This paper investigated the impact of molecular structure on the photooxidation of aromatic SOAs. The effect of the alkyl substitutes on the yield of chamber generated SOA, the chemical composition and the physical properties of SOA were studied. The authors concluded that oxidation of products promote the elevation of SOA yields. They found that the aromatic oxidation increased with increasing alkyl substitute chain length and it also varies with the branching position of an alkyl group on aromatic ring. Using their chamber data, the authors classified aromatics into five groups and ranked them as ortho (o-xylene and o ethyltoluene) > one substitute (ethylbenzene, propylbenzene and isopropylbenzene) > meta (m-xylene and m-ethyltoluene) > three substitute (trimethylbenzenes) > para (p-xylene and p-ethyltoluene).

Overall, all data of this paper need statistical evaluation providing uncertainties in processed data or the errors associated with data points in Figures and Tables. The comparison of data from different aromatic systems and the data interpretation should be based on statistical significance. The explanation of SOA yields and the processed data from analytical instrument are very empirical and needs better interpretation with rationales based on kinetic mechanisms in both the gas phase and aerosol phase. The authors should provide better atmospheric imprecation of the observation and the outcomes of this study based on SOA formation in ambient environments and the classification of aromatic hydrocarbons in current air quality models.

The statistical evaluation is addressed in the revised manuscript and a detailed analysis of the statistical evaluation are included in the replies to Comment 5, 10 and 11. Additional mechanism discussion is included in the revised manuscript to provide interpretation for the results. See replies to Comment 9 and 12 for details.

The implications of this work to ambient environments and air quality models is further enhanced as described in Comment 13.

A point by point reply to the reviewer is listed below.

Comments:
1. Page 6, line 5-10. The SOA formation has been performed at very dry conditions (RH<0.1%), which is very different from ambient environment. The SOA formation can be affected by humidity. Particularly, heterogeneous reactions is sensitive to aerosol water content because some of carbonyls and epoxides can be hydrated with available water in aerosol and oligomerized. Such reactions are also influenced by aerosol compositions and the hygroscopic properties of aerosol. In addition to the reactivity of oxidized carbons in aerosol, the aerosol phase water can modulates the reduction of viscosity of aerosol media, which also affect aerosol growth. Thus, the order in aerosol growth determined at very dry conditions may be/may not be different from aerosol growth in higher humidities. The authors should discuss
about the potential influence humidity on aerosol growth and the rank observed in this study.

We agree SOA formation is potentially affected by humidity due to heterogeneous reaction. The current study provides a fundamental relationship among SOA formed from different aromatic isomers under dry conditions.

We add the following sentences in Page 19, Line 7 (end):
Previous studies found that the humidity insignificantly impacts SOA yield from aromatic hydrocarbons (Cocker, et al., 2001) or maintains the SOA yield relationship between isomers (Zhou, et al., 2001). Therefore, it is predicted that the observation found under dry conditions in this study, especially the molecular structure impact on SOA formation from different aromatic isomers could be extended to atmospherically relevant humidity conditions. However, recent studies observe that the hydration of carbonyls and epoxides could lead to further heterogeneous reaction and oligomerization (Jang, et al., 2002; Liggo, et al., 2005; Minerath and Elrod, et al., 2009; Lal, et al., 2012). It is possible that aerosol compositions and the hygroscopic properties could be altered after the heterogeneous reactions, especially under humid conditions. The impact of molecular structure impact on SOA formation under humidity condition needs to be further studied to extend the findings in the current work.

2. 2nd paragraph, Section 2.2. The aerosol samples were evaporated at 600°C followed by impaction. Such high temperature promotes charring of organic compounds and is able to modify chemical compositions of organic compounds. This should be clarified.

The AMS used in our study followed standard measurement techniques for SOA studies. Heating in the HR-TOF-AMS is conducted under vacuum, which minimizes “charring” of the organic species. We add a “under vacuum” at page 7 line 12 (2nd paragraph, Section 2.2) for clarification. The sentence now reads
“The sample was vaporized by a 600 °C oven under vacuum followed by a 70 eV electron impact ionization.”

3. For Table 1, the author need to provide the uncertainty associated with SOA yields. Please provide the unit with variables instead of footnote. All Figures and Tables need the uncertainties or the errors associated with data.

We insert following sentence addressing SOA uncertainty in the methods section at Page 8 Line 9: “The uncertainty associated with 10 replicate m-xylene and NO experiments SOA yield is <6.65%.”

The standard deviations of O/C, H/C, f44, f’43+57+71, OS, and VFRend are now included in the revised manuscript along with the standard deviations shown in the original manuscript for density.

We modified the manuscript to provide the unit with variables instead of footnotes as requested.
4. Section 3.1 SOA yield. (a) For SOA yields, the authors employed the two product model that was derived by Odum et al. The experimentally observed SOA yield varies with VOC/NOx ratios and NO/NO2 ratios. The authors describe that HC/NO ratios range 11.1 to 171. Does the aerosol at HC/NOx = 11 have the same chemical composition of the SOA at HC/NOx=171? Although all experimental conditions fall into the high NOx regions but they are various. Please discuss about the potential effect of NOx on SOA formation within the experimental conditions of this study. (b) The HC/NOx ratios between aromatic systems in Table are not same. Without the rationale for the effect of HC/NOx on SOA yields, the rank of SOA production from different SOA loses its meaning. (b) SOA yields are influenced by the amount of initial hydrocarbon and its reaction rate. When the aerosol is quickly formed, the loss of gaseous oxidized carbons to the reactor wall becomes smaller (Ng et al, 2007). The authors should clarify whether aerosol yields between different systems are not biased due to the potential loss of gaseous compounds to the wall by different reaction rates and initial experimental conditions. (c) In addition to the kinetic reactivity of hydrocarbons, the determination of SOA yields are subjective to the duration of chamber operation. The authors need to explain how the yield of SOA was determined based on reaction time and aerosol growth.

Only ~10% of experiments have HC/NO larger than 60 ppbC:ppb. The majority of experiments have similar NO conditions. NO could impact SOA formation by either forming organic nitrate or reducing peroxide radical concentrations. First, aerosol at HC/NO = 11 ppbC:ppb may have slightly different SOA chemical composition at HC/NOx=171 ppbC:ppb due to a higher organic nitrate fraction. However, less than 5% of organic matter is organic nitrate in this study and therefore exerts little impact on overall yield and chemical composition. Second, lower HC/NO might lead to lower peroxide radical concentrations and therefore lower SOA yield. However, no significant correlations between SOA yield and radical concentrations are observed as discussed at Page 9 Line 11-15 and presented in Table S5. This shows the lack of correlation between radical concentration and yield. Therefore, NO is not a major factor to SOA yield within the experimental HC: NO range used in this study.

Seeded experiments to minimize gaseous compounds wall loss were conducted in our chamber experiment with no significant difference observed between the seeded and non-seeded experiment. This indicates that the gas-phase wall might not be expected to be significant for experiments in our chamber for this aromatic SOA study.

We add the following sentence at line 8 Page 6 in the revised manuscript.

Seeded experiments to minimize wall effects have also been conducted in our chamber experiment with no measurable difference observed between the seeded and non-seeded experiment.

We agree that SOA yields are subjective to the duration of chamber operation. In this work, SOA yield is calculated after 6-8 hours of photooxidation for each experiment. All the precursors studied have similar kOH (Table S1) or similar kOH[Precursor] (Li, et al., 2016). We selected comparable photooxidation time (6-8 hours) for all precursors and therefore the SOA yields are comparable.
5. Page 8, lines 10-15 and Figure 1. It seems that the substitute length also affects the yield of SOA (C8 vs C9). The data should be treated by the statistical evaluation.

We agree with the reviewer that substitute length also affects the yield of SOA (C8 vs C9) as stated on Page 10, line 1-7. However, the difference caused by substitute length is less significant than substitute location as shown in Fig. 1. Differences in SOA yield due to length is within the SOA yield standard deviation (now shown). There may be differences among all one substitute aromatics, which is answered in detail in the reply to Comment 10. We add the following sentence at line 6 Page 10 in the revised manuscript. “However, the differences between xylenes and their corresponding ethyltoluenes are not statistically significant.”

Section 4 further discusses the differences associated with substitute length (original manuscript on Page 10, line 7.): “These differences are explained by the proposed alkyl group dilution effect (Sect. 4).”

6. Figure 2. Where are the actual data point from each aromatic hydrocarbon in Figure 2? Please include the maker for each data point and the uncertainty of data points.

The actual data points from each aromatic hydrocarbon in Figure 2 become obvious after adding standard deviation on both the x-axis and the y-axis as suggested in Comment 3.

7. 1st paragraph, Section 3.2.1. It is difficult to follow the description of mass fragmentation in the text. It would be better to organize mass fragmentation information using Table. How does the aerosol have carboxylic functional groups (CO2). Does carboxylic acid form via gas phase oxidation of hydrocarbons or autooxidaiton in particle phase? What are the precursor structures to produce carboxylic groups.

We add Table S6 to provide the requested peak information.

CO\textsubscript{2} is a common peak from AMS when measuring aerosol. It is less likely that CO\textsubscript{2} is associated with carboxylic functional group in this study. There are other possible CO\textsubscript{2} fragment sources. For example CO\textsubscript{2} may come from the oligomerization of small cyclic furanones described in earlier work (Li, et al, 2016).

8. Line 20-21, page 10. “While m/z 43 (C3H+7 ) and 57 (C4H+9 20 ) are often considered as markers for hydrogen-like organic aerosol . . .”. What does “hydrogen-like organic aerosol” mean? The explanation about mass fragmentation is unclear.

This is a typo. It should read “hydrocarbon-like organic aerosol” instead of “hydrogen-like organic aerosol”. An m/z 57 and m/z 71 in field studies or ambient atmosphere usually associate with hydrocarbon-like organic fragments, C\textsubscript{4}H\textsubscript{5} and C\textsubscript{5}H\textsubscript{11}, respectively (Zhang, et al., 2005; Ng, et al., 2010). However, m/z 57 and m/z 71 in chamber studies, especially for ethyl and propyl substituted aromatics photooxidation, are majorly C\textsubscript{3}H\textsubscript{5}O and C\textsubscript{4}H\textsubscript{7}O\textsuperscript{+}, which are oxygenated organic aerosol (OOA). Ng. et al (2010) developed f\textsubscript{43} (majorly C\textsubscript{2}H\textsubscript{3}O\textsuperscript{+}) and f\textsubscript{44} (CO\textsubscript{2}+) for ambient OOA categorization without m/z 57 and m/z 71 since m/z 57 and m/z 71
are majorly HOA in ambient atmosphere. However, C$_2$H$_3$O$^+$ (m/z 57) and C$_3$H$_5$O$^+$ (m/z 71) should also be included beside C$_2$H$_3$O$^+$ in SOA chamber studies as OOA to compare the oxidation of different aromatic hydrocarbons. We replace Line 20-24, page 10 with sentences below:

While m/z 57 (C$_4$H$_9^+$) and m/z 71 (C$_5$H$_{11}^+$) are often considered as markers for hydrocarbon-like organic aerosol in ambient studies (Zhang et al., 2015; Ng et al., 2010), oxygenated organic aerosol C$_3$H$_5$O$^+$ and C$_4$H$_7$O$^+$ are the major fragments at m/z 57 and m/z 71, respectively, (Fig. S4, Table S6) in current chamber SOA studies, especially during the photooxidation of ethyl and propyl substituted aromatics. Therefore, m/z 57 and m/z 71 are also considered beside C$_2$H$_3$O$^+$ at m/z 43 in SOA chamber studies as OOA to compare the oxidation of different aromatic hydrocarbons.

9. Last paragraph in section 3.2.1. The authors proposed that longer alkyl substitutes may not lower the oxidation per mass as further oxidation but observed the similar f44 and f43+57+71 of toluene suggesting unidentified oxidation. The interpretation of the analytical data is based on partitioning theory because authors’ interpretation focused on gas phase oxidation. The oxidized carbons such as carbonyl are reactive in aerosol phase and can be transformed into oligomeric matter. In the past, the characterization of SOA have shown that oligomeric matter was significantly contributed to aromatic SOA mass. The interpretation of the data here was very empirical. The authors needs to rationale for observed data with better interpretation based on kinetic mechanisms.

We suggest that longer alkyl substitutes may not lower the oxidation per mass by relying on both the observation in section 3.2.1 and the elemental ratio prediction method (alkyl dilution theory) in Discussion part (section 4). Higher OSc (lower H/C and higher O/C) is observed in longer chain single alkyl substitute aromatics than the alkyl dilution theory predicts from toluene data. Oligomerization, as suggested by the reviewer, consists of highly oxidized monomers (e.g. glyoxal) and could therefore also increase the overall OSc. Therefore, we add following sentence at Page 12 Line 17: It is also possible that oligomerization from highly oxidized carbonyls contribute more to the SOA formation from aromatics with long chain alkyl substitute.

Previous theoretical studies predict that alkyl substitute plays a role in oligomerization, e.g. glyoxal favors acetal oligomerization and methylglyoxal prefers aldol condensation to form oligers (Barsanti and Pankow, 2005; Krizner, et al., 2009). However, these two oligomerization kinetic mechanisms produces products with similar formulas. No matter which mechanism is favored, SOA elemental ratio should not be affected. Therefore, the percentage of oligomerization to other reaction mechanisms rather than the difference in oligomerization mechanism seems a better explanation for the observation in SOA formed from long chain single alkyl substitute aromatics.

