The Raman band of  $\text{HgCl}_2$  at  $314\text{ cm}^{-1}$  can only be resolved as the SSA particle is trapped in a dry environment with RH below  $41\% \pm 11\%$ .



observed. This result indicates that the  $\text{HgCl}_2$  molecule in the suspended droplet is stable when the SSA particle is exposed to  $\text{O}_3$ .

**3.2.3 The photochemical reaction of  $\text{Hg(II)}$  in SSA under UVC radiation.** The Raman spectra of a SSA containing  $\text{Hg(II)}$  trapped with RH of 56% exposed to UVC radiation are shown in Fig. S4.† The spectra show similar changes to those in Fig. 4, which indicates the reaction products are similar to those of the pure SSA. However, compared to Fig. 4, a blue-shift in the vibrational band of OH-stretching at  $3457\text{ cm}^{-1}$  is observed, as shown in Fig. S4.† This difference could be caused by the stronger ionic force as the concentration of  $\text{Cl}^-$  in the original solution is higher than that in SSA due to the  $\text{HgCl}_2$  added in the droplet.

In the global Hg cycling system, the reduction from  $\text{Hg(II)}$  to  $\text{Hg(0)}$  mainly occurs in two ways. (1)  $\text{Hg(II)}$  is reduced to  $\text{Hg(I)}$  by thermal reactions, such as  $\text{BrHgO}$  reacting with  $\text{CO}$  to produce  $\text{HgBr}$  and  $\text{CO}_2$ , and  $\text{Hg(I)Br}$  reacting with  $\text{NO}_2$  to produce  $\text{Hg(0)}$  and  $\text{BrNO}_2$ .<sup>90,91</sup> (2) Reduction from  $\text{Hg(II)}$  to  $\text{Hg(0)}$  takes place largely through photolysis of  $\text{Hg(II)}$  on atmospheric surface, e.g., aerosol's surface, air/gas interface.<sup>92–96</sup> The droplets and aerosol particles in air contribute the most reduction surface for  $\text{Hg(II)}$  to be adsorbed. The photoreduction of  $\text{Hg(II)}$  on the atmospheric surface has been recognized as the important source of  $\text{Hg(0)}$  production in air.<sup>92</sup> However, most of these gaseous-phase reduction reactions are only known from computations and have not been validated experimentally. Most surface-based photoreduction reactions have only been studied empirically and lack a solid mechanistic basis. Most  $\text{Hg(II)}$  compounds have been identified in urban and indoor air as  $\text{HgBr}_2$  and  $\text{HgCl}_2$ .<sup>96</sup> It has been reported that  $\text{O}_3$  potentially reduces  $\text{HgCl}_2$  to  $\text{Hg(0)}$  under UV or solar radiation. Tong *et al.* demonstrated that  $\text{O}_3$  can significantly promote the photolysis rate for  $\text{HgCl}_2$  in the presence of UV radiation and speculated that atomic  $\text{O}$ , which results from the photodissociation of  $\text{O}_3$  via solar radiation in the atmosphere, might reduce  $\text{Hg(II)Cl}_2$  into  $\text{Hg(I)Cl}$  radicals.<sup>55</sup> Lyman *et al.* proposed that  $\text{O}_3$  may react with  $\text{HgCl}_2$  directly in air:  $\text{HgCl}_2 + 2\text{O}_3 \rightarrow \text{HgO} + 2\text{O}_2 + 2\text{ClO}$ .<sup>97</sup> Ai *et al.* conducted an experiment to optically trap a single-particle reactor, which is made by mixing carbon nanotubes and mercury halogens in an ozone environment. They hypothesized that the  $\text{Hg(II)}$  could be reduced on the surface of the carbon nanotubes by  $\text{O}(^3\text{P})$  photolyzed by  $\text{O}_3$ .<sup>98</sup>  $\text{HgCl}_2$  can be photolyzed through the UVC radiation at 185 nm. However, in this experiment, no Raman band changes related to  $\text{Hg(II)}$  are observed when the surrounding RH is high. We speculate that is because the photolysis of  $\text{Hg(II)}$  is slower than its oxidation, and most reduced products of  $\text{Hg(I)}$  have been oxidated and returned to  $\text{Hg(II)}$ .<sup>99</sup> If so, a small amount of  $\text{Hg(I)}$  in the suspended droplet would have been produced and balanced with  $\text{Hg(II)}$  via the photo-reduction and photo-oxidation reactions. On the other hand, as the  $\text{ClO}_3^-$  is formed and  $\text{Hg(I)ClO}_3$  is dissolvable in water, the  $\text{Hg(I)}$  can be in the free  $\text{Hg}^+$  state in the aqueous system. As the Raman spectrum is based on vibrational frequencies of molecular bonds, it is unable to detect  $\text{Hg(I)}$  in its ionic state, other techniques may be used to confirm the above speculation.

