

# Reply to comments of Anonymous Referee #1

The authors present a new chemical mechanism (JAMOC), implemented in MECCA. JAMOC includes explicit oxidation steps of organic compounds in the aqueous phase of cloud droplets and thus exceeds previous aqueous phase chemistry mechanisms suitable for box, regional and global modeling. Such extensions are urgently needed as currently, particularly in global models, aqueous phase chemistry modules are largely limited to sulfur(IV) oxidation. However, the current manuscript needs major clarifications and additions to make it a comprehensive and useful extension to previous multiphase model studies. The choice of the newly added reactions is not always evident and also the discussion of the example results in Figure 3 is misleading. In addition, at several places, terminology is confusing or inaccurate. Also previous literature on atmospheric multiphase modeling should be properly discussed. Overall, while I think that this reaction mechanism could possibly become a useful addition to the currently used ones, my comments below need to be carefully addressed prior to possible recommendation for publication.

Thank you very much for the helpful comments and seeing the value of our work to the community. Please find in black the original comments and in red our replies. We significantly extended the introduction, in order to meet your request to discuss previous literature more extensively.

## Major comments

1. Oligomerization has been discussed to be only relevant in the aqueous phase of aerosol particles where organic concentrations may be higher than in cloud droplets (Ervens et al., 2015; Perri et al., 2009; Tan et al., 2009). Thus, it is not clear why they are included in the mechanism. While aqueous phase reactions might also occur in the aqueous phase of aerosol particles, such an extension would also need to include adjustments of Henry's law constants and kinetic reaction rate constants for high ionic strength. Thus, this seems out of the scope of the current study.

We agree with the reviewer that the importance of oligomerization is higher for typical concentrations found in aerosol water (see Tan et al., 2009). At the same time, earlier modelling studies demonstrated that the secondary organic aerosol formation from cloud processing is globally important (Lin et al., 2012). In a future study, we plan to apply JAMOC, including the necessary adjustments, to the aqueous phase of aerosol particles. Thus, we prefer to keep the representation of oligomerization in JAMOC in order to represent clouds as SOA sources.

2. The authors discuss at several places the role of gem-diols and the need of including their phase partitioning between gas and aqueous phases. Is there any indication of the relevance of such processes? Given the very small concentration of water in the gas phase, the stability of gem-diols in the gas phase is likely very small. I assume that their gas phase fraction is likely negligible. Unless the authors can provide literature or estimates on their Henry's law constants that show the opposite, I would not identify the inclusion of their partitioning into atmospheric chemical mechanisms as one of the main gaps in current mechanism developments. In fact, the hydrated glyoxylic acid has been shown to be of such low volatility that it can be involved in new particle formation (Liu et al., 2017)

- Please show the estimated Henry's law constants in Table 1.

- Add references that indicate their potential relevance in the gas phase.

Overall, the dehydration of many gem-diols is slower than the typical lifetime of cloud droplets. Concerning the stability of gem-diols in the gas phase, Kumar et al. (2017) calculated that for the methanediol the shortest lifetime against decomposition by HCOOH-catalysis is larger than  $1E9$  s. We agree with the reviewer that the inclusion of the gem-diol partitioning is not one of the main gaps in current mechanism development. The reason why we discuss this aspect at multiple places is the fact that this aspect clearly separates JAMOC from CLEPS. JAMOC is designed to be used within global models and the transfer of gem-diols into the gas-phase, once the cloud droplet evaporates, is only possible if their partitioning is explicitly represented within JAMOC. Investigating the importance of their gas-phase oxidation is outside the scope of the manuscript. However, we are currently investigating this process on a global scale and

plan further publications with this focus. Therefore, we want to keep the gem-diol partitioning and oxidation in the developed mechanism. In the revised manuscript, we added an explanation of this mechanistic and added a new table summarising all estimated Henry's law constants.

