Role of sea spray aerosol at the air-sea interface in transporting aromatic acids to the atmosphere

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Abstract

Aromatic acids are ubiquitous in seawater and can be transported to the atmosphere via sea spray aerosol (SSA). Despite their importance in affecting the global radiative balance, the contribution of marine aromatic acids and their transport mechanisms through SSA remain unclear. Herein, the distribution of particle size and number concentration of SSA produced in seawater containing nine different aromatic acids (i.e., benzoic acids, benzenedicarboxylic acids, hydroxybenzoic acids, vanillic acid, and syringic acid) was studied using a custom-made SSA simulation chamber; moreover, enrichment of aromatic acids in SSA and their emission flux to the atmosphere were analyzed. Transmission electron microscopy (TEM) images clearly revealed that aromatic acids can be transferred to the nascent SSA. Interestingly, the morphology associated with benzene dicarboxylic acids-coated particles showed that aromatic acids can promote the growth of other surfaces of sea salt, thus making the sea salt core spherical. Aromatic acids showed a significant enrichment behavior at the air-sea interface, which clearly indicated that SSA represent a source of aromatic acids in the atmosphere. Vanillic acid had the largest global emission flux through SSA (962 tons yr⁻¹), even though its concentration in seawater was lower. The calculated results indicated that the global annual flux of aromatic acids was not only affected by the concentration in seawater, but also by their enrichment factor (EF). These data are critical for further quantifying the contribution of organic acids to the atmosphere via SSA, which may provide an estimate of the potential influence of the atmospheric feedbacks to the ocean carbon cycle.
1. Introduction

Aromatic acids are considered to be important environmental pollutants due to their potential toxicity and persistence (Zhao et al., 2019). They are ubiquitous in seawater and the atmosphere, and have an important influence on marine and atmospheric environment. On the one hand, aromatic acids are readily to be captured by marine organisms and enriched in organisms, or be transported to remote area (Fu et al., 2011; Shariati et al., 2021; Wang and Kawamura, 2006), causing adverse health risks to the endocrine system of aquatic organisms and marine ecosystem (Saha et al., 2006). Other than the influence on marine systems, they have also been reported to have negative impact on human health, bioaccumulation, environmental persistence, and climate change (Lu et al., 2021). Studies also probe that the organic acids can change the SSA component, which in turn affects atmospheric processes in cloud condensation nuclei (CCN) or ice nuclei (IN) activities. Organic acids, as potentially important sources of atmospheric aerosols, may play a major role in aerosol climate forcing (Moore et al., 2011; Zhu et al., 2019). Therefore, due to their importance in upper ocean biogeochemistry and their environmental risk, the transport processes and accumulation potential of aromatic acids at the air-sea interface requires a comprehensive understanding.

In various observations, aromatic acids from esters released and biodegraded by algae have been reported to account for the majority of marine aromatic acids (Zangrando et al., 2019; Dekiff et al., 2014). The degradation of ester in seawater results in substantial accumulation of aromatic acids in the environmental receptors (Benson and Fred-Ahmadu, 2020; Zhang et al., 2020; Gugliandolo et al., 2020). For example, phthalic acid in seawater is produced by the consumption of phthalic acid esters during bacterial growth, as described by laboratory studies (Ren et al., 2021). Many studies have confirmed that aromatic acids cannot only be produced by biological processes, but also be considered as potential sources of oil spill, marine plastics, and sunscreen product discharges (Kristensen et al., 2021; Aitken et al., 2018). An important source of marine aromatic acid is the acidic degradation of spilled oil that occurs extensively in human-impacted areas including coastal oceans and harbors (Lu et al., 2021). Importantly, in addition to biodegradation, the presence of phthalic acid in organisms suggests that it may be derived from marine plastics ingestion (Almulhim et al., 2022). Recent laboratory studies have shown that personal-care products, especially sunscreen (e.g., α-hydroxybenzoic acid), are reduced in levels during algal blooms (Franklin et al., 2022), sparking discussions regarding to “missing aromatic acids”. However, the available data does not seem to be a definitive answer for the “missing aromatic acids”.

