

1. Review #1

Summary: Yun Lin and co-workers have used computational methods to study clusters containing sulfuric acid (SA), ammonia (AM), dimethylamine (DMA), succinic acid (SUA) and water. The computational methods used are adequate for the purpose (if a bit outdated in terms of the DFT method used to obtain the thermodynamics - the sampling approach on the other hand is state-of-the-art). The purpose of the study is to assess the possible atmospheric significance of succinic acid in promoting sulfuric acid - based new-particle formation (and indeed a positive conclusion is implied even in the title). The study provides important new insights and information onto e.g. the hydration behaviour of SA-AM-DMA-SUA clusters, and especially on how this changes with the presence of SUA. Some of the conclusions concerning the effect (or lack of C1 ACPD Interactive comment Printer-friendly version Discussion paper effect) of SUA on new-particle formation are somewhat premature, and not always fully supported (indeed, sometimes flat-out contradicted) by the actual data. While the manuscript is publishable, extensive rewriting and reformulation is thus necessary prior to publication in ACP.

Thanks the reviewer for insightful comments.

Major comments:

-Comment #1. Overall, the authors attach way too much importance to the sign of the standard Gibbs free energy change, “delta-G”. (This is unfortunately very common in similar studies.) The free energies are computed using a 1 atm reference pressure (values computed with other reference pressures would be very different for reactions where the number of molecules changes, as in all the clustering reactions studied here). The concentrations of the studied reactants are (except for water), far far below 1 atm, as the authors themselves note. In other words, the reference pressure is essentially (for these reactions) an arbitrary value that has very little to do with the particular reactions being studied. A negative “delta-G” value for a clustering reaction is a necessary, but very far from sufficient, criterion for the cluster to be said to be “stable”. Consider for example a reaction of the type $X + Y \rightleftharpoons XY$ where X is some core cluster, and Y is a molecule present at ppt - levels (i.e. a partial pressure of $1E-12$ atm, typical for e.g. SA, DMA or SUA in this study). If the delta-G for this reaction is for example -3 kcal/mol, we can use the law of mass balance to write (here R is the gas constant, $T=298K$ is the temperature, e.g. p_{XY} is the partial pressure of XY, and p_{ref} is the reference pressure at which delta-G is computed, in this study 1 atm):

$$\frac{p_{XY}/p_{ref}}{(p_X/p_{ref})(p_Y/p_{ref})} = \exp(-\Delta G/RT)$$

We can solve for the ratio (p_{XY}/p_X) , and get $(p_Y/p_{ref}) \exp(-\Delta G/RT)$. Since p_Y/p_{ref} is $1E-12$, and $\exp(-\Delta G/RT)$ is (for a delta-G of about -3 kcal/mol) about 160, we get a value of about $1.6E-10$ for the ratio. In other words less than one in a billion of the molecules of type Y will be bound to XY clusters at equilibrium, despite a negative delta-G value. Clearly, XY is not “stable” in any meaningful sense of the word, C2 and Y does not “stabilise” X, despite the standard free energy of addition of Y to X being negative. Thus, sentences such as (line 285-286) “All free energy changes shown in Figure 5b are negative, confirming that SUA stabilizes the SA⁺ ubase clusters” are absolutely false: negative free energy changes do not in any way confirm “stabilisation”. (Also, even when a molecule Y is very strongly bound to a cluster X, this does not in and of itself prove “stabilisation” of the cluster as a whole: it just proves that it is difficult for molecule Y to evaporate. Other constituents of the cluster may evaporate either more or less easily

due to the presence of Y, and this needs to be separately evaluated - just looking at the Y to X binding is not sufficient to claim “stabilization”). I recommend that the authors search the manuscript for all occurrences of the word “stable” in any form (verb or adjective), and rigorously consider whether or not its use is justified. My guess is that almost none of the claims of “stability” or “stabilisation” are actually really justified (apart from perhaps the discussion on hydration, where the effect of H₂O concentration is properly accounted for) - the binding of SUA to the clusters tends to be quite weak, with the equilibrium strongly on the side of the reactants even for the highest SUA concentrations claimed by the authors. (See Elm et al., 2017, also cited in the manuscript, for a discussion on what is actually required for a cluster to be “stable” given trace-gas concentration levels of the constituent monomers: this typically means ΔG values far below -10 kcal/mol - which is not reached for any of the SUA addition steps in the manuscript.)