As the Raman spectrum in Fig. 7 shows, the Hg–Cl band located at  $314\text{ cm}^{-1}$  is observed after the droplets are stably trapped in a dry environment of RH condition below  $41\% \pm 11\%$ , and this means that instead of free  $\text{Cl}^-$  in the aqueous system, more ionic Cl bonds with  $\text{Hg(II)}$  to form  $\text{HgCl}_2$  molecules. Fig. 8a shows a contour plot of the Raman spectral intensity distribution of the SSA that contains  $\text{Hg(II)}$  as the droplet is trapped in air with the RH of 42%. Two locations were marked, where S is the position on the surface, and C is the position at the center of the trapped droplet. The corresponding Raman spectra of these two locations are shown in the insets in Fig. 8a. Compared with the Raman spectra in position S, the spectra in position C show one more band at  $267\text{ cm}^{-1}$ , which is from  $\text{HgCl}_4^{2-}$ .<sup>100</sup> Instead of directly forming  $\text{HgCl}_2$  inside the suspended aerosol particle, more  $\text{Cl}^-$  bonds to  $\text{HgCl}_2$  to form  $\text{HgCl}_4^{2-}$ . This is attributed to the photoreduction of  $\text{Hg(II)}$ . As discussed in the previous paragraph, when the droplets are suspended under the high RH condition, a small amount of  $\text{Hg(I)}$  in the suspended droplet can be produced and balanced with  $\text{Hg(II)}$  via the photo-reduction and photo-oxidation reactions. The  $\text{Hg(I)}$  is also produced in lower RH conditions as the droplet is trapped under UVC radiation. As only one ionic Cl bonds to each  $\text{Hg(I)}$ , more free  $\text{Cl}^-$  is left in the aqueous system where it bonds with  $\text{HgCl}_2$  to form  $\text{HgCl}_4^{2-}$ .

Fig. 8b shows the time-resolved Raman spectra of a SSA containing  $\text{Hg(II)}$  trapped under UVC radiation with a RH of 42%. As more free ionic Cl binds to  $\text{HgCl}_2$  to form  $\text{HgCl}_4^{2-}$ , a new band at  $267\text{ cm}^{-1}$  appears and increases in strength. After the droplet was exposed to UVC radiation for several minutes, another new band located at  $166\text{ cm}^{-1}$  appears. This band is from  $\text{Hg}_2\text{Cl}_2$ .<sup>101</sup> The formation of  $\text{Hg}_2\text{Cl}_2$  indicates that  $\text{Hg(II)}$  was reduced to  $\text{Hg(I)}$ . After the Raman band of  $\text{Hg}_2\text{Cl}_2$  appears, the trapped droplet becomes unstable and exits the trap in the next several seconds. This is because of the low dissolvability of  $\text{Hg}_2\text{Cl}_2$  in water and the efflorescence of the suspended droplet. A small amount of  $\text{Hg}_2\text{Cl}_2$  is crystallized inside of the SSA. The formation of solid particles affects the stability of the trapping.

In our exploration, the band change shown in Fig. 8b is difficult to observe. To observe the formation of  $\text{Hg}_2\text{Cl}_2$ , the RH is critical and must be in a specific range of 40–44%. This is because the droplets need to be sufficiently large to provide enough force to hold the solid  $\text{Hg}_2\text{Cl}_2$ . The interrogation time must be sufficiently long to capture the weak Raman signal. Meanwhile, the  $\text{Hg}_2\text{Cl}_2$  that forms need to accumulate to a large volume for observation. For example, if the RH is too low due to the efflorescence of the droplets, the droplet's size will be too small to hold the  $\text{Hg}_2\text{Cl}_2$  that crystallizes inside the droplets. If the RH is too high, the  $\text{Cl}^-$  will be oxidized to  $\text{ClO}_3^-$  in the aqueous system.

Nevertheless, based on the results, we hypothesized that under the UVC radiation of 185 nm and 254 nm, a small amount of  $\text{Hg(II)}$  is photolyzed to  $\text{Hg(I)}$ , and the process reaches a balance between the photo-oxidation and photo-reduction in the SSA when the surrounding RH is high. As RH decreases, the SSA undergoes dehumidification, potentially leading to the formation and subsequent crystallization of  $\text{Hg}_2\text{Cl}_2$ .