The survey data of aromatic acids from the atmosphere of remote marine areas provides evidence of aromatic acid transport at the air-sea interface (Fu et al., 2010), revealing a potential route for the disappearance of aromatic acids from the ocean. The particles sprayed directly from the seawater surface into the atmosphere are collectively known as sea spray aerosols (SSAs) (Hasencz et al., 2020), and represent the main component of atmospheric aerosols (Andreae and Rosenfeld, 2008; Malfatti et al., 2019). SSAs can act as carrier agents for the vertical transport of much more than just sea salt and often include organic surfactants in the ocean, thereby further disturbing ecological systems, as well investigated from field and laboratory studies (Franklin et al., 2022; Dall’osto et al., 2022). Recent data have also revealed that the surface activity and octanol-water partitioning coefficients (Kow) of organic compounds may affect their transport efficiency from the water phase to the atmosphere (Olson et al., 2020). The molecular structure may induce changes of organic acids properties (i.e., surface tension, toxicity), which further affect their transport potential and global emission flux (Lee et al., 2017; Rastelli et al., 2017; Frossard et al., 2019; Van Acker et al., 2021b). Although field studies have shown that
aromatic acids can be detected both in the ocean and atmosphere (Boreddy et al., 2017), the detailed mechanisms affecting their transport at the air-sea interface remain to be further investigated. The emission fluxes of aromatic acids during SSA production need to be identified. A recent study has also revealed that the global annual fluxes of perfluoralkyl acids (PFAAs) from SSA to the atmosphere can be estimated by laboratory-derived enrichment factors and reported median concentrations in seawater, suggesting that SSA is an important source of PFAAs in the atmosphere (Sha et al., 2021a). Consequently, it is critical to understand the factors influencing the enrichment behavior of aromatic acids in SSA at the sea-air interface.

Our previous study has revealed that the air-sea transfer of short-chain organic acids through SSA may depend on seawater surface tension, thereby further altering the enrichment behavior of organic acids (Song et al., 2022). However, extrapolation of these results to all of the organic acids may not be warranted as the mechanism for enrichment behavior may be closely related with molecular structure. Moreover, other factors were discussed in our recent review (Hu et al., 2023). In this study, we systematically identified the transport process of aromatic acids from SSA to the atmosphere in natural seawater environment. To this end, we developed a sea spray aerosol simulation chamber using the plunging jet method that provides the closest proxy to natural SSA currently available in a controlled laboratory monitoring environment, and is used to probe the transport process of aromatic acids. In our SSA simulation chamber, seawater and SSA samples were concurrently analyzed for their aromatic acid concentration, allowing better characterization of any enrichment processes involved in the bubble bursting. Finally, the global emission fluxes of aromatic acids via SSA were assessed by combining data on the concentration of aromatic acids in seawater and experimental enrichment factor data, which provides unique insights into the enrichment behavior and atmospheric transport of aromatic acids-containing SSA particles, particularly at the air-sea interface.

### 2. Experimental Section

#### 2.1 Materials

Aromatic acids were purchased from Shanghai Aladdin Bio-Chem Technology, China. Nine aromatic acids were investigated including benzoic acid, phthalic acids (ω-, m-, and p-isomers), hydroxybenzoic acids (ω-, m-, and p-isomers), vanillic acid, and syringic acid. High performance liquid chromatography (HPLC) grade methanol and acetonitrile were supplied by Fisher Scientific, USA. Seawater collected from the coast of the Yellow Sea in Qingdao, China, was transported into the laboratory SSA simulation chamber. The range of pH values measured by a pH meter (PHS-3C, Shanghai Yidian Scientific Instrument, China) in all experiments was 7.58–7.92. Ultrapure water with a resistivity of 18.2 MΩ cm was generated by a Milli-Q purification system (Merck Millipore, France), which can be used to prepare mobile phases for liquid chromatography. Each aromatic acid was first sonicated in natural seawater for 30 min at a concentration of 1 mM.

#### 2.2 Experimental setup

A jet-based laboratory SSA simulation chamber, shown in Fig.S1, was used to mimic the SSA generation. This chamber was a clamshell cuboid box (length 30 cm, width 20 cm, height 40 cm) with a viewable glass window, which has been recently adapted for air-sea transport process studies for continuously generated plunging jets to generate realistic SSA (Liu et al., 2022; Xu et al., 2023; Zhan et al., 2022). All the enrichment experiments were conducted in the SSA simulation chamber filled with approximately 9 L of the seawater. When used, plunging jets were cycled in seawater by the pump at a flow rate of 1 L min⁻¹ through a stainless-steel nozzle. Then the bubble
breaking on the surface of seawater was observed through the glass window attached to the SSA simulation chamber. To avoid any contamination from indoor air, particle free air was supplied by a compressor and zero-air generator (model 111, Thermo Scientific, USA). A mass flow controller (Beijing Sevenstar Electronics, China) was used to adjust the air flow rate entering the SSA simulation chamber, with a flow rate in the range of 3–50 L min\(^{-1}\). The Nafion drying tube (MD-700-06S-3, Perma Pure, USA) is used to control the relative humidity of the aerosol at the sampling port. The relative humidity and temperature were monitored by a 2-channel thermo-hygrometer (Testo HM42, Vaisala, Finland), in the ranges of 30–40% and 20–25 ℃, respectively. Surface tension was measured with approximately 20 mL of seawater samples. The tensiometer (JK99C, Powereach, China) was calibrated with 20 mL of ultrapure water at 25 ℃ and the sheet metal was cleaned with ethanol between each measurement. The surface tension represents collections of at least three independent measurements for each aromatic acid in order to guarantee reproducibility of the data.