Thanks the reviewer for the clarification of the relationship between cluster stability and ΔG . We agree that a negative value of Gibbs free energy is not sufficient to indicate the stability of a cluster. However, the free energy can imply whether the clustering reaction is favorable or not. Hence, although as reviewer 1 stated, XY is not stable, but a more negative free energy of XY implies the preferable the formation of XY. It means that XY has a high possibility to further react or combine with other species. This is why we used and compared the free energy to judge whether the binding ability of SUA or other precursors.

Furthermore, as the reviewer 1 pointed out, the probability that a core cluster and a precursor molecule can bond together is determined by both the free energy and the precursor atmospheric concentration. Therefore, we examined the possibilities of the core molecule (i.e., SA)/clusters (i.e., SA·base) bonding to SUA under real atmospheric condition (i.e., the ratios for core molecule/clusters bonding to SUA) and how are these bonding ratios for SUA compared to the case for other precursors. To get bonding ratios, first, we have carefully reviewed literatures for reliable atmospheric concentrations of the four precursors studied in this work (i.e., SA, SUA, DMA, AM), and their corresponding concentrations in real atmosphere are listed in Table 3 (please refer to the answer to comment #3 for details). Second, with these reliable atmospheric concentrations, we have calculated the bonding ratios associated with several primary clustering reactions, particularly for SA/SA·base bonding to additional SA or SUA, according to the reviewer 1 illustration. The results are presented in Table 6. Also reported in Table 6 are the interaction energies at 0K (ΔH_0), which are indicators of the strengths of hydrogen bonds for individual cluster system.

Based on Table 6, we did some comparisons of thermochemical properties and cluster concentrations at equilibrium between clustering with SUA and with other precursors. We find that, among the clustering reactions between SA and the four precursors (i.e., SA, AM, DMA, and SUA), SA clustering with SUA shows the second lowest free energy (just following SA clustering with DMA). Also, the hydrogen bonding in SA•SUA cluster is as strong as that in SA•DMA and is higher than SA•AM and (SA)₂ clusters. Under real atmosphere, SA mostly likely binds to DMA since the bonding ratio for SA to DMA is highest. But if we are looking at the case of SUA to AM, we can see that equilibrium cluster concentration for SA•SUA is almost at the same order as the low bound of that for SA•AM, although that the atmospheric concentration of ammonium is about three to four orders of magnitude of SUA. The chance for SA bonding to SUA is much higher than that for two SA molecules bonding together. When SA·base further clustering with SUA or SA, we can see that the possibility of SA·base bonding to SUA are actually comparable to that bonding to SA. Our calculations of bonding ratios suggest that SUA might be a competitive candidate and

compete with SA to participate in further clustering of SA•base system under real atmospheric condition.

Finally, we summarize these findings and added some discussions to section “3.4 Atmospheric Implication”. We also added a description about bonding ratio calculations into section 2. “Computational Methods”. Followed we also carefully examined and modified all the statements containing “stable”, “stability”, and “stabilization” in the main text.

Lines 171-175

“In addition, the cluster concentration, [cluster], for addition of another monomer to the SA•base dimer with or without SUA is estimated, by considering the atmospheric concentrations of the various precursors,

$$[cluster] = [SA] \times [X] e^{\left(\frac{-\Delta G}{RT}\right)} \quad (9)$$

where the precursor species X corresponds to AM, DMA, or SUA.”