Therefore, we also add the following sentences in the Discussion section at Page 18 Line 6: It is also possible that oligomerization from highly oxidized carbonyl component might be more favored for long chain single alkyl substituted aromatics.
10. Page 8, line 30 - Page 9, line 2, for Figure 1. It is uncertain whether the one alkyl substituent on the aromatic ring is clearly separated from meta-positioned aromatics without statistical significance.

We agree that one alkyl substituent on the aromatic ring SOA yield is not clearly separated from meta-positioned aromatics in Figure 1. However, it can be concluded from chemical composition and volatilities (VFR) that one alkyl substitute aromatic hydrocarbon is more oxidized than meta-positioned aromatics. The statistical parameter for the curve fitting is provided in the revised Table 2 as mentioned in the reply to Comment #5. It can be clearly seen that MSRE is much larger in one substitute aromatic fitting than meta-aromatic fitting. This indicates potential SOA yield differences within the one substitute aromatic hydrocarbons along with observed SOA chemical composition differences among one substitute aromatic hydrocarbons (Sections 3.3.2 and 3.2.3). Further studies are warranted to provide more information on SOA yield. At this time, we fit one alkyl substitute aromatics and meta-position aromatics separately.

11. Section 3.2.2 (H/C vs O/C). The data points in Figure S6 are scattered. The reviewer is sure how the authors chose the representative point from each diagram. The difference between the representative point from each diagram should be statistically determined. Without statistical assessment, it is hard to conclude the fact but it looks that difference in representative (averaged) points between systems would be insignificant within standard deviation of scattered data except few systems.

The scattering in Fig S6 is majorly due to a few early period data at the low mass loading when aerosol just starts to form, which have higher H/C and lower O/C than the latter time. However, o-xylene data contains data that does not follow the aerosol aging trend observed for the other isomers. Therefore, we delete the obvious outliers in Fig S6-f-o-xylene and Fig S6-6 o-Ethyltoluene and adjust Fig. 3, Fig. 4 and Fig 7 accordingly. We provide standard deviation for H/C, O/C and OS in Fig 4, Fig 7b and supplemental materials as answered in question 3.

12. 2nd paragraph, in page 13. The explanation about H/C and O/C ratio is empirical and need better interpretation in the point of kinetical mechanisms.

The kinetic parameters ($k_{OH}$) for initial oxidation of ortho, meta and para containing aromatic hydrocarbons are all similar (Table S1). This suggests there may not be a significant difference among all these isomers from kinetic perspective. We draw the conclusion in the paper empirically based on what we observed in our measurement.

13. Atmospheric Implication section. The authors need to provide the implication of the observation of this study to ambient environments. It has been known that toluene is the most abundant in urban areas where NOx and humidity are high. Can the observation of this work be applicable to ambient environments? In the current CMAQ, there are two class of aromatics: high and low yield aromatics. Toluene is currently in the group of high yield aromatics. How can the observation of this paper be applied to the current air quality model? Is the classification of five groups of aromatics in this study meaning? All chamber studies have been limited to high concentration of initial conditions (VOC and NOx) due to detection limit of
analytical instrument. What is the implication of this work to the SOA in the low concentration environments (ambient air)?

We address the relevance of NOX conditions in current study to the ambient atmosphere at the beginning of Atmospheric Implication section. Toluene remains the second highest SOA yield (lower than benzene, especially at higher mass loading, see Li, et al., 2016) precursor according to our earlier work (Li, et al., 2016); however, it is not a target aromatic in current studies. This paper focuses on isomer or molecular structure impact on SOA formation. Current work provides sufficient data to distinguish among para, meta and ortho position containing aromatics and therefore is able to subcategorize the previous “low” and “high” yield aromatics (e.g. only one p-xylene data point in Odum, et al., 1997). The five groups of aromatics and their two product modeling curve fitting provide the practical parameter for more detailed SOA modeling. All SOA yield data provided is under more atmospherically relevant NOX conditions than the earlier “high” and “low” yield work improving reliability of fit parameters as inputs to atmospheric models. Study at the lower atmospheric NOX concentrations provides yields twice as high as those from earlier work at very high NOX concentrations greatly impacting the model predictions from model prediction (e.g., CMAQ).

We add following sentences to emphasize the importance of this study to model and ambient environment at Page 19, Line 7:
Moreover, the five subcategories of aromatics and their two product modeling curve fitting parameters in this work at more realistic NOX loadings provide a more precise prediction of SOA formation form aromatic hydrocarbons under atmospheric conditions.

14. For Figure 5, the author could explain why the predicted density of 1,3,5-TMB is much lower than the measured density.

Thank you for this observation. In fact we observe density underestimation in all meta position containing aromatic hydrocarbons including m-xylene, m-ethyltoluene and 1,3,5-TMB. The underestimation is associated with a bias in elemental ratio analysis from AMS as discussed in Li, et al., 2016; Nakao, et al., 2013.

We add following sentences in Page 16 Line 13:
A comparatively large negative error is found in meta containing aromatic hydrocarbons including m-xylene, m-ethyltoluene and 1,3,5-trimethylbenzene. It is noted that there should be more alkyl substitutes in SOA formed from meta position aromatics than other aromatics since meta position alkyl substitutes are more likely to participate into SOA products than other aromatics (Section 3.2.1 and Section 3.2.2). Previous work suggests that the increase of methyl groups could lead to a change in several key organic fragments (e.g., CO+, CO2+ and H2O+) thereby altering the default fragment table for elemental ratio analysis. This agrees with the density underestimation in SOA formed from meta position aromatics and supports the preference of meta position alkyl substitute to SOA products.

15. For Figure 6, the author should provide the uncertainty of VFR(end) values.

We add uncertainty of VFR(end) values in revised manuscript.
16. *In the Table 1, the author should check the M0 value for 1215A, which is very high (M0=1501). It should be 151. Fixed.*

**Reference**


Reviewer #2

The manuscript presents information on yields for the photooxidation of single ringed aromatic structures. The senior author has been measuring yields from aromatic hydrocarbons (AHCs) for more than 15 years and is well versed. In this particular manuscript, the structure of the aromatic hydrocarbon (AHC) has been varied to examine differences in the organic aerosol (OA) yield. Thus, a series of 12 alkyl-substituted C8 and C9 AHCs have been examined. For these experiments, the aerosol yield has been determined using the Odum two-product model. Other OA parameters examined include the ratio of aerosol mass spectrometry (AMS) peaks attributed to OA, the O: C ratio, the oxidation state (OS), density, and volatility. The authors conclude that changes in the OA chemical composition and volatility influence the yield typically by increasing the mass for increased oxidation. The authors also consider the atmospheric implications of this study.

(1) The study addresses an issue of perhaps abstruse importance. The oxidation of alkyl substituted AHCs and the formation of secondary organic aerosol (SOA) has been examined extensively over the last 20 years and this work appears to cover some old territory. Many of these topics were addressed in Odum et al. 1997a, b (authors’ references) and the present manuscript provides a bit more insight.

Odum’s work was very important and provided a practical way to simplify aerosol yields. However, work over last decade has suggested the importance of NOx to SOA formation from aromatic hydrocarbons (e.g., Song, et al., 2005) with increasing aerosol formation observed for aromatics as initial NO levels are decreased. Lowering NOx conditions from the earlier Odum work improves representation of ambient conditions. Therefore, a comprehensive reinvestigation including isomer effects on SOA formation at more realistic hydrocarbon and NOx conditions is needed. Further, as noted by the reviewer, additional instrumentation available provides more insights into the SOA formation.

Higher yields are observed in this work for low NOx conditions than the earlier high NOx conditions (e.g Odum, et al, 1997, Figure 1, M0=40, yield=~0.03 or ~0.06; current work M0=40, yield 0.07-0.12). This paper demonstrates the molecular structure impact of aromatic hydrocarbons on SOA formation including impacts on SOA yield, chemical composition and physical composition. This is the first comprehensive analysis of SOA formation from aromatic isomers since the original Odum work on SOA from aromatics. The previous Odum work provides only very limited experimental work on isomers which is insufficient to determine molecular impact on SOA formation. (e.g., p-xylene and o-xylene have only two experiments)

(2) Admittedly, the AMS was not around and the aerosol density from the volume distribution was considered to be unity in the 1990s. However, this work also represents a step backwards. Whereas Odum et al. 1996, 1997a sought to simplify aerosol yields, this work goes in the other direction and makes an argument (at least implicitly) that the yields should be addressed more precisely, a contention that I don’t feel has been justified (see Table 2).
This work provides yield information with greater precision to dig into the role of molecular structure in SOA formation from aromatic hydrocarbons. We can't agree more with the reviewer that a simplified curve is a more attractive method for the curve fitting. Prior to further simplification, it is necessary to identify the relative importance of aromatic structure (o, m, p; alkyl length) especially when looking to project these findings for additional aromatic isomers. We currently have a paper under review to further demonstrate a novel method to simplify SOA yield from aromatic hydrocarbons, which requires insight on the relative importance of aromatic molecular structure. This work here focuses on molecular structure impact before stepping largely forward to the general trends found in SOA formation from aromatic hydrocarbons. The limited data sets available in earlier years are insufficient to reveal the difference among isomers and therefore might provide some bias on the similarity among aromatic hydrocarbon SOA formation, especially when conducted under high NOx conditions. As demonstrated in the manuscript, the difference among SOA from aromatic isomers, including SOA yield (for example, para position has significantly lower SOA yield compared with ortha and meta position), chemical composition and physical composition, does exist and should not be ignored by oversimplification. Therefore, this work is valuable to understand SOA formation from aromatic hydrocarbons before generalizing SOA aromatic yield trends.

(3) That said, the experiments appear to have been carefully performed and there certainly are enough of them. Unfortunately, the initial conditions are all over the map and makes it very difficult to get a sense of the reproducibility of a given experiment. Virtually, nothing is said about uncertainty.

There is only ~10% of experiments with HC/NO larger than 60 ppbC:ppb. The majority of experiments have similar NO conditions (see reply to referee #1 in Comment 4). Also, the similar impact of NO on radical and organic nitrate formation is demonstrated to be insignificant (see reply to referee #1 in Comment 4). The uncertainty of experiment is 6.6% based on ten repeat m-xylene experiments (please see referee #1 in Comment 3). Uncertainty in all analysis is included in the updated manuscript.

1. My major substantive comments regarding the manuscript are as follows: (1) these experiments hardly qualify as being under low NOx conditions. The removal of RO2 radicals competitively by NO or RO2 determines the regime that the reaction is in. I would consider the low NOx regime as conditions where the RO2 radical-radical reactions become more important than the NO reaction. As a rule of thumb, I would say that this is certainly at no more than 5 ppb of NO for the conditions of these experiments depending, of course, on the specific RO2 radicals from the precursor AHC.

(1) From a kinetics perspective, low NOx is even lower than the 5 ppb (suggested by the reviewer) tending to occur at NO levels in the 15 to 50 ppt range. However, starting with NOx levels < 50 ppt is substantially lower than practical experimental constraints for Teflon environmental chambers due to offgasing of HONO from Teflon surfaces (Carter et al., 2005). These experiments are referred to as low NOx experiments to be consistent with environmental chamber literature over the last decade, which is referring to the relative amount of NOx at the
beginning of the experiment when compared to initial VOC. The NO\textsubscript{x} ranges in this work are more consistent with urban NO\textsubscript{x} loadings than the earlier high NO\textsubscript{x} experiments performed by Odum (Odum, et al., 1997 and 1996) and others. Also, it is not possible to use an initial NO concentration ~5ppb since the low NO\textsubscript{x} concentration leads to a low reactivity of overall reaction and therefore forms less aerosol which is not atmospherically relevant.

Clearly the NO will compete for the HO\textsubscript{2} or RO\textsubscript{2} when there is sufficient NO. In other earlier work we demonstrate that SOA will not be formed until NO\textsubscript{2}/NO>70 (Li, et al., 2015), which indicates that RO\textsubscript{2} majorly react with NO instead of HO\textsubscript{2} or RO\textsubscript{2} even at NO~5ppb under the range of NO we investigated. The important point is that the NO concentration is extremely low (<10ppt) during the majority of the photooxidation experiment (after onset of O\textsubscript{3} formation). NO\textsubscript{x} mainly exists as NO\textsubscript{2} when PM is formed and HO\textsubscript{2}+RO\textsubscript{2} instead of NO+ HO\textsubscript{2}/RO\textsubscript{2} dominates.

In order to clarify the NO\textsubscript{x} condition we actually used, we add the following information on Page 5 at Line 29 after “under low NO\textsubscript{x}” add “(10-138 ppb)”.

(2) The version of SAPRC referenced to Carter and Heo (2013) is specifically geared to ozone prediction, that is, conditions where NO\textsubscript{x} dominates early product generations (two, at most three). As stated in Carter and Heo, (2013; Atmospheric Environment) SAPRC-11 is not geared for PM predictions. For this to be the case, the importance of RO\textsubscript{2} + RO\textsubscript{2} reactions should be adequately predicted as should the SOA mass. That is the point of the model and to predict radical concentrations as they were an end in themselves. The experimental SOA values could then constrain the model. That said, I question how well SAPRC-11 predicts radical concentrations under low NO\textsubscript{x} conditions. (3) Several sections need a complete writing overhaul. Section 3.2.1 is barely comprehensible. Many sections in the Results and Discussion presents data (e.g., S6) as if they were self-interpretive. The manuscript is written for the audience being other scientists in their research group. There is considerable jargon and the writing is highly imprecise hardly worthy of a scholarly journal. (4) To the extent possible, the authors should give a greater physical interpretation of the metrics they present in Section 3. Some are obvious (e.g., Sec. 3.2.2) others far less so (Sec. 3.2.1; 3.4).