2.3 Sample collection and instrumental analysis

Particle size distributions were measured at a relative humidity of ~30% using a scanning mobility particle sizer (SMPS, TSI, Model 3936). The SMPS consisted of a differential mobility analyzer (DMA, Model 3081, TSI, USA) coupled to a condensation particle counter (CPC, Model 3776, TSI, USA), which measured particles with diameters between 14.1 and 710.5 nm. The SMPS measured particles number concentration and geometric mean diameter (GMD) within the SSA simulation chamber at a time resolution of 3 min, with an inlet and sheath gas flow rate of 0.3 L min\(^{-1}\) and 3.0 L min\(^{-1}\), respectively. Before turning on the pump, we confirmed that the particle number concentration was 0–20 # cm\(^{-3}\) during the first 30 min of each experiment to ensure no leaking and no background particles. Each group of number concentration monitoring experiments lasted approximately 1 h.

SSA samples were collected on aluminum foil (25 mm, Jowin Technology, China) with a 14-stage Dekati low-pressure impactor (DLPI+, DeKati, Finland) for 5 h, and then stored at ~20 ℃ until analysis. All samples were analyzed by attenuated total internal reflectance Fourier transform infrared (ATR-FTIR) spectroscopy (Vertex 70, Bruker, Germany) using an automated fitting algorithm and techniques described in a previous study (Xu et al., 2021). In the range of 4000–600 cm\(^{-1}\), the ATR-FTIR spectra of SSA particles were recorded with an average of 64 scans. Seawater sample was collected as the ATR-FTIR spectra of blank aluminum foil to confirm that there was no infrared absorption of target functional groups on the aluminum foil.

In addition, laboratory SSA transmission electron microscopy (TEM) samples were collected using a single particle sampler (DKL-2, Genstar Electronic Technology, China). The SSA particles impacted onto copper grids films (T11023, Tianld, China) with a flow rate 1 mL min\(^{-1}\) for 1 h. TEM images were used to characterize the morphology of coated and uncoated SSA samples.

SSA particles were impacted onto 25-mm diameter quartz fiber filters (QFF, 1851–025, Waterman, UK), which were baked in muffle furnace at 450 ℃ for 3 h. Filters collected on stages 6–10 (0.19–0.94 μm) of low-pressure impactor were extracted with 4 mL ultrapure water under ultrasonication for 30 min. Thermo scientific series of high performance liquid chromatography (HPLC, Vanquish, Germany) coupled with Atlantis C18 reversed phase column (2.1 mm × 150 mm, 3 μm particle size, 100 Å, Waters) at 30 ℃ and mass spectrometer was performed to determine the concentration of aromatic acids in SSA samples and seawater. The mobile phase consisted of 0.1% formic acid aqueous solution (A) and acetonitrile, and was kept at a flow rate of 0.2 mL min\(^{-1}\). An eluent gradient program was used as follows (Witkowski and Gierczak, 2017): starting with 5% A for 5 min, increasing A from 5% to 25% in 2 min, keeping A at 25% for 4 min, increasing A from 25% to 95% in 5 min, and holding constant.
for 2 min, then turning A to 5% in 0.5 min and keeping it at 5% for 6.5 min. ESI was performed in negative ion mode with a capillary voltage of 3.5 kV. The injection volume was set at 10 μL.

Inorganic ions in seawater and SSA samples were filtered through 0.22 μm PTFE filters and analyzed by ion chromatography (Dionex ICS-6000, Thermo Fisher Scientific, USA) coupled with conductivity detection. Cations were isolated on a Dionex IonPac CS12A (4 mm × 250 mm) column, preceded by a guard column, with 20 mM methanesulfonic acid at a flow rate of 1 mL min\(^{-1}\). The sample volume was 10 μL for each injection.

### 2.4 Data analysis

SSA production was used to evaluate the particle yield at the air-sea layer. In a previous laboratory study (Christiansen et al., 2019), the production of aromatic acids in SSA was calculated as:

\[
\text{SSA Production} = N_{\text{Total}} \times Q_{\text{sweep}}
\]

where \(N_{\text{Total}}\) and \(Q_{\text{sweep}}\) are the total number concentration of particles detected from SMPS and the air flow rate through the headspace of the chamber, respectively.

