

Rapid formation of hydroxymethyl hydroperoxide and its vital role in methanesulfonic acid-methylamine nucleation: impacts of urban industrial areas

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Abstract

Organic peroxides are widely recognized as important contributors to secondary organic aerosols formation. Among these, hydroxymethyl hydroperoxide (HMHP) is a common species found in both the gas phase and fine aerosols. Despite its abundance, the molecular-level formation of HMHP through methanesulfonic acid (MSA)-catalyzed hydrolysis of CH₂OO, particularly in the gas phase and at the air-water interface, remains insufficiently examined. Moreover, the role of HMHP in new particle formation (NPF) has not been fully elucidated. Herein, we employ quantum chemical calculations together with Born-Oppenheimer molecular dynamics simulations to investigate HMHP formation from CH₂OO hydrolysis with MSA under both gas phase and interfacial conditions. Our results show that HMHP forms rapidly and stably in both environments. Further analysis using the atmospheric cluster dynamics code reveals that HMHP not only enhances the clustering stability of MSA-methylamine (MA) clusters, but also exerts a direct role in promoting MSA-MA nucleation. Importantly, in regions with elevated HMHP concentrations (3.00×10^{10} - 1.25×10^{11} molecules·cm⁻³), such as Niwot Ridge and Southeastern United States, the HMHP-involved pathways contribute unexpectedly up to 42% and 59% of total nucleation flux at 258.15 K, respectively. These findings provide new insights into HMHP formation pathways and the efficient MSA-MA-HMHP nucleation mechanism, offering a plausible explanation for the frequent and intense NPF events observed in continental regions.

Keywords: hydroxymethyl hydroperoxide; gas phase; interfacial; new particle formation

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1. Introduction

Organic peroxides serve as important reactive intermediates in atmospheric oxidation processes, facilitating free radical chain termination, oxidant cycling, and the formation of aerosols (Wang et al., 2023; Tröstl et al., 2016). Among these, hydroxymethyl hydroperoxide (HMHP), a product of the hydrolysis of the simplest and highest concentrations criegee Intermediates (CH_2OO), has been detected at notable concentrations in the atmosphere (Lee et al., 1993; Nguyen et al., 2015). Atmospheric concentrations of HMHP typically range around 10^9 molecules·cm⁻³ in locations such as Central Portugal (Jackson and Hewitt, 1996), Pabstthum (Grossmann et al., 2003) and Beijing (Yi, 2000). However, in regions like Guangzhou (Zhang et al., 2001) and Niwot Ridge (Hewitt and Kok, 1991), these levels can increase to approximately 10^{10} molecules·cm⁻³. Notably, in the Southeastern United States (Allen et al., 2018), concentrations can even reach up to 10^{11} molecules·cm⁻³. Due to its high abundance, HMHP undergoes subsequent reactions with other atmospheric species, leading to the formation of hydroperoxides, organic peroxides, and secondary organic aerosols, which play a crucial role in influencing air quality, climate forcing, and human health (Sakamoto et al., 2013; Rissanen et al., 2014). So, understanding the source of HMHP and its impact on the atmospheric environment has become a major topic of significant interest in atmospheric chemistry.

HMHP is formed predominantly through the hydrolysis of CH_2OO , a process that has been extensively investigated through both experimental studies (Chao et al., 2015; Stone et al., 2014; Berndt et al., 2015) and theoretical simulations (Wu et al., 2023a; Lin et al., 2016; Wu et al., 2023b). Experimental data (Chao et al., 2015; Ouyang et al., 2013) estimate the rate constant for the direct CH_2OO hydrolysis to form HMHP to be approximately 4×10^{-15} cm³ molecule⁻¹ s⁻¹. In contrast, hydrolysis involving the water dimer ($(\text{H}_2\text{O})_2$) has rate constants between 4×10^{-12} and 8.9×10^{-12} cm³ molecule⁻¹ s⁻¹ at 294-298 K (Lin et al., 2016; Berndt et al., 2015), approximately three orders of magnitude higher than that of the reaction with monomeric water, demonstrating that the presence of the water dimer significantly enhances the hydrolysis rate (Long et al., 2016; Anglada and Solé, 2016; Newland et al., 2015). Additionally, quantum chemical calculations have shown that atmospheric organic amines, acids, and alcohols can catalyze CH_2OO hydrolysis by substantially lowering activation barriers and accelerating reaction rates (Zhang et al., 2018; Chao et al., 2019). However, the role of methanesulfonic acid (MSA) in facilitating CH_2OO hydrolysis remains underexplored, even through the calculations reported by several groups (Li et al., 2024c; Wang et al., 2025) have shown that MSA is a notably effective catalyst in both CH_3CHOO and HNSO_2 hydrolysis. In fact, MSA is a major oxidation product of organosulfur compounds originating from a variety of sources, including biological processes, biomass combustion, industrial emissions, and agricultural activities. As a result, it has been

57 widely detected in atmospheric aerosol particles across diverse geographic regions, spanning from coastal to inland
58 areas (Barnes et al., 2006; Gaston et al., 2010). Notably, in coastal regions characterized by elevated MSA levels,
59 MSA concentrations range from approximately 10% to 250% of gaseous sulfuric acid concentration (Ning et al.,
60 2022; Ning and Zhang, 2022), whereas in continental regions with lower MSA levels, including many inland urban
61 and industrial regions, MSA concentrations are typically on the order of ~10%-50% of gaseous sulfuric acid
62 concentrations (Berresheim et al., 2002; Chen et al., 2020). Furthermore, MSA is characterized by strong acidity
63 and the ability to act as a proton-transfer bridge, suggesting a potentially important role in modulating the hydrolysis
64 of CH₂OO. So, the lack of understanding of the MSA-catalyzed CH₂OO hydrolysis reaction mechanism limits the
65 accurate understanding and evaluation of HMHP formation, especially in regions with high MSA pollution
66 concentrations.

67 Interfacial reactions at the air-water interface complement gas-phase reactions and can significantly influence
68 reaction rates (Xu et al., 2024; Fang et al., 2024). These interfacial reactions may involve distinct mechanisms, with
69 interfacial water molecules not only serving as reactive sites but also participating directly as reactants, proton
70 transfer mediators, and both donors and acceptors (Tang et al., 2024). For example, MSA-assisted HNSO₂
71 hydrolysis at the air-water interface follows two distinct reaction mechanisms, the ion-forming mechanism and the
72 proton exchange mechanism, resulting in the formation of the SFA⁻···H₃O⁺ ion pair. This contrasts with the gas-
73 phase production of SFA formed from the MSA-catalyzed hydrolysis of HNSO₂ (Wang et al., 2025). In addition,
74 the reaction between glyoxal and dimethylamine proceeds 5.70×10^4 times more rapidly at aqueous interfaces
75 compared to the gas phase, where it is less likely to take place (Dong et al., 2024). Therefore, based on gas-phase
76 reactions, it is essential to investigate the MSA-catalyzed CH₂OO hydrolysis reaction mechanisms at the air-water
77 interface, which will enrich the understanding of HMHP sources in regions affected by MSA pollution.

78 MSA, structurally and chemically similar to the atmospheric aerosol nucleating precursor H₂SO₄, is regarded
79 as an important nucleating precursor for NPF (Elm, 2021; Hodshire et al., 2019). With stringent global regulations
80 reducing SO₂ emissions from fossil fuels (Perraud et al., 2015; Li et al., 2024b), MSA's role in NPF has received
81 widespread attention. Previous studies have investigated the effects of organic compounds (HCOOH, CH₃COOH,
82 CH₃OH, CH₂O, CH₃COCH₃, C₂H₅OC₂H₅, HCOOCH₃, H₂C₂O₄) (Zhao et al., 2017; Xu et al., 2017a; Arquero et al.,
83 2017; Xu et al., 2017b), iodic acid-containing (HIO₂, HIO₃) (Ning et al., 2022; Wu et al., 2023c) and bases (NH₃,
84 CH₃NH₂ (MA), (CH₃)₂NH, (CH₃)₃N) (Chen et al., 2016; Wen et al., 2018; Shen et al., 2020; Liu et al., 2022) on
85 MSA-driven binary nucleation, identifying MA as the most effective base in promoting MSA-driven NPF. However,

the binary MSA-MA nucleation mechanism is insufficient to explain the high NPF rates observed under realistic atmospheric conditions. This implies that other potential gaseous precursors may participate in and further enhance binary MSA-MA nucleation (Lee et al., 2019; Zhang et al., 2022). As a result, organic acids (Zhang et al., 2022) and inorganic acids (Hu et al., 2023) have been proved to enhance the formation rate of MSA-MA-driven ternary nucleation in areas with abundant specific pollutants. However, as one of the organic peroxides, the HMHP involved in MSA-MA-driven NPF remains unexplored. This will limit our knowledge of frequent NPF events, especially in urban industrial areas.