3. At several places in the manuscript, it is not clear whether the authors refer to predictions of aqueous phase rates and budgets or to both phases. For example,

- l. 59: R1 is certainly not a major sink in the atmosphere, but only in the aqueous phase

This is indeed correct. In the revised manuscript, it is now explicitly stated that R1 is only the major sink in the aqueous-phase.

- l. 190: is the explicit oxidation of OVOCs only added in the aqueous or also in the gas phase?

Within the development of JAMOC, only the oxidations of the gem-diols and oxalic acid are added to the gas-phase mechanism. For all other OVOCs, the gas-phase oxidation is represented by the Mainz Organic Mechanism (MOM, Sander et al., 2019). In the revised manuscript, this now reads as "With the explicit oxidation of many OVOCs in the aqueous-phase [...]"

4. At several places in the manuscript, misleading or wrong terminology is used, e.g.

- l. 38: 'nitrogen trioxide' is usually referred to as 'nitrate radical'

We changed it to nitrate radical.

- l. 50: 'simulating hydration and dehydration explicitly' implies that the hydration reaction and dehydration reactions are implemented. However, it seems that only the equilibria are included, separately from the gas/aqueous phase partitioning described by Henry's law.

Even though we list only equilibrium constants in Table 1, the model does indeed calculate hydration and dehydration explicitly. The forward and backward rate constants can be found in the supplement in the file aqueous.eqn.

- l. 134: Do you mean 'recombination of alkyl radicals', i.e. the self-reaction of two radicals?

Exactly. For clarification this now reads "Thus, oligomers formed from the self- and cross-reactions of alkyl radicals are not considered in JAMOC." in the revised manuscript.

5. The model studies are performed for a period of 5 days. However, the typical lifetime of a cloud droplet is on the order of 30 min or less (i.e. the time a droplet spends between cloud base and top in up-and downdraft regimes). You should at least mention that the model simulations are highly idealized and should be regarded as a sensitivity study rather than a realistic scenario.

Indeed, cloud droplets are short lived and the simulations presented in this study are highly idealised. Exactly for this reason, we present a "realistic" cloud event box-model study (i.e. cloud droplet lifetime of 1 hour) in our companion paper by Rosanka et al. (2020). Here, we used the highly idealised CAABA simulations to perform a sensitivity study using JAMOC. We now state the highly idealised nature of this simulations and point to Rosanka et al. (2020) for the realistic cloud scenario simulations using CAABA and EMAC.

6. The authors imply that the additional OVOC oxidation in the aqueous leads to the significant decrease in predicted OH concentration. However, the main reason why significantly less OH is observed in the presence of clouds is the decrease in its formation rate in the gas phase from the reaction of NO + HO<sub>2</sub>. These reactants are separated due to their significantly different water-solubility, e.g. (Jacob, 1986; Lelieveld and Crutzen, 1990). Consequently, the lower HO<sub>2</sub> and higher NO levels in the presence of clouds are not due to the different water solubilities of OH and HO<sub>2</sub> but because of the differences in the gas phase photochemical cycles of HO<sub>x</sub> and NO<sub>x</sub>. This should be discussed and analyzed in Section 3.

Thank you for pointing this out. We reformulated the section significantly.

7. Given that currently only a small fraction (~15%) of organics in cloud water can be identified on a molecular basis, (e.g. Herckes et al., 2013), implies that also even the most detailed chemical aqueous phase mechanism is likely largely incomplete in terms of organic species. Thus, also the predicted OH(aq) concentration is likely biased high as by far not all sinks are included. The idea of the general scavenging rate constant as suggested by Arakaki et (2008) is that

it can be applied to parameterize the loss of OH(aq). Thus, it would seem a reasonable ‘short-cut’ to implement water-soluble organic carbon (WSOC) mass as an additional ‘species’ in the mechanism that reacts with OH(aq) to account for missing OH(aq) sources. As the products may be in many cases other WSOC compounds, this reaction could be implemented as  $\text{WSOC} + \text{OH} \rightarrow \text{WSOC} + \text{HO}_2$  ( $k = 3.8 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ ). How would the implementation of this reaction change the results in general, and in particular the OH(aq) level? Is it then in agreement with the ranges as suggested (Arakaki et al., 2013)?

