Short Communication

Ship emission of nitrous acid (HONO) and its impacts on the marine atmospheric oxidation chemistry

Lei Sun\textsuperscript{a,b,1}, Tianshu Chen\textsuperscript{a,1}, Ying Jiang\textsuperscript{a}, Yang Zhou\textsuperscript{c}, Lifang Sheng\textsuperscript{c}, Jintai Lin\textsuperscript{d}, Juan Li\textsuperscript{a}, Can Dong\textsuperscript{a}, Chen Wang\textsuperscript{b}, Xinfeng Wang\textsuperscript{a}, Qingzhu Zhang\textsuperscript{a}, Wenxing Wang\textsuperscript{a}, Likun Xue\textsuperscript{a,⁎}

\textsuperscript{a} Environment Research Institute, Shandong University, Qingdao, Shandong, China
\textsuperscript{b} School of Environmental Science & Engineering, Qilu University of Technology (Shandong Academy of Sciences), Ji'nan, Shandong, China
\textsuperscript{c} Key Laboratory of Physical Oceanography, College of Oceanic and Atmospheric Sciences, Ocean University of China, Qingdao, Shandong, China
\textsuperscript{d} Department of Atmospheric and Oceanic Sciences, School of Physics, Peking University, Beijing, China

HIGHLIGHTS

- The ship emission ratio of $\Delta$HONO/$\Delta$NO\textsubscript{x} was measured for the first time during a cruise.
- A global HONO ship emission inventory was derived for use in chemical transport models.
- Ship emissions of HONO play an important role in the marine atmospheric oxidation processes.

GRAPHICAL ABSTRACT

ABSTRACT

Nitrous acid (HONO) is an important reservoir of the hydroxyl radical (OH) and thus plays a central role in tropospheric chemistry. Exhaust from engines has long been known as a major primary source of HONO, yet most previous studies focused on vehicle emissions on land. In comparison, ship emissions of HONO have been rarely characterized, and their impacts on the tropospheric oxidation chemistry have not been quantified. In this study, we conducted cruise measurements of HONO and related species over the East China Sea. Contrasting air masses from pristine marine background air to highly polluted ship plumes were encountered. The emission ratio of $\Delta$HONO/$\Delta$NO\textsubscript{x} (0.51 ± 0.18\%) was derived from a large number of fresh ship plumes. Using the in-situ measured emission ratio, a global ship emission inventory of HONO was developed based on the international shipping emissions of NO\textsubscript{x} in the Community Emission Data System inventory. The global shipping voyage emits approximately 63.9 ± 22.2 Gg yr\textsuperscript{−1} of HONO to the atmosphere. GEOS-Chem modelling with the addition of ship-emitted HONO showed that HONO concentrations could increase up to 40–100\% over the navigation areas, leading to about 5–15\% increases of primary OH production in the early-morning time. This study elucidates the potentially considerable effects of ship HONO emissions on the marine atmospheric chemistry, and calls for further studies to better characterize the ship emissions of HONO and other reactive species, which should be taken into account by global and regional models.

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https://doi.org/10.1016/j.scitotenv.2020.139355
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1. Introduction

Acting as a reservoir of the hydroxyl radical (OH), nitrous acid (HONO) plays a crucial role in the atmospheric photochemistry. Photolysis of HONO by sunlight produces an OH radical and a NO molecule, which initiate the atmospheric oxidation processes leading to formations of ozone (O₃) and secondary aerosols (Finlayson-Pitts and Pitts, 1999; Kleffmann, 2007). In the ambient atmosphere, HONO is either emitted from various primary sources such as combustion and microbial processes (Kurtenbach et al., 2001; Su et al., 2011), or formed by gas-phase and heterogeneous reactions involving nitrogen oxides (NOₓ = NO + NO₂) (Finlayson-Pitts et al., 2003; George et al., 2005; Pagsberg et al., 1997). Recent studies have indicated that the sources of ambient HONO are rather complicated and have not yet been fully understood (Kleffmann et al., 2005; Stemmler et al., 2006; Zhou et al., 2011). Exploring the sources and environmental consequences of HONO is a fundamental step towards a better understanding of atmospheric chemistry, air pollution and climate change.