Lines 369-386:

“In addition, the relative concentrations of the precursor species involved in clustering also govern the cluster distribution in the atmosphere. The estimated cluster concentrations using equation (9) show a cluster concentration (i.e., $10^{-3} \sim 10^2 \text{ cm}^{-3}$) for SUA•SA•DMA (Table 6), suggesting that SUA likely contributes to the further growth of SA•base clusters. The calculated ratios of $[SA\cdot X\cdot SUA]/[(SA)_2\cdot X]$ (X denotes AM, DMA, water molecule, or none) under atmospherically relevant concentrations are presented in Table 7. The $(SA)_2\cdot AM$ cluster dominates the cluster distribution when SA and SUA concentrations are similar, i.e., $[SA\cdot AM\cdot SUA]/[(SA)_2\cdot AM] = 1:1000$. The estimated cluster ratio of $[SA\cdot DMA\cdot SUA]/[(SA)_2\cdot DMA]$ is in the range from 3:100 to 30:1, indicating that SA•DMA•SUA or $(SA)_2\cdot DMA$ clusters can be prevalent in the atmosphere. The ratios of $[SA\cdot SUA]/[(SA)_2]$ and $[SA\cdot W\cdot SUA]/[(SA)_2\cdot W]$ are $10^5:1$ and $10^4:1$, respectively, and hence the SUA-containing clusters are prevalent for both unhydrated and hydrated SA clusters with one water molecule. While sulfuric acid dimer is believed to be an important precursor for NPF (Zhang et al., 2012), our study shows that SUA, which is one of most abundant dicarboxylic acids in atmosphere, inhibits the formation or further growth of sulfuric acid dimer because of its strong interaction with SA in the unhydrated or hydrated SA clusters. Such an inhibiting effect of SUA on the formation or further growth of sulfuric acid dimer is more efficient than ketodiperoxy acid (Elm et al., 2016b).”

Line 13:

“Dicarboxylic acids are believed to stabilize pre-nucleation clusters and facilitate new particle formation in the atmosphere...” was changed to “Dicarboxylic acids likely participate in the formation of pre-nucleation clusters to facilitate new particle formation in the atmosphere, ...”

Lines 25-27:

“the uptake of SUA competes with the uptake of the second SA molecule to stabilize the SA•base clusters at atmospherically relevant concentrations” was changed to “SUA competes with the second SA molecule to cluster with the SA•base clusters at atmospherically relevant concentrations.”

Lines 249:

We don't have sufficient evidence to prove it and therefore we just removed this statement here: “While the bending of the carbon chain facilitates hydrogen bonding, such bending also induces steric hindrance, which partially cancels out the energy due to additional hydrogen bonding.”

Lines 368-369:

“All free energy changes shown in Figure 5b are negative, confirming that SUA stabilizes the SA·base clusters.” was changed to “The formation of SA•base•SUA by adding a SUA molecule to the SA•base clusters is energetically favorable (Figure 5b), as reflected by large negative free energies.”

Lines 287-289:

“...the free energy changes for the SA·DMA cluster by SUA addition are more negative than that for the SA·AM cluster, suggesting that SUA more efficiently stabilizes the hydrated SA·DMA clusters than the SA·AM cluster” was changed to “...the free energy changes for the SA·DMA cluster by SUA addition are more negative than that for the SA·AM cluster, suggesting that the clustering between SUA and the SA·DMA cluster core is more favorable than the case of SA·AM cluster core.”

Lines 369:

Remove the statement here “SUA is more effective than SA to stabilize the SA·base clusters.”

Lines 381-384:

“Sulfuric acid dimer has been recognized as an important precursor for NPF (Zhang et al., 2012), but our study shows that, as one of most prevalent dicarboxylic acids in atmosphere, SUA inhibits the formation or further growth of sulfuric acid dimer because of its strong interaction with in the unhydrated or hydrated SA clusters.” **was changed to**

“While sulfuric acid dimer is believed to be an important precursor for NPF (Zhang et al., 2012), our study shows that SUA, which is one of most abundant dicarboxylic acids in atmosphere, inhibits the formation or further growth of sulfuric acid dimer because of its strong interaction with SA in the unhydrated or hydrated SA clusters.”

