INTRODUCTION

The atmospheric (photo-) oxidation of volatile organic compounds (VOCs) emitted from natural and man-made sources leads to the formation of secondary species, notably pollutants like ozone and (VOCs) emitted from natural and man-made sources leads to the formation of secondary species, notably pollutants like ozone and secondary organic aerosol (SOA)

Secondary organic aerosol (SOA) represents a major fraction of atmospheric fine particles. Both biogenic and anthropogenic volatile organic compounds (VOCs) can contribute to SOA through (photo-) oxidation. However, the current understanding of their combined, interactive effect on SOA formation and composition is still limited, challenging the accuracy in assessing global SOA budget, sources, and climate effect. Here we combine laboratory experiments and modelling to show that isoprene can suppress SOA formation from photo-oxidation of anthropogenic aromatics (toluene and p-xylene) with the presence of NOx, and similar SOA suppression phenomena are observed when replacing isoprene with propene. We find that the decreased SOA in such mixed-VOC conditions can be largely attributed to OH scavenging effect, resulting in reduced consumption of parent aromatics. However, various changes in SOA oxidation state (i.e., O/C) and oxidation pathways (i.e., more carboxyls formation) are observed following addition of isoprene, and the SOA chemical composition may not be similar to any single parent hydrocarbon, which implies the existence of complex interactions between the degradation chemistry for alkenes and aromatics. Under the conditions of this work, the OH scavenging effect is largely determined by gas-phase chemistry, which is expected to be widespread in binary or more complex systems in ambient air. More broadly, we infer that the global budget of anthropogenic SOA and its corresponding radiative forcing could be affected by biogenic emission of isoprene, particularly in urban environments with appreciable vegetation coverage.

Generating secondary organic aerosol (SOA) from photo-oxidation of volatile organic compounds (VOCs) emitted from natural and man-made sources leads to the formation of secondary species, notably pollutants like ozone and secondary organic aerosol (SOA). This process is crucial for understanding the global budget of SOA, its sources, and its impact on climate change. While significant progress has been made in understanding the formation of biogenic SOA, the contribution of anthropogenic sources is less well understood, particularly in urban environments.

Recent studies have shown that the presence of isoprene, a biogenic VOC, can suppress the formation of SOA from anthropogenic sources, such as aromatic hydrocarbons. This suppression is due to the scavenging of OH radicals by isoprene, leading to reduced consumption of parent aromatics and altered SOA composition. Studies have also demonstrated that the SOA yield in the presence of isoprene is lower than that of the parent aromatics, indicating a suppressed SOA formation process. This suppression is not confined to binary systems but extends to mixed-VOC conditions, where the interaction between biogenic and anthropogenic VOCs plays a significant role.

Understanding the combined effect of biogenic and anthropogenic VOCs is crucial for accurate assessments of SOA's impact on climate and air quality. While previous studies have primarily focused on biogenic SOA formation, the role of anthropogenic sources is gaining increasing attention. The suppression of SOA formation by isoprene highlights the importance of considering both biogenic and anthropogenic sources in future studies to better understand the global SOA budget.

To address these challenges, we combine laboratory experiments and modelling to study the effect of isoprene on SOA formation from mixed-VOC conditions. Our approach allows us to investigate the interactive effect of biogenic and anthropogenic VOCs on SOA formation and composition, providing insights into the complex processes involved. By understanding these interactions, we can improve models that predict SOA formation and its impact on climate and air quality.