Exhaust of engine combustion is an important primary source of HONO in the atmosphere. In particular, vehicular exhaust has long been recognized as a major source of ambient HONO in urban areas (Pitts et al., 1984). In the recent decade, a number of field studies have measured the emission ratios of ΔHONO/ΔNOₓ from vehicles at tailpipe, tunnel and roadside in many metropolitan areas over the world (Kurtenbach et al., 2001; Li et al., 2018; Liu et al., 2017; Rappenglück et al., 2013; Xu et al., 2015; Yun et al., 2017). These studies have confirmed the significant contributions of vehicle emissions to the ambient HONO levels and hence atmospheric oxidizing capacity (Li et al., 2018; Xu et al., 2015). In comparison, little effort has been dedicated to characterize the emission factors of HONO from other transportation tools such as shipping vessels. Sea transport contributes to over 70% of the global trade by value and around 80% by volume, and at the beginning of 2015, the world’s commercial fleet consisted of nearly 90,000 vessels according to the International Maritime Organization (http://www.imo.org/en/Pages/Default.aspx). Ships emit a large amount of air pollutants into the marine atmosphere, among which HONO is a highly reactive component. Ship emissions of HONO have high potential to perturb the atmospheric chemistry processes by supplying not only OH radicals but also NOₓ, thus affect the marine atmosphere and climate change. To the best of our knowledge, the impacts of ship emissions of HONO on the marine atmosphere and climate have not been well quantified to date.

In the present study, we conducted cruise measurements of HONO and related species on the East China Sea. A large number of fresh ship plumes were encountered due to the vessel operation conditions, which provides a unique opportunity to characterize the emission factors of HONO from ship exhausts. In the following sections, we first show the overview of the measurement data and derive the ΔHONO/ΔNOₓ emission ratio from the ship plumes. We then develop a global ship emission inventory for HONO based on the measured emission ratio and the available NOₓ ship emission database. Finally, a global chemical transport model (GEOS-Chem) is utilized to evaluate the impacts of ship emissions on HONO and OH radicals in the marine atmosphere. Overall, this study attempts to fill the knowledge gap of the ship emission of HONO and its effects on the marine atmosphere, and demonstrates the demand to further characterize the ship emissions of highly reactive air pollutants.

2. Methods

2.1. Field measurements

The cruise measurement campaign was conducted in the East China Sea (117°–131°E, 23°–33°N) from April 22 to May 27, 2017. The Scientific Research Vessel “Dong Fang Hong 2” was deployed as the sampling platform. It is equipped with a medium speed diesel engine (MSDE; MAK 8M332C: 1380 KW and 900 RPM; gross tonnage = 3235) and fueled by relatively clean marine diesel oil (i.e., 0# diesel fuel), and has been in service for over twenty years for scientific research surveys in the China Sea (Zhang et al., 2017). According to the International Maritime Organization, the MSDE vessels account for about 21.1% of the total vessel fleet, similar to that of small-speed diesel engine vessels (20.1%) but slightly lower than that of high-speed diesel engine vessels (35.4%). The cabin housing the measurement instruments was located in the front of the funnel to eliminate potential interferences from ship plumes. We started the campaign when the vessel was far from the shore to avoid the impact of vehicle emissions and other sources at the port. In the present study, however, numerous stops were designed along the cruise route for collections of the air and sea water samples. During these stops, the air parcels sampled by the instruments in the cabin were frequently influenced by the freshly emitted ship plumes, which can be further confirmed by checking the concentration spikes of primary pollutants such as CO and NOₓ. Therefore, the air masses sampled in the present study included not only marine air but also a lot of polluted ship plumes. A schematic diagram for the sampling of marine air and ship plume is shown in Fig. S1.

