We thank the reviewers for their insights and helpful suggestions. We feel that their feedback has led to changes that have strengthened the quality of the revised manuscript. Please find below the original reviewer comments in black, our responses in blue, and changes to the manuscript in red.

**Reviewer 1**

This study focuses on the comparison between measured OH, HO2 and the sum of HO2 and RO2 (XO2) radicals and results from 2 different chemical mechanisms, one the RACM (lumped) and the other the MCM (semi-explicit). Measurements were conducted in an isoprene dominated forest in Michigan where a lot of ancillary species as well as ISOPPOOH were detected.

This study seems to suggest that quite a large loss rate (termination reaction) for both HO2 and XO2 is needed for both chemical mechanisms investigated to agree with the measurements. Several hypotheses are made such as fast deposition of the radicals on surfaces, faster than used RO2+RO2 reactions for isoprene RO2, RO2 radicals reaction with alkenes and/or segregation could play a role. The authors suggest that most probably a combination of all the above could explain the discrepancy although a rather large loss rate of about 60% during daytime is needed.

The paper is well written and structured, and the arguments are presented in a clear manner. The study is interesting but as it is not possible to give a clear conclusion on what is causing the discrepancies, I would recommend adding a bit of analysis to try and see if something more can be understood and after that I would recommend the publication.

My first suggestion would be to try and perform an experimental budget with the available data. I understand that it might not be possible for the HO2 and RO2 but it would be possible to perform it for the OH radical. In this way it should be clear if the very low OH observed in the morning hours is an instrument artefact as it is quite dubious. The OH budget would also help (possibly) to clarify if indeed there doesn’t seem to be the need for additional sources of OH radicals as the comparison with the model seems to show. I have to say that isomerization reactions for isoprene-RO2 are more or less a given now so I am also wondering why there seems to be such a large overestimate of the OH radical when the most up to date mechs are used.

As suggested, we have added an experimental budget for OH to the supplement of the revised manuscript. As expected, the experimental budget is not balanced in the morning hours which suggests either a missing OH sink or an error with the measurement at this time. This is consistent with the discussion of the discrepancy between measured and modeled OH concentrations which was attributed to participant activity near the detection cell in the morning or a potential systematic measurement error.

An experimental OH budget based on measured concentrations of OH, HO2, and other species, is also shown in Fig. S3. The imbalance between 7:00 and 12:00 suggests either a missing OH sink or errors with the OH measurement during this time.
Figure S3: Experimental OH radical budget. In panel (a), shades of blue represent reactions that produce OH, and shades of red represent loss rates, including reactions that propagate to RO2 or HO2. Percentages indicate the relative contribution of each respective process in the morning (06:30 to 14:00) and during the evening (14:00 to 21:00) time periods which are indicated by the vertical dashed lines. The net rate of production or loss is shown in panel (b).

The MCM v3.3.1 and RACM2-LIM1 models likely overestimate the OH concentrations due to a lack of adequate peroxy radical sinks that are relevant in this forested environment. This is illustrated by the green line in Figure 4, which represents a RACM-ACC model that features additional loss terms for HO2 and isoprene-RO2. The additional loss terms were discussed in terms of the necessary loss rate required to match the HO2 and XO2 measurements, but the impact of these loss terms on the modeled OH concentrations was only briefly mentioned – the following sentence has been added to page 23 of the revised manuscript to clarify.

The addition of these peroxy radical loss mechanisms reduces the predicted daytime maximum OH concentration by 25% to $1.65 \times 10^6$ cm$^{-3}$, which is within the combined uncertainties of the measurement and the model (Fig. 4a).

My second suggestion concerns the XRO2. I am wondering if it would not make sense to remove the HO2 fraction from it from the LIF measurement and then have a more or less RO2 measurement. I understand it does not make much of a difference since the measurement is compared with the some of HO2 and RO2 but to e able to compare with previous studies it would make it easier if it was RO2 instead of XO2.

We focused on comparing the direct measurements to their modeled counterparts (i.e. comparing XO2 measurement to the modeled sum of HO2 and RO2) but do understand that this makes comparison to previous studies more complicated. As suggested by the reviewer we have added a plot that compares the average modeled RO2 to the average measured RO2 (HO2 measured by LIF subtracted from XO2 measured by ECHAMP) to the supplement of the revised manuscript.

In addition, measured RO2 mixing ratios (HO2 measured by LIF subtracted from XO2 measured by ECHAMP) are compared with modeled RO2 mixing ratios in Fig. S4.
My third suggestion is about the ISOPOOH. I am not aware of many ISOPOOH ambient measurements and although I could imagine a different publication focusing on that, it would be good to extend the discussion about it in this study. One thing that I find a bit odd is that the measured concentration of ISOPOOH is more or less zero (within the uncertainty) for the all time? I can see a bit of an increase but it is rather small. Even the model results after constraining HO2 and RO2 would expect quite a bit more. Could this be an instrument artefact? Is this consistent with previous measurements?