Lines 403-405:

“Addition of SUA to the SA·base cluster systems is energetically favorable at all hydration levels, suggesting that SUA stabilizes the SA·base clusters and their hydrates” was changed to

“Addition of SUA to the SA•base clusters is energetically favorable at all hydration levels, suggesting that SUA likely stabilizes the SA•base clusters and their hydrates..”

Lines 420-422:

“Hence, the uptake of SUA competes with the uptake of another SA to stabilize the SA·base clusters, and the presence of SUA hinders the formation and further growth of SA dimer clusters.” was changed to “Hence, SUA competes with SA for addition to the SA•base clusters, but the presence of SUA hinders further growth of SA dimer clusters.”

Lines 422:

Removed “The hydrated SA·DMA·SUA cluster is capable of binding with additional acid molecules, which not only stabilizes the cluster but also promotes its further growth” since we don't have sufficient evidence.

Table 3. Typical ranges of gas-phase concentrations (molecules cm⁻³) for sulfuric acid, ammonium, dimethylamine, and succinic acid in the atmosphere.

Precursors	Sulfuric acid ^a	Ammonium ^b	Dimethylamine ^c	Succinic acid ^d
number concentration	1x10 ⁵ ~1x10 ⁷	1x10 ⁹ ~1x10 ¹¹	1x10 ⁷ ~1x10 ⁹	1x10 ⁸ ~1x10 ⁹

^a Weber et al. (1999), Nieminen et al. (2009), and Zhang et al. (2012).

^b Seinfeld and Pandis (1998).

^c Kurten et al. (2008).

^d Ho et al. (2007).

1 **Table 6. Gibbs free energy (ΔG , kcal mol⁻¹), interaction energy (ΔH_0 , kcal mol⁻¹), and typical cluster concentration at equilibrium**
 2 **for basic clustering reactions. The right-hand side of clustering reactions is the product clusters in equation (9), and the core**
 3 **clusters and addition molecules in equation (9) are listed here as well.**

Cluster reactions	ΔG	ΔH_0	Cluster		
			Core cluster	Molecule addition	for [Cluster] (cm ⁻³)
SA+SA \leftrightarrow (SA) ₂	-3.72	-13.08	SA	SA	10 ⁻⁷ ~10 ⁻³
SA+SUA \leftrightarrow SA•SUA	-8.61	-17.94	SA	SUA	10 ⁰ ~10 ³
SA+AM \leftrightarrow SA•AM	-6.36	-14.38	SA	AM	10 ⁻¹ ~10 ⁻³
SA+DMA \leftrightarrow SA•DMA	-11.41	-18.38	SA	DMA	10 ¹ ~10 ⁵
SA•SUA+SA \leftrightarrow (SA) ₂ •SUA	-1.02	-11.04	SA•SUA	SA	10 ⁻¹⁴ ~10 ⁻⁹
SA•AM+SA \leftrightarrow (SA) ₂ •AM	-9.46	-19.53	SA•AM	SA	10 ⁻⁹ ~10 ⁻³
SA•AM+SUA \leftrightarrow SA•AM•SUA	-6.20	-16.01	SA•AM	SUA	10 ⁻⁸ ~10 ⁻³
SA•DMA+SA \leftrightarrow (SA) ₂ •DMA	-10.53	-21.16	SA•DMA	SA	10 ⁻⁶ ~10 ⁰
SA•DMA+SUA \leftrightarrow SA•DMA•SUA	-9.86	-19.07	SA•DMA	SUA	10 ⁻³ ~10 ²
(SA) ₂ •DMA+SA \leftrightarrow (SA) ₃ •DMA	-6.10	-15.25	(SA) ₂ •DMA	SA	10 ⁻¹⁶ ~10 ⁻⁸
SA•DMA•SUA+SA \leftrightarrow (SA) ₂ •DMA•SUA	-5.13	-19.07	SA•DMA•SUA	SA	10 ⁻¹⁴ ~10 ⁻⁷

4

-Comment #2.