In general, JAMOC is targeted for global model applications and focuses on the explicit oxidation kinetics of known compounds. Even though most of the products from WSOC oxidation are expected to be other WSOCs, the real products are unknown. If the WSOC oxidation was implemented using the suggested equation ( $\text{WSOC} + \text{OH} \rightarrow \text{WSOC} + \text{HO}_2$ ), the concentration of each species in the WSOC group would stay the same. Within a global model, all WSOC are transferred into the gas-phase, when the cloud droplet evaporates. Here, the same species, which was artificially oxidised in the aqueous-phase, would undergo an additional oxidation. In addition, we estimate that each  $\text{HO}_2$  formed from WSOC oxidation would roughly return one OH. Therefore, this approach is not suited for global model applications.

8. Figures 1 and 2 need to be improved:

- Captions: ‘The chemical aqueous phase mechanism of glyoxal (oxalic acid)’ is not very meaningful. At the minimum, specify that it is the oxidation (formation) pathways by chemical radical processes as represented in JAMOC.

In the revised version of the manuscript, both captions are updated following your suggestions.

- Add the names of the species next to the structures in both figures.

Done.

- Caption Figure 1: what do you mean by ‘Gly<sub>aq</sub> donates all three species’?

In JAMOC, the oligomerisation of glyoxal is implemented such that the mono-hydrate reacts with glyoxal, the glyoxal mono-hydrate, and the glyoxal dihydrate. For visual simplicity, we represent all three species by Gly<sub>aq</sub>. In the caption, it now states “Here, Gly<sub>aq</sub> denotes all three forms of glyoxal (glyoxal, the glyoxal mono-hydrate, and the glyoxal dihydrate), which is consistent with the kinetic data published by Ervens and Volkamer (2010)”.

- Caption Figure 1: Are there any sources of aqueous phase glyoxal known at all?

Glyoxal and the glyoxal mono-hydrate are formed during the  $\text{HO}_2$  elimination of the peroxy radicals formed from the oxidation of glycolaldehyde and the glycolaldehyde mono-hydrate. In the revised version, the caption of Fig. 1 now includes this additional information.

- in cloud water, glyoxal is predominantly present in its dihydrate form and should be represented as such in the figure. The mono hydrate may form if there is limited water available, and the unhydrated is likely not present at all (Ervens and Volkamer, 2010).

We agree that with the reviewer that glyoxal is predominantly present as a dihydrate in cloud droplets. However, within JAMOC all hydration steps are represented and Fig. 1 is supposed to show JAMOC’s representation of glyoxal oxidation (as stated in the updated caption). Additionally, we plan to apply JAMOC (with the right adjustments) to aerosols with lower water availability.

- Figure 2: How is the phase partitioning of oxalic acid represented in JAMOC? Given that oxalate forms numerous salts and complexes in the condensed phase, the representation of the phase partitioning based on Henry’s law is likely not appropriate.

We have to admit that we do not include the formation of metal complexes in the mechanism yet. However, we plan to implement this for iron (Fe) soon in a follow up publication. Within JAMOC, the phase partitioning of oxalic acid is represented based on the Henry’s law. This is necessary, since if the phase transfer is not represented by the Henry’s law, oxalic acid will not be transferred into the gas-phase by EMAC when the cloud droplet evaporates. Additionally, we plan to apply JAMOC to aerosols (with the necessary adjustments). Due to the lower pH in aerosol water, the representation by the Henry’s law is appropriate.

- Figure 2: Is the oxidation of oxalic acid by NO<sub>3</sub> ignored in the mechanism? If so, why?

For the NO<sub>3</sub> oxidation of oxalic acid, no rate constant is known from the literature. Opposite to the oxidation by OH, no structure activity relationship (SAR) is available for the H-abstraction implemented for the NO<sub>3</sub> oxidation within JAMOC. Therefore, within CLEPS, similarity criteria are applied if no rate constant is known. However, this is not available for oxalic acid. Therefore, JAMOC does not include the oxidation of oxalic acid by NO<sub>3</sub>. In the revised manuscript, we added an explanation on the rate constant and branching ratios for the oxidation by OH and NO<sub>3</sub> to the respective section.