In this work, the gaseous and interfacial formation mechanisms of HMHP catalyzed by MSA, as well as its crucial role in MSA-MA nucleation, were explored. Specifically, quantum chemical calculations and Born-Oppenheimer molecular dynamics (BOMD) simulations were employed to elucidate the reaction mechanism of MSA-mediated CH_2OO hydrolysis leading to the formation of HMHP in both the gas phase and at the air-water interface. Subsequently, the potential effect of HMHP in the MSA-MA-dominated NPF process was assessed under a series of atmospheric conditions, utilizing quantum chemical calculation combined with the Atmospheric Cluster Dynamics Code (ACDC). This work will not only enhance the understanding of HMHP formation but also contributes to explaining some unexplained fluxes in NPF, while highlighting the significant impact of HMHP on nucleation processes.

2. Methods

2.1 Quantum Chemical Calculations

To investigate the MSA-assisted CH_2OO hydrolysis in the gas phase, geometric configurations and vibrational frequencies of all relevant configurations, including reactants, pre-reactive complexes, transition states (TSs), post-reactive complexes, and products, were calculated at the M06-2X/6-311++G(2df,2pd) level (Mardirossian and Head-Gordon, 2016; Pereira et al., 2017) using Gaussian 09. Intrinsic reaction coordinate analyses, performed at the same computational level, verified the correspondence of each TS to its associated pre- and post-reactive complexes. Subsequently, the ORCA 4.2.0 package (Neese, 2012) was performed to compute the single point energies using the CCSD(T)-F12/cc-pVDZ-F12 method.

To identify the global minimum energy configurations of $(\text{MSA})_x(\text{MA})_y(\text{HMHP})_z$ clusters, the ABCluster program (Zhang and Dolg, 2015) was utilized to systematically generate initial configurations for various clusters. Subsequently, these structures were further optimized using different levels of theoretical methods. Specifically, the ABCluster program generated $n \times 1000$ ($1 < n \leq 4$) initial structures for each cluster system. Then, the PM6 semi-

empirical method (Partanen et al., 2016) was used for preliminary optimization of these structures. As the M06-2X functional performs well for noncovalent binding and structural predictions of thermochemical and atmospheric aggregates, 100 of the lowest-energy configurations were chosen from $n \times 1000$ ($1 < n \leq 3$) structures and re-optimized using M06-2X/6-31G(d,p). Next, the 10 most stable configurations were re-optimized at the M06-2X/6-311++G(2df,2pd) level, and the configuration with the lowest free energy was identified. Ultimately, single-point energies were evaluated at the DLPNO-CCSD(T)-F12/cc-pVDZ-F12-CABS level (Tsona Tchinda et al., 2022) in ORCA, using geometries optimized for the stable clusters at the M06-2X/6-311++G(2df,2pd) level.

2.2 Rate coefficient calculations

The rate coefficients of MSA-mediated CH_2OO hydrolysis were investigated through a two-step approach. Initially, high-pressure-limit (HPL) rate coefficients were obtained using the VRC-VTST (Zhang et al., 2023; Zhang et al., 2024) approach implemented in Polyrate 2017-C (Meana-Pañeda et al., 2024). Subsequently, MESMER (Master Equation Solver for Multi-Energy Well Reactions) (Glowacki et al., 2012) was employed to calculate the rate coefficients of MSA-catalyzed CH_2OO hydrolysis across the temperature range of 280.0-320.0 K. The rate coefficients for the barrierless transition from separated reactants to pre-reactive complexes were estimated using the Inverse Laplace Transform (ILT) (Kumar et al., 2021) method. Concurrently, RRKM theory was employed to calculate the rate coefficients describing the conversion of the pre-reactive complex into the post-reactive species through the transition state. Further descriptions of the ILT methods and RRKM theory are provided in Parts S1 and S2 of the Supplement, respectively.

2.3 Born-Oppenheimer Molecular Dynamic (BOMD) simulation

BOMD calculations were carried out in CP2K program (Hutter et al., 2014), with exchange and correlation interactions described by the BLYP functional (Becke, 1988). Dispersion effects were incorporated through Grimme's dispersion (Grimme et al., 2010) (BLYP-D3) method. The Goedecker-Teter-Hutter (GTH) (Goedecker et al., 1996) pseudopotential was adopted for the core region, whereas the valence electrons were represented through a Gaussian DZVP basis (Phillips et al., 2005) in conjunction with an auxiliary plane-wave set. The calculations employed a plane-wave energy cutoff of 280 Ry together with a 40 Ry cutoff for the Gaussian basis set. In the gas phase, a supercell with a side length of 15 Å was employed to minimize periodic boundary effects, and the integration was performed with a 0.5 fs time step. A water droplet containing 191 molecules, serving as the interfacial model, was subjected to BOMD pre-optimization for roughly 5.0 ps at 300 K. CH_2OO and MSA were then positioned at the air-water interface, followed by 10 ps of molecular dynamics simulation. A supercell length

of 35 Å was adopted to prevent interactions between periodic images of the droplet, while the dynamics were advanced with a timestep of 1.0 fs. The gas-phase and interfacial simulations were conducted under NVT conditions at ~300 K, with the temperature maintained via a Nosé-Hoover thermostat.

2.4 Atmospheric Clusters Dynamic Code (ACDC) Model

ACDC (McGrath et al., 2012) was utilized to investigate the cluster formation rates, steady-state cluster concentrations, and growth mechanisms within the extensive MSA-MA-HMHP system. Thermodynamic data, calculated at the DLPNO-CCSD(T)-F12/cc-pVDZ-F12-CABS//M06-2X/6-311++G(2df,2pd) level of theory, were employed as input parameters for the ACDC simulations. The temporal progression of cluster concentrations was numerically resolved by solving the birth-death equations, employing the ode15s solver within the MATLAB-R2013a program. The birth-death equations are given below,

$$\frac{dc_i}{dt} = \frac{1}{2} \sum_{j < i} \beta_{j,(i-j)} C_j C_{(i-j)} + \sum_j \gamma_{(i+j) \rightarrow i} C_{i+j} - \sum_j \beta_{i,j} C_i C_j - \frac{1}{2} \sum_{j < i} \gamma_{i \rightarrow j} C_i + Q_i - S_i \quad (1)$$

In this formulation, c_i denotes the concentration of cluster i , β_{ij} represents the collision coefficient of clusters i with j and then $\gamma_{(i+j) \rightarrow i}$ represents the evaporation coefficient of clusters evaporating from $i + j$ to i and j clusters. Q_i represents the potential source for cluster i , while S_i denotes the sink term for cluster i . External losses of cluster i were represented using a fixed condensation sink coefficient of 0.02 s^{-1} (Qiao et al., 2024; Zhang et al., 2022). Besides, the clusters $(\text{MSA})_4 \cdot (\text{MA})_3$ and $(\text{MSA})_4 \cdot (\text{MA})_4$ were selected as boundary clusters in the MSA-MA-HMHP system, as boundary clusters in ACDC are required to be sufficiently stable to allow continued growth.

