In addition to the changes listed in the point-by-point response, the following changes were made:

1. An error was discovered in how the 0-D photochemical modeling was conducted. The revised model results are used in the revision. The conclusions of the paper remain unchanged. The main difference between the original modeling and the revised modeling is that the relative speciation of peroxy radicals predicted by the four chemical mechanisms (RACM2, RACM2-LIM1, MCM 3.2, and MCM 3.3.1) is now very similar as described in the following two paragraphs from section 3.3 of the manuscript:

   “Measured and MCM 3.2 modeled concentrations for 16, 22, and 24 July are shown in Fig. 8. On all three days the relative contributions from the various types of peroxy radicals are comparable. At 15:30 – when concentrations are highest – the modeled peroxy radicals comprised 30% C\textsubscript{5}H\textsubscript{8}(OH)O\textsubscript{2}, 35% HO\textsubscript{2}, 26% CH\textsubscript{3}O\textsubscript{2} and 7% CH\textsubscript{3}C(O)O\textsubscript{2}. The four chemical mechanisms vary little in the predicted relative speciation (SI). The [XO\textsubscript{2}]/[HO\textsubscript{2}]* ratio modeled by MCM 3.2 between 15:00 and 16:00 is 1.4 for 16 and 22 July and 1.45 on 24 July. The measured [XO\textsubscript{2}]/[HO\textsubscript{2}]* ratio is close to unity on 16 and 22 July, and between 1.2 and 1.5 on 24 July. Increasing these measured ratios by 20% to account for the calibration comparison produces adjusted measured [XO\textsubscript{2}]/[HO\textsubscript{2}]* ratios of 1.2 on 16 and 22 July and 1.4 to 1.8 on 24 July. After accounting for the 20% calibration difference, the modeled and measured ratios agree to within the experimental and model uncertainties.

   Measured [XO\textsubscript{2}] mixing ratios are 20 to 30% lower than the MCM 3.2 [XO\textsubscript{2}] on 16 and 22 July but agree more closely on 24 July (measured MODELED ratio varies from 0.8 to 1.15). The comparison between measured [HO\textsubscript{2}]* and modeled [HO\textsubscript{2}]* for these three days exhibits more variability (Fig. 8). Although all four chemical mechanisms predict a very similar relative speciation, there are variations in the absolute peroxy radical concentrations predicted. MCM 3.3.1 concentrations are very similar to those from MCM 3.2, but RACM2 and RACM2-LIM1 predict 26% and 42% higher peak concentrations, respectively. Further details can be found in the SI.”

2. We have revised the following section in order to address the open comment from Dr. Andres-Hernandez:

   “Similarly, XO\textsubscript{2} measurements from two CO-based chemical amplifiers during the airborne African Monsoon Multidisciplinary Analysis (AMMA) campaign differed by factors of 2-4 when the usual relative humidity-dependent calibration (Mihele and Hastie, 1998) was used for the chemical amplifier data, though the performance of one of the instruments was not assessed with in-flight calibrations (Andrés-Hernández et al., 2010). The relative humidity dependence of the chemical amplification technique is addressed in a variety of ways. Most research groups characterize their instrument’s amplification factor (chain length) as a function of relative humidity (RH) which they then apply to their measurements based on the ambient RH. In some cases, because the RH in the amplification chamber can be lower than ambient because of reduced pressure and higher temperatures, the variability in RH can be considered negligible compared to other experimental uncertainties (Andrés-Hernández et al., 2010; Kartal et al., 2010). In one case the need to apply an RH-dependent calibration was disputed (Sommariva et al., 2011) despite strong experimental evidence (Butkovskaya et al., 2007; Butkovskaya et al., 2005; Butkovskaya et al., 2009; Mihele et al., 1999; Mihele and Hastie, 1998; Reichert et al., 2003).”
3. We have changed the method by which the linear fit was determined for figure 6 – only data between 09:00 and 22:00 are now used due to the low signal-to-noise of the nighttime measurements. This is further described on page 5 of this document.

Our responses to the comments from reviewer #1 are below:

This manuscript details the results of an intercomparison carried out in the field which compares total peroxy radicals using a chemical amplification system with HO2* (which comprises HO2 and a fraction of RO2 radicals) measured by the FAGE technique. Although we may expect HO2* and total RO2 to be well correlated, the comparison presented here is the detected sum of ambient RO2 by two instruments which do not measure different RO2 radicals with the same efficiency, and so is a tricky undertaking. The authors have employed a variety of models with differing chemical mechanisms to predict the composition of peroxy radicals present and from there predict the ratio of total RO2 : HO2* for comparison with the observations. I think on the whole, the approach taken to compare these two observations has led to a meaningful comparison and has demonstrated the performance of the new ECHAMP instrument in the field. I recommend publication once the following comments have been addressed:

Abstract: One concerning result is that the XO2:HO2* ratio is periodically less than one, which the authors themselves note is not possible and must indicate a problem with one or both instruments. A ratio of 0.8 is mentioned in the abstract, but no comment on this low ratio is given until the final pages of the manuscript. I suggest the authors are more upfront about this problem and comment on ratios <1 indicating instrumental issues in the abstract and conclusion.

We have added the following sentence to the abstract:

“Time periods in which the ambient ratio was less than one are definitely caused by measurement errors (including calibration differences) as such ratios are not physically possible.”

and to the abstract:

“The measured [XO2]/[HO2*] ratios usually differed from the ratios predicted by zero-dimensional photochemical modeling by less than the combined measurement and modeling uncertainties, though the lowest ratios observed (0.8) are not physically meaningful and therefore must be due to measurement errors.”

Pg 3, line 18: ‘Measurements of OH by laser-induced fluorescence technique can be affected by a sampling related interference which can exceed the actual concentration of OH..’. This interference is very much dependent on the FAGE instrument design. There are several FAGE instruments in operation that do not observe an OH interference and so this statement needs to be qualified to make this clear.

We have edited that sentence as follows:
“Measurements of OH by the laser-induced fluorescence technique can be affected by a sampling-related interference which can exceed the actual concentration of OH (Mao et al., 2012), though the magnitude of this interference and even its presence varies greatly depending on instrument design.”

Pg 3, line 25: ‘∼90%’ is slightly misleading. In many of the FAGE instruments tested, α is not as high as 90% for the β-hydroxy peroxy radicals; α was as low as 17% in the cited ‘Whalley et al., 2013’ paper.

We have edited that section as follows:

“The sensitivity of the LIF-FAGE technique to each type of organic peroxy radical varies with the amount of NO added for the conversion and is instrument-dependent but in general is highest (up to ∼90%) for β-hydroxy peroxy radicals derived from alkenes and lowest for those derived from small alkanes (Fuchs et al., 2011; Lew et al., 2018; Whalley et al., 2013). This RO2 interference can be greatly reduced by use of lower NO concentrations or reaction times, yielding conversion efficiencies for isoprene-RO2 under 20% (Feiner et al., 2016; Fuchs et al., 2011; Tan et al., 2017; Whalley et al., 2013).”

Pg 7, line 16: please provide the typical Li used for the ambient measurements.

The previous sentence has been edited to clarify that equation 2 is not used to calculate ambient measurements:

“Including a sampling loss term, the sensitivity “α” of ECHAMP to individual organic peroxy radicals relative to that of HO2 can be estimated using Equation 2.”

Later in that paragraph and in the SI, the sampling losses are described.

SI, section S3: Could the authors comment on whether the loss rate of radicals is solely dependent on residence time? Does the shape of the sampling cross and the PFA tees (the sampled air has to flow around corners) impact the loss rate? This could perhaps be determined if the transmitted radical signal was plotted against residence time in the 4 lengths of tube. An intercept would indicate additional losses in the cross piece.

Section S3 describes two types of radical loss tests: 1. measuring the transmission of HO2 through four lengths of tubing, and 2. measuring the HO2 signal when sampling through the sampling cross compared to sampling directly at the reaction chamber. As stated in the SI,

“Similarly, the second method – comparing the ECHAMP signal when sampling a radical source through the sampling cross or directly into one of the reaction chambers – indicated overall losses of less than 4% for an HO2 source.”

Further details of the sampling losses (including loss rates onto several types of material) are the subject of a separate manuscript currently under preparation.

Pg 7, line 23: The authors discuss the impact of alkyl nitrate and alkyl nitrite formation on the sensitivity of ECHAMP to individual RO2 species, but could the authors also comment on the expected sensitivity of ECHAMP to RO2 species which are generated
from alkene + NO3 reactions, so contain an NO3-adduct? ROxLIF instruments are expected to have a low sensitivity to these types of RO2 (Whalley et al. ACP, 2018). If a similarly low sensitivity for these RO2 is expected in ECHAMP, could the authors discuss how this may influence the measured vs modelled ratio during the night?

We have not yet conducted experiments in the lab to determine the sensitivity of ECHAMP to RO2 produced from NO3 reactions, though we do expect that when mixed with NO and ethane these peroxy radicals will lead to increases in NO2. We have added the following sentence regarding the sensitivity of ECHAMP to these radicals at the end of section 2.2:

“We estimate an elevated uncertainty of ~50% for the measurements at night as we have not investigated the sensitivity of ECHAMP to peroxy radical produced by ozonolysis and NO3 reactions.”

We are reluctant to comment on the measured vs. modelled ratio during the night for two reasons: 1. The measurements (of both peroxy radicals and the crucial compound NO) had much lower signal-to-noise ratios and higher uncertainties at night, and 2. The measurement height was only 3 meters which complicates the interpretation of the data since the air was usually stagnant.

Pg 8, line 20: Could the authors make it clear which conversion efficiencies were measured and which have been estimated.

The sentence has been re-worded to clarify:

“The conversion efficiencies for other major RO2 radicals are estimated as 5% for CH3O2 and…”

Pg 8, line 22: The authors reference the Fuchs et al., 2011 work on RO2 interferences in FAGE instruments. ‘α’ is very much dependent upon the specific FAGE instrument and experimental conditions used, however. Using α determined using another FAGE instrument would likely bias the HO2* model measurement comparison. The authors need to make it clear how α was estimated for the RO2 species not experimentally tested with the IU-FAGE. Specifically, how was α = 0.7 derived in equation 5 on page 14, line 21?

The sentence on pg 5 has been edited as follows:

“The conversion efficiencies for other major RO2 are estimated as 5% for CH3O2 and the acetyl peroxy radical (CH3C(O)O2), 8% for ethyl peroxy radical (C2H5O2), and 31-55% for RO2 compounds from the OH oxidation of high-molecular-weight hydrocarbons based on comparisons to several other interference tests (Fuchs et al., 2011; Griffith et al., 2016; Lew et al., 2018).”

The value of 0.7 for the “other” category was chosen since light alkanes, which have low values of α, comprise a minor component of the OH reactivity and most other RO2 compounds have α values near 0.7.

We have added the following sentences:
“The α values for ECHAMP are based on the calculated yields of alkyl nitrates and alkyl nitrites as described in section 2.2. For LIF-FAGE, the α value for C₅H₈(OH)₂O₂ was measured and α for CH₃O₂ and CH₃C(O)O₂ are based on measured yields to several similar instruments all of which have measured values less than 5%. An α of 0.7 is assume for the “other” category since most alkenes have α values between 0.5 and 0.9, and small alkanes, which have lower values, account for a small portion of the OH reactivity (Lew et al., in preparation).”