9. Discussion and Figure 3: The extent is not clear to which the reduction of gas phase mixing ratios is due to uptake into the aqueous phase or due to chemical loss in either phase. I suggest showing total mixing ratios (i.e. gas + aqueous) which would give information on dissolution or net loss, respectively.

We updated Fig. 3 according to your comment. We now show total mixing ratios (i.e. gas + aqueous) for both mechanisms used. In addition, we added an extra line displaying the aqueous-phase mixing ratio of  $\Sigma$ OVOCs, methanol, glycolaldehyde, and methylglyoxal for the simulation using JAMOC. The caption was revised accordingly. Following a suggestion of the first referee of our companion paper (Rosanka et al., 2020), we moved the definition of  $\Sigma$ OVOCs from the caption of Fig. 3 into the newly created Appendix A1.

### Minor comments

l. 17: ‘liquid’ should be replaced by ‘aqueous’

We changed it accordingly.

l. 65: It is not clear which reaction is referred to here (‘reaction with ozone with hydroxide’). Do you mean R2, i.e. the reaction of ozone with the superoxide anion radical (O<sub>2</sub>-)?

In its original version, this statement was supposed to imply that the complete mechanism proposed by Staehelin et al. (1984) and Staehelin and Hoigné (1985) is used in JAMOC. However, we agree that this statement might be confusing. Therefore, we updated this paragraph in the revised version of the manuscript.

l. 69: ‘only outgassing depends on Henry’s law constant’ –I don’t understand this. The standard equations used for the description of mass transfer, e.g. Eq-69 in (Sander, 1999), include the Henry’s law constant which is needed to describe the deviation from equilibrium and thus the concentration gradient that drives the uptake or evaporation, respectively, of species.

We removed this statement in the revised manuscript.

l. 112: I disagree with the authors that ‘little is known’ about NO<sub>3</sub> reactions. There are quite extensive data sets available for NO<sub>3</sub> reactions with organic compounds in the aqueous phase, e.g., (Herrmann, 2003; Herrmann et al., 2015, 2010)

After carefully reevaluating the current literature, we agree with the referee’s comment and removed this statement from the revised manuscript.

l. 127: Why was the rate constant of the dimers estimated as being twice as large as that of the monomer? Is there any reference for this? According to the general kinetic theory, the number of collisions of molecules (which determines the rate constant) scales inversely proportional to molecular mass. Thus, the assumption of a higher rate constant for molecules with doubled mass seems counterintuitive. In addition, the rate constant will also depend on the number of available groups at which the radical attacks. However, since dimers (such as the glyoxal dimer, e.g. (Kua et al., 2008)) form cyclic structures, this trend does not justify a higher rate constant either.

Kua et al. (2008) presents theoretical calculations only of the kinetics of dimerization and not kinetics in the reaction with OH. We estimated the rate constant to be twice as large as the one of the monomer, because the dimers have almost double the number of abstractable H-atoms than the monomers. This reasoning is now included in the respective section.

l. 165: Several multiphase model studies have shown that the direct uptake from the gas phase,

Fenton chemistry and H<sub>2</sub>O<sub>2</sub> photolysis are the main OH(aq) sources in cloud droplets (Deguillaume et al., 2004; Ervens et al., 2003; Herrmann et al., 2005). Compared to these sources, what is the relative contribution of photolysis of organic compounds to OH sources in the aqueous phase?

In Rosanka et al. (2020), we apply JAMOC globally by using EMAC and provide the first global in-cloud OH budget (see Table 2 in the initial submission of Rosanka et al., 2020). Here, the formation of OH from the photolysis of OVOCs containing one carbon atom is estimated to be more than four times higher than the photolysis of H<sub>2</sub>O<sub>2</sub>. Additionally, we estimate that the photolysis of OVOCs containing more than one carbon atom to contribute about one third when compared to the photolysis of H<sub>2</sub>O<sub>2</sub>. A further elaboration of this is added to the revised manuscript.

l. 170: What refers the value 2.33 to? Do you mean ‘enhanced compared to the gas phase photolysis rate’?