3. Results and discussion

3.1 The hydrolysis of CH₂OO with MSA in the gas phase

To evaluate the catalytic role of MSA in the CH₂OO hydrolysis reaction, the potential energy surface was investigated both in the presence and absence of MSA and H₂O. As depicted in Fig. 1(a) and (b), the potential energy surfaces with and without water closely match previously reported data (Wang et al., 2021a; Wang et al., 2021b), suggesting that the CCSD(T)-F12/cc-pVDZ-F12//M06-2X/6-311++G(2df,2pd) method is suitable for assessing MSA's catalytic effect. The MSA-catalyzed CH₂OO hydrolysis reaction (Fig. 1(c)) follows a continuous bimolecular process, involving CH₂OO⋯H₂O + MSA and MSA⋯H₂O + CH₂OO. The stabilization energy of MSA⋯H₂O is 9.8 kcal·mol⁻¹ higher than that of CH₂OO⋯H₂O. Considering the atmosphere concentrations of MSA (Li et al., 2024b), H₂O (Anglada et al., 2013), and CH₂OO (Khan et al., 2018), the concentration of MSA⋯H₂O is two orders of magnitude greater than that of CH₂OO⋯H₂O, as shown in Table S3, predicting that the MSA⋯H₂O +

CH₂OO route is the dominant pathway for the MSA-catalyzed reaction. Starting from MSA $\cdots$ H₂O + CH₂OO, the reaction proceeds through the IM_MSA intermediate, which has stabilization energy 6.5 kcal·mol⁻¹ and 0.7 kcal·mol⁻¹ higher than IM and IM_WM, respectively. The increase in stabilization energy is primarily due to the addition of MSA, which reduces ring strain and favours C-O bond formation. Atoms in Molecules (AIM) topological analysis results also show that in IM_MSA ($\rho = 3.74 \times 10^{-2}$), the strength of the density of all electrons (C-O) is 1.64 times stronger than in IM_WM ($\rho = 2.27 \times 10^{-2}$), further stabilizing the intermediate. The reaction then proceeds through the barrierless transition state TS_MSA, forming a nine-membered ring complex HOCH₂OOH $\cdots$ MSA (labeled as IMF_MSA). This process has a 1.4 kcal·mol⁻¹ lower energy barrier compared to the H₂O-catalyzed reaction, due to MSA's enhanced ability to facilitate proton transfer. Additionally, IMF_MSA is more stable by 7.6 kcal·mol⁻¹ compared to IMF_WM.

To further examine the impact of MSA on the CH₂OO hydrolysis reaction, we calculated the effective rate constants for the CH₂OO hydrolysis assisted by H₂O ([H₂O] = 10¹⁵-10¹⁷ molecules·cm⁻³), NH₃ ([NH₃] = 10⁷-10¹¹ molecules·cm⁻³), H₂SO₄ ([H₂SO₄] = 10⁴-10⁸ molecules·cm⁻³) and MSA ([MSA] = 10⁴-10⁸ molecules·cm⁻³) in Table S5. When the MSA concentration ranges from 10⁶ to 10⁸ molecules·cm⁻³, its catalytic effect is substantially stronger than that of NH₃ ([NH₃] = 10⁷-10¹¹ molecules·cm⁻³), with k'_{MSA} being approximately 2-6 orders of magnitude over the temperature range of 280.0-320.0 K. Similarly, for MSA concentrations between 10⁵ and 10⁸ molecules·cm⁻³, MSA exhibits a significantly higher catalytic activity than SA ([SA] = 10⁴-10⁷ molecules·cm⁻³). In this case, k'_{MSA} exceeds k'_{SA} by about 1-3 orders of magnitude. Taken together, these results demonstrate that MSA is a more effective catalyst than both NH₃ and SA under atmospherically relevant conditions. Nevertheless, even under extreme conditions, with MSA at its upper-limit concentration ([MSA] = 10⁸ molecules·cm⁻³) and H₂O at its lower-limit concentration ([H₂O] = 10¹⁶ molecules·cm⁻³), k'_{MSA} is approximately five orders of magnitude smaller than k'_{WM} , indicating that the catalytic efficiency of MSA remains lower than that of H₂O.

3.2 The hydrolysis of CH₂OO with MSA at the air-water interface

The mechanism of MSA-catalyzed CH₂OO hydrolysis at the air-water interface remained unclear. To elucidate this process, BOMD simulations were employed to investigate possible reaction pathways at the aqueous interface. Based on analogies with CH₂OO reactions involving other atmospheric species (Ding et al., 2024; Li et al., 2023; Cheng et al., 2025), three potential routes were considered: (i) MSA interacting with adsorbed CH₂OO at the air-water interface, (ii) CH₂OO interacting with adsorbed MSA at the air-water interface, or (iii) the MSA-CH₂OO complex reacting at the air-water interface. Because both CH₂OO and MSA exhibit high reactivity toward interfacial

water, their lifetimes at the droplet interface remain very brief, making pathway (i) and (ii) far less likely than pathway (iii). Therefore, only the reaction of the MSA $\cdots$ CH₂OO complex at the aqueous interface was considered, owing to its high stability in the gas phase.

As displayed in Fig. 2, Fig. S1, and Movie S1, the MSA $\cdots$ CH₂OO complex initially interacts with a water molecule at the aqueous interface. At 0.76 ps, this water molecule forms both a van der Waals interaction with the carbon atom in CH₂OO ($d(\text{C}-\text{O}3) = 4.00 \text{ \AA}$) and a hydrogen bond with the oxygen atom in MSA ($d(\text{O}4-\text{H}2) = 3.27 \text{ \AA}$), thereby generating a nine-membered ring structure (CH₂OO $\cdots$ MSA $\cdots$ H₂O). As the reaction progresses, a hydration transition-state-like ring configuration forms at 1.06 ps, with the C-O3, O4-H2 and O2-H1 bonds shorten to 1.75 $\text{\AA}$, 1.27 $\text{\AA}$ and 0.97 $\text{\AA}$, respectively, while the O3-H2 and O1-H1 bonds elongate to 1.16 $\text{\AA}$ and 1.85 $\text{\AA}$, respectively. By 1.16 ps, the bond lengths of C-O3, O4-H2 and O2-H1 further shorten to 1.43 $\text{\AA}$, 1.05 $\text{\AA}$ and 0.96 $\text{\AA}$, while the O3-H2 and O1-H1 bonds further increase to 1.61 $\text{\AA}$ and 1.93 $\text{\AA}$, respectively, indicating the formation of the hydrogen-bonded CH₂OOH₂O $\cdots$ MSA complex. During this process, the interfacial water molecule acts as a reactant, with MSA serving as a proton transfer bridge. Notably, due to the higher abundance of interfacial water molecules compared to MSA, the formation of HMHP at the interface primarily proceeds through the direct hydration of CH₂OO. However, the source of HMHP from MSA-catalyzed CH₂OO hydrolysis at the interface occurs at a significantly faster rate than its corresponding gas-phase formation, with a computed ratio (r_1) of 1.01×10^2 at 298.0 K, which is detailed computational results are provided in Part S3. Traditionally, the loss of CH₂OO in the troposphere has been primarily attributed to its hydrolysis. Therefore, it is crucial to examine the rate ratio r_2 (equation (S4)) between interfacial MSA-catalyzed CH₂OO hydrolysis and the corresponding gas-phase process mediated by water. At 298.0 K, r_2 is 13.4, indicating that the formation of HMHP from interfacial MSA-mediated CH₂OO hydrolysis is much closer to that catalyzed by water in the gas phase. These results indicate that interfacial MSA-catalyzed CH₂OO hydrolysis is a significant source of HMHP formation in MSA-polluted areas under relatively humid conditions. Consequently, when evaluating the comprehensive sources of HMHP in MSA-rich regions, it is essential to consider the formation involving MSA-catalyzed hydrolysis of CH₂OO at the air-water interface.

3.3 The significance of HMHP in MSA-MA nucleation

The role of HMHP on MSA-MA-driven ternary nucleation was assessed. Initial assessments focused on the potential interaction sites of HMHP with MSA and MA, followed by an analysis of the conformation and stability of the (MSA)_x(MA)_y(HMHP)_z ($0 \leq y \leq x + z \leq 3$) clusters. The MSA-MA nucleation mechanism involving

230 HMHP was then examined, along with the effects of temperature and precursor concentrations on the MSA-MA-
231 HMHP system. Finally, the atmospheric implication of HMHP for MSA-MA nucleation were calculated for urban
232 industrial areas.