Related to the previous point, the most interesting part of the data in terms of evaluating the role of SUA on new-particle formation are the energetics for the formation of the clusters shown in Figure 8, i.e. the clusters with two SA molecules. From table 2, it seems that the SA addition free energies to the (SA)(SUA)(DMA)(W)_n clusters vary between +2 and -6 kcal/mol, with the value for the unhydrated cluster being -5.14 kcal/mol. This is significantly HIGHER (more positive) than the free energies of SA addition to the (SA)(DMA) cluster or the (SA)₂DMA cluster (without any SUA), which are about -18 and -9 kcal/mol (respectively) according to the coupled-cluster (DLPNO-CCSD(T)) study of Myllys et al (<https://pubs.acs.org/doi/suppl/10.1021/acs.jpca.5b09762>), and even more negative according to older DFT - or RIC2- based - studies. (In the absence of data at the exact level used by the authors of this study a direct comparison is impossible - the authors might want to compute at least the unhydrated SA(2)DMA and possibly (SA)₃DMA clusters just to check.) Thus, the data presented here indicates - in direct contrast to the manuscript title - that the presence of SUA actually HINDERS sulfuric acid - DMA nucleation rather than enhancing it. I strongly urge the authors to reformulate their title in light of this observation (as well as the comments on stability presented above). Also claims such as SUA “promoting subsequent growth” seem unlikely to be true, if the net effect of SUA is to decrease rather than increase the uptake of further SA molecules.

To justify the point brought up by the reviewer, we did additional computations for unhydrated (SA)₂(DMA) and (SA)₃(DMA) clusters using the same level of theory in this study. Most stable configurations were added into Figure 6 as below. The addition free energy of SA to the (SA)(DMA) cluster or the (SA)₂(DMA) cluster to form (SA)₂(DMA) and (SA)₃(DMA) are -10.5 and -6.1 kcal/mol at the PW91PW91/6-311++G(2d, 2p) level of theory, respectively (see Table 6). These SA addition free energy values are lower than the case of SA addition to the (SA)(SUA)(DMA) (i.e., -5.14 kcal/mol). Therefore, we agree with the reviewer that the presence of SUA actually hinders SA-DMA nucleation rather than enhancing it. As such, we made the revision as follows.

lines 305-315:

“The role of SUA in the subsequent growth of the SA•base clusters was examined by comparing the differences in free energies for adding SA to SA•DMA and SA•DMA•SUA. Computations were also performed for the unhydrated (SA)₂•DMA, (SA)₃•DMA and (SA)₂•DMA•SUA clusters (Table 6). The optimized clusters containing more than one SA molecules are depicted in Figure 6. The free energies of adding SA to SA•DMA and to (SA)₂•DMA are -10.5 and -6.1 kcal mol⁻¹, respectively. The free energy for adding SA to SA•DMA•SUA is -5.14 kcal mol⁻¹, higher than that of SA addition to SA•DMA. However, with hydration (i.e., (SA)₂•DMA•SUA•(W)_x), the free energies for adding SA to SA•DMA•SUA•(W)_x clusters are negative except for the hydration with six water molecules (Table 2), indicating that SA addition to SA•DMA•SUA is also energetically favorable. In addition, addition of SA to SA•DMA•SUA results in formation of strong hydrogen bond (i.e., -19.1 kcal mol⁻¹). Therefore, the clusters containing both the base and organic acid (e.g., SA•DMA•SUA) are capable of further binding with acid molecules to promote the subsequent growth.”

Title:

Based on the update results, we have changed the title to “**Interaction between Dicarboxylic Acid and Sulfuric Acid-Base Clusters**”.