Indeed. In general, gas-phase photolysis rates are available. In order to account for scattering effects within cloud droplets, we apply an enhancement factor of 2.33 to the gas-phase photolysis rates within JAMOC. In order to clarify this, we adjusted the text to: “In order to account for scattering effects within cloud droplets, an enhancement factor of 2.33, the same as used in EMAC’s standard aqueous-phase mechanism for the photolysis of H<sub>2</sub>O<sub>2</sub>, is applied to each gas-phase photolysis rate.”

l. 183/184: Why is the relative humidity 70%? Shouldn’t it be 100% in clouds?

For the cloud scenario simulated, we increased the relative humidity to 100 %. We performed both simulations again and updated Fig. 3.

l. 186: I think it should read ‘a stable cloud droplet population’

We changed this in the revised manuscript.

l. 186: A liquid water content of 3 x 10<sup>-1</sup>g L<sup>-1</sup> does not seem an appropriate liquid water content as it would result in 10000 droplets /cm<sup>3</sup> (with diameter of 40 micrometers). Was indeed this LWC used in the model or is it a typo and should read 0.3 g/m<sup>3</sup>?

Thank you for pointing this out. We checked our simulation setup and the LWC we used is indeed 0.3 g/m<sup>3</sup> and the reported value in the manuscript is a typo. We changed it in the revised manuscript.

l. 222: ‘reaction rates’ should be ‘rate constants’ or ‘rate coefficients’. A reaction rate is rate at which a concentration changes, i.e.  $d[C]/dt = -k[C]$  whereas  $k$  is the rate constant and  $[C]$  is the reactant concentration (Seinfeld and Pandis, 2006).

We agree with the reviewer. We changed it accordingly.

l. 228: I am confused by this terminology. JAMOC stands for ‘Jülich Aqueous phase mechanism of organic compounds’, i.e. it is a chemical mechanism which is usually just a list of reactions and their parameters. Such a mechanism can then be implemented into a model that simulates, e.g. the formation of clouds and processing of chemical species? Thus, rain-out is a process in a model, in which JAMOC comprises one module. Please clarify what you mean by ‘model’ and ‘mechanism’, respectively.

By representing the phase-transfer of organic species, which are not explicitly oxidised in JAMOC, allows to represent their rain-out when using global models. We agree with the reviewer that the current wording is confusing. We updated this statment and it now reads: “Phase-transfer of soluble VOCs into cloud droplets is considered in JAMOC even when their oxidation is not explicitly represented (see Sect. 2.2). This allows their removal from the atmosphere by rain-out when JAMOC is connected to a global model (e.g. using EMAC, see Rosanka et al., 2020).”

#### Technical comments

l. 39: ‘react’ should be ‘reacts’

l. 49: ‘extend’ should be ‘extended’

l. 182: ‘modells’ should be ‘models’

Thank you for spotting these three mistakes. We adjusted them.