233 3.3.1 Cluster conformational analysis

234 The formation of stable clusters is primarily dictated by the pronounced coupling of nucleation precursors
235 (Ning et al., 2024; Li et al., 2024a; Li et al., 2024b). The electrostatic potential (ESP)-mapped molecular van der
236 Waals (vdW) surface was computed to characterize the binding potential of HMHP with MSA and MA and to locate
237 the corresponding interaction sites (Fig. 3(a)). The analysis revealed that HMHP exhibits a positive ESP value
238 ($+54.09 \text{ kcal}\cdot\text{mol}^{-1}$) at the H atom of its -OH group, indicating its role as a hydrogen bond (HB) donor, while the
239 oxygen atom of the -COOH shows a negative ESP of $-32.68 \text{ kcal}\cdot\text{mol}^{-1}$, indicating its role as a hydrogen bond
240 acceptor. Thus, HMHP possesses dual functionalities as a HB donor and acceptor, allowing its association with the
241 nucleating precursors MSA and MA, thereby generating HBs that stabilize the MSA-MA-HMHP clusters. Based
242 on the ESP analysis, the most stable configurations of $(\text{MSA})_x(\text{MA})_y(\text{HMHP})_z$ clusters were obtained (Fig. S2), and
243 the actual binding sites within the ternary clusters were found to be consistent with the predictions from the ESP
244 analysis.

245 To further access the non-covalent interactions within the $(\text{MSA})_x(\text{MA})_y(\text{HMHP})_z$ clusters, the reduced density
246 gradient (RDG) analysis was performed on representative clusters, including $(\text{MSA})_1(\text{MA})_1$,
247 $(\text{MSA})_1(\text{MA})_1(\text{HMHP})_1$ and $(\text{MSA})_1(\text{MA})_1(\text{HMHP})_2$ cluster (Fig. 3(b)). In the RDG analysis, the blue region
248 corresponds to strong attractions, similar to HBs, within the system. When compared to the $(\text{MSA})_1(\text{MA})_1$ cluster,
249 the $(\text{MSA})_1(\text{MA})_1(\text{HMHP})_1$ cluster exhibits two additional blue spikes, suggesting that HMHP forms strong
250 interactions, particularly HBs, with MSA and MA. Moreover, the dominant blue spike in the
251 $(\text{MSA})_1(\text{MA})_1(\text{HMHP})_2$ cluster is shifted further leftward compared to that in the $(\text{MSA})_1(\text{MA})_1(\text{HMHP})_1$ cluster,
252 indicating the introduction of HMHP molecules strengthens the HB interactions within the system. Besides, in
253 clusters involving HMHP (Fig. S2), HMHP facilitates the formation of cage-like clusters with MSA and MA through
254 a spatial network of HBs. Notably, proton transfer is observed in ternary clusters involving HMHP. Taken together,
255 these findings suggest that HMHP forms stable clusters with MSA and MA via both HBs and proton transfer, thereby
256 enhancing the stability of the MSA-MA-HMHP clusters.

257 3.3.2 Cluster stability analysis

258 To evaluate the effect of HMHP on the thermodynamic stability of MSA-MA-clusters, the Gibbs free energy

(ΔG) (Table S7) and total evaporation rates ($\Sigma\gamma$) (Table S10) of clusters in the MSA-MA-HMHP system were calculated. The ΔG of HMHP-involved clusters are consistently lower than those of the corresponding pure MSA-MA clusters, indicating that the participation of HMHP enhances cluster thermodynamic stability. Consistent with the ΔG analysis, HMHP-containing clusters also exhibit markedly reduced $\Sigma\gamma$ values. In addition to thermodynamic stability, cluster growth tendencies play a crucial role in atmospheric cluster formation. From a kinetic perspective, cluster formation is governed by the competition between molecular collisions and evaporation processes. When the ratio of the collision frequency between a cluster and a monomer at concentration C to the total evaporation rate, expressed as $\beta C/\Sigma\gamma$, is higher than 1, the cluster is considered capable of sustained growth. Accordingly, as shown in Fig. 4, the $\beta C/\Sigma\gamma$ ratios were evaluated under various atmospheric conditions to assess the influence of HMHP on the growth behavior of MSA-MA clusters. Among the clusters examined, $(\text{MSA})_2$, $(\text{MSA})_2 \cdot (\text{MA})$, $(\text{MSA})_2 \cdot (\text{MA})_2$, $(\text{MSA})_3 \cdot (\text{MA})_2$, $(\text{MSA}) \cdot (\text{MA}) \cdot (\text{HMHP})$ and $(\text{MSA})_2 \cdot (\text{MA})_2 \cdot (\text{HMHP})$ display sufficient stability against evaporation, enabling further growth through collisions with additional monomers. Overall, these results suggest that HMHP is likely to participate in the nucleation process of MSA-MA clusters by enhancing their thermodynamic stability and increasing their potential for sustained cluster growth.

3.3.3 Cluster formation pathways

To track the detailed nucleation pathways of HMHP involved in the formation of MSA-MA clusters, ACDC simulation were conducted at four different temperatures (238.15 K, 258.15 K, 278.15 K and 298.15 K), with concentrations set as $[\text{MSA}] = 1.0 \times 10^6 \text{ molecules} \cdot \text{cm}^{-3}$, $[\text{MA}] = 2.5 \times 10^8 \text{ molecules} \cdot \text{cm}^{-3}$ and $[\text{HMHP}] = 1.0 \times 10^{11} \text{ molecules} \cdot \text{cm}^{-3}$. The results are presented in Fig. 5(a). At 238.15 K, the primary growth of the MSA-MA-HMHP system can be divided into two distinct routes. One pathway involves MSA-MA clustering alone, while the other incorporates HMHP molecules in the cluster formation process. In the HMHP involving pathway, HMHP initially interacts with the heterodimer $(\text{MSA})_2 \cdot (\text{MA})_2$, resulting in the formation of a $(\text{MSA})_2 \cdot (\text{MA})_2 \cdot (\text{HMHP})$ trimer. This trimer then grows out of the simulated system. Following this, the $(\text{MSA})_2 \cdot (\text{MA})_2 \cdot (\text{HMHP})$ trimer collides with an MSA monomer, forming the cluster $(\text{MSA})_3 \cdot (\text{MA})_2 \cdot (\text{HMHP})$. Subsequently, HMHP evaporates from this cluster, leaving behind the $(\text{MSA})_3 \cdot (\text{MA})_2$ cluster. Finally, the $(\text{MSA})_3 \cdot (\text{MA})_2$ trimer interacts with an MA monomer, resulting in the $(\text{MSA})_3 \cdot (\text{MA})_3$ cluster, which eventually exits the system. These processes illustrate HMHP's dual function: it acts both as a "catalyst" that promotes the formation of MSA-MA clusters and as a "participant" in the assembly of critical clusters. This dual role is also observed at 258.15 K. However, at 278.15 K, HMHP acts only as a "participant" in the formation of critical clusters. With increasing temperature, the contribution of HMHP to

cluster formation diminishes, and at 298.15 K (Fig. S3) cluster formation occurs exclusively through the pure binary MSA-MA pathway. The higher evaporation coefficient of clusters at elevated temperatures leads to their faster evaporation. Together, these findings underscore the significant and direct role of HMHP in MSA-MA nucleation, effectively facilitating cluster formation across different temperatures.

To further investigate the influence of temperature (T) on HMHP's participation role in the MSA-MA nucleation process, we analyzed the contribution of different clustering pathways to the outward flux under varying T . As shown in Fig. 5(b), with increasing T , the dominance of HMHP-involved pathways gradually diminishes. At 238.15-258.15 K, the HMHP-involved pathway accounts for 63%, whereas at 278.15 K, the contribution decreases to 21%. These results highlight the significant role of low temperatures in facilitating HMHP's participation in MSA-MA nucleation. In addition to T , the contribution of nucleating precursor flux also significantly affects trace substances' involvement in the MSA-MA nucleation process. The contribution of HMHP to the MSA-MA system was found to be relatively insensitive to [MA], as illustrated in Fig. S3. In contrast, it is strongly influenced by [MSA], as depicted in Fig. 5(c). As [MSA] increases from 10^4 to 10^8 molecules·cm⁻³, the contribution of HMHP-involved pathways decreases from 65% to 40%. At the low [MSA] (10^4 - 10^6 molecules·cm⁻³), HMHP involvement remains substantial, accounting for 65% of the pathway. Overall, this study reveals that HMHP's contribution to MSA-MA nucleation is most pronounced under conditions of low T , and low concentrations of both MSA and MA.