A package of commercial equipment was utilized for the field measurements of HONO and related species. Specifically, HONO was measured by a long path absorption photometer (LOPAP; QUMA GmbH) at a time resolution of 1 min (Kleffmann et al., 2002). Automatic zero checks using ultrapure nitrogen (purity of 99.999%) were performed regularly twice per day, and manual calibrations by nitrite standard solution were conducted every three days (Li et al., 2018). NOₓ was measured by a Cavity Attenuated Phase Shift (CAPS) spectroscopy instrument (Teledyne API, Model T500U), which is designed to be highly selective for NO₂ and free from possible interferences from other NOₓ species. NO and NO₂ were measured by a chemiluminescence analyzer (Teledyne API Model T200U), with an externally placed molybdenum oxide converter for the conversion of NO to NO₂. O₃ was measured by a UV photometric analyzer (TEI, Model 49C). CO was measured using a non-dispersive infrared analyzer (Teledyne API Model 300EU) with automatic internal zero checks every 4 h. Particle number concentrations and size distribution in the range of 5–10,000 nm were measured by a Wide-range Particle Spectrometer (WPS; MSP Model 1000XP). JNOₓ was measured by a filter radiometer (Meteorologie Consult GmbH). Meteorological parameters including temperature, relative humidity (RH), and winds were monitored by a commercial weather station. The measurement principles, accuracies, and quality assurance and control protocols for all of these instruments can be referred to Xue et al. (2016) and Li et al. (2018).

2.2. GEOS-Chem model

The GEOS-Chem global chemical transport model (Version 9–02) was used to simulate the impacts of ship HONO emissions on the global distributions of HONO and OH. The model was run with the standard NOₓ–NOᵧ–VOC–O₃-aerosol chemistry (Mao et al., 2013) at a horizontal resolution of 2° × 2.5° with 47 vertical layers (including ten layers of ~0.13 km thickness below 850 hPa), as driven by the GEOS-FP assimilated meteorological fields. The global anthropogenic emissions were based on the Emission Database for Global Atmospheric Research (EDGAR) v4.2 inventory for CO and NOₓ and Community Emission Data System (CEDS) for non-methane volatile organic compounds (NMVOCs) (Hoyle et al., 2018), which were further replaced by the regional inventories over Asia (INTEX-B), Europe (EMEP), Canada (CAC; http://www.ec.gc.ca/pdb/cac/cac_home_e.cfm) and China (MEIC; http://www.meicmodel.org/) (Auvray and Bey, 2005; Zhang et al.,...
The international shipping emissions were adopted from the CEDS inventory, which has the updated fuel consumption that matches the bottom-up estimates from the International Maritime Organization (2014), including monthly emissions of CO, NOx, and SO2 (Hoesly et al., 2018). Monthly biomass burning emissions were from the GFED3 inventory (van der Werf et al., 2010). Other natural emissions were parameterized based on the modelled meteorology. See Yan et al. (2016) for the detailed configurations of the model. The model was run with and without ship emissions of HONO, and differences between the two simulations are regarded as the effects of the ship-emitted HONO.

Since the CEDS inventory has been updated to 2014 to reflect the actual emissions in recent years, we performed a half-year spin-up simulation and used the simulation results for July 2014 (summer scenario) for further analyses. Though inconsistent with the observation time period, the results of different sensitivity tests can represent potential impacts of ship HONO emissions on the marine atmospheric chemistry.

3. Results and discussion

3.1. Ship emission ratios of $\Delta$HONO/$\Delta$NOx

The time series of measured air pollutants and weather conditions during the vessel cruise are shown in Fig. 1. Clearly, contrasting air masses from very clean marine background air to highly polluted air parcels were alternately encountered, as indicated by the significant variation in the concentration levels of primary pollutants such as NOx. The fresh ship plumes and pristine marine background air were carefully selected for further analyses. The ship plumes were mainly sampled during the ship stops with obvious concentration spikes for primary pollutants (such as NOx and HONO) but lower O3 levels. Considering the absence of other pollution sources in the sea, such high levels of primary pollutants are believed to be due to the emissions of the research vessel. The detailed selection criteria of ship plumes include (a) only the data when the vessel stopped and the plume moved through the sampling inlet were considered (Fig. S1); and (b) concentration spikes of HONO and NOx as well as reduction in O3 concentrations were observed. A total of 2751 min measurements were identified as the ship plume data, indicating the frequent influence of ship plumes. The averaged mixing ratios ($\pm$SD) of NO, NO2 and HONO for the ship plumes were 73.6 ± 171, 27.6 ± 22.4 and 0.39 ± 0.72 ppbv, with maximum values of 1767, 254 and 7.54 ppbv, respectively (see Table 1). The large range of SD values is mainly caused by the different engine loads during the ship voyage and the stopping time. The marine background air was sampled in the open sea far away from the coast, with extremely low levels of primary pollutants. The mean marine background concentrations ($\pm$SD) of NO, NO2 and HONO were 20 ± 10, 110 ± 40 and 2 ± 2 pptv, respectively. These values are representative of the global background conditions, and will be subtracted from the ship plume data to derive the $\Delta$HONO/ $\Delta$NOx emission ratios.