2) SAPRC-11 is geared to predict O\textsubscript{3} formation under low NO\textsubscript{x} conditions especially for aromatic hydrocarbons as described by Carter and Heo (2013). Literally, the NO\textsubscript{x} range we used in current work is within the used range of NO\textsubscript{x} when the model is updated to SAPRC-11. In fact, the aromatic experiments used to develop the SAPRC-11 update are included in this work. We agree that it could not well predict SOA formation since the gas phase products are not well demonstrated as suggest by Carter and Heo (2013), especially for those associated with gas to particle partitioning. However, SAPRC-11 should be sufficiently good to predict gas phase radical concentration, which is closely associated with ozone formation.

3) Section 3.2.1 extends the traditional f\textsubscript{44} vs f\textsubscript{43} (C\textsubscript{2}H\textsubscript{5}O\textsuperscript{+}) chemical composition analysis by including fragments (C\textsubscript{3}H\textsubscript{5}O\textsuperscript{+} m/z 57 and C\textsubscript{4}H\textsubscript{7}O\textsuperscript{+} m/z 71) from longer alkyl substitute other than methyl since longer alkyl substitutes are included in the isomers investigated. The goal of
Section 3.2.1 is to provide insights into the SOA formation mechanism from different isomers as discussed in the later part of Section 3.2.1.

Some changes are already made in Section 3.2.1 (please see reply to referee #1, Comment 8). Also, we have added the following sentences on Page 11 at Line 3 to Section 3.2.1:

This work extends the traditional $f_{44}$ vs $f_{43}$ ($C_2H_3O^+$) chemical composition analysis by including oxidized fragments ($C_3H_5O^+$ m/z 57 and $C_4H_7O^+$ m/z 71) of the longer (non-methyl) alkyl substitutes. Therefore, $f_{44}$ vs $f_{43}+f_{57}+f_{71}$ is plotted instead of $f_{44}$ vs $f_{43}$.

We demonstrate the calculation of H/C and O/C in Section 3.2.1 and Section 4 to improve the physical interpretation of the metrics used.

Sec. 3.2.1: The physical interpretation is improved after the revision described in 3) comment and the referee #1’s Comment 8. Sec. 4: The physical interpretation stated at the beginning of Section 4 as “Methyl dilution theory (Li, et al. 2015a) is extended to alkyl substitute dilution theory in order to investigate the influence of longer alkyl substitutes compared with methyl group substitutes.” Additionally, the following sentence has been added to clarify the physical interpretation at Line 19 Page 17:

A robust prediction of SOA H/C and O/C trends for longer ($C_2^+$) alkyl substituted aromatics based on the methyl substituted aromatics will suggest a similarity in the role of methyl and longer alkyl to SOA formation; an underestimation or overestimation will indicate different oxidation pathways for aromatics with differing alkyl substitute length.

We also update the Fig 7 a & b according to referee #1’s Comment 3 about standard deviation. The implication from the difference between the measurement and prediction from the aromatics is updated correspondingly in the later part of Sect 4.
Some comments and suggestions:

2. P5, L27. *The goal of the research states the obvious. Perhaps more insight will motivate the reader to actually read the paper.*

We add the following sentence on Page 6 Line 2:

The effects of molecular structure impact on SOA yield, chemical composition (H/C, O/C, OS, f44, f42, f57 and f71) and physical properties (density and VFR) are demonstrated. Alkyl substitute dilution conjecture is further developed from methyl dilution theory (Li, et al., 2016).

3. P6, L7. *UV-350 bulbs have considerable radiation in the UVB which accentuates the photolysis of carbonyl compounds to a considerable extent which accelerates PM formation by increasing the radical concentrations. Thus, the two-product parameters developed (Table 2) may not be applicable for predictions of ambient AHC PM (Tables 2 and S3)*

We agree that UV-350 bulbs do not provide the higher wavelength region which affects photolysis of certain carbonyl compounds. However, UV impacts on different carbonyl are different. The photolysis rate ratios with blacklights will be much lower in the chamber than in the atmosphere if carbonyls have action spectra similar to the α-dicarboxyls; however, blacklight photolysis rate ratios will be higher if carbonyls have action spectra more like that of acrolein (Carter, et al., 1995). The photolysis of carbonyl compounds are more likely to impact the radical concentration (e.g. OH) and may further impact the overall SOA formation by change the kinetic reactivity. Therefore, the light source impact on carbonyl photolysis turns out to be the influence of radical concentration on SOA formation. The difference in radical concentration between chamber and atmosphere is demonstrated in Li, et al., 2015.

Further work is needed to adjust the SOA yield concluded from current chamber studies to better predict the SOA formation under atmospheric conditions. The current work provides the fundamental data for further investigation. Therefore, we add to the paper the statement “Moreover, the five subcategories of aromatics and their two product modeling curve fitting parameters in this work at more realistic NOx loadings provide a more precise prediction of SOA formation form aromatic hydrocarbons under atmospheric conditions” in Section 5 for the atmospheric application as in referee #1 Comment 13. The current study is more focused on the isomer impact on SOA formation. The results and implications of the current study remain reasonable since all the precursors are studied under comparable conditions (see kOH discussion in the reply to referee #1 Comment 4 last part).

4. P6, L18. *Provide a chemical name for the standard OEKANAL.*

OEKANAL is a Sigma-Aldrich Grade (purity) for 1, 2, 3-trimethylbenzene and is not a chemical name. It is followed by “1, 2, 3-trimethylbenzene Sigma-Aldrich”.

5. P6, L23. *How is mixing achieved in this large chamber: fan, diffusion, other?*
The mixing prior to commencing an experiment is achieved by fans. See Carter, et al., 2005 for more details “The two reactors are connected to each other through a series of custom solenoid valves and blowers. The system provides for rapid air exchange prior to the start of an experiment ensuring, that both reactors have identical concentrations of starting material. Each reactor can be premixed prior to the start of an experiment by Teflon coated fans located within the reactor.” During the experiment the vibration on the chamber walls due to air circulation on the outside of the chamber provide sufficient mixing during the experiment.

6. **P7, L10.** Calling a bunch of peaks attributed to innumerable organic compounds completely fragmented by 70 eV electrons as a chemical composition stretches the concept of molecules beyond recognition. I would hardly call this metric a chemical composition in any traditional sense. Perhaps the word “effective” could be incorporated to indicate that this is simply a parameterized metric.

We agree that the chemical composition is derived from peaks from numerable organic compounds completely fragmented. There might be some difference in between the traditional definition and what is widely used nowadays to describe AMS chemical composition, which is a measure of the bulk chemical composition of the aerosol. We keep our manuscript consistent with recent publications using AMS results (eg., Crippa, et al., 2013; Lambe, et al., 2015).

7. **P7, L23.** The authors should reference the 2013 Atmospheric Environment article by Carter and Heo rather than the CARB report. The article went through peer-review and should be more reliable.

Done.

8. **P8, L11.** The sentence is unneeded; include the information in the caption.

Done.

9. **P9, L3.** To use the word “claimed” in a pejorative fashion is particularly bad form. If you believe the statement in Odum et al. is wrong, simply state it.

Fixed. Changed “claimed” to “stated”.

10. **P9, L14.** Delete the sentence. The supplement does nothing to support the sentence other than to simply repeat itself and refer to a paper in preparation. Nothing is gain by including the sentence in the paper or in the supplement.

We keep this sentence to clarify the differences in the kinetics is insignificant (e.g., k_{OH}[OH], [HO2], …) and therefore the molecular structure of the isomers is driving the difference in SOA formation. The referenced paper is now published.

11. **P9, L23.** Why is the assumption needed? The two-product model is just a fitting exercise anyway.
Similar products are expected to be formed from the aromatic isomers. Fixing the $K_{om,2}$ value provides for similar treatment of the high volatility products allowing us to focus on the low volatility products most important to SOA formation under atmospheric conditions.

12. **P9. It might be worthwhile to examine partitioning using a volatility basis set (VBS) to see if any insight could be gained beyond the standard two-product fit which at this point is rather dated. This might provide a more useful metric for describing the partitioning of the AHC products.**

We agree that VBS is an attractive way to describe the SOA yield. However, VBS is fundamentally based on gas-particle partitioning theory which is the same as two-product model. The application of VBS only provide similar result in a different format. VBS presents the contribution of products with different volatility using bins and here two-product model use $K_{om}$. Therefore, we keep our analysis using the traditional two-product model.

13. **Sec 3.2.1. This section suffers from a lack of an understandable interpretation of the various fragments from the AMS output and their combinations into the combined metrics (e.g. Eq 1). An annoying aspect of this section is the comparison with other work before any interpretation is provided (e.g., P11, L5, 16; P12 L6, 25, . . ..). How do we even know that the conditions are applicable between these experiments and the ones being compared to?**

The interpretation is improved according to the reply to Comment 3). We cited other’s work to provide the AMS result found in other chamber work for selected isomer species. The initial hydrocarbon and NOx conditions used in other’s work are not completely the same as ours. Therefore, the AMS data is not exactly the same. However, we demonstrate the AMS data we use are reasonably in-line with earlier studies to contextualize the results and demonstrate that further discussion of the AMS data is reasonable.

14. **P13, L7. LV-OOA and SV-OOA are presented both undefined and without context.**

We change “LV-OOA and SV-OOA” to “low volatility oxygenated organic aerosol (LV-OOA) and semi-volatile oxygenated organic aerosol (SV-OOA)” The definition of LV-OOA and SV-OOA can be found in detailed in Ng, et al 2011.

15. **P13, L8. The sentence as written belongs in the introduction. The intent of the sentence needs a rationale from the data. It is not self-evident.**

We added a sentence to the introduction part to emphasize this part as a reply to Comment 2 (P5, L27). The sentence referred to here is to transition from evolution data to average data. We will delete it here and rewrite it as a sentence below in blue. We mention in the manuscript that “The evolution trend agrees with Fig. S3 (Sect. 3.2.1).” This means that the evolution trend is not significant during the photooxidation similar to what is mentioned in Section 3.2.1 for $f_{44}$ vs. $f_{43+57+71}$. Therefore, average value is sufficient to describe H/C and O/C. We add the following sentence in P13, L8 after “The evolution trend agrees with Fig. S3 (Sect. 3.2.1).” to
provide the rationale.

…, which means no significant H/C and O/C evolution is observed in the current study. Therefore, average H/C and O/C with standard deviation provided is used to explore the impact of molecular structure on SOA chemical composition.


Fixed. Inserted “than H/C and O/C”

17. P16, L1. Eliminate first name for Borrás.

Fixed

18. P17, L15. Use of the term “theory” (in any scientific sense) strikes me as somewhat pretentious. I would consider it as more of a conjecture.

Good point. We use “Alkyl Dilution Conjecture” according to reviewer’s suggestion; we keep “methyl dilution theory” as published in (Li, et al., 2016)

19. P18, L8-16. What does the term “extremely low” mean? Provide a value for comparison. I would characterize most of this part as speculation. Also, experimental limitations in Forstner et al., 1997 (their specific quartz filter configuration) suggested that the furan-type compounds were in the particle phase but were almost certainly in the gas phase. The metrics in the present paper refer only to particle phase OA. The reference should be avoided here.

We have now added the actual value for OSₙ and delete the reference in the revised manuscript. This part has been slightly modified after considering standard deviation/uncertainty as mentioned in response to Referee #1’s Comment 3.

20. Section 5. I seriously doubt that any of these parameterizations would appear in any wide-used air quality model. I would consider the work mainly for academic purposes.

The SOA yield parameters are widely used in current model (e.g. CMAQ see Carlton, et al., 2010; GEOS-Chem see Heald, et al., 2011, WRF-CHEM model Li, et al., 2011 ). Current work provides improved SOA yield parameters than previous work under high-NOx conditions (e.g. Odum. et al., 1997).


Fixed

23. **P25, L8.** Replace the ACPD manuscript with newly published ACP paper.

Done

24. **Table 1, S2 and text.** How many AHCs were studied: the text say twelve AHCs (P5, L7), Table 1 gives ten AHCs, and Tables 1 and S2 together give fourteen. Which is the right number? (For good measure, Figure 6 shows eleven precursors.)

Twelve is the right number. Table 1 and Table S2 combine to give the 12 unique AHCs used in this study.

25. **Table 3.** What’s the point of the table if the p value are greater than 0.05. Certainly, the p-value for VFR and k (OH) is not zero.

It is included in note below the table that “Alpha (α) level used is 0.05. If the p-value of a test statistic is less than alpha, the null hypothesis is rejected”. It the p value is greater than 0.05, the correlation found is not trusted within the α level. In another word, the larger the p-value the less confidence in the correlation provided. Certainly, the p-value for VFR and k (OH) is not absolute zero. However, it is <0.0005. Therefore, we change “0” into “0.000” for significant figure purposes.

26. **Figure 1.** The final points control the shape of the curve. The data below 20 ug m⁻³ would be of most interest for atmospheric applications.