In conclusion, the suppression of SOA formation by isoprene underscores the importance of considering both biogenic and anthropogenic sources in future studies. This work contributes to a broader understanding of SOA formation in mixed-VOC environments, which is essential for accurate assessments of SOA's impact on climate and air quality.
SOA formation by mixing two anthropogenic VOCs (n-dodecane and 1,3,5-trimethylbenzene) in the presence of NOx and sunlight, and their experiments were conducted in a large-scale outdoor chamber, which is close to the real atmospheric conditions. By contrast, Voliotis et al.\textsuperscript{32} found that the volatility of SOA from anthropogenic and biogenic VOC mixtures (α-cresol and α-pinene) was in between those measured in single-VOC systems, and their experimental conditions were representative of typical ambient daytime photochemistry with moderate relative humidity (50 ± 5%) and temperature (24 ± 2 °C). Kari et al.\textsuperscript{33} further conducted chamber experiments by injecting both α-pinene and real-world exhaust from gasoline direct-injection (GDI) engine, and they found that the presence of vehicle exhausts may change the oxidation pathways of α-pinene and result in suppression (by more than 50%) of biogenic SOA mass yield. These recent laboratory studies suggested that SOA formation from mixed-VOC condition was governed by the molecular interactions of the oxidation products from different hydrocarbons, which was different and more complex than the SOA derived from single-VOC experiments. However, there are still rather limited laboratory studies on the interactive effect of VOC mixtures from different sources—particularly the effects of biogenic VOC on anthropogenic SOA formation and composition—implying a significant knowledge gap regarding SOA formation chemistry. Given the fact that the real atmosphere is a complex mixture of thousands of organic and inorganic species from various sources\textsuperscript{1,24,34}, an improved understanding of such interaction effects on SOA formation is much needed\textsuperscript{18,31,35}, which consequently motivated the current work.

Aromatics such as toluene and p-xylene are well known as an important class of VOCs with a high SOA yield from anthropogenic emissions such as vehicles and industry\textsuperscript{36,37}, while isoprene is the most abundant biogenic hydrocarbon with global emission of ~500 Tg year\textsuperscript{-1}.\textsuperscript{38} In this study, we combine chamber experiments and photochemical modelling to investigate the effect of isoprene on anthropogenic SOA formation from toluene and p-xylene. We find a significant suppression of aromatic SOA by isoprene, and further elucidate the underlying chemical mechanisms through additional investigation on analogous reactants, gas-phase and SOA modelling, and SOA composition analysis. Finally, the potential implications on global SOA budget, sources and climate effect from photochemical interaction between anthropogenic and biogenic VOC are discussed.

**RESULTS AND DISCUSSION**

**Experimental evidence of anthropogenic SOA suppression by isoprene**

A series of chamber experiments on SOA formation from toluene/NOx and p-xylene/NOx photo-oxidation in the absence and presence of isoprene or propene were performed (see "Methods"), and the initial conditions of each experiment are given in Table 1. The time-dependent SOA mass concentrations and total particle number concentrations are shown in Fig. 1, which is divided into

| Table 1. Summary of experimental initial conditions carried out at CAPS-ZJU chamber as well as the chemical parameters of SOA derived from HR-ToF-AMS. |
|---|
| **Expt ID** | **Toluene/NOx** | **p-xylene/NOx** | **Isoprene/NOx** | **Propene/NOx** | **NOx (ppbv)** | **H2O2 min\(^{-1}\)** | **f\(_{43}\) (%)** | **f\(_{44}\) (%)** | **H/C** | **O/C** | **OM/OC** |
| ZJU095 | 144 | – | – | 33 | 0.40 | – | – | – |
| ZJU098 | 144 | – | – | 30 | 0.40 | – | – | – |
| ZJU104 | 144 | – | – | 28 | 0.39 | 9.4 | 14.3 | 1.56 | 0.64 | 1.99 |
| ZJU130 | 144 | – | – | 39 | 0.38 | 8.7 | 12.8 | 1.56 | 0.60 | 1.93 |
| Isoprene/NOx | – | – | 165 | 32 | 0.36 | – | – | – |
| ZJU178 | – | – | 165 | 31 | 0.36 | 9.8 | 7.1 | 1.78 | 0.36 | 1.64 |
| ZJU189 | – | – | 165 | 31 | 0.36 | 9.8 | 7.1 | 1.78 | 0.36 | 1.64 |
| Propene/NOx | ZJU191 | – | – | 152 | 34 | 0.36 | – | – | – |
| ZJU193 | – | – | 152 | 33 | 0.36 | – | – | – |
| Toluene/isoprene/NOx | ZJU164 | – | – | 165 | 33 | 0.37 | 9.3 | 9.9 | 1.66 | 0.48 | 1.78 |
| ZJU183 | – | – | 38 | 33 | 0.36 | 9.3 | 7.0 | 1.71 | 0.34 | 1.60 |
| Toluene/propene/NOx | ZJU176 | 144 | – | – | 152 | 32 | 0.36 | 8.6 | 8.1 | 1.69 | 0.37 | 1.63 |
| p-xylene/NOx | ZJU220 | – | 110 | – | 53 | 0.38 | 10.7 | 9.8 | 1.63 | 0.45 | 1.74 |
| ZJU232 | – | 110 | – | 50 | 0.37 | 10.9 | 10.8 | 1.61 | 0.49 | 1.79 |
| p-xylene/isoprene/NOx | ZJU224 | – | 110 | 120 | 51 | 0.38 | 11.4 | 11.3 | 1.63 | 0.55 | 1.88 |
| p-xylene/propene/NOx | ZJU230 | – | 110 | – | 52 | 0.37 | 10.8 | 11.1 | 1.61 | 0.49 | 1.79 |