The reviewer is correct that there are not many measurements of ambient ISOPOOH available. The data in this manuscript were first presented in Vasquez et al. (2018). The measured mixing ratios for the sum of ISOPOOH isomers were between approximately 10-250 ppt, with an average maximum of approximately 100 ppt. We have added an inset in Figure 7a for better visibility. The measured mixing ratios of ISOPOOH were similar to that measured during the SOAS campaign (Kaiser et al., 2016). This has been clarified on pages 21 and 22 of the revised manuscript.

Measurements of isoprene hydroxy hydroperoxides (ISOPOOH) produced from the reaction of isoprene-based RO2 radicals with HO2 can provide an additional test of the model chemistry at this site. Figure 7 shows the average ISOPOOH mixing ratio measured during PROPHET-AMOS between July 22 and July 27 by the Caltech low-pressure GC-CIMS instrument (Vasquez et al., 2018). The measured mixing ratios were similar to that observed during the SOAS campaign (Kaiser et al., 2016).
Figure 7: Measured and modeled mixing ratios of (a) isoprene hydroxy hydroperoxides (ISOPOOH) and (b) isoprene hydroxy nitrates (IHN). Measurements of ISOPOOH are an average from July 23-27 (Vasquez et al., 2018) and measurements of IHN are an average of July 6-31. The solid lines represent modeled mixing ratios from MCM-ACC models, the dashed line represents predictions of the same model constrained to measured values of HO₂ and measurements of XO₂ scaled to the modeled isoprene RO₂ composition.

The modeled ISOPOOH is sensitive to the rate of ISOPOOH deposition. As described in the original manuscript, the model runs shown in Figure 7 include a literature-based ISOPOOH deposition term, but this term could vary significantly from forest to forest or with meteorological conditions. Due to this and also to the limited amount of ISOPOOH data Figure 7 is not intended to present a direct comparison between measured and modeled ISOPOOH, but instead to illustrate that the measured ISOPOOH is more consistent with the measured HO₂ and XO₂ radical concentrations, regardless of how ISOPOOH deposition is treated in the model. While the model constrained to the peroxy radical concentrations still overestimates the measurements, the model overestimation of the measured ISOPOOH in this study is similar to that observed during the SOAS campaign (Kaiser et al., 2016), where a large dilution rate was needed to bring the modeled ISOPOOH into agreement with the measurements. This has been clarified on page 21 of the revised manuscript.

Constraining the model to the measured concentrations of HO₂ and isoprene-RO₂ (assuming the same relative distribution of RO₂ radicals predicted by the models) improves the agreement (Fig. 7a), although the model still overestimates the measured concentrations. This overestimate of the measured ISOPOOH is similar to that observed during the SOAS campaign (Kaiser et al., 2016), where a large dilution rate was needed to bring the modeled ISOPOOH into agreement with the measurements.

Are there other products that show up which would compensate the production rate of RO2 that, as mentioned, was rather high, and the reacted isoprene must go somewhere.

In addition to ISOPOOH, measurements of isoprene hydroxy nitrates (IHN) produced from the reaction of isoprene peroxy radicals with NO were also overpredicted by the model. Constraining the model to the measured peroxy radical concentrations also brought the modeled IHN into
better agreement with the measurements, again suggesting that the measured isoprene hydroxy nitrates is more consistent with the measured peroxide radical concentrations. This result has been added to the revised manuscript (pages 21 and 22), and we have included a plot of the modeled and measured IHN in Figure 7b.

Similarly, the model also overestimates the concentrations of isoprene IHN produced from the reaction of isoprene peroxy radicals with NO and measured using iodine adduct CIMS (Xiong et al., 2015). Constraining the model to the measured peroxy radical concentrations improves the agreement with the measurements (Fig. 7b). It is also worth noting that the model does not account for losses of IHN due to reactive uptake onto aerosol and subsequent hydrolysis in the aerosol phase (Jacobs et al., 2014; Morales et al., 2021; Wang et al., 2021). Knowledge and incorporation of such loss rates in the model could better constrain the modeled IHN concentrations but the effect is expected to be small in comparison to the adjustment in the modeled output when constrained to measured RO2. (Wei et al., 2021; Mayhew et al., 2022) These results suggest that the measured HO2 and XO2 concentrations are consistent with the measured ISOPOOH and IHN concentrations and that the models are overpredicting the concentrations of HO2 and isoprene-based peroxy radicals, either through an overestimation of their production or an underestimation of their loss.

I noticed that on few occasions the subscripts are not correct so I would recommend checking and fixing that.

Thank you for noticing these mistakes. Subscript formatting errors have been corrected in the revised manuscript.

I also recommend adding the work by J. Medeiros et al. (2022) which is consistent with LIM1.

Thank you for the suggestion. We have added this reference to the introduction section discussing laboratory measurements of isoprene oxidation.

References