The enrichment factor (EF) of aromatic acid was then calculated using Eq. (2), where \([X]_{\text{SSA}}\) and \([\text{Na}^+]_{\text{SSA}}\) are the concentrations of aromatic acids and Na\(^+\) in SSA, \([X]_{\text{SW}}\) and \([\text{Na}^+]_{\text{SW}}\) denote the concentrations in seawater (Sha et al., 2021a).

\[
EF = \frac{[X]_{\text{SSA}}}{[\text{Na}^+]_{\text{SSA}}} / \frac{[X]_{\text{SW}}}{[\text{Na}^+]_{\text{SW}}}
\]

An equation to estimate organic acids global flux from SSA emission was developed by Sha et al. (2021b). Both the laboratory-derived EF and the concentration of Na\(^+\) data were obtained to quantify the global emission flux of aromatic acid.

\[
[X]_{\text{SSA}} = k_{\text{SSA}} \times [\text{Na}^+]_{\text{SSA}}
\]

\[
[X]_{\text{SSA}} = EF \times \frac{[X]_{\text{SW}}}{[\text{Na}^+]_{\text{SW}}} \times [\text{Na}^+]_{\text{SSA}}
\]

\[
\text{flux}_X = k_{\text{SSA}} \times \text{flux}_{[\text{Na}^+]}
\]

In these equations, \(k_{\text{SSA}}\) represents the concentration of aromatic acid transferred to the atmosphere per μg of Na\(^+\). In addition, \(\text{flux}_{[\text{Na}^+]}\) means SSA annual production. Note that this quantification assumes a constant concentration of each aromatic acid reported from the literature and no atmospheric aging process of aromatic acids.

### 3. Results and discussion

#### 3.1 Surface tension of seawater containing aromatic acids

The variation of seawater surface tensions containing different aromatic acids at room temperature is given in Fig. 1. Surface tension measured in seawater ranged from 73.59 to 73.84 mN m\(^{-1}\) with a median of 73.75±0.06 mN m\(^{-1}\). The surface tension of seawater containing benzoic acid is 73.67±0.03 mN m\(^{-1}\) and certainly lower than that of seawater, indicating that benzoic acid slightly reduces the surface tension of seawater. Likewise, it is generally observed that surfactants in seawater will decrease the surface tension of seawater (Cravigan et al., 2020; Liu and Dutcher, 2021; Pierre et al., 2022; Keene et al., 2017; Enders et al., 2023).

The surface tension of both benzene dicarboxylic acids and hydroxybenzoic acids is also lower than that of...
seawater, indicating that the effect of carboxyl and hydroxyl groups on the surface tension of seawater is similar. However, differences in the positions of the functional groups of the compounds lead to some differences between the surface tension of seawater containing the two types of aromatic acids. Seawater containing p-phthalic acid (73.92±0.14 mN m\(^{-1}\)) had the highest surface tension among the isomers of benzene dicarboxylic acids. The seawater surface tension varied with the position of carboxyl group, in the order of p-phthalic acid > m-phthalic acid > o-phthalic acid. It is important to note that benzene dicarboxylic acid used here is amphiphilic, which may be the main cause of results observed above. The current findings on the surface tension of seawater containing aromatic acids are also consistent with recent studies that the surface propensity of monocarboxylic acids leads in general to a reduction of surface tension compared to seawater, while dicarboxylic acids give higher values (Ozgurel et al., 2022). The theory applies to hydroxybenzoic acids as the surface tension of seawater containing hydroxybenzoic acid was lower than that of seawater devoid of aromatic acids, with surface tension depressions ranging from 0.01 to 0.17 mN m\(^{-1}\). Unlike the order of seawater surface tension containing benzene dicarboxylic acid, the most obvious inhibition of surface tension was found with the addition of p-hydroxybenzoic acid (73.58±0.10 mN m\(^{-1}\)), while o-hydroxybenzoic acid had the weakest effect on seawater surface activity. Similar to the hydroxybenzoic acid, vanillic acid and syringic acid inhibited the surface tension of seawater to varying degrees. The results showed that the hydrophobic group (methoxy group) enhanced the surface activity of seawater, in the order of syringic acid > vanillic acid > p-hydroxybenzoic acid. As illustrated in Fig. 1D, the seawater surface activity increased with the number of methoxy groups. Our data demonstrated strong functional groups connections to the seawater surface tension containing aromatic acids.