All experiments were carried out at ~35 °C, in dry conditions (RH < 10%) and in the absence of seed particles; the initial VOC concentrations were measured and comprised over 95% NOx; AMS measurements were not available for ZJU095/098/178; the elemental ratios of H/C (hydrogen to carbon), O/C (oxygen to carbon) and OM/OC (organic matter to organic carbon) were calculated using the Improved-Ambient (I-A) method\textsuperscript{40}; all AMS-derived parameters including f\(_{43}\) and f\(_{44}\) (fraction of m/z = 43 or 44) were calculated using averaged data of 420–480 min.
The SOA mass yields at the end period (7–8 h) of each experiment were calculated according to the wall-loss corrected SOA mass concentrations and reacted hydrocarbons, with the detailed results summarised in Supplementary Table 1. Note that the reacted amount of VOC (ΔVOC) from most of the experiments was obtained from direct VOC measurements (further scaled by initial concentrations), except for four single-hydrocarbon experiments without VOC measurements where simulated VOC decay from SAPRC-18 was used (see more details of gas-phase modelling in the next section). As for mixed-VOC experiments, two yield values (Y_max and Y_only) were calculated based on alternative assumptions to quantify the aerosol mass yield. Y_max was calculated following the assumption that both reacted aromatics and alkenes contributed to SOA mass, while Y_only was calculated by conservatively assuming that only aromatics contributed to SOA mass. For single-hydrocarbon experiments, Y_only alone was calculated because only one parent VOC was in the system.

In the single-hydrocarbon experiments, the average mass yield of toluene SOA was ~22%; while the isoprene SOA yield was 0.78–1.18%; these were both within typical ranges in previous studies. We found that the addition of isoprene or propene into toluene/NO_x could significantly lower the Y_max (by at least ~50% depending on conditions), particularly under high isoprene conditions (ZJU164). This was consistent with the fact that propene is usually not considered to be an SOA precursor and the isoprene SOA yield was also relatively small (0.78–1.18% in this work). Even if it was assumed that toluene was the only contributor to SOA mass in these mixed-VOC experiments, the
SAPRC-11 and SAPRC-18 provided reasonable predictions for O3 found in the toluene-only experiments. Similar results were also found in the p-xylene experiments, where the addition of isoprene or propene clearly lowered the SOA mass yield regardless of whether $Y_{mix}$ or $Y_{only}$ was calculated.

**Modelling gas-phase chemistry and SOA formation**

Detailed photochemical modelling was performed to further investigate the reasons behind the observed SOA suppression. Considering that gas-phase chemistry initiates SOA formation, simulations were conducted using five well-established gas-phase chemical mechanisms i.e., SAPRC-07/11/18, CB6 and MCMv3.3.1. As for single-hydrocarbon experiments, Supplementary Figs. 1 and 2 show that almost all the mechanisms were able to explain the measured data to some extent, though some differences among the five mechanisms were observed. This provided a good baseline for mechanistic modelling of more complex conditions. More details about the mechanism performance for the single-hydrocarbon experiments are provided in the Supplementary Discussion.

Here, we focus on the gas-phase results from mixed-VOC experiments (Fig. 2). For toluene/isoprene/NOx experiments, both SAPRC-11 and SAPRC-18 provided reasonable predictions for O3 formation, NO oxidation and hydrocarbon decay, while the other three mechanisms slightly underestimated the O3 formation rate and NO oxidation, particularly under lower isoprene condition (ZJU183). All the mechanisms provided reasonable predictions for the toluene/propene/NOx experiment (ZJU176), though CB6 slightly underestimated the O3 formation. As for the two p-xylene experiments with the addition of isoprene (ZJU224) or propene (ZJU230), the general trends and levels of gas-phase species were correctly captured by SAPRC-11, SAPRC-18 and MCMv3.3.1, though all the mechanisms showed small discrepancies.