We then deduced the emission ratio (ER) of $\Delta$HONO/$\Delta$NOx for the above identified fresh ship plumes. Based on the huge number of instantaneous ship plume data, the average ($\pm$SD) ER was 0.46 ± 0.31% with a median value of 0.41% (see Table 2). Furthermore, 90% of the individual ER values (i.e., 5–95% percentiles) are concentrated in the range of 0.08–0.97%. The scatter of ER values case by case should be due to the differences in the engine operation conditions and/or the ambient atmospheric conditions (e.g., age of plumes), which is also the case for the vehicle emission ratios on land (Xu et al., 2015). We also calculated the ER of $\Delta$HONO/ $\Delta$NOx for the more strictly defined ship plumes, which should last for at least 15 min with concurrent high concentrations of HONO and NOx. With this criterion, a total of 37 typical plumes were screened out, covering 1202 min measurement data. These plumes further removed the possible interference to the measurement data in short durations and ensured the high HONO and NOx levels were indeed affected by the ship plumes. The averaged ($\pm$SD) ER for

Fig. 1. Time series of measured species and meteorological parameters during the East China Sea voyage from April 22 to May 27, 2017. The black dots represent the ship plume data selected for analyses. Note that only data collected when the vessel stopped and the plume passing through the sampling inlet were selected as ship plumes in this study.
these typical plumes was calculated as $0.51 \pm 0.18\%$ with a median value of 0.43%, which are quite comparable to those derived from the instantaneous data case. In the following sections, the ER of 0.51% was adopted to derive the ship emission inventory of HONO, which can be considered as an overall average estimation of ship HONO emission.

An interesting finding is the discrepancy between the ΔHONO/ΔNOx ratios derived from the daytime and nighttime ship plumes. The ER derived from the daytime ship plume data (7:00–19:00 local time, LT; including 1447 min data) was 0.53% ± 0.37%, which is significantly higher than that (0.37% ± 0.21%) obtained from the nighttime data (19:00–7:00 LT; containing 1304 min data) (p < 0.01). Fig. S2 (a) and S2(b) show that the ΔHONO/ΔNOx ratios generally increase with the measured JNO2 levels, (except that JNO2 value is in the range of $3-4 \times 10^{-3} \text{s}^{-1}$), indicating the significant influence of the solar radiation. Therefore, the higher daytime ΔHONO/ΔNOx ratios should be attributed to the rapid photo-enhanced heterogeneous conversion of NOx to HONO on the surface of freshly-emitted particles such as soot. Laboratory studies have proposed the photo-enhanced uptake of NOx on aerosol surfaces as a potential daytime source of HONO (George et al., 2005; Monge et al., 2010), and our results provide the direct evidence of this process in the ambient atmosphere.

In general, the ocean-going vessels, such as container ships, cargo carriers and tankers, usually burn heavy fuel oil (HFO) in the open waters, which release more SO2 and metals. Whereas in harbors or specific emission regulation areas, the ferries, bulk carriers, and other smaller ships mostly burn marine diesel oil (MDO) instead. Diesch et al. (2013) showed that the ship gross tonnage was directly proportional to the engine power. The emission ratios of NOx of larger vessels (gross tonnage >30,000 tons) were 1.5 times higher than that of the smaller vessels (gross tonnage <5000 tons). In addition, Celik et al. (2020) pointed out that more efficient fuel combustion resulted in more NOx emissions owing to the higher combustion temperature. Another important impact factor is fuel quality. According to Zhao et al. (2020), the emission of NOx could be reduced by approximately 20% when the research vessel burned MDO instead of HFO under different engine loads. The research vessel in this study is a smaller vessel (gross tonnage = 3235 tons) and uses MDO as the fuel. Thus, we believe that the emission of NOx under this circumstance is relatively low.