Pg 12, line 12, fig 3: Add the limit of detection of XO₂ to the figure. Also make it clear in the figure caption which instrument measured HO₂⁺RO₂⁺

The limit of detection (LOD) depends on the relative humidity and the variability in the ambient ozone concentration as described in Wood et al. 2017. To exactly determine the LOD at any given time requires operating both reaction chambers in background mode, precluding simultaneous knowledge of the exact LOD and the ambient concentrations. Rather than add an estimated limit of detection to the figure, we have added the following text to the caption:

“The sum of [RO₂] and [HO₂] was measured by the ECHAMP instrument, with a detection limit typically between 1 and 2 ppt (signal-to-noise ratio of two).”

Pg 12, section 3.3: The authors acknowledge that comparing the 30 min averaged ECHAMP measurements to a single FAGE measurement made during the 30 minute bin is not ideal. I worry that this approach could introduce bias into the comparison, given that the peroxy radical concentrations will generally be increasing throughout the morning hours and then decreasing during the afternoon and evening. Are the FAGE HO₂⁺ measurements made at the midpoint of each 30 minute bin? Does the gradient XO₂ vs HO₂⁺ vary if the FAGE measurement falls at the start of a 30 minute bin? I think the authors need to explore the robustness of this averaging approach used for the ECHAMP data to satisfy the reader that the two measurements are comparable at the times they are taken.

We have majorly revised the paragraph below. We have also changed the averaging time used for the linear regression:

“A bi-variate linear regression of the XO₂ and HO₂⁺ measurements between 09:00 and 22:00 yields the relationship [XO₂] = (1.08 ± 0.05) [HO₂⁺] – (1.4 ± 0.3) ppt (Fig 6.). The regression is restricted to this window of time because of the degraded precision of the ECHAMP measurements at night due to the higher relative humidity. The [XO₂]/[HO₂⁺] slopes were highest on the last two days of measurements – 24 and 25 July, with slopes of 1.25 and 1.08, respectively, or 1.5 and 1.3 after adjusting for the calibration difference. These two days were characterized by the highest mixing ratios of peroxy radicals, O₃, isoprene, and the anthropogenic VOCs ethene and ethyne. The lowest [XO₂]/[HO₂⁺] ratios were observed on 13 July during which a passing thunderstorm led to low concentrations during mid-day with higher values before and after the storm. The higher [XO₂]/[HO₂⁺] ratios observed later in the field campaign may simply be the result of a change in sensitivity in one of the instruments. These linear are difficult to interpret, however, since the XO₂ measurements are 30 minute averages and the HO₂⁺ measurements are 1-minute averages taken every 30 minutes. A regression of the binned data shown in Fig. 5 gives the relation [XO₂] = 1.0 ± 0.14 [HO₂⁺] + (1.5 ± 1.6) ppt; accounting for the calibration difference gives an adjusted slope of 1.2. The [XO₂]/[HO₂⁺] ratio using the binned data was highest between 9:45 and 10:45 (Fig. 5), but was between 0.9 and 1.1 between 14:45 and 19:15. This overall temporal trend is apparent in several days (Fig. 4). Applying a 30-min offset to the XO₂ data largely removes this trend and leads to
fewer time periods when [XO$_2$]/[HO$_2^*$] was less than 1.0, but such an offset does not agree with the synchronized time-base of both measurements. The two instruments’ different averaging times and precision levels preclude further assessment and conclusions regarding possible time offsets.”

*Page 13, line 6 - 8: the authors report the highest XO$_2$:HO$_2^*$ ratio on days when isoprene and ethene concentrations were most elevated. This is unexpected, given the high sensitivity of FAGE to alkene-derived RO$_2$ species. Could the authors comment on this finding?*

We have edited that sentence:

“These two days were characterized by the highest mixing ratios of O$_3$, isoprene, and the anthropogenic VOCs ethene and ethyne. The high [XO$_2$/[HO$_2^*$]] ratios observed those days may simply be the result of a change in sensitivity in one of the instruments.”

*Page 13, line 10: The data in figure 5 has already been binned and then averaged over 9 days. Does the linear regression on the figure 5 data provide a reduced uncertainty relative to the data presented in figure 6? Errors on the fit should be included. I may have misunderstood, but don’t both linear regressions use the same data (just one if further averaged into a diurnal)? Does the change in the regression slope as the data is averaged further suggest that the binning approach is biasing the correlation?*

The binning is useful because there are occasional gaps in the time series (e.g., the morning of 14 July). Without the binning, the morning data is slightly “underrepresented” because of that gap. We have changed the caption as follows (including the fit errors):

“Figure 6. Correlation of ambient [XO$_2$] measured by ECHAMP with [HO$_2^*$] measured by IU-LIF-FAGE. The linear fit is for data between 09:00 and 22:00, indicated by the points with green circles. The equation of the fit is [XO$_2$] = (1.08 ± 0.05) [HO$_2^*$] – (1.4 ± 0.3) ppt.”

*Page 13, line 23: Although I appreciate that the authors do not know the reason why the measurements diverge on the 22nd, the possible explanation ‘a transient interference in the HO$_2^*$ measurement when sampling ambient air.’ is rather vague. Could the authors elaborate on what they think this transient interference may be or what it may be related to?*

We agree that the explanation of a “transient interference” is vague, but feel that any possible reason offered at this point would be too speculative. We note that since HO$_2^*$ is measured as OH after conversion by reaction with NO, any interference in the OH measurement would affect the HO$_2^*$ measurements as well.

*Pg 13, line 25 –Pg 14, line 4: I suggest moving this paragraph to the start of section 3.3. It is important that the α of two instruments to different RO$_2$, and how the ratio is expected to change as ambient RO$_2$ types vary, is set out at the beginning of this section.*

We agree and have made that paragraph the 2nd paragraph of section 3.3 in the revision.
Section 3.3: in general, there is a lot to consider when comparing HO2* and XO2 measured and modelled. The ratio varies with RO2 type present and calibration differences also need to be considered. A table detailing the measured HO2*, XO2 and XO2:HO2* and the 4 modelled HO2*, XO2 and XO2:HO2* on the individual days and campaign average would help to clarify the text.

We hope that the majorly revised paragraph quoted earlier (starting with “A bi-variate linear regression...”) has clarified these issues. Furthermore, the results from the 4 models are shown in the SI.

Figure 4: The caption on the figure is obscuring the top x-axis
This has been fixed in the revision.

Figure 5: There does not seem to be a measured ratio for each 30 min point? Between the hours of 4 – 8, there are only 3 points?
Because of the low signal-to-noise ratios for the nighttime measurements (especially by ECHAMP), the ratio of the measured XO2/[HO2*] varies greatly at night, from 0.3 to 2.1, and so some of those points were off-scale (the graph’s axis was from 0.8 to 1.8). For the revision, we only show the ratio for time periods between 08:00 and 22:00, with the following revised caption:

“…The upper plot shows the [XO2]/[HO2*] ratio - both measured by the two instruments and modeled using the MCM 3.2 chemical mechanism. The measured ratio is only shown for time periods between 08:00 and 22:00 due to the poor signal-to-noise ratios for the night-time measurements.”

Figure 4 – Figure 8: It is unclear whether the ECHAMP data has been corrected for the calibration comparison or not? This should be clear in each figure caption
We have added the following sentence to section 3.3 to clarify:

“No adjustments have been made to either of the datasets in Fig. 4 (or any other figures) to account for the calibration difference.”

Reviewer #2’s comments:

The paper describes the measurement of peroxy radicals (HO2, RO2) with two different techniques. The LIF-FAGE technique by Indiana University was originally designed to measure solely HO2 radicals by chemical conversion with NO to OH, which is then detected by LIF. However, different experimental studies (including a study by authors of this paper) have shown that the technique is also sensitive to specific RO2 radicals with different sensitivities when the instrument is tuned for maximum HO2-to-OH conversion efficiency. The measured quantity is called HO2*. The new ECHAMP technique, a chemical amplifier using ethane instead of CO, is designed to measure the sum of HO2 and all RO2 species. Due to different amplifier chain lengths for different radical species, the resulting quantity XO2 is a proxy for the total peroxy radical concentration. Comparing the measurements by the two techniques sounds like comparing apples with oranges. The present paper demonstrates that such a comparison can be done in a
meaningful way, if the instruments are carefully characterized and additional information about the peroxy radical speciation is available (here from box model calculations constrained by measured trace gases). The direct comparison of the conceptually different calibration methods (photolysis of water vapor vs. photolysis of acetone) and the field comparison show that the measurement techniques yield consistent data within the specified experimental uncertainties. These findings suggest that the two described methods can also be used for meaningful tests of atmospheric chemistry models, if the measured peroxy radicals (HO2*, XO2) are appropriately simulated by the model by taking RO2-specific weighting factors of the instruments into account. This requirement should be explicitly stated in the conclusions.

Furthermore, recent progress in the measurement of HO2 by LIF-FAGE instruments should be mentioned. It has been shown that the interference by RO2 can be avoided by reducing the concentration of NO that is used for conversion to OH (e.g. Fuchs et al., 2011; Whalley et al., 2013; Feiner et al., 2016; Tan et al., 2017).

This section has been re-written as follows:

‘The sensitivity of the LIF-FAGE technique to each type of organic peroxy radical varies with the amount of NO added for the conversion and is instrument-dependent but in general is highest (up to ~90%) for β–hydroxy peroxy radicals derived from alkenes and lowest for those derived from small alkanes (Fuchs et al., 2011; Lew et al., 2018; Whalley et al., 2013). This RO2 interference can be greatly reduced by use of lower NO concentrations or reaction times, yielding conversion efficiencies for isoprene-RO2 lower than 20% (Feiner et al., 2016; Fuchs et al., 2011; Tan et al., 2017; Whalley et al., 2013).’

Overall, the paper is thoroughly and well written. It is suitable for ACP, but could have been submitted to AMT as well. The authors and editor should consider whether the paper should appear as a "Technical note" in ACP. I recommend publication after the following minor comments have been addressed.

We agree that the paper could have been suitable for AMT as well. We chose to submit to ACP because we think that the comparison of the measured concentrations with those by the models provided information beyond that of an instrument assessment and provided information on our community’s understanding of HOx chemistry in low-NOx, high biogenic VOC environments, which has historically been problematic.

(1) Introduction: as the topic of the paper is an instrumental intercomparison, I suggest to provide a more complete list of previous intercomparisons. For instance, Mount et al. (JGR vol.102, no.D5, p6437, 1997), Zenker et al. (JGR vol 103, no Dll, p13,615, 1998), Ren et al. (JGR vol 108, no D19, 4605, 2003), Fuchs et al. (AMT 5, 1611–1626, 2012), Onel et al. (AMT 10, 4877–4894, 2017), Sanchez et al. (Atmos. Env. 174, 227–236, 2018).

We have added the suggested references.

(2) In the experimental section, the authors point out that the use of ethane instead of CO offers advantages. Safer operation is obviously a plus. However, I don’t understand
why the choice of ethane reduces the sensitivity on relative humidity. Is this due to the reduced chain length? Is there evidence for water influence on the OH+CO reaction? To my knowledge, the water effect has been attributed to the reaction HO2+NO (e.g., Mihele et al. 1999, Butkovskaya et al., 2007). Why is the amplification factor lower, if ethane is used? Another advantage of ethane could be mentioned, although it is probably not relevant in a forest environment. Ethane avoids possible interferences from ClOx, which can lead to amplification in CO/NO systems (Perner et al., J. Atmos. Chem. 34, 9, 1999).