Overall, all the chemical mechanisms were able to predict the gas-phase processing under single or mixed-VOC conditions, particularly the decay process of hydrocarbons under mixed-VOC conditions. Among the five chemical mechanisms, SAPRC-18 modelling results seem to show better agreement with experimental data probably because it considers latest advancement of chemical mechanism development. The simulation results showed that a lower amount of toluene or p-xylene reacted after the addition of isoprene or propene, consistent with the experimental data (Fig. 2 and Supplementary Fig. 2). This is attributed to OH competition by the highly reactive alkene species, consequently resulting in lower OH levels (Supplementary Fig. 3). The lower consumption of toluene or p-xylene due to the addition of isoprene or propene provides a qualitative explanation for the suppression of SOA mass concentration. As indicated by Song et al. that the SOA from m-xylene/NOx photo-oxidation is strongly dependent on OH levels in the system, where the addition of propene can decrease OH and slow down the reaction rate of m-xylene. Jaoui et al. also found that the addition of...
isoprene to complex mixture conditions (α-pine/toluene/NOx) can decrease the amount of reacted toluene and consequently lower the SOA formed.

It has been recognised that the gas-phase autooxidation of peroxy radicals (RO2) can produce highly oxygenated organic molecules (HOMs), which can further contribute to new particle formation and SOA formation43,44. More recently, Heinritzi et al.25 found that the addition of isoprene to the α-pine oxidation system suppresses nucleation and growth, because the RO2 radicals derived from isoprene and α-pine are competitive and result in reduced C19–20 dimers and increased C14–15 dimers. Thus, such RO2 termination processes observed in the isoprene and α-pine mixtures are considered as a critical step that determines the HOMs distribution and subsequent nucleation and growth of SOA. Similar RO2 termination processes are expected to exist in our complex interactions between the degradation chemistry for alkenes and aromatics. Here we further explore the possible connection between SOA formation and RO2 fate under our mixed-VOC conditions. The RO2 removal rates (RO2 + NO and RO2 + HO2 pathways) and concentrations predicted by the MCMv3.3.1 were summarised in Supplementary Fig. 4. For all the experiments in general, the removal pathway of RO2 was overwhelmingly dominated by RO2 + NO in the early period, and gradually shifted to RO2 + HO2 once the NO levels approached ~0. Compared to the base conditions (toluene/NOx or p-xylene/NOx system), the RO2 removal was delayed due to the addition of isoprene, while the addition of propene resulted in an earlier RO2 removal. Such variation of RO2 fate was consistent with the normalised NO decay signal, and may explain the delayed SOA removal. Such variation of RO2 fate was consistent with the connection between SOA formation and RO2 fate under our conditions of these experiments, although the chemical systems in our chamber experiments were simplified compared to the complexity of real atmospheric conditions.

### SOA chemical composition from AMS measurements

The evolution trends of SOA chemical composition are further investigated in mass spectra (see Supplementary Discussion), Van Krevelen diagram (O/C vs. H/C space) and triangle plots (f43 vs. f44 space; f43 and f44 are AMS organic signal fraction at m/z = 43 and 44, respectively) from AMS measurements. Figure 4a shows the changes in H/C and O/C ratios as a function of OH exposure for each experiment in Van Krevelen diagram. This space has been widely used to interpret the possible involved reactions and addition of functional groups during SOA formation46–48, according to the slopes of the straight lines (the slopes of k = 0, −0.5, −1, and −2 are shown in dash lines in Fig. 4a). Specifically, the replacement of an aliphatic carbon (CH2) with a carbonyl group (C(=O)) would result in a slope of −2 in the Van Krevelen diagram, suggesting the ketones or aldehydes formation in the SOA. By contrast, the replacement of a hydrogen with an alcohol group (OH) would only lead to O/C increase, as shown by a slope of 0 in the O/C vs. H/C space. Similarly, the addition of functional group like carboxylic acid (C(=O)OH) during SOA formation would result in a slope of −1, while carboxylic acid formed in the SOA by fragmentation would result in a slope of −0.5 in the evolution of H/C and O/C ratios48. The triangle plots (f43 vs. f44 space) in Fig. 4b can also indicate similar information of SOA evolution processes with less noisy data than those of elemental ratios49,50. Note that the AMS ion signals of m/z 43 and 44 primarily originate from carboxyls and carboxylic acids respectively, with dominant oxygenated fragment ions of C3H5O+ (at m/z = 43) and CO2+ (at m/z = 44)49,51,52. As shown in Fig. 4, most of the experiments showed clear evolving trends in both Van Krevelen space and triangle plots as measured vs. predicted SOA for all chamber experiments. The experiments were divided into a toluene and b p-xylene systems. The measured SOA mass concentrations were from SMPS measurements (with wall-loss correction) over 7–8 h. The two-product fitted yield curves (see Supplementary Fig. 5) of specific chamber experiments were derived from their respective toluene/NOx, isoprene/NOx, or p-xylene/NOx, and modelled VOCs from SAPRC-18 were used to derive the reacted amount of VOC (ΔVOC) for these experiments.