The higher Mₒ allows one to improve the parameters used to fit the overall aerosol formation trends, especially that for α₂. The final points are therefore controlling the shape of the curve used to fit of the high volatility products, in this case α₂. The identified curves reasonably represent the lower organic mass loadings (< 20 ug m⁻³) as seen in the quality of the fit where α₁ and Kₒ,1 (lower volatility products) dominate the shape of the curve. Curve fitting with and without α₂ and Kₒ,2 are presented below.
27. **Figures 2, 3, 7. Am I missing something? Why are there no symbols for the values; is there that much uncertainty? The use of colors for the precursors in Table 2 is particularly annoying. Are not words sufficient without colors?**

The exact locations of these values show up after adding error bar to each value (see Referee #1 Comment 6’s reply). I think you mean Figure 2 instead of Table 2. The colors we use categorize all the 12 isomers into different subgroups (e.g. all xylenes are in green as what we labeled in the upper right). We think these colors help the audience to understand what kind of molecular structure impact it is (location vs. length). We would like to keep the colors to help us demonstrate the findings.

28. **In Figure 4, what are the estimated uncertainties in the model-generate radical concentrations. Carter and Heo, 2013 suggests that these could be substantial.**

We are unsure of what the reviewer is requesting. Figure 4 provides “Oxidation state (OSc) of SOA formed from different aromatic hydrocarbon” which nothing about model generated radical concentrations. Table S4 lists model-generated radical concentrations. Generally speaking, the [OH] is fitted through precursor measurement from GC-FID and therefore [OH] has little uncertainty (<~5%). SAPRC-11 adjusted photoreactive product quantum yield parameters are used to minimize average biases in Rate (Δ(O3-NO)) (Carter and Heo, 2013). The uncertainties of radical prediction is minimized since O3 prediction relies on radical predictions. However, the uncertainties associated with SAPRC-11 is not a focus for current work. We provide the radical prediction provide by SAPRC-11 to rule out the impact of kinetic difference during the aromatic hydrocarbon photooxidation in order to emphasize the molecular structure impact.

29. **Table S3. How about the C9-trimethyl compounds studied? Why not put the parameters obtained from the TMB compounds in the table? It would also be informative to include the data for toluene (which must have been studied at some point) for comparison with the other single position substituents, ethylbenzene and n-,i-propylbenzene.**

Table S3 is used to support Fig. S1 to demonstrate the length effect among all C₈ and C₉. We don’t study triple alkyl substitute that contain longer chains and therefore there is no need to list C9-trimethyl compounds in Table S3. (The fitting parameters for trimethylbenzenes can be found in Table 2.) We don’t include toluene since it is not in the range of C₈ and C₉ aromatics; instead, we refer to our earlier work (Li, et al., 2016) at Page 10 Line 5.

30. **Table S4. The value of this table for predicting radical concentrations is very limited as noted above. It may be useful for urban NOX conditions but not where RO2 + RO2 is the dominant source of the aerosol. It doesn’t surprise me that the p-values for virtually all comparisons in S5 are no different than the null hypothesis.**

We agree that SOA formation is tied to peroxyde radical reactions. It should be noted that peroxyde radical reaction is associated with NO, precursor concentration and other radicals (e.g.
OH) as is ozone formation. We maintain that SAPRC could predict radical concentrations sufficiently well for how they are used in this paper (see response to comments above comment 1-2) and comment 28). The insignificant correlations between yield and radicals are not due to the limitation of the model but the similarity in kinetics among all the isomer precursor we studied. We actually found pretty good radical (e.g. HO2/RO2) correlation with yield in our earlier work (Li, et al., 2015). Therefore, we prefer to keep the radical discussion as part of the supplement supporting the manuscript.

31. SI Table 3 should be Table S6. Use lower case k in the table

Fixed

Reference


In this manuscript, the authors Li et al. studied formation of secondary organic aerosol from aromatic hydrocarbons, quantified the formation yields and investigated the chemical and physical properties in order to infer the differences in underlying mechanisms between different molecular structures. The hydrocarbons studied spanned from C7 to C9 and from 1 to 3 alkyl substitutes on the aromatic ring. These hydrocarbons are important for urban areas, and have not been well studied compared to benzene and toluene. The authors presented a variety of measurements (yields, aerosol mass spectrometer, density, volatility) conducted with state of the art instrumentation. The experiments are well designed and the manuscript is generally well written with relatively minor typos. However, I believe that the data interpretation is weak and it is very difficult to draw conclusions based on the data presented, especially with regards to SOA yields. I recommend major revisions before the manuscript can be considered for publication in ACP.

Major comments:

1. The biggest weakness of this manuscript is that the SOA yields are quite varied and many of the conclusions drawn by the authors are not very convincing. In Fig. 1, there are no error bars shown for the individual yields and Mo, and for the empirical fits. For example, the authors conclude from this figure that ortho compounds yields are similar and regression was performed on the combined data set. However, one could easily argue that o-xylene yields and m-ethyltoluene yields are similar and should be fitted to the same curve. Also, one of the major conclusions about ortho-compounds having higher yields is very weak. At atmospherically relevant Mo (Mo < 40 ug/m3), the yields are all very similar. Any differences in mechanisms do not translate to significant differences in formation yields (smaller than experimental uncertainties in yields). Therefore, one can argue that molecular structure plays a relatively minor role.

We insert following sentence in method at Page 8 Line 9 to provide error bar for yields, as what replied to reviewer 1 in Comment 3: The uncertainty associated with 10 replicate m-xylene and NO experiments SOA yield is <6.65%.

We agree that yields are hard to distinguish by only looking limited datasets at a low mass loading (eg. M_o < 40 ug/m³). Our current fitting parameters already provide a good estimation for SOA formation under low mass loadings.

The argument about the minor role of molecular structure is not true as significant differences are suggested by the physical properties and chemical composition trend. The categorization of aromatic hydrocarbons for yield fitting are based on the measured data and the molecular structures such as the relative position of substitutes and the number of substitutes. Our yield
categorization agrees with the observation found in later physical properties and chemical composition trend part. For example, o-xylene is found to be more oxidized than m-ethyltoluene according to chemical composition (Fig. 2-4) and therefore it is reasonable to be categorized into different group.

2. Another critique I have about this manuscript is the lack of mechanistic insights revealed by the experiments. I read the earlier paper on carbon dilution theory by the same authors, and found that the AMS data were very useful in identifying the general effect: methyl substituents divert OH oxidation from the ring, leading to less oxidized SOA components. A similar approach is used here but much less effectively. It is unclear from the proposed theory why the position of the alkyl substituents matters. Again, the authors claim that SOA from ortho-compounds is more oxidized than that from para- and meta-compounds. Why is that the case? The authors can make this manuscript much stronger if they can propose mechanisms along with careful experimental work to support them. Since the theme of the paper is molecular structure, I think these explanations are very important and deserve more attention. Can the authors propose any mechanisms (does not have to 100% proven, only needs to be plausible) that may be consistent with the results? AMS data are not suitable for resolving isomer-specific differences. More speciated measurements (such as LC or GC/MS) will provide more insights.

We provide the mechanism leading to the difference in SOA formation from meta, para and ortho position in Section 3.2.2 Page 11, Line 20-Page 12 Line 10. Some mechanisms are also added as mentioned in the reply to Referee #1’s Comment 12.

“"A robust prediction of SOA H/C and O/C trends for longer (C2+) alkyl substituted aromatics based on the methyl substituted aromatics will suggest a similarity in the role of methyl and longer alkyl to SOA formation; an underestimation or overestimation will indicate different oxidation pathways for aromatics with differing alkyl substitute length.” as described in the reply to Referee #2’s Comment 1-(4). The proposed theory is to demonstrate the impact of alkyl substitute length. The impact of position is filtered out since aromatics with similar alkyl location are used for the prediction as stated in Page 17 Line 21-24.

Following sentences is added on Page 19 Line 1:

“It is possible due to the alkyl substitute location impact on the further oxidation of five-membered bicyclic radicals. Different carbonyl compounds can form as the ring opening products from the dissociation of five-membered bicyclic radical. It is assumed that oligomerization of these carbonyl compounds can contribute to SOA (Li, et al., 2016). Aromatic hydrocarbons with para position alkyl substitute tend to form more ketone like dicarbonyl compounds than other aromatics. Ketone might contribute less to oligomerization formation compared with aldehyde as suggested in Li, et al (2016)”.

The following sentence is added on Page 19 Line 2:

It might be due to a higher percentage of carbonyl with alkyl substitute formed during the
oxidation of meta containing aromatics (e.g. methylglyoxal, 2-methyl-4-oxopent-2-enal), which contributes to oligomerization and thereby SOA formation.

More speciated measurements are warranted in future studies

3. Similar to my previous comment (and other reviewers’ comments), the AMS does not really provide information about specific chemical composition. For example, the statement in Section 3.2.2 “SOA components from all isomers are located in between slope= −1 and slope= −2 lines suggesting that SOA from these aromatic hydrocarbons is composed primarily of acid (carbonyl acid and hydroxycarbonyl) and carbonyl (ketone or aldehyde) like functional groups”. The elemental ratios does not suggest that the SOA *contain* these functional groups, but rather they suggest that the SOA composition *evolve* as if they were adding these function groups. This is a very important difference that should be noted throughout the manuscript. The AMS elemental ratios help elucidate the bulk composition and evolution, not the specific chemical composition and mechanism, as suggested by the manuscript. I believe that AMS was useful in showing O/C changes from benzene to toluene to xylenes, but the specific isomeric differences in this paper are not convincingly argued in this manuscript.

We agree that elemental ratios suggest that the SOA composition *evolve* as if they were adding these function groups. We will delete the following sentences:

“SOA components from all isomers are located in between slope=-1 and slope=-2 lines (Fig. S6) suggesting that SOA from these aromatic hydrocarbons is composed primarily of acid (carbonyl acid and hydroxycarbonyl) and carbonyl (ketone or aldehyde) like functional groups. The elemental ratio of SOA from p-xylene photooxidation was nearly located on the acid line (slope=-1)”

All the isomers start from the same precursor location in the Van Krevelen graph and therefore the difference in aerosol phase composition indicates the difference in oxidation. We agree that AMS is not specially for the detection of detailed species. However, the overall chemical composition provides the oxidation state of SOA and can be used to interpret the different extend of oxidation and related mechanisms.

4. It is not clear to me why the authors chose to conduct the experiments under low NOx conditions. These highly substituted aromatic hydrocarbons are emitted in urban areas, and have lifetimes around 1 day or so. Even in rural areas, the NO levels are quite high, resulting in at least 50% oxidation by RO2+NO pathway (see Ortega et al., ACP, 2014). One could argue that the experiments conducted are still experiments with NOx (unlike HO2 dominated experiments). However, the HC/NOx ratios are not fixed (ranges over 1 order of magnitude), bringing into question the relative role of NOx. Why aren’t the experiments conducted with a fixed HC/NOx ratio, or with NOx so high that it is not limiting?

The low NOx we mentioned here is compared with earlier work as described on Page 4 Line 9. The NOx range we use is comparable to urban atmosphere. Range is clarified in revised
manuscript (See reply to Referee #2 Comment 1-(1)) and is not the major driver of the differences in SOA formation between the isomers. There might be a difference in the so called NO range when RO$_2$+HO$_2$ is dominated (See reply to Referee#2 Comment 1-(1)).

NO is depleted very fast at the beginning of the photooxidation and HO$_2$ and RO$_2$ reaction is dominating the major period of the photooxidation(see Reply to Referee #2 Comment 1-(1)).

Other comments:

5. Pg 5 line 2: what is the significance of the furanone relative abundances? Does it point to a particular propensity of fragmentation and/or SOA formation from one particular molecular structure?

The abundances of these products are

1.4 (± 0.39)% of 3-methyl-2,5-furanone is observed in toluene oxidation (Forstner et al., 1997);

7.4 (±3.8) % 3-ethyl-2,5-furanone in ethylbenzene oxidation (Forstner et al., 1997).

The reference here is to give an example of molecular structure impact on aromatic oxidation products. The observed results in earlier work are not directly comparable to our work and therefore the exact abundance is not included in the manuscript.

6. Pg 9 line 20: The parameters are for fitting purposes only, and are semi-empirical at best. Because the Odum 2-product equations are non-linear, alpha and K are coupled parameters (i.e. one can use a higher alpha and lower K and still get a decent fit to the experimental data). Therefore I suggest the authors not derive insights into relative volatilities from the fitted parameters.

The semi-empirical two product model is based on the fundamental theory gas-particle partitioning. $\alpha_1$ and K have different impact on the curve and the pair we reported here is based on a best fit. It is partially empirical because two lumped groups are assumed. This model fitting parameters can’t tell us which detailed species are favored in SOA from different aromatic. However, the difference among $\alpha$ and K provide the general information about the high volatility and low volatility products. Therefore, we would like to keep these implications from the yield parameters.

7. Pg 14 line 2-11: Is it possible that photolysis of aromatic carbonyls is playing a role in affecting the H/C of SOA?

It is possible. However, aldehydes higher than formaldehyde appear to react dominantly with OH radicals (Atkinson and Arey, 2003).
8. Table 3: It seems to me that the correlations are quite weak (around 0.5 at best) and the p-
value is always greater than 0.05, sometimes much greater. From a statistics point of view, the
correlations are inconclusive. I suggest rewording in main text to call these “weakly
correlated” (“. . . best correlated. . .” is misleading).

Done.

Use “correlated” instead of “best correlated”. on Page16 Line 7

Delete “strong” before “correlation” on Page16 Line 25

Delete “well” before “correlated” on Page16 Line 2

9. Sect 3.3.2: Can the authors isolate the effect of kinetics and molecular structure by
comparing VFR at the same extent of reaction?

All experiments are conducted under comparable kinetics and similar extent of reaction. (See
kOH discussion in the reply to referee #1 Comment 4 last part).