3.4 Atmospheric implications of HMHP to MSA-MA system

Building on the above findings, low T coupled with low [MSA] and [MA] appear to favor an enhanced role of HMHP in MSA-MA nucleation. To assess the atmospheric significance of these variations, we quantified the contributions of MSA-MA cluster growth pathways, with and without HMHP involvement (Fig. 6), under favorable conditions of temperature ($T = 258.15$ K) and precursor concentrations ([MSA] = 1.00×10^4 molecules·cm⁻³, [MA] = 2.50×10^7 molecules·cm⁻³). Indeed, substantial variability in atmospheric HMHP concentrations has been observed across diverse environments worldwide. For example, levels range from 2.50×10^9 to 6.25×10^9 molecules·cm⁻³ in Central Portugal, Pabstthum, and Beijing. Higher concentrations, between 1.15×10^{10} to 3.00×10^{10} molecules·cm⁻³, have been observed in Guang Zhou and Niwot Ridge, while the Southeastern United States exhibit the highest concentrations, reaching up to 1.25×10^{11} molecules·cm⁻³. As shown in Fig. 6, in low [HMHP] regions such as Pabstthum and Beijing, HMHP-involved pathways account for only 11% and 12% of total NPF, respectively. In contrast, in environments characterized by high HMHP concentrations, such as the southeastern United States (1.25×10^{11} molecules·cm⁻³) and Niwot Ridge (3.00×10^{10} molecules·cm⁻³), HMHP-involving

nucleation pathways become dominant. Under these conditions, HMHP acts both as a “catalyst”, facilitating the formation of MSA-MA clusters, and as an “participant” in the assembly of critical clusters (Figs. S5 and S6). These two roles contribute up to 59% and 42%, respectively, to the overall nucleation process. These results highlight that HMHP exerts a markedly stronger influence on MSA-MA nucleation at elevated concentrations, particularly in urban industrial regions, where its contribution to NPF can be substantial.

Previous studies have revealed that SA-MA and SA-A nucleation mechanisms are widely regarded as key contributors to new particle formation in urban industrial regions (Yin et al., 2021; Liu et al., 2021). To underscore the importance of MSA-MA-HMHP nucleation in urban industrial regions, the cluster formation rates (J) of the MSA-MA-HMHP system have been compared with those of the SA-MA and SA-A systems (Qiao et al., 2024) (Fig. S7). The results show that, over the temperature range of 238.15 K-298.15 K, the J of MSA-MA-HMHP system is 1-5 orders of magnitude higher than that of SA-MA system at equivalent precursor concentrations ($[SA] = 1.00 \times 10^6 \text{ molecules}\cdot\text{cm}^{-3}$ and $[MA] = 1.00 \times 10^8 \text{ molecules}\cdot\text{cm}^{-3}$). Similarly, under the conditions of $[SA] = 1.00 \times 10^6 \text{ molecules}\cdot\text{cm}^{-3}$ and $[A] = 1.00 \times 10^{10} \text{ molecules}\cdot\text{cm}^{-3}$, the J of MSA-MA-HMHP systems slightly exceeds that of SA-A system by approximately 5-6 orders of magnitude. These comparisons suggest that HMHP plays a key role in enhancing MSA-MA nucleation, particularly in urban industrial environments.

4. Conclusions

The hydrolysis of CH_2OO by MSA, occurring in both the gas phase and at interfaces and resulting in hydroxymethyl hydroperoxide (HMHP) formation, was investigated. In addition, the role of HMHP in promoting MSA-MA nucleation was elucidated.

Quantum chemical calculations suggest that the activation energy for HMHP formation via MSA-catalyzed gaseous hydrolysis of CH_2OO is relatively low, observed to be $2.9 \text{ kcal mol}^{-1}$. At a concentration of $[MSA] = 10^6$ - $10^8 \text{ molecules}\cdot\text{cm}^{-3}$, MSA exhibits a dominant catalytic effect compared to other catalysts, such as NH_3 ($[\text{NH}_3] = 10^7$ - $10^{11} \text{ molecules}\cdot\text{cm}^{-3}$), within the temperature range of 280.0-320.0 K. Notably, the computed effective rate constant for MSA is larger by about 2-6 orders of magnitude compared with that of NH_3 . Simultaneously, when $[MSA] = 10^5$ - $10^8 \text{ molecules}\cdot\text{cm}^{-3}$, k'_{MSA} is found to be 1 to 3 orders of magnitude larger than k'_{SA} , which corresponds to $[SA] = 10^4$ - $10^7 \text{ molecules}\cdot\text{cm}^{-3}$. BOMD simulations further demonstrate that the MSA-catalyzed CH_2OO hydrolysis at the gas-liquid interface follows a ring-based reaction mechanism on a picosecond timescale. Remarkably, HMHP formation through MSA-catalyzed CH_2OO hydrolysis occurs rapidly and stably at the interface.

HMHP participates in MSA-MA-driven ternary nucleation through its bifunctional hydrogen-bonding capacity,

346 which allows direct association with both MSA and MA. Thermodynamic analysis shows that MSA-MA-HMHP
347 trimers possess lower Gibbs free energy than MSA-MA dimers, indicating enhanced cluster stability. Cluster
348 formation pathways analysis reveals a temperature-dependent role: at 238.15 K and 258.15 K, HMHP functions as
349 both catalyst and participant, whereas at 278.15 K it acts solely as a participant due to increased evaporation rates.
350 Under low temperature, low [MSA] and [MA], and high [HMHP] conditions, particularly in Niwot Ridge and the
351 southeastern United States, HMHP-involved pathways contribute up to 59% and 42% of total nucleation flux. These
352 results predict that HMHP substantially enhances MSA-MA-driven NPF in urban industrial regions, helping to
353 explain previously unaccounted NPF sources and improve nucleation models.

354 Overall, this work deepens the understanding of hydroperoxide formation in MSA-polluted regions, especially
355 at the gas-liquid interface. It also reveals the potential contribution of other organic peroxides to NPF, offering a
356 plausible explanation for part of the unaccounted particle fluxes in urban industrial regions.

357 **Data availability**

358 Data will be made available on request.

359 **Authorship contributions**

360 RL: methodology, investigation, data curation, writing (original draft). ZL: writing (review), data curation,
361 methodology, investigation. CZ: writing (review), data computation. JY: data curation, data computation. HC:
362 writing (editing), data curation, visualization, investigation. XL: writing (review and editing). NH: data curation,
363 project administration, writing (review and editing). RW: funding acquisition, writing (review and editing). TZ:
364 writing (review and editing), funding acquisition.