**Fig. 3** Measured vs. predicted SOA for all chamber experiments. The experiments were divided into a toluene and b p-xylene systems. The measured SOA mass concentrations were from SMPS measurements (with wall-loss correction) over 7–8 h. The two-product fitted yield curves (see Supplementary Fig. 5) of specific chamber experiments were derived from their respective toluene/NOx, isoprene/NOx, or p-xylene/NOx experiments; these were used for SOA prediction, particularly for toluene/NOx, p-xylene/NOx, and isoprene/NOx, ZJU224 that showed some bias outside of the 1:1 line. This suggested that the suppression of SOA mass concentrations observed in the mixed-VOC experiments could be attributed to the decreased consumption of the parent hydrocarbons alone. As discussed earlier, established gas-phase chemistry can account for the OH scavenging effect upon SOA yields induced by isoprene or propene in our mixed-VOC experiments, associated with lower consumption of the parent aromatic hydrocarbons. Gas-phase chemistry and yield-based approaches can provide an overall explanation of SOA formation suppression under the conditions of these experiments, although the chemical systems in our chamber experiments were simplified compared to the complexity of real atmospheric conditions.
the OH exposure increased. As for toluene/NOx base experiments (ZJU104/130), the H/C and O/C ratios of SOA fall along the slope around \(-0.5\), while \(f_{44}\) remained mostly unchanged as \(f_{44}\) increased in the triangle plots. This suggests that the chemical evolution of toluene SOA (ZJU104/130) is probably dominated by the formation of carboxylic acid, particularly through the fragmentation process. By contrast, the addition of isoprene or propene into toluene/NOx base condition leads to some changes of SOA evolving trends. For example, adding low isoprene of \(\sim 38\) ppbv (ZJU183) showed both increase of \(f_{33}\) and \(f_{44}\) in the SOA evolution process, suggesting the simultaneous formation of carbonyl compounds and carboxylic acids in this case. ZJU183 further showed a trajectory along the slope of \(-0.81\) in Van Krevelen space, consistent with the above deduction that carbonyl compounds are formed in addition to carboxylic acids. On the other hand, adding high isoprene of \(\sim 165\) ppbv (ZJU164) did not show a clear evolving trajectory, with the H/C and O/C ratios distributing in a narrow range. This case is similar to the pattern of isoprene/NOx (ZJU189), as both experiments have similar level of initial isoprene and relatively low SOA loadings. The addition of \(\sim 152\) ppbv propene (ZJU176) showed a trajectory of H/C and O/C ratios along the slope around \(-1\), while \(f_{33}\) remained mostly unchanged as \(f_{44}\) increased in the triangle plots. This suggests that the formation route of carboxylic acid in ZJU176 may shift to some extent from fragmentation to functionalization due to the addition of propene. Besides, both \(f_{44}\) and O/C represent the oxidation state of SOA. The addition of isoprene or propene both reduced the magnitude of the increase in \(f_{44}\) and O/C during an experiment. Table 1 further summarises the typical chemical parameters of SOA at the end of each experiment. A decrease in \(f_{33}\) and O/C, as well as an increase in H/C were observed due to the addition of isoprene or propene, which agreed with the results shown in Van Krevelen space in Fig. 4a. Such phenomena indicated that the oxidation state of SOA composition was reduced under mixed-VOC conditions, probably because of (1) reduced OH exposure due to the OH scavenging effect and (2) the aforementioned changes in the oxidation pathways and chemical composition of SOA.