## References

- Arakaki, T., Anastasio, C., Kuroki, Y., Nakajima, H., Okada, K., Kotani, Y., Handa, D., Azechi, S., Kimura, T., Tsuchi, A., and Miyagi, Y.: A General Scavenging Rate Constant for Reaction of Hydroxyl Radical with Organic Carbon in Atmospheric Waters, *Environmental Science & Technology*, 47, 8196–8203, <https://doi.org/10.1021/es401927b>, PMID: 23822860, 2013.
- Deguillaume, L., Leriche, M., Monod, A., and Chaumerliac, N.: The role of transition metal ions on HO<sub>x</sub> radicals in clouds: a numerical evaluation of its impact on multiphase chemistry, *Atmospheric Chemistry and Physics*, 4, 95–110, <https://doi.org/10.5194/acp-4-95-2004>, 2004.
- Ervens, B. and Volkamer, R.: Glyoxal processing by aerosol multiphase chemistry: towards a kinetic modeling framework of secondary organic aerosol formation in aqueous particles, *Atmospheric Chemistry and Physics*, 10, 8219–8244, <https://doi.org/10.5194/acp-10-8219-2010>, 2010.
- Ervens, B., George, C., Williams, J. E., Buxton, G. V., Salmon, G. A., Bydder, M., Wilkinson, F., Dentener, F., Mirabel, P., Wolke, R., and Herrmann, H.: CAPRAM 2.4 (MODAC mechanism): An extended and condensed tropospheric aqueous phase mechanism and its application, *Journal of Geophysical Research: Atmospheres*, 108, <https://doi.org/https://doi.org/10.1029/2002JD002202>, 2003.
- Ervens, B., Renard, P., Tlili, S., Ravier, S., Clément, J.-L., and Monod, A.: Aqueous-phase oligomerization of methyl vinyl ketone through photooxidation – Part 2: Development of the chemical mechanism and atmospheric implications, *Atmospheric Chemistry and Physics*, 15, 9109–9127, <https://doi.org/10.5194/acp-15-9109-2015>, 2015.
- Herckes, P., Valsaraj, K., and Collett, J.: A review of observations of organic matter in fogs and clouds: Origin, processing and fate, *Atmospheric Research*, 132-133, 434–449, <https://doi.org/10.1016/j.atmosres.2013.06.005>, 2013.
- Herrmann, H.: Kinetics of Aqueous Phase Reactions Relevant for Atmospheric Chemistry, *Chemical Reviews*, 103, 4691–4716, <https://doi.org/10.1021/cr020658q>, PMID: 14664629, 2003.
- Herrmann, H., Tilgner, A., Barzaghi, P., Majdik, Z., Gligorovski, S., Poulain, L., and Monod, A.: Towards a more detailed description of tropospheric aqueous phase organic chemistry: CAPRAM 3.0, *Atmospheric Environment*, 39, 4351 – 4363, <https://doi.org/https://doi.org/10.1016/j.atmosenv.2005.02.016>, URL <http://www.sciencedirect.com/science/article/pii/S1352231005001937>, FEBUKO and MODMEP: A Combined Study of Aerosol-Cloud Interaction by Field Experiments and Model Development, 2005.
- Herrmann, H., Hoffmann, D., Schaefer, T., Bräuer, P., and Tilgner, A.: Tropospheric Aqueous-Phase Free-Radical Chemistry: Radical Sources, Spectra, Reaction Kinetics and Prediction Tools, *ChemPhysChem*, 11, 3796–3822, <https://doi.org/https://doi.org/10.1002/cphc.201000533>, 2010.
- Herrmann, H., Schaefer, T., Tilgner, A., Styler, S. A., Weller, C., Teich, M., and Otto, T.: Tropospheric Aqueous-Phase Chemistry: Kinetics, Mechanisms, and Its Coupling to a Changing Gas Phase, *Chemical Reviews*, 115, 4259–4334, <https://doi.org/10.1021/cr500447k>, PMID: 25950643, 2015.
- Jacob, D. J.: Chemistry of OH in remote clouds and its role in the production of formic acid and peroxymonosulfate, *Journal of Geophysical Research: Atmospheres*, 91, 9807–9826, <https://doi.org/https://doi.org/10.1029/JD091iD09p09807>, 1986.
- Kua, J., Hanley, S. W., and De Haan, D. O.: Thermodynamics and Kinetics of Glyoxal Dimer Formation: A Computational Study, *The Journal of Physical Chemistry A*, 112, 66–72, <https://doi.org/10.1021/jp076573g>, PMID: 18067276, 2008.
- Kumar, M., Anglada, J. M., and Francisco, J. S.: Role of Proton Tunneling and Metal-Free Organocatalysis in the Decomposition of Methanediol: A Theoretical Study, *The Journal of Physical Chemistry A*, 121, 4318–4325, <https://doi.org/10.1021/acs.jpca.7b01864>, PMID: 28513154, 2017.