We have added the following text to briefly clarify the important issue of RH-dependence:

“The cause of the RH-dependence of the CO-based amplification chemistry is the RH-dependence of the main radical termination step: the reaction of HO2 with NO to form HNO3 (Butkovskaya et al., 2007;Butkovskaya et al., 2005;Butkovskaya et al., 2009;Mihele et al., 1999;Reichert et al., 2003), with a smaller contribution from the RH-dependent wall losses of HO2. These two RH-dependent radical termination steps affect the ethane-based amplification chemistry as well, but the most important terminations steps are from the formation of ethyl nitrite and ethyl nitrate – neither of which depends on relative humidity.”

(3) Page 9: “For this project, [O3] was instead quantified by the ECHAMP CAPS NO2 sensors after conversion to NO2 by reaction with excess NO”. A few details should be explained: is the flow in the calibrator laminar or turbulent? Where is the NO added (upstream, downstream of the calibrator)? Is the NO2 measured after it has been passed through the FAGE cell or is it measured in the air that bypasses the inlet of the FAGE cell? How much NO is added and how large is the resulting NO2 mixing ratio?

We have edited the following section in order to provide more information on this quantification:

“For this project, [O3] was instead quantified by the ECHAMP CAPS NO2 sensors after conversion to NO2 by reaction with excess NO. This was accomplished by having the IU calibration source overflow the ECHAMP inlet. ECHAMP was operated without the ethane flowing, so that each reaction channel sampled 1 LPM of air from the cal source into which 80 sccm of 21 ppm NO was added. This resulted in a diluted concentration of 1.7 ppm NO, which is high enough to react with 99% of the O3 formed during the transit from the inlet to the CAPS detectors. This produces a very precise measurement of the sum of [O3] and [NO2] (1σ precision of 22 ppt for 10 second averages). The accuracy of this ozone determination is thus ultimately traceable to the CAPS NO2 calibration (see SI). Typical [O3] values measured were between 0.4 and 2.0 ppb.”

(4) Page 9, line 18: is the water vapor correction based on laboratory characterization of the LIF-FAGE instrument, or on theoretical calculations using published data for the OH fluorescence lifetime and cross sections for quenching?

It is based on laboratory characterizations. The new sentence:

“The sensitivity of the instrument is corrected for fluorescence quenching by water vapor as per laboratory characterization”

(5) Model constraints: was atmospheric CO measured? Which formaldehyde data were used (GC-FID or DNPH)?
CO was not measured but was estimated based on published emission ratios of CO with benzene. Formaldehyde was only measured using the DNPH cartridges.

The relevant sentences were edited as follows:

“…cartridges to measure carbonyls, including formaldehyde (which was not measured by the GC-FID system), acetaldehyde and…”

and

“Measured VOC concentrations (every 90 min) were interpolated on to this 30 min time resolution. Carbon monoxide was not measured but instead estimated based on emission ratios of CO with benzene (Warneke et al., 2007).”

I see large gaps in the measured time series of NO in the first half of the campaign. Was NO (when available) used as a constraint, or was NO calculated by the model using NO2 as a constraint?

The comparison to the models is heavily focused on the three days when there measurements of NO, XO2 (by ECHAMP) and HO2* (by LIF-FAGE) as described in the text:

“Due to gaps in the NO data because of problems with the Thermo chemiluminescence sensor, there are only three days for which we have model results and measured peroxy radical concentrations by both ECHAMP and LIF-FAGE – on the 16th, 22nd, and 24th of July. The model was run for these three days, and also a diurnal profile for the entire campaign was run using diurnal average concentrations of constrained species.”

The box model was constrained with 30 minute average mixing ratios. As peroxy radicals show a strong non-linear dependence on NO, using 30 minute average values as constraint can lead to systematic bias in the model results. I would like to see the model results that are averaged to 30 minutes after the model has been run at the much higher time resolution of the NOx measurements.

The time resolution of the model is limited by the 90-minute frequency of the VOC measurements which we have interpolated to values every 30-minutes. Thus we are unable to run the model at higher time resolution.

(6) Figure 4 - 6: Is it meaningful to adjust the result of the linear regression for the calibration difference (section 3.1)? This would only make sense, if the calibration would be done for the same peroxy radical speciation as encountered during the measurement days in the field.

We have intentionally included in the text both the “raw” regression/ratio results and those corrected for the calibration difference. Since both ECHAMP and LIF-FAGE are both sensitive (high α) to HO2 and isoprene RO2, we do think that “correcting” the comparisons for the 20% calibration difference helps to frame the discussion of the differences between the two measurements.

(7) Figure 1: what is the scale of the map?
The caption has been updated to address this:
“The arrow represents a distance of 1 km.”

(8) Figure 3: what is causing the noise and spikes on the NO data? Is it measurement precision or atmospheric variability from nearby NO sources?

The data shown in figure 3 are the 5-minute averaged NO concentrations which have a 1σ precision of approximately 100 ppt. The “spikes” in the figure are actually of 15 to 60 minute duration, and thus are from atmospheric variability (mostly during the early morning).

(9) Figure 3 and 4: vertical dotted lines = midnight?

Yes. We have updated the figure captions to clarify:

“The vertical grid lines indicate midnight for odd-numbered days, in local time.”

(10) Figure 5: the shown error bars (1sigma precisions) seem too large compared to the variability of the shown data

In the caption for figure 5 we had erroneously described the error bars as indicative of the 1σ precision of the measurements when they actually just describe the distribution of the measured concentrations. We have changed that sentence in the caption to the following:

“The error bars indicate the ± one standard deviation of the measured concentrations in each 30-minute time bin during those nine days.”
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Peroxy Radical Measurements by Ethane - Nitric Oxide Chemical Amplification and Laser-Induced Fluorescence / Fluorescence Assay by Gas Expansion during the IRRONIC field campaign in a Forest in Indiana


1Department of Chemistry, University of Massachusetts, Amherst MA, 01003, United States
2Department of Chemistry, Indiana University, Bloomington IN 47405, United States
3School of Public and Environmental Affairs, Indiana University, Bloomington, IN 47405, United States
4IMT Lille Douai, Université Lille, Département Sciences de l’Atmosphère et Génie de l’Environnement (SAGE), F-59000 Lille, France
5Department of Chemistry, Drexel University, Philadelphia PA, 19104, United States

* now at Momentive Performance Materials, Inc., Tarrytown, NY 10591, United States
** now at Department of Chemistry, University of Colorado, Boulder CO 80309, United States
*** now at Cooperative Institute for Research in Environmental Sciences, University of Colorado, Boulder, CO 80309, USA

Correspondence to: Ezra Wood (Ezra.Wood@drexel.edu)
**Abstract.** Peroxy radicals were measured in a mixed deciduous forest atmosphere in Bloomington, Indiana, USA, during the Indiana Radical, Reactivity and Ozone Production Intercomparison (IRRONIC) during the summer of 2015. Total peroxy radicals (\([XO_2] = \{HO_2\} + \Sigma[RO_2]\)) were measured by a newly developed technique involving nitric oxide (NO) – ethane (\(C_2H_6\)) chemical amplification followed by NO\(_2\) detection by cavity attenuated phase shift spectroscopy (hereinafter referred to as ECHAMP). The sum of hydroperoxy radicals (\(HO_2\)) and a portion of organic peroxy radicals (\(\{HO_2^*\} = \{HO_2\} + \Sigma \alpha_i[RO_iO_2], 0<\alpha<1\)) was measured by the Indiana University Laser-Induced Fluorescence / Fluorescence Assay by Gas Expansion instrument (LIF-FAGE). Additional collocated measurements include concentrations of NO, NO\(_2\), O\(_3\), and a wide range of volatile organic compounds (VOCs); and meteorological parameters. \(XO_2\) concentrations measured by ECHAMP peaked between 13:00 to 16:00 local time, with campaign average concentrations of 41 ± 15 ppt (1σ) at 14:00. Daytime concentrations of isoprene averaged 3.6 ± 1.9 ppb (1σ) whereas average concentrations of NOx (\([NO] + [NO_2]\)) and toluene were 1.2 ppb and 0.1 ppb, respectively, indicating a low impact from anthropogenic emissions at this site.

We compared ambient measurements from both instruments and conducted a calibration source comparison. For the calibration comparison, the ECHAMP instrument, which is primarily calibrated with an acetone photolysis method, sampled the output of the LIF-FAGE calibration source which is based on the water vapor photolysis method and, for these comparisons, generated a 50-50% mixture of \(HO_2\) and either butane or isoprene-derived RO\(_2\). A bivariate fit of the data yields the relation \([XO_2]_{ECHAMP} = (0.88 \pm 0.02) ([HO_2]+[RO_2])_{IU\text{-cal}} + (6.6 \pm 4.5) \text{ ppt}\). This level of agreement is within the combined analytical uncertainties for the two instruments’ calibration methods.

A linear fit of the daytime (09:00 – 22:00) 30-minute averaged \([XO_2]\) ambient data with the 1-minute averaged \([HO_2]^*\) data (one point per 30 minutes) yields the relation \([XO_2] = \{1.08 \pm 0.05\} [HO_2^*] - \{1.4 \pm 0.3\}\). Day to day variability in the \([XO_2]/[HO_2^*]\) ratio was observed. The lowest \([XO_2]/[HO_2^*]\) ratios between 13:00 and 16:00 were 0.8 on 13 and 18 July, whereas the highest ratios of 1.1 to 1.3 were observed on 24 and 25 July – the same two days on which the highest concentrations of isoprene and ozone were observed. Although the exact composition of the peroxy radicals during IRRONIC is not known, 0-dimensional photochemical modeling of the IRRONIC dataset using the RACM2, RACM2-LIM1, MCM 3.2, and MCM 3.3.1 chemical mechanisms all predict afternoon \([XO_2]/[HO_2^*]\) ratios of between 1.2 to 1.5.

Differences between the observed ambient \([XO_2]/[HO_2^*]\) ratio and that predicted with the 0-D modeling can be attributed to deficiencies in the model errors in the two measurement techniques, or both. **Time periods in which the ambient ratio was less than one are definitely caused by measurement errors (including calibration differences) as such ratios are not physically meaningful.** Although these comparison results are encouraging and demonstrate the viability of using the new ECHAMP technique for field measurements of peroxy radicals, further research investigating the overall accuracy of the measurements and possible interferences from both methods is warranted.
1. Introduction

Peroxy radicals in the atmosphere comprise hydroperoxy (HO$_2$) and organic peroxy radicals (RO$_2$, R = organic group). The most important sources of peroxy radicals are the reactions of oxidants (OH, O$_3$, and NO$_3$) with volatile organic compounds (VOCs), photolysis of oxygenated VOCs, and decomposition of peroxyacetyl nitrate (PAN) (Atkinson, 2000). Chemistry involving "ROx" radicals ([ROx] ≡ [OH] + [RO$_2$] + [H$_2$O]) leads to the formation of ozone (O$_3$), oxygenated VOCs, and secondary aerosol particles (Atkinson, 1997; Atkinson and Arey, 2003; Claeys et al., 2004; Kroll and Seinfeld, 2008; Ng et al., 2008). The chemical identity and concentrations of peroxy radicals can provide important information on atmospheric oxidation processes such as ozone production, the removal efficiency of primary pollutants, and radical budgets. This information is ultimately required to formulate pollution control strategies and to evaluate the impacts of atmospheric chemistry on health and global climate. It is therefore crucial to understand the concentrations and chemistry of RO$_x$ radicals in the atmosphere.