10. The authors alluded to the carbon dilution theory they proposed in an earlier publication
(Li et al., 2015a) many times throughout the manuscript without explaining the theory. This
theory is relatively novel and is not well known. Therefore I suggest adding a short section to
explain the theory early on in the manuscript and show how this will be applied to the molecular
structures studied in this work.

Following sentence is added in revised manuscript on Page 17 Line 17 to add a short section to
explain carbon dilution theory:

Carbon dilution theory proposed by Li et al (2016) successfully explain that methyl group
impacts remain similar in SOA elemental ratios as in the aromatic precursor. The chemical
composition of SOA formation from alkyl substituted aromatics is predicted by simply adding
the alkyl substitute into the chemical composition of SOA formed from pure aromatic ring
precursor (benzene).

We also add explanation in the revised manuscript to show how the theory is applied to this
work. Please see reply to referee #2 1-(4)

Technical comments:

11. Pg 2 Line 5: eight to nine carbon should be “C8- to C9-” or “eight- to nine-carbon”

Done. Changed “eight to nine carbon” to “eight- to nine-carbon”

12. Pg 2 Line 23 IPPC should be IPCC
Fixed.

13. Pg 2 Line 26 What does growth potential mean?

Changed “have larger growth potential than biogenic aerosol sources” into “are more likely to increase”

14. Pg 3 Line 14 Toluene and C8 aromatics do not dominate aromatic SOA, because they are not SOA themselves, but are precursors to SOA. I suggest rewording this sentence.

Changed “Toluene and C8 aromatics dominate anthropogenic SOA” into “Toluene and C8 aromatics dominate the anthropogenic SOA precursors”.

15. Pg 5 line 12: references to mass loading (Shilling et al. 2009 and Pfaffenberger et al 2013) and NO effect (Eddingsaas et al. 2012) on SOA formation are for α-pinene, not for aromatics. The authors should clarify that point.

Add “(α-pinene)” after literature in revised manuscript to clarify the SOA precursor.

16. Pg 6 line 13-20: are the chemicals used without further purification?

Yes.

17. Pg 7 line 8-9: “Volume fraction remaining” should be all capitalized

Done.

18. Pg 7 line 13-14: clarify that fx is the *mass* fraction of organic signal at m/z = x

Done.

19. Pg 7 line 19: “squirrel” should be capitalized

Done. Change to “Squirrel 1.56D / Pika 1.15D”

20. Pg 8 line 10: organic mass concentration should be Mo (o for organics), not M0. M0 would suggest that it is an initial mass concentration

Fixed.

21. Pg 10 line 17: what does the n stand for? It might be more useful to use n to denote the carbon number of the alkyl substitute. So for m/z 57 is derived from an ethyl-substituted aromatic, so n would be 2, and the formula would be C_{(n+1)}H_{(2n+1)}O+.
Add “n = carbon number of the alkyl substitute” after C\text{\(n\)}H\text{\(2n\text{-1}\)}O⁺.

The formula should be C\text{\(n\)}H\text{\(2n\text{-1}\)}O⁺ while using n to denote the carbon number of the alkyl substitute.

21. Pg 10 line 21: “hydrogen-like organic aerosol” should be “hydrocarbon-like organic aerosol”; it seems quite obvious to me that it is not possible to have a C3H7 or C4H9 fragment from aromatic compounds here. Even isopropyl benzene does not yield C3H7 upon EI fragmentation (see NIST spectra). I suggest removing that explanation to make it more concise.

Fixed. “hydrogen-like organic aerosol” should be “hydrocarbon-like organic aerosol”.

We want to keep the explanation to clarify the different between chamber conditions and ambient.

22. Pg 11 line 25: This section is very hard to follow. Is there a proposed mechanism of how an oxidation product of aromatic compounds can produce m/z 43 that is also consistent with proposed carbon dilution theory? It would be very beneficial here to use a figure to illustrate the key points.

We add Figure in supplemental materials to demonstrate the proposed mechanism. (New Fig. S7)

23. Pg 12 line 1: do the authors mean ROOH here? The bicyclic peroxides formed in aromatic oxidation are internally bridged, and are therefore ROOR, not ROOH (or hydroperoxides). I presume that in the presence of NOₓ, RO₂+HO₂ is negligible and ROOH is not formed. Also, is there a reference for peroxides not yielding CO₂⁺ from the AMS literature?

Change “bicyclic hydroperoxides” to “bicyclic peroxides” RO₂+HO₂ dominated since ozone is formed (see Reply to Comments 4).

CO₂⁺ can come from carbonates, cyclic anhydrides and lactones (McLafferty and Turecek, 1993). This indicates that the CO₂⁺ should come from a fragments with -O-C-O- structure. Neutral CO₂ should be formed before EI to generate CO₂⁺. Thermal decarboxylation is a possible pathway to form CO₂ from compounds such as aliphatic acid (McLafferty and Turecek, 1993). We hypothesize that it is impossible to form a neutral CO₂ if CO₂⁺ comes from -C-O-O-.

We do not find a reference to support that peroxides not yielding CO₂⁺ from the AMS.

We deleted “More importantly, the difference in f₄₄ implies that substitute location influences the further reaction pathway to form CO₂⁺ since CO₂⁺ is not readily available from bicyclic hydroperoxides.” Add following sentences to demonstrate the source of CO₂⁺ on page 11 Line 26.

CO₂⁺ are generally formed during MS electrical ionization from carbonates, cyclic anhydrides
and lactones (McLafferty and Turecek, 1993) indicating that the CO$_2^+$ is associated with -O-C-O- structure. Within the AMS, the CO$_2^+$ is also associated with decarboxylation of organic acids during heating followed by electrical ionization of the CO$_2$. We hypothesize that CO$_2^+$ formation from bicyclic peroxides is insignificant since CO$_2$ loss is not expected to come from -C-O-O- structure during thermal decomposition. Therefore, it is the reaction products of bicyclic peroxides that lead to the formation of CO$_2^+$ and the difference in $f_{44}$.

24. Pg 13 line 8: “The current study concentrated on” should be “The current study concentrates on”

Fixed.

25. Pg 17 line 20: the chemical formula C2H2n+1 seems wrong

Fixed. It should be C$_n$H$_{2n+1}$

26. Figure 2: there should not be a continuous scale for molecular structure; Also for all the figures, the experiment numbers should be noted as such. Otherwise the numbers do not mean anything to readers

Used color legend instead of color scale in revised manuscript.

27. Figure 4: can the bars be color coded to correspond to those in other figures (e.g. Figs. 5 and 6)?

Done
Impact of Molecular Structure on Secondary Organic 
Aerosol Formation from Aromatic Hydrocarbon 
Photooxidation under Low NO$_x$ Conditions

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Abstract

The molecular structure of volatile organic compounds (VOC) determines their oxidation pathway, directly impacting secondary organic aerosol (SOA) formation. This study comprehensively investigates the impact of molecular structure on SOA formation from the photooxidation of twelve different eight- to nine-carbon aromatic hydrocarbons under low NO$_x$ conditions. The effects of the alkyl substitute number, location, carbon chain length and branching structure on the photooxidation of aromatic hydrocarbons are demonstrated by analyzing SOA yield, chemical composition and physical properties. Aromatic hydrocarbons, categorized into five groups, show a yield order of ortho ($o$-xylene and $o$-ethyltoluene) > one substitute (ethylbenzene, propylbenzene and isopropylbenzene) > meta ($m$-xylene and $m$-ethyltoluene) > three substitute (trimethylbenzenes) > para ($p$-xylene and $p$-ethyltoluene). SOA yields of aromatic hydrocarbon photooxidation do not monotonically decrease when increasing alkyl substitute number. The ortho position promotes SOA formation while the para position suppresses aromatic oxidation and SOA formation. Observed SOA chemical
composition and volatility confirm that higher yield is associated with further oxidation. SOA chemical composition also suggests that aromatic oxidation increases with increasing alkyl substitute chain length and branching structure. Further, carbon dilution conjecture developed by Li., et al (2015a) is extended in this study to serve as a standard method to determine the extent of oxidation of an alkyl substituted aromatic hydrocarbon.

**Key Words**

Secondary Organic Aerosol; Aromatic Hydrocarbon; Molecular Structure; Alkyl Substitute; SOA Yield; Chemical Composition; Volatility

1. Introduction

Organic aerosols are critical to human health (Dockery, et al., 1993; Krewski, et al., 2003; Davidson et al., 2005), climate change (IPPC, 2007) and visibility (Pöschl 2005; Seinfeld and Pandis, et al., 2006). Global anthropogenic secondary organic aerosol (SOA) sources are underestimated by current models (Henze, et al., 2008; Matsui, et al., 2009; Hallquist, et al., 2009; Farina, et al., 2010) and are more likely to increase have larger growth potential than biogenic aerosol sources due to the increase of known anthropogenic emissions (Heald, et al., 2008). Therefore, it is crucial to explore SOA formation mechanism from anthropogenic precursors.

Aromatic hydrocarbons are major anthropogenic SOA precursors (Kanakidou, et al., 2005; Henze, et al., 2008; Derwent, et al., 2010). C₈ (ethylbenzene, xylenes) and C₉ (ethyltoluenes and trimethylbenzenes) aromatics are important aromatic hydrocarbons in the atmosphere besides toluene and benzene (Monod, et al., 2001; Millet, et al., 2005; Heald, et al., 2008; Kansal, et al., 2009; Hu, et al., 2015). The major sources of C₈ and C₉ aromatic hydrocarbons are fuel evaporation (Kaiser, et al., 1992; Rubin, et al., 2006; Miracolo, et al., 2012), tailpipe exhaust (Singh, et al., 1985; Monod, et al., 2001; Lough, et al., 2005; Na, et al., 2005; Correa and Arbilla, et al., 2006) and solvent use (Zhang, et al., 2013). C₈ aromatic hydrocarbons (ethylbenzene and xylenes (ortho, meta and para) are categorized as hazardous air pollutants
(HAPs) under the US Clean Air Act Amendments of 1990

Toluene and C8 aromatics dominate the anthropogenic SOA precursors and SOA yield from all C9 aromatics is currently predicted to be equal to that of toluene (Bahreini, et al., 2009). The chemical composition of aromatic SOA remains poorly understood with less than 50% of aromatic hydrocarbon photooxidation products identified (Forstner, et al., 1997; Fisseha, et al., 2004; Hamilton et al., 2005; Sato et al., 2007). Aromatic hydrocarbon photooxidation mechanisms remain uncertain except for the initial step (~90% OH-addition reaction) (Calvert, et al., 2002). Hence, understanding the atmospheric reaction mechanisms of C8 and C9 aromatic hydrocarbons and properly quantifying their SOA formation potential presents unique challenges due to the variety in their molecular structure and the electron density of the aromatic ring.

Volatile organic compound (VOC) structure impacts the gas phase reaction mechanism (Ziemann and Atkinson, 2012) and kinetic reaction rate (eg. $k_{OH}$ Atkinson, 1987) thereby influencing the resulting SOA properties and mass yield. Molecular structure impacts on SOA formation from alkanes have been previously studied (Lim and Ziemann, 2009; Ziemann, 2011; Lambe, et al., 2012; Tkacik, et al., 2012; Yee, et al., 2013; Loza, et al., 2014). It is generally observed that SOA yield decreases from cyclic alkanes to linear alkanes and to branched alkanes. The relative location of the methyl group on the carbon chain also affects SOA yield (Tkacik, et al., 2012). It is further found that the SOA yield and structure relationship is influenced by C=C groups (Ziemann, 2011). Understanding the SOA yield and structure relationship of aromatic compounds in a similar way is necessary due to the atmospheric importance of aromatic hydrocarbons.

Previously, aromatic studies categorized SOA yield solely based on substitute number (Odum, 1997a, b). However, those chamber experiments were conducted at high NOx conditions, which are well above levels present in the atmosphere. Song, et al (2005, 2007) found that initial HC/NOx ratios significantly impact SOA yields during aromatic photooxidation with yields increasing as NOx levels decreased. Ng et al. (2007) shows there is no significant yield difference between one substitute (toluene) and two substitute (m-xylene) aromatics in the absence of NOx. The current work focuses on molecular structure impact on SOA formation
at more atmospherically relevant NOx and aerosol loadings. Li et al (2015a) demonstrated the methyl group number impact on SOA formation under low NOx conditions. Also, aromatic compounds with para position alkyl groups have been observed to form less SOA under various NOx conditions than their isomers in previous studies. Izumi and Fukuyama (1990) found that p-xylene, p-ethyltoluene and 1, 2, 4-trimethylbenzene have low SOA formation potential under high NOx conditions. Song, et al (2007) observed that p-xylene has the smallest SOA yield among all xylenes in the presence of NOx. The relative methyl position to -OH in dimethyl phenols also impacts SOA yield in the absence of NOx (Nakao, et al., 2011), while Song et al.(2007) observed no significant SOA yield difference between o-xylene and p-xylene under NOx free conditions. Moreover, previous studies mainly focused on the carbon number effect on SOA formation (Lim and Ziemann, 2009; Li, et al, 2005a, 2016) and seldom addressed the substitute carbon length impact on VOC oxidation and hence SOA formation. Different percentages of similar compounds are found when the substitute carbon length on the aromatic ring changes (Forstner, et al., 1997; Huang, et al., 2007; Huang, et al., 2011). For example, a higher-lower percentage of 3-methyl-2, 5-furanone is observed in toluene than that of 3-ethyl-2, 5-furanone in ethylbenzene (Forstner, et al., 1997). Further, the branching structure on the aromatic substitute might impact the reaction pathway. It is possible that fragmentation is more favored on branched substitute alkoxy radicals than n-alkane substituents similar to alkanes (Atkinson, et al., 2003).