The ship emission related experimental studies can be performed in stationary platforms (at harbours or ports and mobile platforms in research vessels/aircrafts measurements), and in the exhaust funnels. Merico et al. (2016) reported the observed NO peak concentration of ~102 ppbv at the harbour area (50 m from the water), which was comparable to the NO levels in this study. In contrast, Zhao et al. (2020) measured the emission gases directly from the chimney and the mean NO concentrations (MGO: 1438 ± 289 ppm; HFO: 1448 ± 272 ppm) were much higher than that found in this study. Table S1 compares our measured ship emission ratios of ΔHONO/ΔNOx against the results obtained for on-road vehicle emissions from many metropolitan areas in the world. The different values of ER implied the effects of other factors such as different types of engines, fuels, and working conditions on the HONO emissions in the real atmosphere (Huang et al., 2018; Kurtenbach et al., 2001). In general, the emission ratios derived in the present study are well within the reported range of ER for vehicle emissions (e.g., 0.10–1.70%), suggesting the similarity of HONO emission factors from engine exhaust. Thus, results in this study are representative of the emission ratios from ship engine exhausts. To improve the representativeness of this study, we divided the ER into three categories: 0.51% for this study; 1.7% for the maximum ratio and 0.03% for the minimum ratio.

### 3.2. A global ship HONO emission inventory

A global ship emission inventory of HONO was then developed based on the average in-situ measured ER of ΔHONO/ΔNOx (0.51%) and the CEDS ship emission inventory for NOx (Hoesly et al., 2018). Fig. 2 shows the spatial distribution of ship HONO emissions. The ship-emitted HONO primarily concentrated over the navigation areas (i.e., the shipping routes and international ports), with much stronger emission intensity in the Northern Hemisphere. The estimated mean (±SD) annual emission of HONO from global shipping was 63.9 ± 22.2 Gg yr$^{-1}$, which was much smaller than those of NOx, SO2 and CO. The Northwest Pacific (i.e., west of 180°E and north of 15°N), Northeast Pacific (i.e., east of 180°E and north of 15°N), Northern Atlantic (i.e., north of 15°N), Indian Ocean & Red Sea, and the Mediterranean & offshore areas of continental Europe are the major emission zones, with annual emissions of 8.1 ± 2.8, 14.1 ± 4.9, 24.7 ± 6.8, 8.7 ± 3.0, and 49.0 ± 17.2 Gg yr$^{-1}$, respectively. Monthly emissions were also estimated and results are summarized in Table S2. Overall, the ship related HONO emissions in the second half of the year (July–December; 5.4–5.7 Gg mon$^{-1}$ with a total of 33.5 ± 11.7 Gg) are slightly higher than those in the first half (January–June; 4.4–5.5 Gg mon$^{-1}$ with a total of 30.4 ± 10.6 Gg). This pattern was the same as NOx emissions, which reflects seasonal variations of in shipping business or other ship-related activities.

The HONO emission inventory derived this study is subject to large uncertainties. Potential uncertainties mainly arise from uncertainties of the HONO/NOx emission ratio and the CEDS ship emission inventory
for NOx. The emission ratio was derived from only one vessel despite a huge amount of ship plume data. Therefore, we estimated ship HONO emissions based on previous reported values of ER to improve its representativeness. The ship HONO emissions estimated from the maximum ratio (1.7%) and the minimum ratio (0.03%) of HONO/NOx were listed in Table S2. In addition, the uncertainty of ship NOx emissions may result from the uncertainties of the CEDS inventory as well. Hoesly et al. (2018) have shown that IEA-reported consumption is incomplete and additional fuel consumption is needed to match the bottom-up estimates from the International Maritime Organization in CEDS. However, more research is needed to improve the reliability of CEDS ship emission inventory. Furthermore, nearly 70% of the ship emissions occurred within 400 km of coastlines along the main trade routes (Corbett et al., 1999). Therefore, the global inventory may underestimate the emissions of short-distance ships near the continents. Despite these deficiencies, this first piece of ship emission inventory of HONO may still be considered by modelling studies before a more refined inventory is in place.