Comparison of measured radical concentrations to those produced by photochemical models is a common exercise used to assess our understanding of atmospheric chemistry. Discrepancies of a factor of two or more between measured and modeled OH concentrations have been reported in biogenic VOC-rich forest environments (Lelieveld et al., 2008; Lu et al., 2012; Pugh et al., 2010), suggesting that our knowledge of atmospheric photochemistry is deficient. Similarly, discrepancies between measured and modeled peroxy radicals have suggested the presence of unknown sources or sinks of peroxy radicals (Griffith et al., 2013; Wolfe et al., 2014). These findings have fueled research into the oxidation mechanisms of biogenic VOCs, especially isoprene (e.g., Wennberg et al., 2018). Although much has been learned in the past decade, the atmospheric fate of biogenic VOCs remains incompletely understood.

Some past model-measurement comparisons are difficult to interpret because of measurement errors that have recently been discovered. Measurements of OH by the laser-induced fluorescence technique can be affected by a sampling-related interference which can exceed the actual concentration of OH (Mao et al., 2012), though the magnitude of this interference and even its presence varies greatly depending on instrument design. Similarly, many previous measurements of HO$_2$ by chemical conversion to OH through the HO$_2$ + NO → OH + NO$_2$ reaction using both the LIF-FAGE and the perCIMS techniques have been shown to have been affected by a variable contribution from organic peroxy radicals (Fuchs et al., 2011; Hornbrook et al., 2011) and the LIF-based measurements subject to this interference are now referred to as HO$_2^*$ ([HO$_2^*$$] = [HO$_2$] + \alpha\Sigma[R,O$_2$], 0 < \alpha < 1). The sensitivity of the LIF-FAGE technique to each type of organic peroxy radical varies with the amount of NO added for the conversion and is instrument-dependent but in general is highest (up to ~90%) for β-hydroxy peroxy radicals derived from alkenes and lowest for those derived from small alkanes (Fuchs et al., 2011; Lew et al., 2018; Whalley et al., 2013). This RO$_2$ interference can be greatly reduced by use of lower NO concentrations or reaction times, yielding conversion efficiencies for isoprene-RO$_2$ under 20% (Feiner et al., 2016; Fuchs et al., 2011; Tan et al., 2017; Whalley et al., 2013).
Discrepancies between measured and model-predicted OH and XO2 concentrations can be caused by a combination of measurement errors, missing or incorrect chemistry in models and erroneous model constraints. Measurement errors can be evaluated by the comparison of atmospheric measurements by multiple techniques. Several HOx intercomparison projects have been conducted in the past few decades (Eisele et al., 2003; Fuchs et al., 2010; Fuchs et al., 2012; Hofzumahaus et al., 1998; Mount and Williams, 1997; Onel et al., 2017; Ren et al., 2003; Ren et al., 2012; Sanchez et al., 2018; Schlosser et al., 2009; Zenker et al., 1998). There have been few intercomparisons, however, of total peroxy radical ([HO2] + ∑[RO2]) measurements and these have produced mixed results. For example, excellent agreement between the matrix isolation electron spin resonance (MI-ESR) and the ROx LIF-FAGE techniques was observed in a chamber study involving HO2, CH3O2, and C4H7O2 produced by the oxidation of methane and 1-butene (Fuchs et al., 2009). An earlier comparison of XO2 measurements between a CO-based chemical amplifier (PERCA) and MI-ESR showed overall agreement of within 10% (Platt et al., 2002). In contrast, XO2 measurements in a forest from two similar CO-based chemical amplifiers differed by more than a factor of three (Burkert et al., 2001). This disagreement was attributed to sampling losses on a rain cover. Similarly, XO2 measurements from two CO-based chemical amplifiers during the airborne African Monsoon Multidisciplinary Analysis (AMMA) campaign differed by factors of 2-4 when the usual relative humidity-dependent calibration (Mihele and Hastie, 1998) was used for the chemical amplifier data, though the performance of one of the instruments was not assessed with in-flight calibrations (Andrés-Hernández et al., 2010).

The relative humidity dependence of the chemical amplification technique is addressed in a variety of ways. Most research groups characterize their instrument’s amplification factor (chain length) as a function of relative humidity (RH) which they then apply to their measurements based on the ambient RH. In some cases, because the RH in the amplification chamber can be lower than ambient because of reduced pressure and higher temperatures, the variability in RH can be considered negligible compared to other experimental uncertainties (Andrés-Hernández et al., 2010; Kartal et al., 2010). In one case the need to apply an RH-dependent calibration was disputed (Sommariva et al., 2011) despite strong experimental evidence (Butkovskaya et al., 2007; Butkovskaya et al., 2005; Butkovskaya et al., 2009; Mihele et al., 1999; Mihele and Hastie, 1998; Reichert et al., 2003). Due to the paucity of XO2 measurement intercomparisons and these new questions regarding the impact of relative humidity on the traditional chemical amplifier technique, further intercomparisons involving different instruments are required before we have enough confidence in the measurements to interpret model-measurement discrepancies as arising from unknown chemistry in models.

This paper presents measurements of XO2 in a mixed deciduous forest by the new Ethane CHEmical AMPlifier (ECHAMP) technique (Wood et al., 2017), which is a variation of the traditional chemical amplification or “PERCA” method (Cantrell and Stedman, 1982; Hastie et al., 1991; Wood and Charest, 2014). Measured XO2 concentrations at this high isoprene, low NOx environment are described along with supporting measurements of ozone (O3), nitrogen oxides (NOx), biogenic and anthropogenic VOCs, and meteorology. We compare measurements of XO2 by ECHAMP with collocated ambient measurements of HO2⁺ by the Indiana University LIF-FAGE technique. We also describe calibration comparison experiments in which ECHAMP, which was calibrated by an acetone photolysis calibration method, quantified radical
concentrations produced by the LIF-FAGE calibration source which is based on the more common water photolysis method. Ozone formation rates are also calculated based on measured XO$_2$ and NO concentrations.

2 Experimental Section

2.1 Site description

The measurements were carried out at the Indiana University Research and Teaching Preserve (IURTP) field laboratory during the Indiana Radical, Reactivity and Ozone Production Intercomparison (IRRONIC) campaign over the time period of 9 July – 8 August 2015. The IURTP is located in a mixed deciduous forest 1 km from the perimeter road for Indiana University in Bloomington (Fig. 1). Instrument inlets and related instrumental accessories were set atop a 3 meter scaffolding platform in a clearing behind the IURTP building. The height of the scaffolding was several meters below the forest canopy. The major analytical instruments and gas cylinders were housed inside the building.

2.2 ECHAMP Measurements of Total Peroxy Radicals (XO$_2$)

XO$_2$ concentrations were quantified using a newly developed analytical technique, which involves chemical amplification by ethane (C$_2$H$_6$) - nitric oxide (NO) followed by nitrogen dioxide (NO$_2$) detection using cavity attenuated phase shift spectroscopy (hereinafter referred as ECHAMP: Ethane CHemical AMPlifier) (Wood et al., 2017). This instrument can be thought of as a descendent of “traditional” chemical amplifiers, also known as PERCA, in which ambient air is mixed with carbon monoxide and nitric oxide and the resulting “amplified” NO$_2$ measured by the luminol technique (Cantrell and Stedman, 1982; Clemitshaw et al., 1997; Kartal et al., 2010; Mihele and Hastie, 2000). Our initial peroxy radical sensor (Wood and Charest, 2014) relied on the original CO/NO amplification chemistry but utilized a modern, highly sensitive NO$_2$ detection method: cavity attenuated phase shift spectroscopy (CAPS) (Kebabian et al., 2007; Kebabian et al., 2008). The major modification made for the ECHAMP method used for the measurements described in this study is that ethane (C$_2$H$_6$) replaces CO as a reagent. This results in greatly improved deployability thanks to the relative safety of C$_2$H$_6$ compared to CO, a smaller dependence of the sensitivity on relative humidity, but at the expense of lower amplification factors. The cause of the RH-dependence of the CO-based amplification chemistry is the RH-dependence of the main radical termination step: the reaction of HO$_2$ with NO to form HNO$_3$ (Butkovskaya et al., 2007; Butkovskaya et al., 2005; Butkovskaya et al., 2009; Mihele et al., 1999; Reichert et al., 2003), with a smaller contribution from the RH-dependent wall losses of HO$_2$. These two RH-dependent radical termination steps affect the ethane-based amplification chemistry as well, but the most important terminations steps are from the formation of ethyl nitrite and ethyl nitrate – neither of which depends on relative humidity. Details of the experimental technique are described elsewhere (Wood et al., 2017) but its deployment at the IURTP is described here. The ECHAMP inlet was attached to scaffolding at a height of 3 m. Ambient air was sampled at a flow rate of 5.5 standard liters per minute (SLPM) into a 0.4 cm inner diameter (ID) glass sampling cross internally coated with halocarbon wax (Halocarbon Products Corp., series 1500) and externally coated with PTFE tape. The sampled air then
entered two identical reaction chambers (0.4 cm ID × 61 cm, FEP tubing) at a flowrate of 0.87 SLPM - see schematic in Wood and Charest (2014). The total residence time in the sampling cross before entering the reaction chambers was approximately 18 ms.

At any given point in time, one reaction chamber operated in “amplification” (ROx) mode while the other operated in “background” (Ox) mode. In “ROx” mode, the air was immediately mixed with NO and C\(_2\)H\(_6\) in the “upstream” reagent addition port and, 0.1 second later, mixed with nitrogen (N\(_2\)) in the “downstream” reagent addition port, effecting the following radical propagation reactions:

\[
\begin{align*}
\text{RO}_2 + \text{NO} & \rightarrow \text{RO} + \text{NO}_2 \\
\text{RO} + \text{O}_2 & \rightarrow \text{HO}_2 + \text{products} \\
\text{HO}_2 + \text{NO} & \rightarrow \text{OH} + \text{NO}_2 \\
\text{OH} + \text{C}_2\text{H}_6 & \rightarrow \text{H}_2\text{O} + \text{C}_2\text{H}_5 \\
\text{C}_2\text{H}_5 + \text{O}_2 + \text{M} & \rightarrow \text{C}_2\text{H}_6\text{O}_2 + \text{M} \\
\text{C}_2\text{H}_6\text{O}_2 + \text{NO} & \rightarrow \text{C}_2\text{H}_3\text{O} + \text{NO}_2 \\
\text{C}_2\text{H}_3\text{O} + \text{O}_2 & \rightarrow \text{CH}_3\text{CHO} + \text{HO}_2
\end{align*}
\]

Reactions R3 through R7 repeat several times, leading to the formation of NO\(_2\) that is subsequently measured by a CAPS sensor. In background (Ox) mode, the N\(_2\) and C\(_2\)H\(_6\) flows were switched: sampled air was mixed with NO and N\(_2\) upstream and C\(_2\)H\(_6\) downstream. During this sampling mode, sampled radicals are removed by a combination of reactions R1, R2, R3 and finally the reaction of OH with NO to form HONO. The flowrates of NO, N\(_2\) and C\(_2\)H\(_6\) were each maintained at 45 sccm using mass flow controllers (MKS model 1179A and Alicat MC series). Cylinder concentrations of NO and C\(_2\)H\(_6\) (Indiana Oxygen) were 21.1 ppm and 30\%, respectively, leading to concentrations in the reaction chamber of 0.9 ppm and 1.4\% respectively. Both upstream and downstream injections were delivered with PFA tubing (0.16 cm i.d. × 6 m). Each reaction chamber alternated between ROx and Ox mode every 45 seconds on an anti-synchronized schedule using four solenoid valves controlled by Labview software (National Instruments). After the downstream reagent addition, the air from each reaction chamber flowed through 1 m of 0.32 cm ID FEP tubing, a particulate matter filter (United Filtration Systems, Inc., DIF BN60), and another 6 m of tubing before entering identical CAPS monitors located inside the laboratory. The CAPS NO\(_2\) measurements during “ROx” mode are from ambient NO\(_2\), NO\(_2\) from the reaction of NO and O\(_3\) in the reaction chamber and transport tubing, and NO\(_2\) from the chemical amplification reactions involving HO\(_2\) and RO\(_2\) (R1 through R7). In “Ox” mode, the CAPS measures NO\(_2\) from the first two categories above and NO\(_2\) produced by R1 and R3 but not from the amplification reactions (R3 to R7), as ethane is not added until all radicals are removed by formation of HONO.