365 **Competing interests**

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

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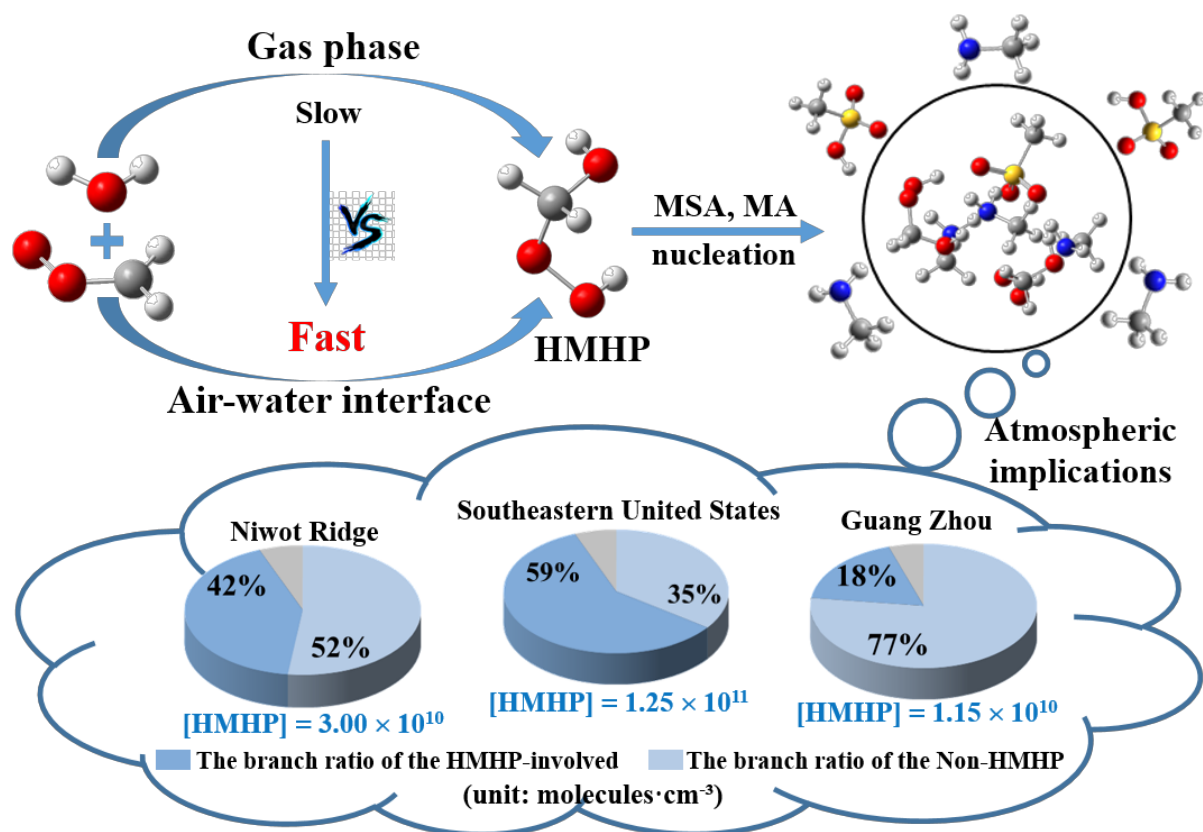
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MSA: methanesulfonic acid MA: methylamine HMHP: hydroxymethyl hydroperoxide

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Graphic Abstract

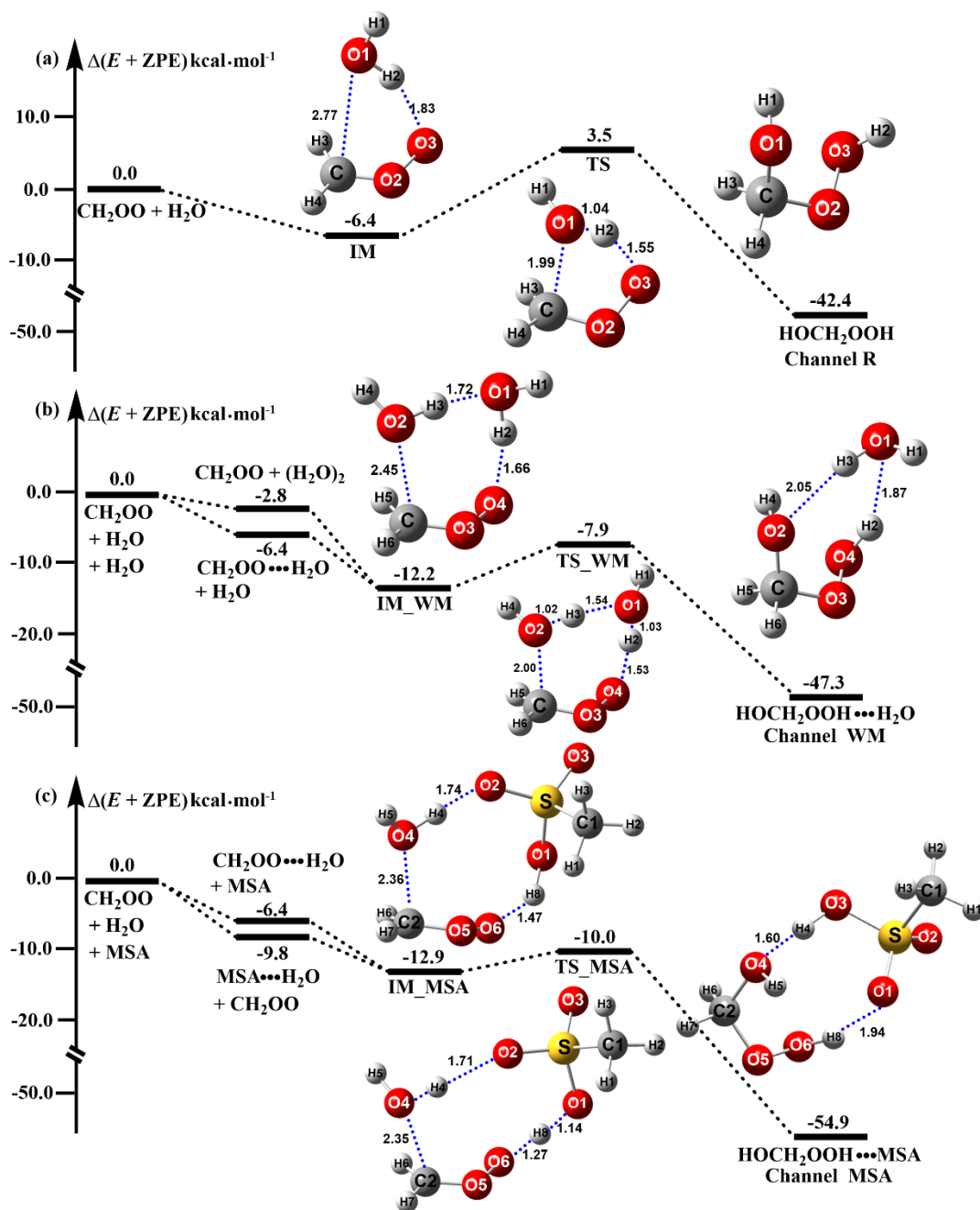


Fig. 1 Potential energy surface for the hydrolysis of CH_2OO without (a) and with H_2O (b) and $\text{CH}_3\text{SO}_3\text{H}$ (c) at the CCSD(T)-F12/cc-pVDZ-F12/M06-2X/6-311++G(2df,2pd) level