Few studies comprehensively consider the overall alkyl effect on SOA formation from aromatic hydrocarbons, including the substitute number, position, carbon chain length and branching structure, especially under low NOx conditions. It is valuable to understand the relationship between aromatic hydrocarbon molecular structures and SOA physical and chemical characteristics. The effects of OH exposure (Lambe, et al., 2011, 2015), mass loading (Shilling, et al., 2009 (α-pinene); Pfaffenberger, et al., 2013 (α-pinene)) and NO condition (Ng, et al., 2007; Eddingsaas, et al., 2012 (α-pinene)) on SOA physical and chemical characteristics are previously discussed. However, few studies address the molecular structure effect of the precursor on SOA chemical composition, especially under atmospherically relevant conditions. Sato et al (2012) shows the chemical composition...
difference between ethylbenzene, \( m\)-xylene, \( o\)-xylene, 1, 2, 4-trimethylbenzene and 1, 3, 5-trimethylbenzene under high absolute NO\(_x\) conditions and hypothesizes that ketones prevent further oxidation during aromatic photooxidation compared with aldehydes. The SOA products detected in Sato’s study are mainly small volatile compounds which are less likely to partition into the particle phase (Chhabra, et al., 2011). Therefore, the study of Sato, et al. (2012) indicates that further oxidation or oligmerization might contribute to SOA formation during aromatic photooxidation. Less SOA characterization data on propylbenzene and ethyltoluene compared with trimethylbenzene is available. However, Bahreini, et al. (2009) suggests that the sum of the propylbenzene and ethyltoluene is on average a factor of 4–10 more abundant than trimethylbenzene.

This work examines twelve aromatic hydrocarbons, all of which are isomers with eight or nine carbons, to investigate the impact of molecular structure on SOA formation from aromatic hydrocarbon photooxidation under low NO\(_x\) (10-138 ppb). Here, we investigate the substitute number, substitute position, alkyl carbon chain length and alkyl branching impacts on aromatic hydrocarbon oxidation. The effects of molecular structure impact on SOA yield, chemical composition (H/C, O/C, OS\(_c\), f\(_{44}\), f\(_{43}\), f\(_{57}\) and f\(_{71}\)) and physical properties (density and VFR) are demonstrated. Alkyl substitute dilution conjecture is further developed from methyl dilution theory (Li, et al., 2016).

2. Method

2.1 Environmental chamber

The UC Riverside/CE-CERT indoor dual 90 m\(^3\) environmental chambers were used in this study and are described in detail elsewhere (Carter et al., 2005). Experiments were all conducted at dry conditions (RH<0.1%), in the absence of inorganic seed aerosol and with temperature controlled to 27±1°C. Seeded experiments to minimize wall effects have also been conducted in our chamber experiment with no measurable difference observed between the seeded and non-seeded experiment. Two movable top frames were slowly lowered during each experiment to maintain a slight positive differential pressure (~0.02” H\(_2\)O) between the
reactors and enclosure to minimize dilution and/or contamination of the reactors. 272 115 W Sylvania 350BL blacklights are used as light sources for photooxidation.

A known volume of high purity liquid hydrocarbon precursors (ethylbenzene Sigma-Aldrich, 99.8%; n-propylbenzene Sigma-Aldrich, 99.8%; isopropylbenzene Sigma-Aldrich, analytical standard; m-xylene Sigma-Aldrich, 99%; o-xylene Sigma-Aldrich, 99%; p-xylene Sigma-Aldrich, 99%; m-ethyltoluene Sigma-Aldrich, 99%; o-ethyltoluene Sigma-Aldrich, 99%; p-ethyltoluene Sigma-Aldrich, ≥95%; 1, 2, 3-trimethylbenzene Sigma-Aldrich, OEKANAL analytical standard; 1, 2, 4-trimethylbenzene Sigma-Aldrich, 98%; 1, 3, 5-trimethylbenzene Sigma-Aldrich, analytical standard) was injected through a heated glass injection manifold system and flushed into the chamber with pure \( \text{N}_2 \). NO was introduced by flushing pure \( \text{N}_2 \) through a calibrated glass bulb filled to a predetermined partial pressure of pure NO. All hydrocarbons and NO are injected and well mixed before lights are turned on to start the experiment.

2.2 Particle and Gas Measurement

Particle size distribution between 27 nm and 686 nm was monitored by dual custom built Scanning Mobility Particle Sizers (SMPS) (Cocker et al., 2001). Particle effective density was measured with an Aerosol Particle Mass Analyzer (APM-SMPS) system (Malloy et al., 2009). Particle volatility was measured by a Dekati® Thermodenuder Volatility Tandem Differential Mobility Analyzer (VTDMA) (Rader and McMurry, 1986) with a 17 s heating zone residence time (Qi, et al., 2010a). The heating zone was controlled to 100 °C in this study with Volume Fraction Remaining (VFR) calculated as \( \frac{D_{p, \text{after TD}}}{D_{p, \text{before TD}}} \). Particle-phase chemical composition evolution was measured by a High Resolution Time of Flight Aerosol Mass Spectrometer (HR-ToF-AMS; Aerodyne Research Inc.) (Canagaratna et al., 2007; DeCarlo et al., 2006). The sample was vaporized by a 600 °C oven under vacuum, followed by a 70 eV electron impact ionization. \( f_x \) in this study is calculated as the mass fraction of the organic signal at \( m/z=x \). For example, \( f_{44}, f_{43}, f_{57}, \) and \( f_{71} \) are the ratios of the organic signal at \( m/z \) 44, 43, 57 and 71 to the total organic signal, respectively (Chhabra et al., 2011; Duplissy et al., 2011). Elemental ratios for total organic mass, oxygen to carbon (O/C),
and hydrogen to carbon (H/C) were determined using the elemental analysis (EA) technique (Aiken et al., 2007, 2008). Data was analyzed with ToF-AMS analysis toolkit Squirrel. Evolution of SOA composition (Heald, et al., 2010; Jimenez, et al., 2009) refers to SOA chemical composition changes with time. $f_{44}$ and $f_{51-77}$ evolution and H/C and O/C evolution refer to the change of $f_{44}$ and $f_{51-77}$ with time and the change of H/C and O/C with time, respectively.

The Agilent 6890 Gas Chromatograph – Flame Ionization Detector was used to measure aromatic hydrocarbon concentrations. A Thermal Environmental Instruments Model 42C chemiluminescence NO analyzer was used to monitor NO, NO$_2$-NO and NO$_3$. The gas-phase reaction model SAPRC-11 developed by Carter and Heo (2012) was utilized to predict radical concentrations (·OH, HO$_2^-$, RO$_2^-$ and NO$_3^-$).

3. Result

3.1 SOA yield

Photooxidation of twelve C$_8$ and C$_9$ aromatic hydrocarbons were studied for low NO$_x$ conditions (HC/NO ratio 11.1-171 ppbC:ppb). SOA yields for all aromatic hydrocarbons were calculated according to Odum, et al. (1996) as the mass ratio of aerosol formed to parent hydrocarbon reacted. Experimental conditions and SOA yields are listed (Table.1) along with additional m-xylene, o-xylene, p-xylene and 1, 2, 4-trimethylbenzene experimental conditions from previous studies (Song, et al, 2005; Song, et al, 2007; Li, et al, 2015a, 2016) (Table S2). The uncertainty associated with 10 replicate m-xylene and NO experiments SOA yield is <6.65%. SOA yield as a function of particle mass concentration ($M$, $M_0$), shown in Fig. 1, includes experiments listed in both Table 1 and Table S2. Each symbol represents a different aromatic hydrocarbon. It is observed that both alkyl substitute number and position affect SOA yield. The SOA yield of two-substitute C$_8$ and C$_9$ aromatic hydrocarbons depends more on the substitute location than substitute length. This means that the yield trend of o-xylene is analogous to that of o-ethyltoluene. Similarly, the yield trends for meta and para position substituted C$_8$ and C$_9$ aromatic hydrocarbons will be analogous to each other. Ortho isomers
(o-xylene and o-ethyltoluene, marked as solid and hollow green circles, respectively) have the highest SOA yield for similar aerosol concentrations while para isomers (p-xylene and p-ethyltoluene, marked as solid and hallow blue diamonds, respectively) have the lowest SOA yield level. Lower SOA yield for para isomers are consistent with previous observation by Izumi and Fukuyama (1990). Izumi and Fukuyama (1990) also suggest that 1, 2, 4-trimethylbenzene yields are lower than for other aromatic hydrocarbons. The current study does not show a significant SOA yield difference between 1, 2, 4-trimethylbenzene and 1, 3, 5-trimethylbenzene. It is difficult to compare 1, 2, 3-trimethylbenzene yields with the former two trimethylbenzenes since 1, 2, 3-trimethylbenzene mass loading is much higher than the former two.

Aromatic hydrocarbons having only one substitute (ethylbenzene, n-propylbenzene and isopropylbenzene) or three substitutes (1, 2, 3-trimethylbenzene, 1, 2, 4-trimethylbenzene and 1, 3, 5-trimethylbenzene) tend to have yields similar to the meta position two alkyl aromatics. Odum, et al (1997b) categorized SOA yield formation potential solely based on substitute number and claimed stated that aromatics with less than two methyl or ethyl substitutes form more particulate matter than those with two or more methyl or ethyl substitutes on the aromatic ring. However, Odum’s work was conducted for high NOx conditions and had insufficient data to compare isomer yield differences (e.g., only two low mass loadings for o-xylene data). The strong low yield (two or more substitutes) and high yield (less than two methyl or ethyl substitutes) trends for high NOx conditions (Odum, et al., 1997) are not observed for low NOx aromatic experiments in this study. Rather, high yield is observed only for benzene (Li, et al., 2015a, 2016) while low yield is seen for substituted aromatic hydrocarbons. Similar SOA yield trends from different C8 and C9 aromatic isomers are further confirmed by comparing yields at similar radical conditions (Table S4, Fig. S2). It is also found that molecular structure exerts a greater impact on SOA yield than reaction kinetics (supplemental material, Table S5). A two product model described by Odum, et al. (1996) is used to fit SOA yield curves as a function of \( M_{2} \). The twelve aromatics are categorized into five groups to demonstrate the alkyl group number and position effect on SOA formation. The five groups include one substitute group (1S), ortho position two alkyl group (ortho), meta
position two alkyl group (meta), para position two alkyl group (para) and three substitute group (3S). Fitting parameters ($\alpha_1$, $K_{om,1}$, $\alpha_2$ and $K_{om,2}$; Table 2) in the two product model are determined by minimizing the sum of the squared residuals. The lower volatility partitioning parameter ($K_{om,2}$) is the same for all yield curve fits by assuming similar high volatile compounds are formed during all aromatic hydrocarbon photooxidation experiments. The ortho group is associated with a much higher $K_{om,1}$ compared with other aromatic groups, indicating aromatic hydrocarbon oxidation with an ortho position substitute forms much lower volatility products than other isomers. $K_{om,1}$ are also slightly higher in the meta group and one substitute groups than in the three substitute and para substitute groups.

A slight SOA yield difference remains within each group (Fig. S1&Table. S3), indicating the influence of factors other than alkyl group position. Generally, lower yields are found in aromatics with higher carbon number substitute alkyl groups, such as when comparing propylbenzene (i- and n-) with ethylbenzene or toluene (Li, et al., 2005a, 2016), m-ethyltoluene with m-xylene and p-ethyltoluene with p-xylene, respectively. These differences are explained by the proposed alkyl group dilution effect (Section 4). However, the differences between xylenes and their corresponding ethyltoluenes are not statistically significant.

### 3.2 Chemical composition

#### 3.2.1 $m/z_{44}$ vs $m/z_{43+57+71}$

The ratio of alkyl substitute carbon number (H:C >1) to the aromatic ring carbon number impacts SOA composition since the H:C ratio on the alkyl substitute is larger than 1 and the H:C ratio on aromatic ring itself is no more than 1. $m/z$ 43 ($C_2H_3O^+$ and $C_3H_7^-$) combined with $m/z$ 44 ($CO_2^-$) are critical to characterize oxygenated compounds in organic aerosol (Ng, et al., 2010; Ng, et al., 2011). $C_2H_3O^+$ is the major contributor to $m/z$ 43 in SOA formed from aromatic hydrocarbons having only methyl substitute (Li, et al., 2015a, 2016) while $C_3H_7^-$ fragments are observed in this work for SOA from propylbenzene and isopropylbenzene (Fig. S4, Table S6). The $C_nH_{2n-1}O^+$ ($n$-carbon number of the alkyl substitute) fragment in SOA corresponds to a $C_nH_{2n+1}^+$ alkyl substitute to the aromatic ring. $C_3H_7O^+$ ($m/z$ 57) and $C_4H_7O^+$
(m/z 71) are important when investigating SOA from ethyl or propyl substitute aromatic precursors. While m/z 57 (C₄H₉+) and m/z 71 (C₅H₁₁+) are often considered as markers for hydrocarbon-like organic aerosol in ambient studies (Zhang et al., 2015; Ng et al., 2010), oxygenated organic aerosol C₄H₅O⁺ and C₅H₇O⁺ are the major fragments at m/z 57 and m/z 71, respectively. (Fig. S4, Table S6) in current chamber SOA studies, especially during the photooxidation of ethyl and propyl substituted aromatics. Therefore, m/z 57 and m/z 71 are also considered beside C₂H₃O⁺ at m/z 43 in SOA chamber studies as OOA to compare the oxidation of different aromatic hydrocarbons. While m/z 43 (C₃H₇+) and m/z 57 (C₄H₉+) are often considered as markers for hydrogen-like organic aerosol (Zhang, et al., 2015; Ng, et al., 2010), C₃H₅O⁺(m/z 57) and C₄H₇O⁺ (m/z 71) are the major fragments (Fig. S4) from SOA originating from ethyl and propyl substituted aromatics, consistent with Sato, et al., (2010) and Mohr, et al., (2009). Fig S4 lists all fragments found at m/z 43, 44, 57 and 71 and Fig S5 shows the fraction of each m/z in SOA formed from all aromatic hydrocarbons studied. The m/z 43+m/z 44+m/z 57+m/z 71 accounts for 21.2%–29.5% of the total mass fragments from all C₈ and C₉ aromatics studied, suggesting similar oxidation pathways. Only a small fraction (<=0.7%) of m/z 71 (C₄H₅O⁺) or m/z 57 (C₅H₇O⁺) was observed in ethyltoluenes and trimethylbenzenes, respectively.