### 3.2 Transfer of individual aromatic acids to submicron SSA particles

#### 3.2.1 SSA size distribution in the different seawater system containing individual aromatic acids

The size distribution of SSA generated from the SSA simulation chamber was measured to study the influence of molecular structure of aromatic acids on SSA production. Fig. 2A shows the size distribution of sea salt particles and benzoic acid-added particles with a peak number concentration near 186.26 nm, which could be fitted to a lognormal-mode distribution as observed in previous studies (Quinn et al., 2017; Saliba et al., 2019; Xu et al., 2022). The number size distribution showed no significant changes in response to SSA particles after adding benzoic acid, with an SSA production of 1.08×10\(^7\) part s\(^{-1}\). To explore the response of SSA formation to the number of carboxyl groups in aromatic acids, we measured the particle size distribution in the system after adding different benzene dicarboxylic acids. The number concentration of SSA particles with added benzene dicarboxylic acids was much higher than with added benzoic acids. As shown in Fig. 2B, p-phthalic acid promoted a much higher SSA particles number concentration, which could be observed visually through the SSA simulation chamber window as an increase in bubble bursting. The observed bursting phenomenon is in accordance with the conclusion that higher seawater surface tension could promote the SSA production. The most common explanation is that p-phthalic acid has two hydrophilic carboxyl groups at both ends, and the increase of particle number concentration cannot be attributed to the inhibition effect of surfactants on bubbles bursting. Obviously, the presence of benzene dicarboxylic acid increases the particle number concentration, as described in a recent study (Dubitsky et al., 2023). For the particle size distribution, the size distribution of particles with added benzene dicarboxylic acid was much narrower in contrast to the system with added benzoic acids. The morphology of SSA
particles seems to change after the addition of benzene dicarboxylic acid, which is very similar to the SSA chamber studies described by Lv et al. (2020) and Lee et al. (2020a), and further indicating that the number and position of carboxylic groups in aromatic acids could significantly affect the formation of SSA.

Further, different functional groups were investigated to determine their effect on particle generation. The distributions show a decrease in the particle number concentrations when adding hydroxybenzoic acids, compared with the system with added benzoic acids, leading to the conclusion that hydroxybenzoic acids as surfactants could inhibit SSA production (Fig. 2C). Compared with the seawater with added benzene dicarboxylic acids, the particles number concentration with added hydroxybenzoic acids decreased significantly. The hydroxybenzoic acids enhanced the seawater surface activity, which would increase the bubble lifetime and then, decrease the SSA production. Importantly, Fig. 2D shows that the particle number concentration decreased proportionally to the increase of the number of methoxy groups, providing further ground that organic matter with hydrophobic functional groups had preferential atomization ability. Previous studies also showed that particles number concentration and GMD increased with the seawater surface tension (Guzman et al., 2014; Liu et al., 2022), further indicating that the types of functional groups and the seawater surface tension both affect SSA generation.

Integrated particle size, SSA production and mass concentration based on SMPS are shown in Fig. 3. The mode diameter of SSA particle containing m-hydroxybenzoic acid was the largest among all the hydroxybenzoic acid isomers, suggesting that the m-hydroxybenzoic acid was more likely attached to sea salt particles. In all of the benzene dicarboxylic acids position isomers, p-phthalic acid-containing particles gave the largest mode diameter while SSA particles containing o-phthalic acid expressed the smallest mode diameter. Rather, the SSA particle diameters of vanillic acid and syringic acid did not change significantly, with a deviation of less than 3.75%. For SSA production, the rate of p-phthalic acid-added particles (1.54×10^7 part s^-1) is higher than those of m-phthalic acid-added (1.33×10^7 part s^-1) and o-phthalic acid-added particles (1.05×10^7 part s^-1). In addition, the SSA production of particles when adding o-hydroxybenzoic acid (1.13×10^7 part s^-1) was obviously higher than those of other isomers (m-hydroxybenzoic acid: 0.90×10^7 part s^-1, p-hydroxybenzoic acid: 0.98×10^7 part s^-1) shown in Fig. 3, because the surface tensions of seawater when adding m-hydroxybenzoic acid and p-hydroxybenzoic acid were lower than that of seawater. These differences in SSA production can likely be attributed to the selective transfer of aromatic acids at the air-sea phases. As described in previous studies, the bubble formation and breakout were controlled by seawater surface tension (Lee et al., 2020b; Liu and Dutcher, 2021). Furthermore, the mass concentration was increased with the SSA production, which is similar to previous findings (Sha et al., 2021a).