Compared to the p-xylene/NOx base condition (ZJU220/232), the addition of isoprene (ZJU224) or propene (ZJU230) leads to a more discrete pattern in the evolving trend in the H/C vs. O/C space in Fig. 4a. This phenomenon is similar to the p-xylene/NOx base experiments, p-xylene/NOx base experiments (ZJU220/232) showed increase of both \(f_{33}\) and \(f_{44}\) in the SOA evolution process, suggesting the simultaneous formation of carbonyl compounds and carboxylic acids in p-xylene case. As observed in the \(f_{33}\) vs. \(f_{44}\) space in Fig. 4b, the addition of isoprene (ZJU224) resulted in less increase of \(f_{44}\) as \(f_{33}\) increased, indicating relatively more carbonyl compounds formation due to the addition of isoprene. This is consistent with the results from toluene/isoprene/NOx experiment of low isoprene concentration (ZJU183) in the earlier discussion, where more carbonyl formation was evidenced by the changed slope in Van Krevelen space and increased \(f_{33}\) trends in triangle plots. The addition of propene (ZJU230) showed a changed trajectory in the low OH exposure in
triangle plots, and reason for this phenomenon is still not clear and requires further investigation. Overall, we find that the addition of isoprene or propene could not only suppress the aromatic SOA mass and yield, but also change the oxidation state and chemical composition of SOA. The addition of isoprene into aromatic/NO₂ photo-oxidation may reduce the magnitude of oxidation state increase during the experiment and result in SOA formation with more carbonyl compounds, though these are qualitative and the extent of such changes could depend on the addition amount of isoprene. The current AMS analysis also indicates the complexity of SOA composition under such mixed-VOC conditions, which implies that a detailed molecular characterisation of SOA composition is required in future studies.

Atmospheric implications

Aromatics are well-known as an important class of anthropogenic VOCs from vehicle and industrial emissions, and many previous chamber studies⁵¹,⁵² have confirmed their high SOA yields in the presence of NO₂ and sunlight. Therefore, it is expected that aromatic hydrocarbons can contribute to urban fine particles through photochemical processes. The present study shows that the aromatic SOA mass yields were suppressed by the addition of isoprene or propene, which could be attributed to the OH scavenging effect that lowers consumption of the parent aromatics. Considering the reasonably good performance of the five different chemical mechanisms (SAPRC-07/11/18, CB6, and MCMv3.3.1) in simulating our experiments, we further demonstrated that the suppression of aromatic SOA could be quantitatively explained by the current gas-phase chemistry and yield-based SOA predictions. Such OH scavenging effect proposed by McFiggans et al.²⁵ could be widespread in binary or more complex mixture conditions. Our current experiments are performed under high-NO₂ conditions that are representative of the daytime photochemistry, and the OH production in such reaction system is limited due to the radical recycles (e.g., HO₂ + NO → OH + NO₂). However, it still requires more investigations to examine whether the OH scavenging effect holds in reaction systems with surplus OH (e.g., low-NO₂ conditions), where adding isoprene may have limited influences on the OH consumption. Note that the product scavenging effect (removal of oxidation products with high SOA-formation potential) further proposed by McFiggans et al.²⁵ might also exist in our mixed-VOC conditions; this could further prevent the multi-generation oxidation process, as evidenced by the decreased O/C of SOA due to the addition of isoprene in our toluene experiments. Unfortunately, due to limitations over measurements of oxidised products, this cannot be examined in greater depth in the present study.

The findings of this study imply that the global budget of anthropogenic SOA might be significantly affected by biogenic emissions such as isoprene, particularly in some urban environments with large coverings of land vegetation where both anthropogenic and biogenic emissions are mixed. More broadly, this study further revealed the critical importance of the interaction between aromatics and short-chained alkenes on SOA formation: combining aromatics (higher SOA yield and lower OH rate constant (k_H₂O)) and alkenes (lower SOA yield and higher k_NO₂) in the presence of NO₂ and sunlight could result in a lower SOA mass. Accounting for such interactive effects can provide new insights for improving the SOA module of chemical transport models, which will allow us to achieve an improved understanding of SOA sources under real-world conditions. Additionally, anthropogenic SOA is well-recognised as an important component for evaluating the impact of human activities on climate change. If the suppression by isoprene on the global budget of anthropogenic SOA is considered and parameterised in current climate models, the corresponding magnitude of the impact of human activities on radiative forcing might also change. Given that both anthropogenic and biogenic VOC emissions will continue to change over time, it is also expected that the aerosol-climate interaction will present a highly dynamic system in the future; this highlights the importance of better understanding SOA formation derived from VOC mixtures of different sources.