3.3. Impacts on the atmospheric oxidation processes

Influences of ship HONO emissions on the ambient HONO and OH radicals in the marine atmosphere were determined by the GEOS-Chem model. Model simulated meteorology (2-m temperature, relative humidity, and winds) was evaluated against the high qualities GEOS-FP output which uses the most recent validated GEOS system. Fig. S3 shows that the spatial distributions of different meteorological variables are comparable with the GEOS-FP output data. In addition, O3, NOx and HONO were evaluated against observational data. Fig. S4, Table S3 and S4 show that the model performed relatively well in reproducing the observed levels of O3 and HONO. Fig. S5 shows that the model cannot capture observed high NOx, probably due to the low model resolution and the positive biases of the measurement (Sun et al., 2019; Xu et al., 2015). Overall, the model performed reasonably well in reproducing the HONO measurement. Fig. S6 shows the global distributions of HONO, OH, and primary OH sources (POH: the OH production rate from photolysis of HONO, O3, H2O2, HNO3 and OVOCs, as well as ozonolysis reactions of unsaturated VOCs) from the Base Scenario without ship HONO emissions. The model predicted higher HONO levels in the polluted continental areas (e.g., East Asia and western Europe), with maximum values of 1–2 ppbv for Eastern China. Such levels are comparable to the in-situ determined concentrations in these continental regions (e.g., Li et al., 2018 and references therein). In comparison, the much lower HONO mixing ratios (i.e., 2–50 pptv with an average of 9.8 ± 8.8 pptv) simulated in the East China Sea and near ocean areas are actually higher than the measured values (2 ± 2 pptv for the marine background air) in this study. The overestimation of HONO is mostly due to the low model resolution, i.e., one grid may cover both coastal and offshore areas. For OH and POH, the model calculated higher levels in the polluted continents and their adjacent coastal regions, and moderate levels (1–4 × 10^6 molecules cm^-3 and 0.5–2 × 10^6 molecules cm^-3 s^-1) in the vast oceans.

Figs. 3 and 4 presents the detailed impacts of ship emissions on the ambient 24-h average HONO and OH in the global marine atmospheres. With the inclusion of ship HONO emissions, the model predicts significant enhancements of HONO concentrations in the navigation areas. The enhancements in the absolute concentration of HONO reach up to 10 pptv in the North Sea, 3 pptv in the east of North America and are about 0.5–1.5 pptv in the Northwestern Pacific, Northeast Pacific, North Atlantic, Red Sea and some famous seaports. Generally, the enhancements of HONO in percentage are 40–100% along the major shipping routes. This indicates ship emission presents a major source of HONO in the global marine atmosphere. HONO concentration enhancements mostly occurred in the nighttime, reaching up to 80–100% in the major shipping routes (Fig. S7). In comparison, the enhancement was much lower during the daytime due to the fast photolysis.