The concentrations of peroxy radicals were calculated by dividing the difference between the two CAPS sensors’ NO\(_2\) measurements (\(\Delta\text{NO}_2\)) between “ROx” and “Ox” modes by an experimentally determined amplification factor F:

\[
[\text{RO}_2] + [\text{HO}_2] = \frac{\Delta[\text{NO}_2]}{\text{CAPS A} - \text{CAPS B}} / F
\]
The RH-dependent amplification factor \( F \) was measured using the acetone photolysis method described by Wood and Charest (2014). Briefly, methyl peroxy (\( \text{CH}_3\text{O}_2 \)) and peroxyacetyl (\( \text{CH}_3\text{C}(\text{O})\text{OO} \)) radicals (50 – 400 ppt) were produced by the photolysis of acetone vapor and reacted with excess NO to form \( \text{NO}_2 \) which was quantified using a CAPS \( \text{NO}_2 \) sensor. The accuracy of this calibration method ultimately depends on the accuracy of the CAPS \( \text{NO}_2 \) measurement (see supplementary information (SI)) and knowledge of the products of the reaction of \( \text{CH}_3\text{O}_2 \) and \( \text{CH}_3\text{C}(\text{O})\text{OO} \) with NO but does not depend on measurements of actinic flux.

The amplification factor \( F \) was measured to be 28 at 0% relative humidity (RH) and decreased to 6 at 90% RH (Wood et al., 2017). The RH was typically between 50 and 75% during the afternoon, corresponding to values of \( F \) between 20 and 11. These values are based on laboratory calibrations performed before and after the field project. During the field campaign, we attempted to use a variation on the calibration method described by Wood and Charest (2014). Rather than flow air through the headspace over pure acetone to produce dilute acetone vapor, we instead flowed air through the headspace of dilute (1%) aqueous acetone in an attempt to obviate the need to dilute the resulting acetone vapor (i.e., by reducing the vapor pressure of the acetone per Raoult’s Law). Inconsistent calibrations resulted, however, and subsequent laboratory tests demonstrated that the use of aqueous acetone sometimes produced compounds that absorb blue light and therefore interfered with the CAPS \( \text{NO}_2 \) measurement which is based on absorption of light at 450 nm with a bandpass of 10 nm (full width at half maximum). Because field calibrations were unsuccessful, we have increased the measurement uncertainty accordingly (see below). The acetone vapor photolysis calibration results obtained in the laboratory also agreed with our prototype \( \text{H}_2\text{O} \) photolysis method as described in Wood et al (2017). Further details on the calibration are described in the SI.

Individual peroxy radicals are not detected with equal sensitivity by ECHAMP due to the formation of organic nitrates and organic nitrites in the reaction chambers:

\[
\begin{align*}
\text{RO}_2 + \text{NO} & \rightarrow \text{RO} + \text{NO}_2 \quad \text{R8a} \\
\text{RO}_2 + \text{NO} + \text{M} & \rightarrow \text{RONO}_2 + \text{M} \quad \text{R8b} \\
\text{RO} + \text{O}_2 & \rightarrow \text{R'O} + \text{HO}_2 \quad \text{R9a} \\
\text{RO} + \text{NO} + \text{M} & \rightarrow \text{RONO} + \text{M} \quad \text{R9b}
\end{align*}
\]

Including a sampling loss term, the sensitivity “\( \alpha \)” of ECHAMP to individual organic peroxy radicals relative to that of \( \text{HO}_2 \) can be estimated using Equation 2:

\[
\alpha_{\text{RO}_2} = \frac{S_{\text{RO}_2}}{S_{\text{HO}_2}} = L_i(1-Y_i)(k_{\text{RO}_2}[\text{O}_2]/(k_{\text{RO}_2}[\text{O}_2] + k_{\text{RO}_6}[\text{NO}])) 
\]

where \( S_{\text{RO}_2}/S_{\text{HO}_2} \) is the sensitivity of ECHAMP to individual \( \text{RO}_2 \) compounds relative to that of \( \text{HO}_2 \), \( L_i \) is the fractional sampling transmission of an individual organic peroxy species “\( \text{RO}_2 \)” through the short inlet into the reaction chambers.
(relative to that of HO₂), Y is the alkyl nitrate yield (Y = R8b/(R8a + R8b)), and the remaining terms in parentheses account for alkyl nitrite (RONO) formation. Alkyl nitrate yields increase with carbon backbone number, from less than 0.1% for CH₃O₂ to 8% for isoprene to over 25% for C10 and larger alkyl peroxy radicals (Lockwood et al., 2010; Orlando and Tyndall, 2012). Alkyl nitrite (RONO) formation accounts for less than 4% loss for most organic peroxy radicals and is likely negligible for alkene-derived peroxy radicals due to the rapid decomposition of beta hydroxy alkoxy radicals (Atkinson, 1997), but can sequester a calculated 10% of CH₃O₂ (Wood et al., 2017). Sampling losses are limited to the 18 ms transit time in the halocarbon wax–coated sampling cross to the tee in which the NO and C₂H₆ are added. Mihele et al. (1999) measured effective first order wall loss rate constants of 3 to 7 s⁻¹ for HO₂ onto ¼” OD PFA tubing, depending on RH, and ~0.5 s⁻¹ for CH₃O₂ and C₂H₅O₂. Though this would suggest losses in our inlet of up to 12% for HO₂ and 1% for the alkyl peroxy radicals, laboratory tests on our inlet have demonstrated losses of less than 2% for HO₂ in our inlet and loss rate constants onto various fluoropolymers much lower than presented in Mihele et al. (1999) as described in the SI.

At an RH of 50%, the theoretical 1σ precision of the ECHAMP measurements, limited by only the precision of the CAPS NO₂ measurements and the amplification factor, was 0.8 ppt for a 90-second average. The atmospheric variability of O₃, which after reaction with NO accounts for most of the NO₂ observed by the CAPS sensors, led to an additional contribution to the noise due to the slightly different time responses of the two CAPS sensors. The observed precision during sampling was typically 2.5 ppt (1σ) for 90-second averaging (Wood et al., 2017), leading to a detection limit of 5 ppt for 90-second averaging and 1.6 ppt for 15 minute averages at a signal-to-noise ratio of two. At night, although variability of O₃ was negligible, high RH values of over 95% and the resulting low values of F led to detection limits of between 2 ppt and 8 ppt for 90 second average measurements.

We assign an uncertainty of 27% (2σ) to the ECHAMP measurements during the IRRONIC project, comprising the uncertainty in the NO₂ calibration of the CAPS sensors (5%), the uncertainty in the relative humidity - dependent amplification factor (usually 16%, but increased to 25% because post-deployment laboratory calibrations were used instead of the unsuccessful field calibrations using aqueous acetone), and the variable sensitivity to speciated peroxy radicals. We estimate an elevated uncertainty of ~50% for the measurements at night as we have not investigated the sensitivity of ECHAMP to peroxy radicals produced by ozonolysis and NO₃ reactions. These uncertainties are more fully described in Wood et al. (2017). Except where noted otherwise, all ECHAMP XO₂ measurements presented are 15-minute averages.

2.3 Laser-Induced Fluorescence Measurements HO₂*

HO₂* was measured by the Laser-Induced Fluorescence / Fluorescence Assay by Gas Expansion (LIF-FAGE) technique described in detail elsewhere (Griffith et al., 2013a; Dusanter et al., 2009). Briefly, air is sampled through a pinhole into a low pressure chamber and mixed with NO which converts HO₂ into OH. OH radicals are excited by 308 nm radiation from a tunable dye laser and the subsequent fluorescence detected with a time-gated microchannel plate photomultiplier
Concentrations of generated peroxy radicals are measured with a UV-absorption spectrometer. The conversion efficiencies for other major RO2 are estimated as 5% for CH3O2 and the acetyl peroxy radical (CH3C(O)O2), 8% for ethyl peroxy radical (C2H5O2), and 31-55% for RO2 compounds from the OH oxidation of high-molecular-weight hydrocarbons based on comparisons to several other interference tests (Fuchs et al., 2011; Griffith et al., 2016; Lew et al., 2018). These conversion efficiencies are average values weighted over the distribution of isomers where applicable.