This work extends the traditional f₄₄ vs f₄₃ (C₂H₃O⁺) chemical composition analysis by including oxidized fragments (C₄H₅O⁺ m/z 57 and C₅H₇O⁺ m/z 71) of the longer (non-methyl) alkyl substitutes. Therefore, f₄₃ vs f₄₄+f₅₇+f₇₁ is plotted instead of f₄₄ vs f₄₃. Fig S3 shows the evolution of f₄₄ and f₄₃,f₅₇,f₇₁ in SOA formed from the photooxidation of different aromatic hydrocarbons at low NOx conditions. f₄₄ and f₄₃,f₅₇,f₇₁ ranges are comparable to previous chamber studies (Ng, et al., 2010; Chhabra, et al., 2011; Loza, et al., 2012; Sato, et al., 2012). Only slight f₄₄ and f₄₃,f₅₇,f₇₁ evolution during chamber photooxidation is observed for the C₈ and C₉ isomers hence only the average f₄₄ and f₄₃,f₅₇,f₇₁ will be analyzed in this work.

A modification is applied to the mass based m/z fraction in order to compare the mole relationship between m/z 44 and m/z 43+m/z 57+m/z 71(Eq-1).

\[ f'_{43+57+71} = \frac{44}{43} f_{44} + \frac{44}{57} f_{57} + \frac{44}{71} f_{71} \]  

Eq-1
The average $f_{44}$ vs $\Gamma_{43-57-71}$ for all $C_8$ and $C_9$ isomers (Fig. 2) are located around the trend line for methyl group substituted aromatic hydrocarbons (Li, et al., 2015a, 2016), implying a similarity in the SOA components formed from alkyl substituted aromatic hydrocarbons. A decreasing trend in oxidation from upper left to lower right is included in Fig 2, similar to what Ng, et al (2011) found in the $f_{44}$ vs $f_{43}$ graph, especially while comparing similar structure compounds. The methyl group location on the aromatic ring impacts $f_{44}$: $\Gamma_{43-57-71}$. Decreasing $f_{44}$ and increasing $\Gamma_{43-57-71}$ trends are observed from $p$-xylene to $o$-xylene to $m$-xylene and from 1, 2, 4-trimethylbenzene to 1, 2, 3-trimethylbenzene to 1, 3, 5-trimethylbenzene. The $\Gamma_{43-57-71}$ may partially depend on the relative position between the alkyl substitute and the peroxide oxygen of the bicyclic hydroperoxide. For instance, allylically stabilized five-membered bicyclic radicals are the most stable bicyclic peroxide radical formed from aromatic hydrocarbon photooxidation (Andino, et al., 1996). Two meta position substitutes connected to the aromatic ring carbon with -C=O yield higher fractions of $C_nH_{2n-1}O^+$ fragments than the para and ortho position, which have at most one substitute connected with -C=O. (Fig. S6). $CO_2^+$ are generally formed during MS electrical ionization from carbonates, cyclic anhydrides and lactones (McLafferty and Turecek, 1993) indicating that the $CO_2^+$ is associated with -O-C=O- structure. Within the AMS, the $CO_2^+$ is also associated with decarboxylation of organic acids during heating followed by electrical ionization of the $CO_2$. We hypothesize that $CO_2^+$ formation from bicyclic peroxides is insignificant since $CO_2$ loss is not expected come from -C=O-O- structure during thermal decomposition. Therefore, it is the reaction products of bicyclic peroxides that lead to the formation of $CO_2^+$ and the difference in $f_{44}$. More importantly, the difference in $f_{44}$ implies that substitute location influences the further reaction pathway to form $CO_2^+$ since $CO_2^+$ is not readily available from bicyclic hydroperoxides. This indicates that the alkyl groups are more likely to contribute to SOA formation at the meta position than the ortho and para position. The para position substituted aromatics form the least SOA as they exclude the possibility that para position alkyl substitutes are further oxidized to other less volatile components instead of $C_nH_{2n-1}O^+$. $p$-Xylene displays the high $f_{44}$ similar to benzene (Li, et al., 2015a) implying that the para position substitute exerts the least dilution effect on $CO_2^+$ formation pathway among all isomers. Bicyclic hydroperoxides formed from the OH-addition reaction pathway and their
dissociation reaction products are both used to explain the substitute location impact on \( f_{44} \) and \( \Gamma_{43,57,71} \) relationship. However, the existence of longer alkyl substitutes diminishes the alkyl substitute location impact. SOA \( f_{44} \) and \( \Gamma_{43,57,71} \) in ethyltoluenes are all analogous to \( m \)-xylene. One substitute \( C_8 \) and \( C_9 \) aromatic hydrocarbons have similar \( f_{44} \) and \( \Gamma_{43,57,71} \) with slightly lower \( f_{44} \) and \( \Gamma_{43,57,71} \) compared to toluene (Li, et al., 2015a, 2016). Longer alkyl substitutes may not lower the average oxidation per mass as further oxidation of the longer chain alkyils may render other oxidized components not included in Fig. 2. Their lower total \( f_{44} + \Gamma_{43,57,71} \) (Fig. S5) further supports the possibility of oxidation of the longer alkyl substituents. It is also possible that oligomerization from highly oxidized carbonyls contribute more to the SOA formation from aromatics with long chain alkyl substituent. Elemental ratio (Section 3.2.2) and oxidation state (Section 3.2.3) are further used to evaluate the impact of increasing alkyl group size on SOA formation.

### 3.2.2 H/C vs O/C

Elemental analysis (Aiken, et al., 2007, 2008) serves as a valuable tool to elucidate SOA chemical composition and SOA formation mechanisms (Heald, et al., 2010; Chhabra, et al., 2011). Fig. S6-S7 shows H/C and O/C evolution in SOA formed from the photooxidation of different aromatic hydrocarbons under low NOx (marked and colored similarly to Fig. S3). H/C and O/C ranges are comparable to previous chamber studies (Chhabra, et al., 2011 (\( m \)-xylene and toluene); Loza, et al., 2012 (\( m \)-xylene); Sato, et al., 2012 (benzene and 1, 3, 5-trimethylbenzene)). SOA components from all isomers are located in between slope=-1 and slope=-2 lines (Fig. S6) suggesting that SOA from these aromatic hydrocarbons is composed primarily of acid (carbonyl acid and hydroxycarbonyl) and carbonyl (ketone or aldehyde) like functional groups. The elemental ratio of SOA from \( p \)-xylene photooxidation was nearly located on the acid line (slope=-1). The SOA elemental ratio for \( C_8 \) and \( C_9 \) aromatic isomers are located near the alkyl number trend line found by Li, et al (2015a, 2016) for methyl substituents, indicating a similarity between SOA from various alkyl substituted hydrocarbons. SOA formed is among the low volatility oxygenated organic aerosol (LV-OOA) and semi-volatile oxygenated organic aerosol (SV-OOA) regions (Ng, et al., 2011). The evolution
trend agrees with Fig. S3 (Section 3.2.1), which means no significant H/C and O/C evolution is observed in the current study. Therefore, average H/C and O/C with standard deviation provided is used to explore the impact of molecular structure on SOA chemical composition.

The current study concentrated on experimentally averaged H/C and O/C to explore the impact of molecular structure on SOA chemical composition. Average H/C and O/C locations are marked with aromatic compound names in Fig. 3. All H/C and O/C are located around the predicted values for C₈ and C₉ SOA (dark solid circle) based on the elemental ratio of benzene SOA (Li, et al., 2015a, 2016). This confirms the presence of a carbon dilution effect in all isomers. Ortho position aromatic hydrocarbons (o-xylene or o-ethyltoluene) lead to a more oxidized SOA (higher O/C and lower H/C) than that of meta (m-xylene or m-ethyltoluene) and para (p-xylene or p-ethyltoluene) aromatics. SOA formed from 1, 2, 4-trimethylbenzene and 1, 2, 3-trimethylbenzene is more oxidized than that from 1, 3, 5-trimethylbenzene. It is noticed that 1, 2, 4-trimethylbenzene and 1, 2, 3-trimethylbenzene both contain an ortho position moiety on the aromatic ring. This indicates that the ortho position aromatic hydrocarbon is readily oxidized and this ortho position impact on oxidation extends to triple substituted aromatic hydrocarbons.

Substitute length also plays an important role in aromatic hydrocarbon oxidation. Overall, SOA from a one-substitute aromatic with more carbon in the substitute is located at a more oxidized area of the O/C vs. H/C chart (lower right in Fig. 3.) than those multiple substitute aromatic isomers with the same total number of carbon as the single substituted aromatic. SOA from isopropylbenzene is located in a lower position of the chart and to the right of propylbenzene indicating that branch carbon structure on the alkyl substitute of aromatic hydrocarbons leads to a more oxidized SOA. Lines in Fig. S7-S8 connect the O/C and H/C of resulting SOA to that of the aromatic precursor. Most SOA components show a slight H/C increase and a dramatic O/C increase from the precursor, which is consistent with results observed for methyl substituted aromatics (Li, et al., 2015a, 2016). However, H/C barely increases (1.33 to 1.34) from the propylbenzene precursor to its resulting SOA and there is even a decreasing trend from isopropylbenzene to its SOA. This indicates that a high H/C component loss reaction such as alkyl part dissociation during photooxidation is an important
reaction to SOA formation from longer carbon chain containing aromatic hydrocarbons. The carbon chain length of propylbenzene increases the possibility of alkyl fragmentation. The branching structure of isopropylbenzene facilitates fragmentation through the stability of tertiary alkyl radicals. Elemental ratio differences between xylenes and ethyltoluene can be attributed to the alkyl dilution effect, similar to the methyl dilution theory by Li, et al. (2015a, 2016). m-Ethyltoluene is an exception as it is more oxidized than m-xylene after accounting for the alkyl dilution effect. This suggests that the meta substituted longer chain alkyl is more readily oxidized since the substitute is at the end of ring-opening products. Prediction of elemental ratios from toluene and xylenes are discussed later (Section 4) to further quantify the carbon length and branching effect on SOA formation from aromatic hydrocarbons.

3.2.3 OS_c

Oxidation state (OS_c ≈ 2O/C-H/C) was introduced into aerosol phase component analysis by Kroll et al. (2011). It is considered to be a more accurate metric for describing oxidation in atmospheric organic aerosol than H/C and O/C (Ng et al., 2009; Canagaratna, et al., 2015; Lambe, et al., 2015) and therefore well correlated with gas-particle partitioning (Aumont, et al., 2012). Average OS_c of SOA formed from C_8 and C_9 aromatic isomers ranges from -0.53 to -0.20 and -0.82 to -0.22, respectively (Fig. 4), implying that the precursor molecular structure impacts the OS_c of the resulting SOA. An OS_c decrease with alkyl substitute length is observed in one-substitute aromatic hydrocarbons from toluene (toluene OS_c=-0.049; Li, et al. 2015a, 2016) to propylbenzene. However, OS_c provides the average oxidation value per carbon not considering whether these carbons start from an aromatic ring carbon or an alkyl carbon. Alkyl carbons are associated with more hydrogen than aromatic ring carbons, thus leading to a lower precursor OS_c and therefore lower SOA OS_c Dilution conjecture theory in Section 4 will be used to further explore the carbon chain length effect on aromatic hydrocarbon oxidation by considering the precursor H:C ratio. Single substitute aromatic hydrocarbons generally show higher OS_c than multiple substitute ones, consistent with the yield trend of Odum, et al (1997b). However, it is also found that ortho position moiety
containing two or three substitute aromatic hydrocarbons have analogous or even higher OS,
to single substitute aromatic hydrocarbons (\(\alpha\)-xylene -0.402±0.03 to ethylbenzene
-0.402±0.033; 1,2,4-trimethylbenzene -0.425±0.072 and \(\alpha\)-ethyltoluene -0.492.
481±0.030 to propylbenzene -0.435±0.111). This suggests that both substitute number and
position are critical to aromatic hydrocarbon oxidation and therefore SOA formation. OS,
trends also support that the meta position suppresses oxidation while the ortho position
promotes oxidation when the OS, of xylenes (\(\alpha\)-xylene>\(p\)-xylene>(insignificant) m-xylene),
ethtoluenes (\(\alpha\)-ethyltoluene>\(p\)-ethyltoluene>(insignificant) m-ethyltoluene) and especially,
trimethylbenzenes (1, 2, 4-trimethylbenzene (ortho moiety containing)> (insignificant)1, 2,
3-trimethylbenzene (ortho moiety containing)>1, 3, 5-trimethylbenzene (meta moiety
containing)) are compared separately. Further, SOA formed from isopropylbenzene shows the
highest OS, among all C\(_9\) isomers, nearly equivalent to that of ethylbenzene. This
demonstrates that the branching structure of the alkyl substitute can enhance further oxidation
of aromatic hydrocarbons.