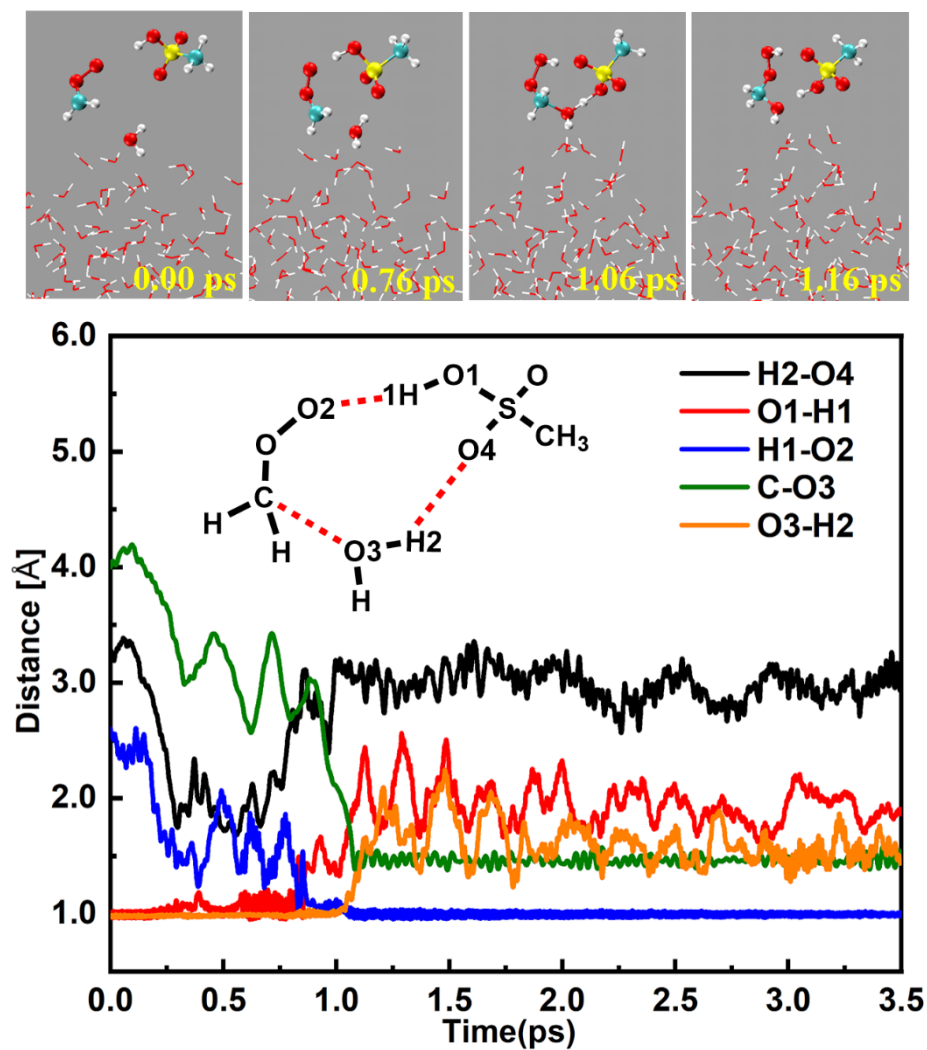


Fig. 2 BOMD simulation trajectories and snapshots for $\text{CH}_3\text{SO}_3\text{H}$ -mediated CH_2OO hydration at the water microdroplet interface

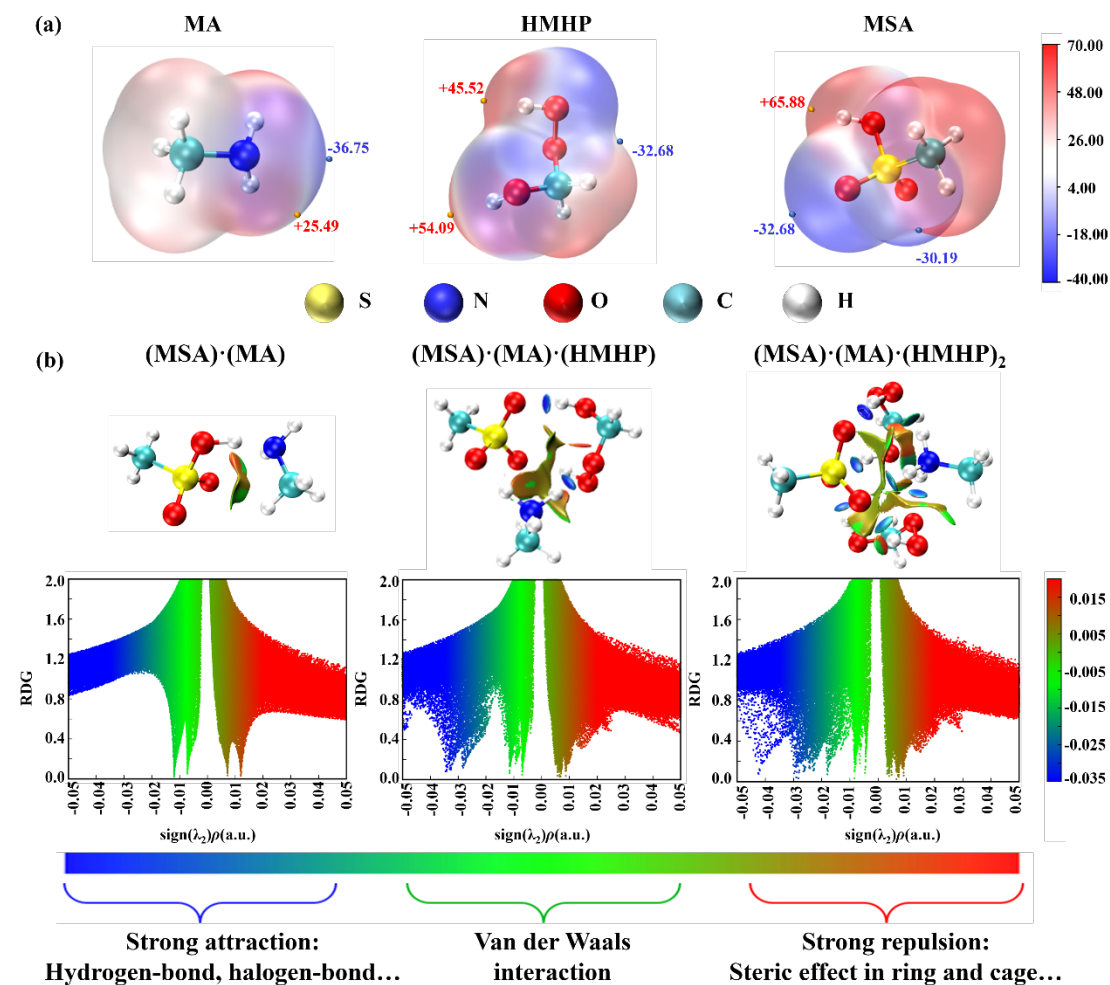


Fig. 3 (a) Electrostatic potential (ESP)-mapped molecular van der Waals (vdW) surface of MA, HMHP, and MSA calculated at M06-2X/6-311++G(2df, 2pd) + aug-cc-pVTZ level of theory. The orange and blue dots indicate the positions of ESP maximums and minimums (in kcal·mol⁻¹), with the corresponding values labeled in red and blue numbers, respectively. (b) The reduced density gradient (RDG) color-filled maps and isosurface of $(\text{MSA})_1 \cdot (\text{MA})_1$, $(\text{MSA})_1 \cdot (\text{MA})_1 \cdot (\text{HMHP})_1$, and $(\text{MSA})_1 \cdot (\text{MA})_1 \cdot (\text{HMHP})_2$ clusters

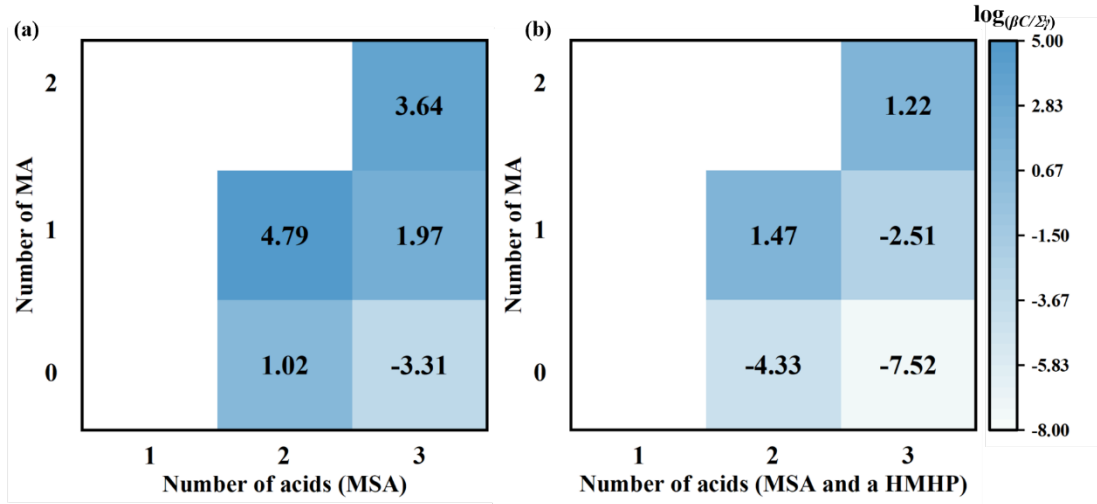


Fig. 4 The $\log(\beta C / \Sigma \gamma)$ of $(\text{MSA})_x(\text{MA})_y(\text{HMHP})_z$ ($0 \leq y \leq x + z \leq 3$) clusters at 238.15K. (a) without HMHP monomer, (b) containing a HMHP monomer. The $\beta C / \Sigma \gamma$ is the ratio of collision frequency between the cluster and monomer molecule at the concentration C to the total evaporation frequency at $C = 2.5 \times 10^8 \text{ molecules} \cdot \text{cm}^{-3}$