The LIF-FAGE instrument was calibrated using a portable calibrator in which quantified amounts of OH/RO2 and NOx were produced through the photolysis of water vapor by a low-pressure mercury lamp at 184.9 nm (Dusantner et al., 2008). Humid air containing either isoprene (80 ppb) or n-butane (1.4 ppm) entered the rectangular calibrator (1.27 × 1.27 × 30 cm). Light from a low-pressure mercury lamp (UVP Inc., model 11sc1) illuminated a ~3 cm3 photolysis volume through a quartz window. The flow rate of air was maintained at 45 SLPM. A mixture with equal concentrations of HO2 and either C3H6(OH)O2 (from isoprene) or C4H6O2 (from butane) were produced when isoprene or butane were added to the calibration gas upstream of the photolysis region, respectively. Ozone actinometry was used to quantify the product of the actinic flux and the exposure time (“Ft”) in the calibrator (Dusantner et al., 2008). Concentrations of generated peroxy radicals are calculated by the following equation:

\[
[\text{HO}_2] + [\text{RO}_2] = \frac{\left[\sigma_1 [\text{H}_2\text{O}] \sigma_2 [\text{O}_2] \phi_{\text{HO}_2} \phi_{\text{RO}_2}\right]}{\left[\sigma_1 [\text{H}_2\text{O}] \sigma_2 [\text{O}_2] \phi_{\text{HO}_2} \phi_{\text{RO}_2}\right]}
\]

where [O2] is the concentration of ozone generated by the photolysis of O2; \(\sigma_{\text{HO}_2}\) and \(\sigma_{\text{RO}_2}\) are the absorption cross sections of H2O and O2 at 184.9 nm, respectively, and \(\phi_{\text{HO}_2}\) and \(\phi_{\text{RO}_2}\) are the photolysis quantum yields, both equal to two (Waschita et al., 1971). A value of 7.14 × 10^{-20} cm² molecule⁻¹ (base e) was used for \(\sigma_{\text{HO}_2}\) (Cantrell et al., 1997; Hofzumahaus et al., 1997; Lanzendorf et al., 1997). The effective value of \(\sigma_{\text{RO}_2}\) depends on the O2 optical depth and the operating conditions of the mercury lamp and was determined to be 1.20 × 10^{-20} cm² molecule⁻¹ (Dusantner et al., 2008; Lanzendorf et al., 1997). The water vapor mixing ratio was measured by IR absorption spectrometry using a LI-COR 6262 monitor. Ordinarily the ozone mixing ratio is determined using a calibrated photodiode installed in the calibrator (Griffith et al., 2013). The conversion factor (calibration) that converts the photodiode reading to an O3 mixing ratio is determined from separate experiments in which a range of O3 concentrations produced by the calibrator are measured with a UV-absorption O3 sensor. For this project, \([\text{O}_3]\) was instead quantified by the ECHAMP CAPS NO2 sensors after conversion to NO2 by reaction with excess NO. This was accomplished by having the IU calibration source overflow the ECHAMP inlet. ECHAMP was operated without the ethane flowing, so that each reaction channel sampled 1 LPM of air from the calibration.
source into which 80 sccm of 21 ppm NO was added. This resulted in a diluted concentration of 1.7 ppm NO, which is high enough to react with 99% of the O₃ formed during the transit from the inlet to the CAPS detectors. This produces a very precise measurement of the sum of [O₃] and [NO₂] (1σ precision of 22 ppt for 10 second averages). The accuracy of this ozone determination is thus ultimately traceable to the CAPS NO₂ calibration (see SI). Typical [O₃] values measured were between 0.4 and 2.0 ppb. Linking the IU FAGE HO₂* calibration to the ECHAMP NO₂ measurement has ramifications for the intercomparison of the IU calibration source and the ambient measurements as discussed in the relevant sections below.

The sensitivity of the instrument is corrected for fluorescence quenching by water vapor as per laboratory characterization. This amounted to a correction of approximately 20% at a water mixing ratio of 1%. The limit of detection of HO₂* was 0.8 ppt (30 s average, signal-to-noise ratio of two). The overall accuracy of the HO₂* measurements was ±36% (2σ). On all days except 22 July, HO₂* data were collected for 1 minute every 30 minutes and OH was measured during the rest of the 30 minute cycle. On 22 July, OH was not measured and instead the FAGE instrument measured HO₂* continuously.

2.4 Supporting Measurements

Ambient NO₂ was measured using a separate CAPS monitor (Aerodyne Research) (Kebabian et al., 2007; Kebabian et al., 2008). The standard 450 nm bandpass filter used by the CAPS monitor was replaced with a 470 nm bandpass filter to eliminate any interference by glyoxal and methyl glyoxal (Kebabian et al., 2008). This reduced the sensitivity by approximately a factor of three but still provided high signal-to-noise ratios (>100) for the ambient measurements. O₃ was measured with a UV absorbance monitor (2B Technologies model 202). NO was measured using a Thermo Fisher chemiluminescence sensor (Model 42i Trace Level). NO, NO₂, and O₃ data were averaged to 1 minute. Additional details regarding the calibrations and baseline measurements for the NO, NO₂, and O₃ measurements can be found in the SI.

A wide variety of biogenic and anthropogenic VOCs including isoprene and its oxidation products (methyl vinyl ketone and methacrolein), monoterpenes, non-methane hydrocarbons (C₂–C₅ and C₆–C₁₂), including aromatics, and oxygenated VOCs (alcohols, aldehydes and ketones) were measured during IRRONIC. An online GC-FID-FID was used to measure 57 NMHCs (Badol et al., 2004). Ambient air was sampled through a NAFION membrane and NMHCs were trapped at a temperature of -30 °C inside a quartz tube filled with Carbosieve SIII and Carbopack B. A thermodesorption unit (Perkin Elmer, ATD 400) was used to inject the sample into two columns (PLOT alumine and CPSil 5CB) to separate C₂–C₆ and C₆–C₁₂ compounds. Two FID detections provided limits of detection of 10–60 pptv at a time resolution of 90 min.

A second online GC-FID instrument was used to measure ethanol, isopropanol, methylethylketone and a few monoterpenes (α-pinene, 3-carene) (Roukos et al., 2009). A sampler unit (Markes International, air server Unity 1) allowed continuous sampling of ambient air through a trap held at 12 °C and filled with Carbopack B and Carbopack X. After thermodesorption,
the GC separation was performed using a high-polarity CP-Lowox column (Varian, France). Limits of detection reached with this instrument were in the range 10–90 pptv for a time resolution of 90 min. Offline sampling was performed on multisorbent cartridges to measure > C9 anthropogenic compounds (alkanes and aromatics) and monoterpenes (pinenes, terpinenes, limonene, ocimene, terpinolene, camphene, myrcene, borneol, camphor, cumene), and on DNPH (DiNitroPhenyldyrazine) cartridges to measure carbonyls, including formaldehyde (which was not measured by the GC-FID system), acetaldehyde and higher compounds. The cartridge measurements were integrated over 2-h sampling periods. Technical details can be found in (Ait-Helal et al., 2014; Detournay et al., 2011; Detournay et al., 2013).

Zero-dimensional photochemical modeling of this field campaign data was performed using the Framework for 0-Dimensional Atmospheric Modeling (F0AM) which was constrained by the 30 minute average mixing ratios of the supporting measurements (Wolfe et al., 2016). Measured VOC concentrations (every 90 min) were interpolated on to this 30 min time resolution. Carbon monoxide was not measured but instead estimated based on emission ratios of CO with benzene (Warneke et al., 2007). F0AM was executed using four different chemical mechanisms: two versions of the Regional Atmospheric Chemistry Mechanism (RACM2 and RACM2-LIM1) and the Master Chemical Mechanism (MCM 3.2 and 3.3.1). RACM2 groups various compounds based on similar rates of reaction resulting in 363 reactions from 17 stable inorganics, 4 inorganic intermediates, 55 stable organics, and 43 intermediate organics (Goliff et al., 2013). RACM2-LIM1 incorporates the revision to the isoprene oxidation mechanism (Peeters et al., 2009) that includes the Leuven Isoprene Mechanism (LIM) including a 1,6 H-shift and a 1,5 H-shift for isoprene peroxy radicals. MCM is a near-explicit chemical reaction model resulting in approximately 17000 reactions from 6700 radical species from methane and 142 non-methane species. Similar to the LIM1 mechanism, MCM 3.3.1 was updated to include revisions to the isoprene oxidation mechanism resulting in HOx recycling from peroxy radical H-shift isomerization as well as NOx recycling and updated ozonolysis rate constants.

3 Results and Discussion

3.1 Calibration comparisons between ECHAMP and IU calibration source

On 24 and 26 July the IU calibration source was positioned so that its output overflowed the ECHAMP inlet. Figure 2 compares the response of ECHAMP to variable concentrations of peroxy radicals generated by the IU calibrator. Concentrations of peroxy radicals were varied by adjusting the mixing ratio of water or by changing the intensity of the UV lamp. H2O mixing ratios varied from 0.1 to 1.4%, corresponding to relative humidities between 5 and 45% and F values between 28 and 17. A bivariate fit (York et al., 2004) between the ECHAMP measurements and the concentrations calculated by eq. 1 results in the relation ECHAMP = (0.88 ± 0.02) × (IU cal source) + (6.6 ± 4.5) ppt with an R² of 0.99. If both instrument’s calibrations were perfectly accurate, however, the slope would not be expected to equal unity because the two instrument’s calibration methods do not produce the same type of peroxy radicals. ECHAMP is calibrated with the acetone photolysis method, which produces an equimolar mixture of CH3O2 and CH3C(O)O2 radicals (Wood and Charest,
Because a calculated 10% of both of these radicals will be converted to CH\textsubscript{3}ONO in the reaction chambers and not be detected, ECHAMP is expected to be 11% (1/0.9) more sensitive to HO\textsubscript{2} than to CH\textsubscript{3}O\textsubscript{2} and CH\textsubscript{3}C(O)O\textsubscript{2}. Moreover, ECHAMP is expected to be between 7 to 12% less sensitive to RO\textsubscript{2} from butane and isoprene than to HO\textsubscript{2} because of the respective alkyl nitrate yields for both peroxy radicals: 8% for butane and 7 – 12% for isoprene (Atkinson et al., 1982; Lockwood et al., 2010; Patchen et al., 2007; Paulot et al., 2009). Thus if both instruments’ calibrations were perfectly accurate, then the expected slope for the calibration comparison using butane (i.e., 50% HO\textsubscript{2} and 50% C\textsubscript{4}H\textsubscript{9}O\textsubscript{2}) would be 1.07 (i.e., 1.11 × 0.96) and the expected slope when using isoprene would be between 1.07 and 1.04 depending on the isoprene alkyl nitrate yield. These values differ from the observed slope of 0.88 by 18 to 22%.

The 2σ analytical uncertainty for the IU calibration source and ECHAMP measurements are 36% and 27%, respectively. Because the IU calibration source’s O\textsubscript{3} mixing ratios were determined by ECHAMP, however, a portion of these two uncertainties is correlated. The uncertainty bars in Fig. 2 have been reduced to remove this component of the uncertainty - to 23% for IU (Dusanter et al., 2008) and 26.6% for ECHAMP. The 18 to 22% difference between the observed slope of 0.88 and the expected slope of 1.04 to 1.07 is within the adjusted uncertainties of both the ECHAMP measurements and the IU calibration source. Moreover, that ECHAMP evidently has near identical sensitivity to these two types of organic peroxy radicals demonstrates that differences in the mechanisms for converting RO\textsubscript{2} to HO\textsubscript{2} between β-hydroxy and alkyl peroxy radicals do not appear to affect their detection by ECHAMP.

The excellent linearity of Fig. 2 is notable because the calibrations were performed over a range of relative humidity values, each of which requires a different amplification factor to be used by ECHAMP. If the RH-dependence of the ECHAMP calibration had been ignored and only the dry calibration factor been used instead, the comparison would have been inferior as indicated by the squares in Fig. 2, for which a linear fit (not shown) gives the relation ECHAMP = 0.69(IU \textsuperscript{20}cal source) + 10.8 ppt. This serves as evidence that RH-dependent calibrations are indeed needed for producing accurate results from chemical amplifiers, including traditional CO and NO\textsubscript{2}-based instruments (e.g., PERCA).

### 3.2 Ambient concentrations of total peroxy (XO\textsubscript{2}) radicals, trace gases, and meteorological parameters

Ambient concentrations (15-minute averages) of XO\textsubscript{2}, isoprene, ethene, O\textsubscript{3}, NO, and NO\textsubscript{2}, along with meteorological parameters are shown in Fig. 3.