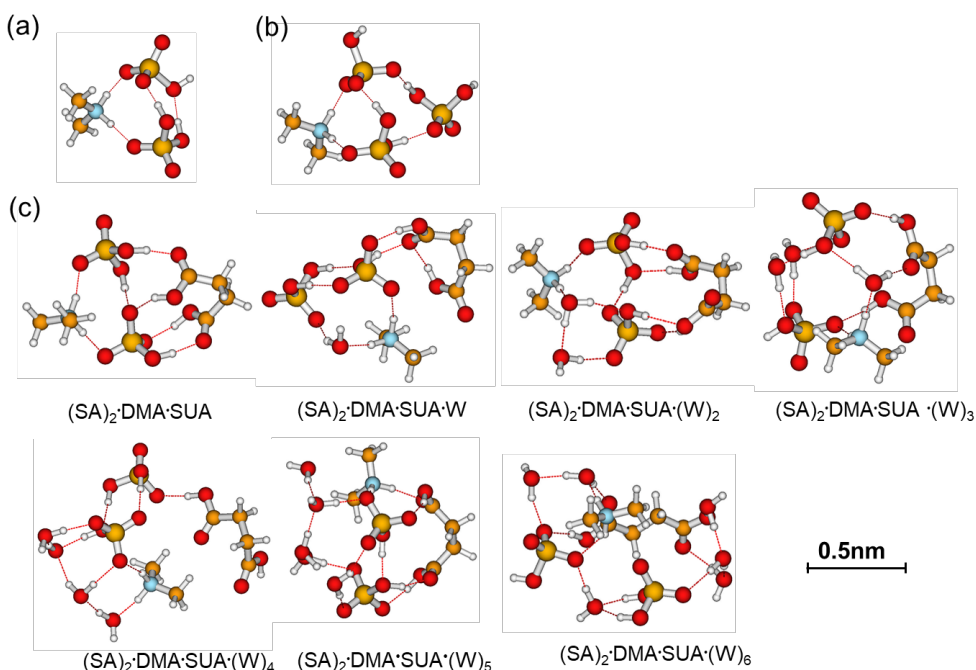


Fig. 6. Most stable configurations of (a) unhydrated $(SA)_2 \cdot DMA$, (b) $(SA)_2 \cdot DMA$, and (c) the hydrated $(SA)_2 \cdot DMA \cdot SUA$ clusters. The hydration is with 0-6 water molecules (W).

-Comment #3. The authors present somewhat misleading concentration ranges and ratios for SUA and SA. The SA value of $1E5$ molecules per cm^3 corresponds to fairly clean conditions - in polluted environments the SA concentration can easily be a couple orders of magnitude higher. The SUA concentration of $1E7$ quoted is from Los Angeles, which I presume corresponds to fairly polluted conditions (with SA certainly exceeding $1E5$!) - in clean conditions the SUA concentration is very likely much lower. The total organic acid concentration range of $1E8$. . $1E9$ quoted is valid, BUT the majority of these will be simple monocarboxylic acids - dicarboxylics such as SUA will only account for a small fraction of the total. Thus, the $[SUA]/[SA]$ range used in the paper (1 to 10 000) is obtained by combining minimum values for SA with maximum values for SUA - in reality, the ratio may well be below one in most places, and I find it hard to believe that values of 10 000 will be found anywhere in the atmosphere. Using the range 1 to 10 000 is fine for answering the question “could SUA possibly play a role in NPF anywhere in the atmosphere, even in the best case?”, but then this should be stated openly, instead of implying that the range used is representative for most areas of the world. Again, together with the two previous issues, this indicates that the statement in the manuscript title is exaggerated at best, and false at worst.

In fact, we did claim in the original text that the value of $1E5$ molecules/ cm^3 is the lower limit of the atmosphere SA, so it does correspond to fairly clean conditions, as reviewer 1 pointed out.