(a) Cluster formation pathway ($[MSA] = 1.0 \times 10^6$; $[MA] = 2.5 \times 10^8$; $[HMHP] = 1.0 \times 10^{11}$ molecules·cm⁻³)

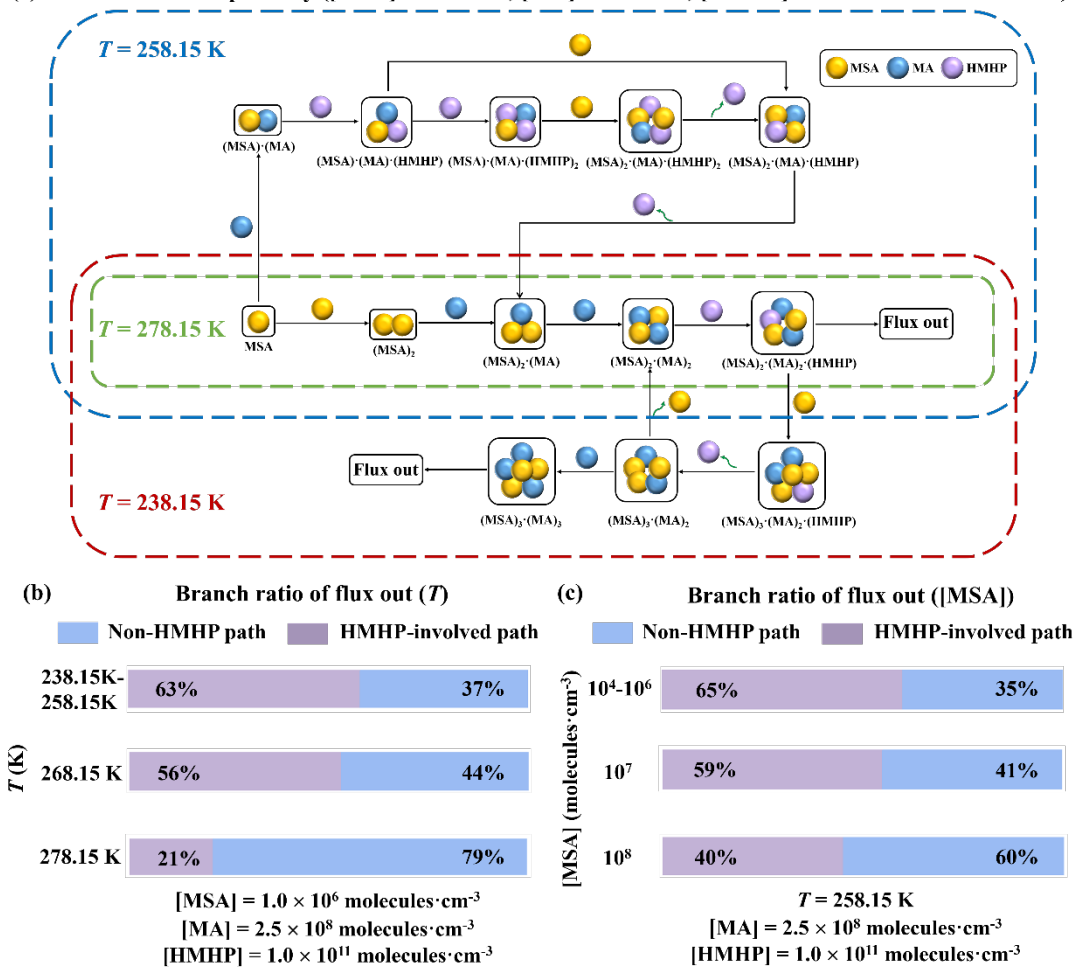


Fig. 5 Nucleation mechanism of the HMHP-MSA-MA system. (a) Cluster formation pathway at different temperatures; (b) the branch ratio of outward flux under varying T (238.15 K-298.15K); and (c) the branch ratio of outward flux under varying $[MSA]$ (1.0×10^4 - 1.0×10^8 molecules·cm⁻³)

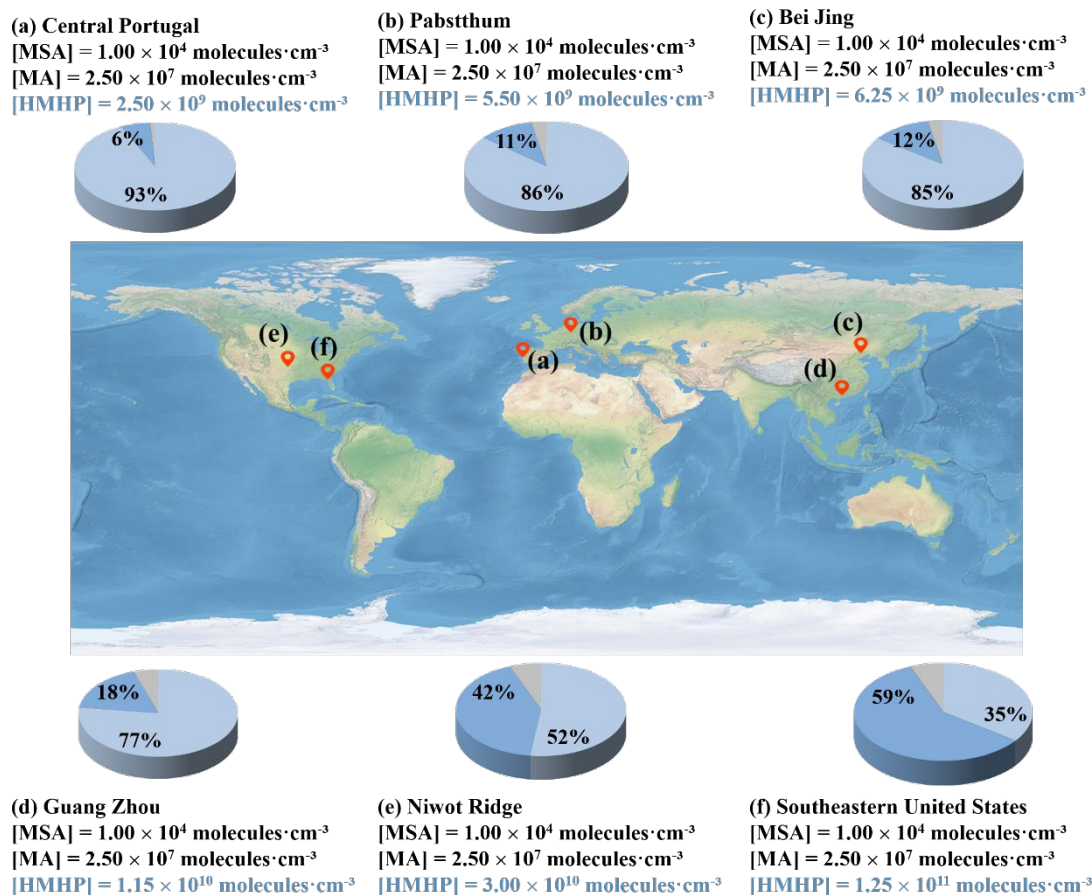


Fig.6 The branch ratio of the MSA-MA-HMHP (blue pie) and MSA-MA (light blue pie) growth pathways based on field data in different regions with a different $[HMHP]$. The data recorded in blue are from field observations, and those in black are set to be a median in this study. $[MA] = 2.5 \times 10^8 \text{ molecules} \cdot \text{cm}^{-3}$. The map is from ©Google Maps (<https://www.google.com/maps>, last access: 15 July 2025)

Table 1 The rate ratio (r_1) between the interfacial MSA-catalyzed CH₂OO hydrolysis reaction and the corresponding reaction in the gas phase as well as the rate ratio (r_2) between the MSA-catalyzed CH₂OO hydrolysis reaction at the air-water nanodroplet interface and the hydrolysis of CH₂OO facilitated by H₂O in the gas phase at 298.0 K

Rate ratio	r_1	r_2
	1.01×10^2	13.4