The 15-min average XO\textsubscript{2} concentrations in the daytime ranged from below the detection limit of ~5 ppt to 77 ppt. Among the VOCs measured, the daytime concentrations of low-molecular weight total alkanes (C\textsubscript{2}-C\textsubscript{5}) were the highest (average mixing ratio ± 1 standard deviation: 5.7 ± 3.9 ppb) followed by isoprene (3.6 ± 1.9 ppb), total C2-C5 alkenes (1.1 ± 0.3 ppb), high-molecular-weight alkanes (C\textsubscript{6}-C\textsubscript{14}, 0.3 ± 0.2 ppb), toluene (0.1 ± 0.1 ppb) and monoterpenes (0.1 ppb). NO concentrations typically peaked at 0.2 to 0.8 ppb between 09:00 - 11:00 and were almost always below 0.2 ppb between
12:00 and 21:00, whereas NO\textsubscript{2} concentrations in the daytime ranged between 0.3 to 3 ppb. \textit{O}_{3} concentrations varied between 0 to 71 ppb (av. 35.0 ± 8.4 ppb).

Measured XO\textsubscript{2} concentrations during IRRONIC exhibited a diurnal profile characterized by low mixing ratios (often below detection limit) between 0:00 – 07:00, increasing values from 07:00 to 13:00, peak values between 13:00 and 16:00, followed by a decrease in the late afternoon, similar to past measurements in other forests (Burkert et al., 2001; Hewitt et al., 2010; Mihele and Hastie, 2003). XO\textsubscript{2} mixing ratios were generally positively correlated with concentrations of isoprene, total alkenes, and ozone (Fig. 3). The highest XO\textsubscript{2} concentrations of over 60 ppt were measured during the afternoon of 24 and 25 July, coinciding with the highest average concentrations of isoprene (4.4 ppb), total alkenes (1.8 ppb), and \textit{O}_{3} (61 ppb), and the lowest average concentration of NO (0.1 ppb). The lowest daytime concentrations of XO\textsubscript{2} were observed on 13 July and 15 July, which were also characterized by lower isoprene and ozone mixing ratios and higher NO\textsubscript{2} mixing ratios.

We compare our XO\textsubscript{2} concentrations with reported XO\textsubscript{2} and HO\textsubscript{2}\textsuperscript{*} concentrations from other forests. The observed daytime XO\textsubscript{2} mixing ratios (campaign daytime average 26 ppt) at the IRRONIC site at Indiana are similar to those reported in a tropical rain forest in Malaysia (range 2-68 ppt) (Hewitt et al., 2010), in a northern Michigan forest during several intensive campaigns (range 8-65 ppt) (Griffith et al., 2013; Mihele and Hastie, 2003), and in a tropical forest over South America (campaign av. 42 ppt) (Lelieveld et al., 2008). XO\textsubscript{2} concentrations at Indiana never exceeded 80 ppt, in contrast to studies in which measured peroxy radical mixing ratios sometimes exceeded 150 ppt (Burkert et al., 2001; Wolfe et al., 2014).

Measurements of peroxy radical and NO concentrations enable ozone production rates to be calculated directly rather than rely on photochemical models. Using the measured concentrations of peroxy radicals and NO, calculated ozone production rates at the IURTP were at most 9 ppb/hr and described more in the SI.

### 3.3 Comparisons of Ambient Peroxy Radical Mixing Ratios

Figure 4 compares ambient [XO\textsubscript{2}] measurements by ECHAMP (30-minute averages) with the [HO\textsubscript{2}\textsuperscript{*}] measurements by LIF-FAGE (1-minute average every 30 minutes) during 13-25 July. Only data from days in which both instruments were operational are shown. No adjustments have been made to either of the datasets in Fig. 4 (or any other figures) to account for the calibration difference. Although in general it is preferable to compare measurements with equal time averaging, the precision of ECHAMP during this campaign – typically 2.5 ppt (1σ) for the 1.5 minute average measurements – necessitated this averaging. The diurnal profiles of both measurement sets, divided into 30-minute bins, are displayed in Fig. 5. Both figures indicate that the ECHAMP and LIF-FAGE measurements are in general well correlated and follow the same diurnal trend, though closer inspection reveals significant day to day and even hour to hour variability in the ratio.
The “true” [XO\textsubscript{2}]/[HO\textsubscript{2}^*] ratio, i.e., the ratio that would be produced by the two instruments’ measurements if they were calibrated to the same source and operated exactly as expected without any uncharacterized interferences or losses, depends on the composition of the peroxy radicals. As described in Section 2 (Experimental Methods), for both ECHAMP and LIF-FAGE, the sensitivity of the instrument to individual RO\textsubscript{2} compounds depends on the R-group and is characterized by the parameter “α”, which is the instrument’s sensitivity to each RO\textsubscript{2} relative to its sensitivity to HO\textsubscript{2}. For ECHAMP α is determined largely by the fraction of RO\textsubscript{2} that is converted to alkyl nitrates (RONO) and alkyl nitrates (RONO) following reaction with NO at atmospheric pressure. For LIF-FAGE, α is mostly determined by how quickly each RO\textsubscript{2} is converted sequentially to HO\textsubscript{2} and then OH following reaction with NO after the expansion of the sampled gas into the low-pressure region of the instrument (Fuchs et al., 2011; Lew et al., 2018). Air in which CH\textsubscript{3}O\textsubscript{2}, CH\textsubscript{3}C(O)O\textsubscript{2}, and small (<C\textsubscript{5}) alkyl peroxy radicals have a large contribution to the total peroxy radical concentration would thus produce a relatively high [XO\textsubscript{2}]/[HO\textsubscript{2}^*] value, since ECHAMP is sensitive to those peroxy radicals (α>0.9) whereas the LIF-FAGE HO\textsubscript{2}^* measurement is not (α<0.1). In contrast, air with a relatively high fraction of alkene-derived RO\textsubscript{2} (e.g., isoprene Peroxy radicals), for which both ECHAMP and LIF-FAGE HO\textsubscript{2}^* α values are near one, would be expected to lead to lower [XO\textsubscript{2}]/[HO\textsubscript{2}^*] values (i.e., closer to unity).

A bi-variate linear regression of the measured XO\textsubscript{2} and HO\textsubscript{2}^* concentrations between 09:00 and 22:00 yields the relationship [XO\textsubscript{2}] = (4.08 ± 0.05) [HO\textsubscript{2}^*] – (1.4 ± 0.3) ppt (Fig 6). The regression is restricted to this window of time because of the degraded precision of the ECHAMP measurements at night due to the higher relative humidity. The [XO\textsubscript{2}]/[HO\textsubscript{2}^*] slopes were highest on the last two days of measurements – 24 and 25 July, with slopes of 1.25 and 1.08, respectively, or 1.5 and 1.3 after adjusting for the calibration difference. These two days were characterized by the highest mixing ratios of peroxy radicals, O\textsubscript{3}, isoprene, and the anthropogenic VOCs ethene and ethyne. The lowest [XO\textsubscript{2}]/[HO\textsubscript{2}^*] ratios were observed on 13 July during which a passing thunderstorm led to low concentrations during mid-day with higher values before and after the storm. The higher [XO\textsubscript{2}]/[HO\textsubscript{2}^*] ratios observed later in the field campaign may simply be the result of a change in sensitivity in one of the instruments. These linear regressions are difficult to interpret, however, since the XO\textsubscript{2} measurements are 30 minute averages and the HO\textsubscript{2}^* measurements are 1-minute averages taken every 30 minutes.

A regression of the binned data shown in Fig. 5 gives the relation [XO\textsubscript{2}] = 1.0 ± 0.14 [HO\textsubscript{2}^*] + (4.5 ± 1.6) ppt; accounting for the calibration difference gives an adjusted slope of 1.2. The [XO\textsubscript{2}]/[HO\textsubscript{2}^*] ratio using the binned data was highest between 9:45 and 10:45 (Fig. 5), but was between 0.9 and 1.1 between 14:45 and 19:15. This overall temporal trend is apparent in several days (Fig. 4). Applying a 30-min offset to the XO\textsubscript{2} data largely removes this trend and leads to fewer time periods when [XO\textsubscript{2}]/[HO\textsubscript{2}^*] was less than 1.0, but such an offset does not agree with the synchronized time-base of both measurements. The two instruments’ different averaging times and precision levels preclude further assessment and conclusions regarding possible time offsets.

To further investigate the effect of this different averaging on the comparison, on 22 July the IU-LIF-FAGE instrument operated in HO\textsubscript{2}^*-only mode (i.e., with no time devoted to measuring OH). We compare the resulting 1-minute and 15-minute averaged HO\textsubscript{2}^* measurements to the 1.5 minute and 15-minute averaged XO\textsubscript{2} measurements (Fig. 7).
Between 15:00 and 17:00, the HO$_2$ measurements increased from 50 to 70 pptv and decreased back to 50 pptv while the XO$_2$ measurements were relatively invariant at 40 pptv. Ignoring the difference between the average mixing ratios, this difference in the temporal profile of the two instruments’ measurements result could only be “real” if there were changes in the peroxy radical relative composition on this two-hour time scale, e.g. a simultaneous increase in HO$_2$ and a decrease in alkyl peroxy radicals, such that [HO$_2$] actually did increase while the mixing ratio of total peroxy radicals was almost constant. Measurements of VOC composition and NOx do not support such a fast change in peroxy radical composition, suggesting that these observations were more likely the result of an instrumental issue. Currently we are unable to identify the exact cause of this observation, but possible explanations are a transient interference in the HO$_2$ measurements. Due to gaps in the NO data because of problems with the Thermo chemiluminescence sensor, the model was run for these three days, and also a diurnal profile for the entire campaign was run using diurnal average concentrations of constrained species. From these model results we calculate a “true” [XO$_2$]/[HO$_2$] ratio, i.e., the ratio that would be produced by the two instruments’ measurements if they were calibrated to the same source and operated exactly as expected without any uncharacterized interferences or losses, depends on the composition of the peroxy radicals. As described in Section 2 (Experimental Methods), for both ECHAMP and LIF-FAGE, the sensitivity of the instrument to individual RO$_2$ compounds depends on the R-group and is characterized by the parameter “α”, which is the instrument’s sensitivity to each RO$_2$ relative to its sensitivity to HO$_2$. For ECHAMP α is determined largely by the fraction of RO$_2$ that is converted to alkyl nitrates (RONO) and alkyl nitrates (RONO) following reaction with NO at atmospheric pressure. For LIF-FAGE, α is mostly determined by how quickly each RO$_2$ is converted sequentially to HO$_2$ and then OH following reaction with NO after the expansion of the sampled gas into the low-pressure region of the instrument (Fuchs et al., 2011; Lew et al., 2018). As which CH$_2$O, CH$_3$CO, and small (<C$_5$) alkyl peroxy radicals have a large contribution to the total peroxy radical concentration, this ratio would thus produce a relatively high [XO$_2$]/[HO$_2$] value, since ECHAMP is sensitive to those peroxy radicals (α≈0.9) whereas LIF-FAGE HO$_2$ measurement is not (α<0.1). In contrast, air with a relatively high fraction of alkene-derived RO$_2$ (e.g., isoprene peroxy radicals), for which both ECHAMP and LIF-FAGE HO$_2$ values are near one, would be expected to lead to lower [XO$_2$]/[HO$_2$] values (i.e., closer to unity). 