**Fig. 8** (a) A contour plot of the spectral intensity distribution of Raman spectra of a SSA containing Hg(II). Two locations are marked where S is on the surface, and C is at the center. The insets show the position-resolved Raman spectra from two marked locations. (b) Raman band variations by the time-resolved Raman spectra as the SSA containing Hg(II) exposed to UVC radiation with RH of 42%. The new emerging peak at  $166\text{ cm}^{-1}$  corresponds to  $\text{Hg}_2\text{Cl}_2$ , which indicates the photo-reduction reaction.

## 4. Conclusion

In this study, we investigated the heterogeneous chemistry and photochemistry of single suspended SSA particles and SSAs containing Hg(II) under different RH, ozone concentration, and UVC radiation using OT-RS.

In Part I, we explored the hygroscopicity, heterogeneous chemical reaction with  $\text{O}_3$ , and photochemical reaction with UVC radiation of single, suspended SSA particles by analyzing their Raman spectra. In this part, we show the RH-dependent single-particle Raman spectra of the SSA when it is suspended in air. Different spectral features show different sizes and states of the droplets. Specifically, the Raman signal is with no/weak OH-stretching band when it is trapped with RH below the ERH of  $39\% \pm 8\%$ . The solute of NaCl will neither be crystalline nor ionic, but NaCl molecules in the aqueous system. The droplet's size is smaller than  $2\ \mu\text{m}$ . The Raman spectra show a strong OH-stretching band under the RH condition of 39–69%. The solute of NaCl in SSA is in the ionic state, and the droplet's size is in the range of 4 to  $20\ \mu\text{m}$ , which are shown in the intensity of the Raman spectra. As the RH increases above  $69\% \pm 5\%$ , the WGM is observed in the Raman spectra. We also observe the heterogeneous chemical reaction between  $\text{O}_3$  and a suspended single SSA particle, and the main reaction product is  $\text{ClO}_3^-$ . This reaction is supposed to happen only on the surface of the droplets when the environmental RH is higher than the ERH of SSA which is  $39\% \pm 8\%$ . Note that this reaction should be very slow in the natural environment because of the low ozone concentration. When the SSA is exposed to UVC radiation, the ionic Cl is oxidized to  $\text{ClO}_3^-$ , and a small amount of  $\text{ClO}_3^-$  can be further oxidized to  $\text{ClO}_4^-$  on the surface of the droplets. In the dry environment, the solute in SSA becomes supersaturated, and becomes NaCl molecules in the trapped SSA particle. Under this condition, no reaction is observed as the SSA is trapped under the UVC radiation.

In Part II, we explored the heterogeneous chemical reaction of the single suspended SSA particle containing Hg(II) with  $\text{O}_3$ , and the photochemical reaction of Hg(II) in SSA with UVC radiation. In this part, we show that the Raman signal response is similar to pure SSA particle when the SSA containing Hg(II) is trapped in air with different RH. When the solute inside the

trapped droplet becomes supersaturated with the RH lower than the ERH of SSA containing Hg(II) of  $41\% \pm 11\%$ , the Raman band of  $\text{Hg}_2\text{Cl}_2$  is observed. However, the size and state changes of SSAs containing Hg(II) are the same as the pure SSA particles. This indicates that the hygroscopicity of SSAs is not affected by the Hg(II). Furthermore, no reduction reaction of Hg(II) to Hg(I) is observed when the SSA containing Hg(II) is exposed to UVC radiation in the high RH condition. This is likely due to the photolysis process of Hg(II) being slower than its oxidation reaction. A small amount of Hg(I) may be produced, but in the free  $\text{Hg}^+$  state, and it cannot be detected by Raman spectroscopy. With the RH of 40–44%, we observe the Raman band at  $166\text{ cm}^{-1}$ , which indicates the formation of  $\text{Hg}_2\text{Cl}_2$ , as the droplet trapped under UVC radiation. Based on these results, the photo-reduction of Hg(II) to Hg(I) is hypothesized to happen in the SSA under UVC of 185 nm and 254 nm radiation. Limited quantities of Hg(I) are produced and balanced with Hg(II) between photo-oxidation and photo-reduction under high RH conditions. With the decrease of RH, the SSA experiences efflorescence, and the  $\text{Hg}_2\text{Cl}_2$  is formed and crystallized inside the SSA.

## Conflicts of interest

There are no conflicts to declare.

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