### 3.2.2 Morphologies of SSA particles

The particle size and number concentration of particulate matter obtained by SMPS were varied with adding different aromatic acids, which may be closely related to the interaction between aromatic acids and sea salt. To test this hypothesis, SSA particles from seawater experimental systems with addition of different aromatic acids were collected and the morphologies of individual particles were characterized (Fig. 4). The TEM images visually show that the cores of the sea salt particles were coated to varying degrees, forming a typical core-shell structure. Furthermore, aluminum foil samples were also characterized by ATR-FTIR for the qualitatively analysis of organic coating. The broad band in the range of 3700–3000 cm^-1 is O–H stretching in water or acids (Diniz et al., 2018). The other absorption band peak was observed at approximately 1643 cm^-1, representing the O–H stretching.
in acids (Jin et al., 2013). The image from Fig. S2A provides the evidence that SSA produced by bubble bursting can transfer acids. The absorbances in the ranges 1900–1670, 1600–1480, 1475–1300 cm\(^{-1}\) are the stretching vibration of C=O in aromatic carboxylic acid, C=C stretching in aromatic ring, C–H symmetrical carboxylate stretching (Andreeva and Burkova, 2017; Diniz et al., 2018; Geng et al., 2009; Koutstaal and Ponec, 1993).

As can be seen from Figs. 4A–B, the sea salt particles had a cubic shape, and the benzoic acid coated sea salt particles formed a core-shell structure. With this perspective, the finding of TEM images is in agreement with previously reported morphology for SSA particles (Unger et al., 2020). Figs. 4C–E clearly show the single particle morphology characteristics of benzene dicarboxylic acids. Notably, it can be seen that the core morphology of salt particles had changed significantly, where the cubic structure has changed into a ball structure. The core of SSA particles containing p-phthalic acid (Fig. 4E) on the TEM grid became more round compared with SSA particles containing m-phthalic acid (Fig. 4D) and o-phthalic acid (Fig. 4C). As previously shown by a different study, these morphologies may suggest that o-phthalic acids-containing SSA form round particles via promoting the growth of other surfaces of sea salt particles (Ballabh et al., 2006). The organic coating of sea salt is a well-known process, and has been shown specifically from phytoplankton bloom to produce more organics that likely further lead to spherical structures (Ault et al., 2013). Moreover, the organic coating was getting thicker, in the order of p-phthalic acid > m-phthalic acid > o-phthalic acid. This order in the organic coating clearly explained the result discussed above that the GMD of p-phthalic acid-containing particles was markedly larger than those of other isomers of benzene dicarboxylic acids in SSA particles. According to the spectrum in Fig. S2B, for SSA particles containing benzene dicarboxylic acids, the very weak absorption band featuring range of 1000–650 cm\(^{-1}\) represents the C–H out of plane bending in the present study and in the literature (Chang et al., 2022; Lin et al., 2014; Świslocka et al., 2013). The in-situ ATR-FTIR identified SSA particles functional groups provide the key information on the transfer of aromatic acids from seawater to the atmosphere.

Figs. 4F–H show the morphology images of SSA particles containing hydroxybenzoic acids, where the core kept the crystalline phase of sea salt with its cubic structure and the coatings mainly consisted of hydroxybenzoic acids. Notably, the organic coating of m-hydroxybenzoic acid was the thickest among all the isomers. This is consistent with the result that the GMD of m-hydroxybenzoic acid was the largest measured by SMPS. The TEM images of Figs. 4I–J show that individual nascent SSA containing vanillic acid and syringic acid also have the cubic sea salt core, despite their coating thickness is similar. Both characteristic absorption peaks of hydroxybenzoic acid, vanillic acid and syringic acid are observed in Figs. S2C–D, consistent with the results for benzene dicarboxylic acids discussed above. Besides, we expressed herein that sea salt particles surrounded by the aromatic acid coatings would be also inferred from TEM images and ATR-FTIR spectra.

### 3.3 Enrichment of individual aromatic acids to submicron SSA particles

The enrichment factors (EFs) of different aromatic acids in SSA are characterized to visually demonstrate the influence of different functional groups on the accumulation degree of aromatic acids from seawater to the atmosphere through SSA. As shown in Fig. 5, all aromatic acids exhibit varying degrees of enrichment in SSA particles. The EF ranged from 5.97 to 24.69, with the largest and smallest being of m-hydroxybenzoic acid and o-phthalic acid, respectively. The reason for the difference in EFs may be related to the difference in surface tension of seawater and the difference in octanol-water partitioning coefficients (log(K\(_{\text{ow}}\))) (Olson et al., 2020). In detail, ...
The EF of benzoic acid was approximately 10, indicating that benzoic acid can be significantly transferred to the atmosphere. In combination with the TEM images, not only is the \( p \)-phthalic acid coating the thickest, but its EF is also the largest among its three isomers. Interestingly, the EFs show the \( \log(K_{ow}) \) dependence for the three isomers of benzene dicarboxylic acids, in the order of \( p \)-phthalic acid > \( m \)-phthalic acid > \( o \)-phthalic acid. A negative correlation between EF and the \( \log(K_{ow}) \) of surfactants has been observed (Olson et al., 2020), and the detailed data are available on the website (https://comptox.epa.gov/dashboard/) and summarized in Table S2. The EFs order of benzene dicarboxylic acids are reversed from that of surfactants, probably due to the amphiphilicity of \( o \)-phthalic acid.