Because the composition of the peroxy radicals during IRRONIC is not exactly known, we examine the predicted speciation generated by zero-dimensional photochemical modeling of the IRRONIC dataset using two versions of the Regional Atmospheric Chemistry Mechanism (RACM2 and RACM2-LIM1) and the Master Chemical Mechanism (MCM 3.2 and 3.3.1). A full comparison of the modeled and measured concentrations is beyond the scope of this paper; we use these model outputs mainly to inform the discussion of the relative speciation of total peroxy radicals and its relation to the expected and measured [XO$_2$]/[HO$_2$] ratio. A fuller description of the photochemistry at this site, including OH reactivity measurements, will be described in a companion paper (Lew et al., in preparation).

The accuracy of the model results is, of course, subject to how comprehensive and accurate the supporting measurements and underlying chemical mechanisms are, but nonetheless help to frame the interpretation of the two instruments’ measurements. Due to gaps in the NO data because of problems with the Thermo chemiluminescence sensor, there are only three days for which we have model results and measured peroxy radical concentrations by both ECHAMP and LIF-FAGE – on the 16th, 22nd, and 24th of July. The model was run for these three days, and also a diurnal profile for the entire campaign was run using diurnal average concentrations of constrained species. From these model results we calculate the expected values measured by ECHAMP and LIF-FAGE based on each instrument’s relevant values for α:

$$\text{ECHAMP } [\text{XO}_2]_{\text{EXPECTED}} = [\text{HO}_2] + 0.9([\text{CH}_3\text{O}_2]) + 0.92([\text{C}_3\text{H}_6\text{O}_2\text{O}_2]) + 0.9([\text{CH}_3\text{C}(\text{O})\text{O}_2]) + 0.9(\text{Other})$$ (4) 

$$\text{LIF-FAGE } [\text{HO}_2^*]_{\text{EXPECTED}} = [\text{HO}_2] + 0.05([\text{CH}_3\text{O}_2]) + 0.83([\text{C}_3\text{H}_6\text{O}_2\text{O}_2]) + 0.05([\text{CH}_3\text{C}(\text{O})\text{O}_2]) + 0.7(\text{Other})$$ (5) 

The “Other” category includes all types of peroxy radicals, e.g., from monoterpenes, methy vinyl ketone, ethene, etc. The α values for ECHAMP are based on the calculated yields of alkyl nitrates and alkyl nitrates as described in section 2.2. For LIF-FAGE, the α value for C$_3$H$_6$(OH)O$_2$ was measured and α for CH$_3$O$_2$ and CH$_3$C(O)O$_2$ are based on measured yields from several similar instruments, all of which have measured values less than 5%. An α of 0.7 is assumed for the “other” category since most alkenes have α values between 0.5 and 0.9, and small alkanes, which have lower values, account for a small portion of the OH reactivity (Lew et al., in preparation).
The comparison between measured [XO\*]/[HO\*\_] ratio modeled by MCM 3.2 and measured using all days when there were both XO and HO\*\_ measurements. Between 10:00 and 18:00 the modeled [XO\*]/[HO\*\_] ratio varied between 1.2 and 1.5, whereas the measured ratio varied between 0.9 and 1.4, with a greater amount of variability from hour to hour. Increasing the observed ratio by 20% to account for the calibration comparison (section 3.1) gives an adjusted measured ratio of between 1.1 and 1.7. The highly variable ratios during nighttime mainly reflect the lower signal to noise ratios of both instruments when peroxy radical concentrations were low (less than ~5 ppt).

Measured and MCM 3.2 modeled concentrations for 16, 22, and 24 July are shown in Fig. 8. On all three days the relative contributions from the various types of peroxy radicals are comparable. At 15:30 – when concentrations were highest – the modeled peroxy radicals comprised 30% C\(_2\)H\(_5\)(OH)\(_2\), 25% HO\_, 26% CH\(_3\)O\(_2\), and 7% CH\(_3\)C(O)O\(_2\). The four chemical mechanisms vary little in the predicted relative speciation (SI). The [XO\*]/[HO\*\_] ratio modeled by MCM 3.2 between 15:00 and 16:00 is 1.4 for 16 and 22 July and 1.45 on 24 July. The measured [XO\*]/[HO\*\_] ratio is close to unity on 16 and 22 July, and between 1.2 and 1.5 on 24 July. Increasing these measured ratios by 20% to account for the calibration comparison produces adjusted measured [XO\*]/[HO\*\_] ratios of 1.2 on 16 and 22 July and 1.4 to 1.8 on 24 July. After accounting for the 20% calibration difference, the modeled and measured ratios agree to within the experimental and model uncertainties.

Although all four chemical mechanisms predict a very similar relative speciation, there are variations in the absolute peroxy radical concentrations predicted. MCM 3.3.1 concentrations are very similar to those from MCM 3.2, but RACM2 and RACM2-LIM1 predict 26% and 42% higher peak concentrations, respectively. Measured [XO\_] mixing ratios are 20 to 30% lower than the MCM 3.2, [XO\_] on 16 and 22 July but agree more closely on 24 July (measured/modelled ratio varies from 0.8 to 1.15). The comparison between measured [HO\*\_] and modeled [HO\*\_] for these three days exhibits more variability (Fig. 8). Further details can be found in the SI.

Observations of [XO\*]/[HO\*\_] ratios less than one were observed during parts of 13, 17, and 18 July and even after increasing by 20% to account for the calibration comparison do not seem reasonable or in some cases even possible. These observations were most likely caused by issues with one or both instruments. Two possible causes that warrant investigation in subsequent field measurements are discussed below:

1. Error in the ECHAMP calibration, especially for RH values greater than 45%. Although the calibration comparison presented in section 3.1 show that the ECHAMP and LIF-FAGE instrument’s calibrations agreed to within measurement uncertainties, that is not necessarily true for RH values higher than those used during those calibration tests. The highest RH value during the calibration comparisons was 45%, whereas the daytime minimum RH values between 12:00 and 16:00, when measured [XO\_] and [HO\*\_] were both highest, were typically between 45% and 65% (Fig 1). Furthermore, we cannot prove that the ECHAMP calibration was invariant from day to day. We include potential sampling losses to be a part of the overall ECHAMP calibration.
2. Interferences in the LIF-FAGE measurement. The comparison of high temporal resolution in Fig. 7 revealed differences in the temporal profile of the LIF-FAGE and ECHAMP sensor. If these were caused by an interference in the LIF-FAGE measurement when sampling ambient air, then it would follow that the two instruments would agree when sampling a calibration source but differ when sampling ambient air.

As discussed earlier, the RH-dependence of the sensitivity of chemical amplifiers has recently been questioned (Sommariva et al., 2011). Had we ignored the RH dependence for ECHAMP’s amplification factor and simply used the value under dry conditions, the daytime XO$_2$ values would have been roughly 50% lower than those presented in this paper, leading to unrealistically low [XO$_2$]/[HO$_2$*] ratios of ~0.5.

4. Conclusions

The results of this comparison of the IU calibration source and the ambient measurements of peroxy radicals by ECHAMP and LIF-FAGE provide encouraging first results that the newly developed ECHAMP method can be used for ambient measurements of total peroxy radicals. The ECHAMP measurements, based on the acetone photolysis method, and the IU water vapor photolysis calibration source agreed within 12%, within the experimental uncertainties. The measured mixing ratios of XO$_2$ and HO$_2$* were usually lower than the concentrations predicted by the RACM2, RACM2-LIM1, MCM v. 3.2, and MCM v. 3.3.1 chemical mechanisms. The measured [XO$_2$]/[HO$_2$*] ratios usually differed from the ratios predicted by zero-dimensional photochemical modeling by less than the combined measurement and modeling uncertainties, though the lowest ratios observed (0.8) are not physically meaningful and therefore must be due to measurement errors.

An attribute of these comparison exercises is that the two instruments operate on very different measurement principles and the calibration methods differ greatly. Although the calibration comparison was favorable, due to the time required to conduct successful calibrations with the acetone photolysis method and its overall inconvenience (Wood and Charest, 2014) we have discontinued its use. For subsequent field measurements we have used the water vapor photolysis method and another method based on methyl iodide photolysis (Anderson et al., 2019; Clemitshaw et al., 1997; Liu and Zhang, 2014). All three calibration methods do indicate that a humidity-dependent calibration must be used for both CO-based and ethane-based chemical amplifiers.

Data Availability

Data are available upon request from the corresponding author (Ezra.Wood@drexel.edu)
Author contributions.

EW and PS designed the research project. SK, BD, and EW were responsible for the ECHAMP measurements and supporting measurements of NO, NO₂, and ML, BB, PR, and PS were responsible for the LIF-FAGE measurements and photochemical modeling. SD, SS, TL, and NL were responsible for the measurements of VOCs. SK and EW conducted the analysis and wrote the paper with feedback from all co-authors.

Competing Interests. The authors declare that they have no conflicts of interest.

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Figure 1. Map of the sampling site. The star symbol represents the Indiana University Research and Teaching Preserve (IURTP) in Bloomington, Indiana, USA. The arrow represents a distance of 1 km.
Figure 2. Results of the calibration comparison in which ECHAMP measured the total peroxy radical concentration in the output of the IU calibration source. The error bars indicate $2\sigma$ uncertainties of the ECHAMP measurements and IU calibration source, adjusted for the fact that the IU actinometry was based on the ECHAMP NO$_2$ calibration. The slope of the dotted line is unity.
Figure 3. Time series data of measured chemical and physical parameters during IRRONIC. Except where noted, all measurements are in ppb. The sum of $[\text{RO}_2]$ and $[\text{HO}_2]$ was measured by the ECHAMP instrument, with a detection limit typically between 1 and 2 ppt (signal-to-noise ratio of two). The vertical grid lines indicate midnight for odd-numbered days, in local time.
Figure 4. Concentrations of ambient total peroxy radicals (\( \text{XO}_2 \)) by ECHAMP and \( \text{HO}_2^* \) by IU-LIF-FAGE. 30-minute averaged measurements are shown for ECHAMP \( \text{XO}_2 \). For \( \text{HO}_2^* \), measurements are 1-minute averages every 30 minutes. The vertical grid lines indicate midnight for odd-numbered days, in local time.
Figure 5. Lower plot: Mean diurnal profile of ECHAMP XO\textsubscript{2} and IU-LIF-FAGE HO\textsubscript{2}* measurements for the 9 days in which both instruments were operational. The HO\textsubscript{2}* values are displayed with a 6 minute horizontal offset for clarity. The error bars indicate the ± one standard deviation of the measured concentrations in each 30-minute time bin during those nine days. The upper plot shows the [XO\textsubscript{2}]/[HO\textsubscript{2}*] ratio - both measured by the two instruments and modeled using the MCM 3.2 chemical mechanism. The measured ratio is only shown for time periods between 09:00 and 22:00 due to the poor signal-to-noise ratios for the night-time measurements.
Figure 6. Correlation of ambient $[\text{XO}_2]$ measured by ECHAMP with $[\text{HO}_2^*]$ measured by IU-LIF-FAGE. The linear fit is for data between 09:00 and 22:00, indicated by the points with green circles. The equation of the fit is $[\text{XO}_2] = (1.08 \pm 0.05) [\text{HO}_2^*] - (1.4 \pm 0.3)$ ppt.
Figure 7. Time series comparing IU LIF-FAGE HO$_2^*$ and ECHAMP XO$_2$ measurements from 22 July, 2015 when the IU LIF-FAGE instrument was run in HO$_2^*$-only mode.
Figure 8. Peroxy radical mixing ratios measured by ECHAMP and LIF-FAGE and modeled by MCM v3.2.
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