The current AMS analysis also showed some interesting and highly variable phenomena related to the chemical characteristics of SOA, particularly the changed oxidation state (i.e., O/C) and oxidation pathways (i.e., more carbonyls formation) under mixed-VOC conditions, which are still not fully understood at present. This implies that SOA formation under mixed-VOC conditions likely undergoes highly complex chemical processes between the degradation chemistry for the alkenes and aromatics. However, parameterisation of such interactive effect in models requires additional work, and it relies on the ability of modelling the transformation from gas-phase species to SOA with fundamental understanding and representation rather than adjustments to fit chamber data. In addition to the daytime photochemistry, nighttime NO₃ oxidation can also be a dominant pathway of SOA under certain circumstances, and SOA formation from VOC mixtures through NO₂ oxidation is relevant to the real atmosphere but not yet understood. We suggest that more comprehensive mixed-VOC experiments with high quality data and molecular characterisation of SOA composition should be conducted to further explore the interaction between anthropogenic and biogenic VOC mixtures and their combined effect on SOA formation, which should cover a wide range of experimental conditions that are more representative of real-world conditions.

METHODS

Experiment summary

The CAPS-ZJU (Complex Air Pollution Study-Zhejiang University) smog chamber is a 3 m³ Teflon-FEP chamber housed in a well-controlled enclosure, with 60 controlled ultraviolet (UV) lamps providing the light source. A total of 15 smog chamber experiments across a broad range of scenarios were performed to support the objectives of this study, with the initial concentrations of VOCs and NO₂ in each experiment given in Table 1. Specifically, toluene/NO₂ and p-xylene/NO₂ experiments were used as two base conditions, as toluene and p-xylene are well recognised as typical precursors of anthropogenic SOA. Isoprene was the major focus of this study, and three experiments (ZJU164/183/224) were conducted with different levels of isoprene added into the chamber. The main aim of this study was to address the main issue of how isoprene affects anthropogenic SOA formation. In addition, since propene is a typical reactive hydrocarbon and usually not regarded as an SOA precursor, two experiments (ZJU176/230) were performed by adding propene into the base conditions instead of isoprene, assisting explain the isoprene addition case considering some chemical similarities between the two short-chained reactive alkenes. The abbreviations ‘tol’, ‘p-xy1’, ‘iso’, and ‘prop’ for toluene, p-xylene, isoprene, and propene, respectively, are used elsewhere. Note that some parallel single-hydrocarbon experiments were conducted under similar conditions to confirm the reproducibility of the results. Each experiment was performed in a 7–8 h photochemical window at equivalent light intensity (Uₑq) of 0.36–0.40 min⁻¹. All the experiments were conducted under dry conditions (relative humidity (RH) < 10%) and in the absence of seed aerosol. We choose low RH (< 10%) in our chamber experiments to avoid and exclude the complex influence (e.g., multiphase chemistry) under high humidity condition. Upon turning on the UV lights, the temperature inside the chamber increased from ~20 to ~35 °C within 2 h due to the heat released from the lights, and then stabilised at ~35 °C until the end of the experiment. All experiments had similar temperature profiles, which ensured that the experiments were comparable and repeatable. The background air was supplied by a Zero Air generator (Aadco 737-15, USA). After each experiment, the chamber was continually flushed with clean air for at least 12 h to ensure the particle number concentration below 10 cm⁻³ and the reactive gas concentration below 1 ppbv. More details of the chamber characterisation and experimental methods can be found in our recent publication.⁶⁴
Instrumentation and analysis
The CAPS-ZJU chamber facility was equipped with a suite of state-of-the-art instruments to characterise the chemical composition, mass concentration, size distribution, effective density, gas concentration, and other physicochemical properties of particles. Specifically, the NO2 and O3 concentrations were measured using commercial gas analysers (Thermo Scientific 42i and 49i, USA). The gaseous hydrocarbons were measured by a gas chromatography-mass spectrometer (GC-MS; Agilent 7890B–5977A, USA) coupled with an automated pre-concentrator (Entech 7100A, USA). The particle size distribution (14–700 nm) was determined by a scanning mobility particle sizer (SMPS, TSI 3936) with a long differential mobility analyser (DIMA, TSI 3081) in combination with a butanol condensation particle counter (CPC, TSI 3776). An aerosol particle mass analyser (APM, Kanomax 3601) was used to measure particle effective density by coupling the DMA, APM, and CPC. As summarised in Supplementary Table 2, the effective density of SOA measured by DMA-APM-CPC ranged from 1.27 to 1.36 g cm⁻³ depending on the reaction system, which was also close to the density range of typical SOA reported in previous literature. Then, the aerosol mass concentration was derived from the particle number size distribution. Size-dependent aerosol wall-loss correction was performed to account for the particles deposited onto the walls, following the method applied in our previous work.