Notably, the correlation between the EF containing the isomers of hydroxybenzoic acid and the corresponding organic coating was also strong, among which the EF of \( m \)-hydroxybenzoic acid (EF = 24.7±8.2) was the highest. Meanwhile, the GMD of SSA particles containing \( m \)-hydroxybenzoic acid was also the highest. One possible reason could be that the \( \log(K_{ow}) \) for \( m \)-hydroxybenzoic acid (1.50) supports preferential transfer of \( m \)-hydroxybenzoic acid to SSA particles. The fundamental role of \( \log(K_{ow}) \) in the transfer of organic matter from aqueous phase to particulate phase has been proved (Olson et al., 2020; MCCord et al., 2018; De Maagd et al., 1999). However, the enrichment behavior of \( o \)-hydroxybenzoic acid is weaker than those of others. Therefore, \( \log(K_{ow}) \) is not the only factor affecting the EF, which should be combined with the octanol-air partitioning coefficients (\( \log(K_{oa}) \)). Just as \( o \)-hydroxybenzoic acid is easily transferred to the seawater surface microlayer according to \( \log(K_{ow}) \), it is not easily transferred to the atmosphere according to \( \log(K_{oa}) \). Compared with \( p \)-hydroxybenzoic acid, vanillic acid has one more hydrophobic group (methoxy) to its molecular structure. Accordingly, the EF of vanillic acid (13.3±2.5) in SSA is slightly higher than that of \( p \)-hydroxybenzoic acid (12.1±2.5). The EFs of vanillic acid and syringic acid depend on their surface activities. Syringic acid has strong surface activity, making it to be more easily transported to the atmosphere through SSA than vanillic acid. The above EF data indicates that aromatic acids have significant transport potential at the air-sea interface.

We hypothesized that the enrichment of aromatic acids is largely related to the bridging effect of \( Ca^{2+} \) as well and, as a result, the EFs of \( Ca^{2+} \) were calculated. The observed selectivity in cation transfer shown in Fig. 6 has the same trend as the cation binding affinity of aromatic acids. Moreover, another reason for the effective transfer of \( m \)-hydroxybenzoic acids to SSA particles is their binding to \( Ca^{2+} \), as evidenced by the significant enrichment of \( Ca^{2+} \) (Fig. 6). It is known that \( Ca^{2+} \) has the potential to transfer organics (Shaloski et al., 2015; Hasenecz et al., 2019). The fact is that aromatic acids can form complexes with cations which are then transferred to SSA by bubble bursting. The strong binding abilities of divalent cations to phenolic −OH and aromatic C=C have been reported (Jayaratne et al., 2016). Thus, the enrichment of major cations acts also as a standard for assessing the abundance of organic matter. In Fig. 6, the enrichment of \( Ca^{2+} \) was the most obvious, followed by those of \( Mg^{2+} \) and \( K^{+} \) among all the experiments. In SSA, \( Ca^{2+} \) always exhibited high enrichment (EF > 1), while the EFs for \( K^{+} \) and \( Mg^{2+} \) were more lousy and below 1. The above observation of the enrichment capacity of \( Ca^{2+} \) is consistent with recent studies (Salter et al., 2016; Unger et al., 2020; Lee et al., 2021; Van Acker et al., 2021). Interestingly, the greater the EF for \( Ca^{2+} \), the larger the EF for the corresponding aromatic acids. Hence, the current results further confirm that \( Ca^{2+} \) bridging is important in complexes formation and transport of organics.
3.4 Estimation of SSA contribution to global aromatic acids emissions

The $K_{SSA}$ for aromatic acids ranged from 0.0002 to 0.623 (Table S3). These values were strongly influenced by the concentration of aromatic acid in seawater, with differences of 1–4 orders of magnitude. For example, the seawater concentrations of syringic acid, $p$-hydroxybenzoic acid, benzoic acid are 0.3, 4.58, 34 ng L$^{-1}$, respectively. Such large variations highlight the importance of concentration variations in seawater of aromatic acid in assessing the SSA contribution. More than that, benzoic acid and $p$-hydroxybenzoic acid concentrations in the ocean were 205 ng L$^{-1}$ and 8.66 ng L$^{-1}$, respectively (Zhao et al., 2019). Therefore, it is necessary to estimate their atmospheric fluxes. Notably, for syringic acid, $K_{SSA}$ is equivalent to that of $o$-hydroxybenzoic acid, while the concentration of syringic acid in seawater is obviously lower. As a result, EFs of aromatic acids also contribute to SSA emission flux. Furthermore, the enrichment process is much more complex than controlled laboratory monitoring experiments, including factors such as wind speed and temperature at the seawater surface. Hence, it is necessary to further study the environmental enrichment mechanism of aromatic acids in SSA to reduce uncertainties in the estimation of aromatic acid emissions.