As HONO photolysis is a major source of OH, ship emissions of HONO could serve as an additional OH supplier and hence enhance the atmospheric oxidation processes. Our modelling results showed an enhancement of 20–40% in the OH production rate from HONO photolysis over the majority of shipping routes after including shipping emissions (Fig. S8). As shown in Fig. 4a and b, ship emissions of HONO also resulted in significant increases (i.e., 1–4 × 10^4 molecules cm^-3 s^-1 or 5–15% in percentage) of the total primary source of OH (Ptot) in the navigation areas during the early morning period (6:00–7:00 LT). This suggests that ship-emitted HONO may play an important role in the jump
start of the atmospheric photochemistry in the marine atmosphere. In comparison, the impact of ship-emitted HONO on OH production is minor during the daytime (Fig. S9), indicating that OH productions are dominated by other pathways (i.e., \( \text{HO}_2 + \text{NO} \) and so on), and the accumulation of HONO during the nighttime has been photolyzed at dawn. We also examined the impacts of ship HONO emissions on the tropospheric \( \text{O}_3 \) in the marine atmosphere, and found that they are almost negligible (<1%; figures not shown). This is because \( \text{O}_3 \) formation is \( \text{NO}_x \)-limited in the pristine marine atmosphere, and adding OH radicals is insensitive to \( \text{O}_3 \) formation and might even lead to \( \text{O}_3 \) destruction via the \( \text{O}_3 + \text{H} \) reaction cycles. In order to improve the representativeness of the simulation results, two sensitivity tests were performed by setting the ER to 1.7% and 0.03%, representing the maximum and minimum ratios in ship plume, respectively (according to the literature results in Table S2). The two test results showed significant differences. HONO concentration reached up to 40 pptv in the North Sea and 100–200% over the major shipping routes when we set ER to 1.7% (Fig. S10(a) and S10(b)). Similarly, the ship HONO emissions promoted the OH production, which increased \( 3 \times 10^4 \) molecules cm\(^{-3}\) s\(^{-1}\) or 20–40% for the POH (Fig. S11). However, when ER was set to 0.03%, HONO concentrations and POH did not show much difference, suggesting the impacts can be ignored.

The aforementioned analyses elucidate the significant contributions of ship emissions to the ambient HONO and OH sources in the marine air. Once OH is produced, it immediately involves in the atmospheric oxidation cycles and oxidizes a variety of reduced substances such as VOCs and DMS. One should keep in mind that the present analysis may be subject to some uncertainty due to limitations of the methodology data representativeness. Nevertheless, we believe that the current estimation should represent a lower limit for the importance of ship HONO emissions given the following reasons. First, as mentioned above, the CEDS emission may not cover all the shipping vessels, especially for those with short travel voyages (Hoesly et al., 2018; Liu et al., 2016). Second, the marine diesel oil used by the research vessel in this study is a typical clean fuel and emits relatively low levels of air pollutants compared to common cargo ships. Third, our analysis suggests the additional formation of HONO from the heterogeneous reactions of \( \text{NO}_2 \) on the surface of particles, both of which are major constituents of ship emissions, and this process was not considered in the GEOS-Chem simulations. Obviously, more efforts are urgently needed to test more ships, fuels, operation conditions, etc., and to compile more accurate ship emission inventories for a better understanding of the influences of global shipping on the marine atmosphere and climate.
4. Summary

We present the direct measurements of emission ratio of ΔHONO/ΔNOx from a large number of fresh ship plumes. A global ship emission inventory of HONO was compiled for the first time based on the measured emission ratio and the available emission data for NOx, and is ready for use in chemical transport models. GEOS-Chem modelling analysis illustrates the potentially important impacts of ship emissions on ambient HONO, primary OH sources, and thus oxidation capacity in the marine atmosphere. Despite the inherent uncertainty from the field measurements and ship emission database, this study is a good initiative to revisit the roles of ship emissions in the marine atmospheric chemistry. Further studies are urgently needed to better characterize the ship emissions of HONO and other reactive species and to thoroughly evaluate their impacts on atmospheric chemistry and climate change.

CRediT authorship contribution 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.

Acknowledgements

This work was funded by the National Natural Science Foundation of China (Project No.: 91544213, 41505111 and 11922051), Shandong Provincial Science Foundation for Distinguished Young Scholars (ZR2019JQ09), the Qilu Youth Talent Program of Shandong University, the Jiangsu Collaborative Innovation Center for Climate Change, the Taishan Scholars (ts201712003) and the Young Scientist Cooperation Fund of Qilu University of Technology (Shandong Academy of Sciences) (Grant No. 2017BSHZ2020). We appreciate Penggang Zheng and Rui Li for their help in setting up the instruments. The model simulations were performed at the Supercomputing Center of Shandong University in Weihai.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.scitotenv.2020.139355.

References


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