However, as the reviewer suggested, since many studies report that the gas-phase concentration of sulfuric acid in atmosphere typically ranges from $1 \times 10^5 \sim 1 \times 10^7$ molecules cm^{-3} (Weber et al., 1999; Nieminen et al., 2009; Zhang et al., 2012), we decided to use this concentration range in our calculations to represent the typical atmospheric condition of SA. Regarding the SUA concentration, we misused the particle-phase concentration as the gas-phase concentration for SUA as reviewer 2 pointed out, therefore we did additional literature review for a reliable data source. However, there is limited studies reporting gas-phase concentration of SUA in atmosphere (most related studies only report particle-phase concentration since dicarboxylic acids are normally semi-volatile and tend to partition most portion into particle phase). One available study to report gas-phase concentration is by Limbeck et al. [2001], based on which the atmospheric concentration of SUA in gas phase is set as 1×10^7 molecules cm^{-3} (Please refer to the answer to comment #1 of reviewer 2 for details). We have tabulated the typical atmospheric concentrations of four precursors and added it to the manuscript as Table 3.

According to the reviewer 1' comment, with the updated concentrations of SUA and SA, we have adjusted SUA/SA ratio, ranging from 1 to 10,000 to 0.1 to 100, used to compute the concentration ratios between $\text{SUA} \cdot \text{SA} \cdot \text{X}$ and $(\text{SA})_2 \cdot \text{X}$ Clusters, with $\text{X} = \text{W}, \text{AM}, \text{and DMA}$ (Table 6). Of course, in the revised manuscript, the extreme condition still is taken into account (such as the condition with the lowest possible level of sulfuric acid but overestimated level of succinic acid), but it is just for testing the role of SUA on the NPF process, and hence it is believed to be reliable.

Table 7. Concentration Ratios between $\text{SUA} \cdot \text{SA} \cdot \text{X}$ and $(\text{SA})_2 \cdot \text{X}$ Clusters, with $\text{X} = \text{W}, \text{AM}, \text{and DMA}$.

SUA/SA	X=(None)	X=W	X=AM	X=DMA
0.1:1	3.80E+02	5.30E+01	4.11E-04	3.19E-02
1:1	3.80E+03	5.30E+02	4.11E-03	3.19E-01
10:1	3.80E+04	5.30E+03	4.11E-02	3.19E+00
100:1	3.80E+05	5.30E+04	4.11E-01	3.19E+01

With the updated Table 7, modifications on the manuscript include the description in section 2. "Computational Methods" and the related discussions in section "3.4 Atmospheric Implication" as below.

Lines 167-171:

"We considered a concentration of $10^8 \sim 10^9$ molecules cm^{-3} for SUA, according to Ho et al. (2007). For comparison, the concentrations of sulfuric acid, ammonia, and dimethylamine in the atmosphere are typically in the range of $10^5 \sim 10^7$, $10^9 \sim 10^{11}$, $10^7 \sim 10^9$ molecules cm^{-3} (Zhang et al., 2012), as listed in Table 3. Hence, the SUA/SA ratio typically ranges from 0.1 to 100."

Lines 373-381:

"The calculated ratios of $[\text{SA} \cdot \text{X} \cdot \text{SUA}] / [(\text{SA})_2 \cdot \text{X}]$ (X denotes AM, DMA, water molecule, or none) under atmospherically relevant concentrations are presented in Table 7. The $(\text{SA})_2 \cdot \text{AM}$ cluster dominates the cluster distribution when SA and SUA concentrations are similar, i.e., $[\text{SA} \cdot \text{AM} \cdot \text{SUA}] / [(\text{SA})_2 \cdot \text{AM}] = 1:1000$. The estimated cluster ratio of $[\text{SA} \cdot \text{DMA} \cdot \text{SUA}] / [(\text{SA})_2 \cdot \text{DMA}]$ is in the range from 3:100 to 30:1, indicating that $\text{SA} \cdot \text{DMA} \cdot \text{SUA}$ or $(\text{SA})_2 \cdot \text{DMA}$ clusters can be prevalent in the atmosphere. The ratios of $[\text{SA} \cdot \text{SUA}] / [(\text{SA})_2]$ and

$[SA \cdot W \cdot SUA] / [(SA)_2 \cdot W]$ are $10^5:1$ and $10^4:1$, respectively, and hence the SUA-containing clusters are prevalent for both unhydrated and hydrated SA clusters with one water molecule.”