On the basis of $K_{SSA}$ and modeled annual SSA, the global emission fluxes of aromatic acids from seawater to the atmosphere through SSA can be estimated as eq. 4. The flux of Na$^+$ was obtained by using the annual emission of SSA calculated by Textor et al. (2006) and Jonas et al. (2021). Estimate of aromatic acids fluxes using the above Na$^+$ global emission is included in Table S4 for reference, and the comparison between aromatic acids fluxes is also shown in Fig. 7. For example, the laboratory-derived annual flux of benzoic acids was 27 tons yr$^{-1}$ for SSA emission of approximately $3.65 \times 10^{12}$ kg yr$^{-1}$, but increased to 71 tons yr$^{-1}$ when SSA emission were about 9.7 kg yr$^{-1}$. Therefore, the SSA emission used in the global annual emission of aromatic acids may be one possible reason of observed differences. Similarly, the atmospheric fluxes measured for aromatic acids are of similar order of magnitude with those for perfluorocarboxylic acids in a previous laboratory study based on field samples (Sha et al., 2021b). It follows that using lab-derived global annual fluxes gives results that are close to the modeling results. As illustrated in Fig. 7, vanillic acid has the highest global emission among all aromatic acids studied, although it has lower concentration in seawater than others. This demonstrates that the EF plays a very important role in the global emission fluxes of organic matter, in addition to concentration. Moreover, an in-depth quantification of the organic matter global emission flux transferred to the atmosphere through SSA will help estimating the CCN influence of organic contribution.

4. Conclusion

Based on our experimental data, we highlight that aromatic acids can be transferred from seawater to the atmosphere through bubble bursting. The air-sea transfer efficiency of aromatic acids was evaluated by simulating the SSA generation with a plunging jet and collecting the SSA particles. First and foremost, aromatic acids transferred from seawater to the SSA particles were confirmed as possible by combining infrared spectra and TEM images. TEM images intuitively show that aromatic acids are coated with sea salt particles to form a core-shell structure, of which further proved that there exists a good correlation between EF of organic acids and organic coating. Our data confirm that the enrichment of aromatic acid was enhanced by the increase of hydrophobic functional groups, while the corresponding number concentration of SSA particles decreased. As a whole, the transfer capacity of aromatic acids may depend on their functional groups and on the bridging effect of cation, as well as their concentration in seawater, as these factors influence the global emission flux of aromatic acids via
SSA. Overall, our research helps to close an existing knowledge gap in studying the global annual flux of aromatic acids from SSA particles to the atmosphere. Though the current results deepen the understanding of aromatic acids transport, much detailed work is still needed to explore how different aromatic acids affect the marine carbon cycle.
CRediT author statement


Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgments

This work was supported by National Natural Science Foundation of China (22076099, 22376121). We would like to thank Xiaoju Li from State Key laboratory of Microbial Technology of Shandong University for help and guidance in TEM.

Appendix A. Supplementary data

Supplementary material
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Fig. 1. Measured surface tension values of natural seawater and aromatic acid-containing seawater: benzoic acids (A), benzene dicarboxylic acids (B), hydroxybenzoic acids (C), $p$-hydroxybenzoic acid, vanillic acid, and syringic acid (D). The dark spots represent the mean values of at least three measurements and the boxes represent the ranges of 25th–50th–75th percentiles.
Fig. 2. The number concentration distribution of sea salt particles and SSA particles containing benzoic acids (A), benzene dicarboxylic acids (B), hydroxybenzoic acids (C), \( p \)-hydroxybenzoic acid, vanillic acid, and syringic acid (D). The inset depicts the number concentration of SSA particles with diameters of 50–300 nm.
Fig. 3. SSA production, particle size, mass concentration distribution of aromatic acids. The symbol size represents the geometric mean diameter of SSA particles, and the symbol color indicates the particle mass concentration.
Fig. 4. Particle morphology observed using TEM of sea salt (A) and mixed particles composed of aromatic acids-coated sea salt particles (B–K).
Fig. 5. Enrichment factors of benzene dicarboxylic acids (A), hydroxybenzoic acids (B), p-hydroxybenzoic acid, vanillic acid, and syringic acid (C) from seawater to the atmosphere.
Fig. 6. Enrichment factors of K⁺, Mg²⁺, and Ca²⁺ in submicron SSA during the experiment.
Fig. 7. Estimated annual global aromatic acids emission (tons yr\(^{-1}\)) via SSA.