- Lelieveld, J. and Crutzen, P. J.: Influences of cloud photochemical processes on tropospheric ozone, *Nature*, 343, 227–233, <https://doi.org/10.1038/343227a0>, URL <https://doi.org/10.1038/343227a0>, 1990.
- Lin, G., Penner, J. E., Sillman, S., Taraborrelli, D., and Lelieveld, J.: Global modeling of SOA formation from dicarbonyls, epoxides, organic nitrates and peroxides, *Atmospheric Chemistry and Physics*, 12, 4743–4774, <https://doi.org/10.5194/acp-12-4743-2012>, 2012.
- Liu, L., Zhang, X., Li, Z., Zhang, Y., and Ge, M.: Gas-phase hydration of glyoxylic acid: Kinetics and atmospheric implications, *Chemosphere*, 186, 430–437, <https://doi.org/https://doi.org/10.1016/j.chemosphere.2017.08.007>, URL <http://www.sciencedirect.com/science/article/pii/S0045653517312298>, 2017.
- Perri, M. J., Seitzinger, S., and Turpin, B. J.: Secondary organic aerosol production from aqueous photooxidation of glycolaldehyde: Laboratory experiments, *Atmospheric Environment*, 43, 1487 – 1497, <https://doi.org/https://doi.org/10.1016/j.atmosenv.2008.11.037>, URL <http://www.sciencedirect.com/science/article/pii/S1352231008011096>, 2009.
- Rosanka, S., Sander, R., Franco, B., Wespes, C., Wahner, A., and Taraborrelli, D.: Oxidation of low-molecular weight organic compounds in cloud droplets: global impact on tropospheric oxidants, *Atmos. Chem. Phys. Discuss.*, <https://doi.org/10.5194/acp-2020-1041>, 2020.
- Sander, R.: Modeling Atmospheric Chemistry: Interactions between Gas-Phase Species and Liquid Cloud/Aerosol Particles, *Surveys in Geophysics*, 20, 1–31, <https://doi.org/10.1023/A:1006501706704>, URL <https://doi.org/10.1023/A:1006501706704>, 1999.
- Sander, R., Baumgaertner, A., Cabrera-Perez, D., Frank, F., Gromov, S., Grooß, J.-U., Harder, H., Huijnen, V., Jöckel, P., Karydis, V. A., Niemeyer, K. E., Pozzer, A., Riede, H., Schultz, M. G., Taraborrelli, D., and Tauer, S.: The community atmospheric chemistry box model CAABA/MECCA-4.0, *Geoscientific Model Development*, 12, 1365–1385, <https://doi.org/10.5194/gmd-12-1365-2019>, 2019.
- Seinfeld, J. H. and Pandis, S. N.: Atmospheric chemistry and physics – From air pollution to climate change, John Wiley & Sons, 2006.
- Staehelin, J. and Hoigné, J.: Decomposition of ozone in water in the presence of organic solutes acting as promoters and inhibitors of radical chain reactions, *Environmental Science & Technology*, 19, 1206–1213, <https://doi.org/10.1021/es00142a012>, 1985.
- Staehelin, J., Buehler, R. E., and Hoigné, J.: Ozone decomposition in water studied by pulse radiolysis. 2. Hydroxyl and hydrogen tetroxide (HO<sub>4</sub>) as chain intermediates, *The Journal of Physical Chemistry*, 88, 5999–6004, <https://doi.org/10.1021/j150668a051>, 1984.
- Tan, Y., Perri, M. J., Seitzinger, S. P., and Turpin, B. J.: Effects of Precursor Concentration and Acidic Sulfate in Aqueous Glyoxal–OH Radical Oxidation and Implications for Secondary Organic Aerosol, *Environmental Science & Technology*, 43, 8105–8112, <https://doi.org/10.1021/es901742f>, PMID: 19924930, 2009.