Minor comments:

-Comment #4 In the BPMC conformational sampling, what force field was used for the SA, AM, DMA and SUA molecules? (For water TIP3P was apparently used.) Presumably some variant of the AMBER force field, but this should be stated.

According to the reviewer 1’s suggestion, we added following description in the manuscript:

Lines 104-106:

“We employed the Generalized Amber Force Field (GAFF) for AM, DMA and SUA following Wang et al. (2004; 2006). The force field parameters from Loukonen et al. (2010) were adapted for SUA and bisulfate ion.”

-Comment #5. -In reaction equations (e.g. 4, 5), arrows (going in both directions to indicate a reversible reaction) should be used rather than equals signs

Thanks for the reviewer 1’s comment, done as suggested.

-Comment #6. -On line 218, please specify that the proton transfer occurring upon hydration of (SA)(AM)(DMA)(W₅) is the SECOND proton transfer, forming the sulfate dianion (overall, it would be good to separate discussion of first and second proton transfers).

We have modified the statement as **(Lines 220-222)**

“In our study, proton transfer due to hydration occurs in the monohydrate of SA•AM. A second proton transfer also occurs due to hydration, for example, when SA•AM•DMA•(W)₅ clusters are hydrated with one more water molecule (Figures 1a and 3a).”

-Comment #4. -It’s a bit unclear what’s being plotted in Figure 5a, apparently the free energy of the reaction $X + nH_2O \Rightarrow X(H_2O)_n$ where X is some core cluster? This could be explicitly specified. Or even better, plot the stepwise hydration energies instead, as the actual hydration in the atmosphere likely involves addition (or removal) of single water molecules, not e.g. 5 molecules at the same time. (Since the stepwise energies are then referred to in discussing the hydrate equilibria, it would be better to plot them in the first place.)

The original Fig. 5a shows free energies of hydration of nH_2O , not the stepwise hydration energies. To be clear, we plotted stepwise hydration energies instead as the reviewer suggested. The new figure 5a is as below. Modification was also done in manuscript correspondingly.

Lines 261-263:

“The stepwise hydration free energies for the clusters, along with the number of water molecules contained in the clusters, are presented in Figure 5a. The hydration energies are also provided in Table 2.”

Lines 270:

“Figure 5a shows that the stepwise hydration energies are negative at most hydration degrees...”

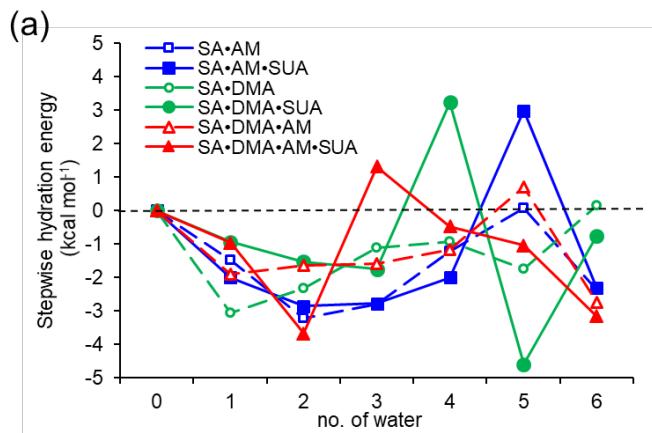


Fig. 5. Stepwise hydration free energies (a) and the relative Gibbs free energy changes due to addition of one SUA molecule to SA•base clusters (b) at T=298.15 K and p=1 atm. The free energy is calculated at the PW91PW91/6-311++G(2d, 2p) level.