The composition of SOA was determined by a high-resolution time of flight aerosol mass spectrometry (HR-ToF-AMS, hereafter AMS; Aerodyne Research Inc., USA), with method described in detail in our previous publications. In brief, the AMS was operated in the high sensitivity V-mode and the high resolution W-mode every 2 min, alternately. The AMS data was analysed by SQUIRREL v1.57 and PIKA v1.16 software written into Igor Pro 6.3.7.2 (WaveMetrics Inc., USA). The elemental ratios (i.e., H/C and O/C) of SOA were determined using the Improved-Ambient (IA) method developed by Canagaratna et al. More details of AMS operation and analysis were described in the Supplementary Method.

Gas-phase chemistry and SOA modelling
The gas-phase chemistry box models used for this study are outlined in Supplementary Table 3. SAPRC and CB (carbon bond) mechanisms are widely used as core components for chemical transport models. The specific versions used in this study were SAPRC-07, SAPRC-11, SAPRC-18 and CB6, and have been applied in our recent work. SAPRC-18 is the latest version of SAPRC mechanism (release in May 2020, see https://intra.cert.ucr.edu/~carter/SAPRC/18/), which has a significant update with alkene chemistry and considers the H-shift isomerizations of peroxy radicals. Notably, all the deposited series mechanisms (SAPRC-07/11/18) used in this study were detailed versions with explicit representation of individual VOC degradation chemistry. There are simplified versions of SAPRC by lumping the VOC species for chemical transport modelling, while the detailed versions of SAPRC are widely used for mechanism evaluation utilising environmental chamber data; more details are available at http://www.cert.ucr.edu/~carter/SAPRC/. CB6 was developed as a lumped mechanism for simplicity, with the least chemical detail among the five mechanisms. The Master Chemical Mechanism (MCMv3.3.1) is a near-explicit mechanism with more detailed reactions (http://mcm.leeds.ac.uk/MCM/). The smog chamber experiments were simulated using the above five mechanisms with the additional CAPS-ZJU chamber-specific auxiliary wall-mechanism, which has been characterised in detail in our recent publication.

The SOA mass yield is defined as the mass of SOA produced per unit of hydrocarbon reacted, which has been widely used to predict SOA formation. In this study, yield curves were obtained from the single-hydrocarbon experiments by fitting the time-dependent SOA mass yield to aerosol mass concentration, which could be used to predict SOA formed by toluene, isoprene, and p-xylene, respectively. Combined with the measured VOC decay, these specific-fitted yield curves were further used to predict SOA formation under mixed-VOC conditions.

DATA AVAILABILITY
The datasets generated during and/or analysed during the current study are available from the corresponding author (likiangweijp@foxmail.com) on reasonable request.

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AUTHOR CONTRIBUTIONS
KL conceived the study; KL, X.Z, and C.L performed experiments; KL performed the modelling; KL, X.Z., and B.Z. analysed data; KL, B.Z., and Z.B. wrote the paper; W.J.B, S.W., H.Y., L.C., C.Geng, W.Y., A.M., and C.George helped to interpret the results and revised the manuscript. All authors contributed to the manuscript preparation and discussions.

COMPETING INTERESTS
The authors declare no competing interests.

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