3.3 Physical property

3.3.1 SOA Density

SOA density is a fundamental parameter in understanding aerosol morphology, dynamics,
phase and oxidation (De Carol, et al., 2004; Katrib, et al., 2005; Dinar, et al., 2006; Cross, et
al., 2007). SOA density ranges from 1.29-1.38 g/cm\(^3\) from aromatic photooxidation under low
NO\(_x\) conditions in this study (Fig. 5). The range is comparable to previous studies under
similar conditions (Esther Borrás and Tortajada-Genaro 2012; Ng, et al; 2007; Sato, et al.,
2010). There is no significant difference in the density of SOA formed from C\(_8\) and C\(_9\)
aromatic hydrocarbon isomers and molecular structure is not observed to be a critical
parameter to determine SOA density. The standard deviation results from differences in initial
conditions (e.g., initial HC/NO) that also determine the oxidation of aromatic hydrocarbons
(Li, et al., 2015b) and thus further affect density. SOA density is best correlated with the O/C
ratio and OS, (0.551 and 0.540, Table 3), consistent with the observation of Pang, et al. (2006)
that SOA density increases with increasing O/C ratio. The density prediction method developed by Kuwata, et al. (2011) based on O/C and H/C is evaluated as

\[ \rho = \frac{12 + H/C + 16 \times O/C}{7 + 5 \times H/C + 15 \times O/C} \]  

Eq-2

The black lines (Fig. 5) are predicted (Eq-2) densities and show a good agreement between predicted and measured SOA densities (-6.01% ~ 7.62%). A comparatively large negative error is found in meta containing aromatic hydrocarbons including \( \text{m-xylene, \text{m-ethyltoluene}} \) and \( \text{1,3,5-trimethylbenzene} \). It is noted that there should be more alkyl substitutes in SOA formed from meta position aromatics than other aromatics since meta position alkyl substitutes are more likely to participate into SOA products than other aromatics (Section 3.2.1 and Section 3.2.2). Previous work suggests that the increase of methyl groups could lead to a change in several key organic fragments (e.g., \( \text{CO}^+, \text{CO}_2^+ \) and \( \text{H}_2\text{O}^+ \)) thereby altering the default fragment table for elemental ratio analysis. This agrees with the density underestimation in SOA formed from meta position aromatics and supports the preference of meta position alkyl substitute to SOA products.

### 3.3.2 SOA Volatility

SOA volatility is associated with reactions such as oxidation, fragmentation, oligomerization and mass loading (Kalberer, et al., 2004; Salo, et al., 2011; Tritscher, et al., 2011; Yu, et al., 2014). SOA volatility in this study is measured as VFR. Initial (<30 minutes after new particle formation) SOA VFRs are around 0.2 for all the aromatic precursors studied and increase up to 0.58 during photooxidation. This suggests that aromatic hydrocarbon oxidation undergoes an evolution from volatile compounds to semivolatile compounds. The VFR trends and ranges are comparable to previous studies (Kalberer et al., 2004; Qi et al., 2010a; Qi et al., 2010b; Nakao et al., 2012). Fig. 6 shows the VFR at the end of aromatic hydrocarbon photooxidation (VFR\textsubscript{end}). A decreasing VFR\textsubscript{end} trend is found as the number of substitutes increase and for meta position (e.g. \text{m-xylene}) or meta position containing (e.g. \text{1,3,5-trimethylbenzene}) aromatic precursors. Strong Correlations among VFR\textsubscript{end} and chemical composition are observed in the aromatic hydrocarbons studied here (Table 3). This is consistent with recent findings that O:C ratio is well correlated to aerosol volatility (Section 3.2.1) and the density prediction method (Eq-2).
3.3.2) (Cappa, et al., 2012, Yu, et al., 2014), thereby affecting the gas-particle partitioning, which in turn relates to SOA yield. It is also observed that VFR	extsubscript{end} is strongly correlated (-0.937) with reaction rate constant ($k_{\text{OH}}$). Higher $k_{\text{OH}}$ is associated with faster reaction rates of initial aromatic precursors and is therefore expected to lead to further oxidation for a given reaction time. However, the inverse correlation between $k_{\text{OH}}$ and VFR	extsubscript{end} indicates that $k_{\text{OH}}$ value represents more than just the kinetic aspects. $k_{\text{OH}}$ increases with increasing number of substitutes on the aromatic ring. Additionally, aromatic hydrocarbons with meta position substitutes have higher $k_{\text{OH}}$ than those with para and ortha (Table S1) position substitutes. This suggests that the precursor molecular structures for aromatics associated with $k_{\text{OH}}$ values determine the extent of oxidation of the hydrocarbons and therefore impact SOA volatility more than simply the precursor oxidation rate.

### 4. Alkyl Dilution Conjecture Theory on SOA formation from aromatic hydrocarbons

The dependence of SOA formation on molecular structure can be partially represented by the alkyl carbon number. Carbon dilution theory proposed by Li et al (2016) successfully explain that methyl group impacts remain similar in SOA elemental ratios as in the aromatic precursor. The chemical composition of SOA formation from alkyl substituted aromatics is predicted by simply adding the alkyl substitute into the chemical composition of SOA formed from pure aromatic ring precursor (benzene). Methyl dilution theory (Li, et al., 2015, 2016) is extended to alkyl substitute dilution conjecture in order to investigate the influence of longer alkyl substitutes compared with methyl group substitutes. A robust prediction of SOA H/C and O/C trends for longer (C2+) alkyl substituted aromatics based on the methyl substituted aromatics will suggest a similarity in the role of methyl and longer alkyl to SOA formation; an underestimation or overestimation will indicate different oxidation pathways for aromatics with differing alkyl substitute length. Fig. 7a and Fig. 7b shows the predicted elemental ratio and OS for SOA formed from longer alkyl substitutes (-C\textsubscript{n}H\textsubscript{2n+1}, n>1) based on methyl only substitute. The elemental ratio of SOA formed from single substitute aromatic
hydrocarbons including ethylbenzene, propylbenzene and isopropylbenzene are predicted by toluene and those of ethyltoluenes are predicted by corresponding xylenes with similar alkyl substitute location. H/C and O/C are generally well predicted by alkyl dilution effect, except for \( \alpha \)-ethyltoluene and iso-propylbenzene. O/C (15%), H/C (1%) and OS\(_c\) (13%) of \( \alpha \)-ethyltoluene are slightly overestimated by only considering alkyl dilution effect. This indicates that \( \alpha \)-ethyltoluene is less oxidized than \( \alpha \)-xylene possibly due to the hindrance effect of the longer alkyl substitute.

However, OS\(_c\) prediction is close to measurement (\( \pm \)15%; Fig. 7b and Dashed line in Fig. 7a) for ethyltoluenes. This suggests that higher carbon number alkyl substitutes may suppress reactions that have little effect on OS\(_c\) but large effect on elemental ratio (e.g. hydrolysis). OS\(_c\) is largely underestimated in SOA formed from single substitute aromatic hydrocarbons, especially for isopropylbenzene (-49%) and ethylbenzene (-25%). This implies that longer alkyl substitutes are more oxidized than the methyl group on toluene. A direct \( \cdot \)OH reaction with the alkyl part of the aromatic is more favored on longer alkyl chains since tertiary and secondary alkyl radicals are more stable than primary alkyl radicals (Forstner, et al., 1997). It is also possible that oligomerization from highly oxidized carbonyl component might be more favored for long chain single alkyl substituted aromatics. The less significant OS\(_c\) underestimation from xylenes (meta and para) is due to the presence of an “inert” methyl group which lowers the average OS\(_c\). The extreme low H/C in Fragmentation on alkyl substitute of isopropylbenzene implies an additional hydrogen loss with the branching alkyl substitute can lead to a higher OS\(_c\) (-0.22\( \pm \)0.04) than propylbenzene (-0.42 \( \pm \)0.11), which possibly occurs while forming 2, 5-furandione (Forstner, et al., 1997) or 3-\( H \)-furan-2-one due to the increased stability of the isopropyl radical compared to the \( n \)-propyl radical. It is also possible that longer carbon chain substitutes might have higher probability to form other cyclic or low vapor pressure products by additional reaction due to their increased length. The similarity in \( \Gamma_{43} \)\(_{43}\) and \( \Gamma_{43}^{57,71} \) but discrepancy (insignificant) in elemental ratio among all single substitute C\(_8\) and C\(_9\) aromatics supports that additional reactions leading to further oxidization of alkyl substitutes can occur.
5. Atmospheric Implication

This study elucidates molecular structure impact on a major anthropogenic SOA source, photooxidation of aromatic hydrocarbons, under atmospherically relevant NOx conditions by analyzing SOA yield, chemical composition and physical properties. These observations, when taken together, indicate the roles of alkyl substitute number, location, carbon chain length and branching structure in aromatic hydrocarbon photooxidation. SOA yield of all C\textsubscript{8} and C\textsubscript{9} aromatic hydrocarbon isomers are comprehensively provided in this study with a focus on the impact of molecular structure. It is demonstrated that aromatic hydrocarbon oxidation and SOA formation should not be simply explained by substitute number. The promoting of SOA formation by the ortho position is found along with confirmation of the suppression effect by the para position during oxidation of aromatic hydrocarbons. It is possible due to the alkyl substitute location impact on the further oxidation of five-membered bicyclic radicals. Different carbonyl compounds can form as the ring opening products from the dissociation of five-membered bicyclic radical. It is assumed that oligomerization of these carbonyl compounds can contribute to SOA (Li, et al., 2016). Aromatic hydrocarbons with para position alkyl substitute tend to form more ketone like dicarbonyl compounds than other aromatics. Ketone might contribute less to oligomerization formation compared with aldehyde as suggested in Li, et al (2016). Meta position alkyl substitutes on aromatic ring lead to a lower extent of aromatic hydrocarbon oxidation. It might be due to a higher percentage of carbonyl with alkyl substitute formed during the oxidation of meta containing aromatics (e.g. methylglyoxal, 2-methyl-4-oxopent-2-enal), which contributes to oligomerization and thereby SOA formation. Evidence is provided to demonstrate Aromatic aromatic oxidation is proved to increase with alkyl substitute chain length and branching structure. Further, carbon dilution theory developed by Li., et al (2015a) is extended to this study. Carbon dilution theory not only serves as a tool to explain the difference in SOA components due to the difference in substitute alkyl carbon number but also acts as a standard to determine the oxidation mechanism based on alkyl substitute structure. Moreover, the five subcategories of aromatics and their two product modeling curve fitting parameters in this work at more realistic NO\textsubscript{x} loadings provide a more precise prediction of SOA formation.
form aromatic hydrocarbons under atmospheric conditions. Previous studies found that the humidity insignificantly impacts SOA yield from aromatic hydrocarbons (Cocker, et al., 2001) or maintains the SOA yield relationship between isomers (Zhou, et al., 2001). Therefore, it is predicted that the observation found under dry conditions in this study, especially the molecular structure impact on SOA formation from different aromatic isomers could be extended to atmospherically relevant humidity conditions. However, recent studies observe that the hydration of carbonyls and epoxides could lead to further heterogeneous reaction and oligomerization (Jang, et al., 2002; Liggio, et al., 2005; Minerath and Elrod, et al., 2009; Lal, et al., 2012). It is possible that aerosol compositions and the hygroscopic properties could be altered after the heterogeneous reactions, especially under humid conditions. The impact of molecular structure impact on SOA formation under humidity condition needs to be further studied to extend the findings in current work. This study improves the understanding of SOA formation from aromatic hydrocarbons and contributes to more accurate SOA prediction from aromatic precursors. Further study is warranted to reveal the detailed oxidation pathway of aromatic hydrocarbons with longer (carbon number >1) alkyl substitutes.

Acknowledgments

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Reference


Dinar, E., Mentel, T., and Rudich, Y.: The density of humic acids and humic like substances (HULIS) from fresh and aged wood burning and pollution aerosol particles, Atmos. Chem. Phys., 6(12), 5213-5224, doi:10.5194/acp-6-5213-2006, 2006.


Mohr et al. (2009)......


Salo, K., Hallquist, M., Jonsson, Å.M., Saathoff, H., Naumann, K.-H., Spindler, C., Tillmann, R., Fuchs, H., Bohn, B., Rubach, F., Mentel, T. F., Müller, L., Reinnig, M., Hoffmann, T., and


Tritscher, T., Dommen, J., DeCarlo, P. F., Gysel, M., Barret, P. B., Praplan, A. P., Weingartner, E., Prévôt, A. S. H., Riipinen, I., Donahue, N. M., and Baltensperger, U.:


Table 1. Experiment conditions

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<th>HC ppm</th>
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<th>NOX µg m⁻³</th>
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<td>1162A</td>
<td>15.8</td>
<td>33.4</td>
<td>80.1</td>
<td>391</td>
<td>46.5</td>
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<td>1162B</td>
<td>14.9</td>
<td>40.0</td>
<td>80.4</td>
<td>399</td>
<td>46.6</td>
<td>0.117</td>
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<td>1, 3, 5-Trimethylbenzene</td>
<td>1153A</td>
<td>65.2</td>
<td>11.0</td>
<td>79.5</td>
<td>309</td>
<td>12.4</td>
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<td>1153B</td>
<td>35.3</td>
<td>20.4</td>
<td>80</td>
<td>381</td>
<td>19.6</td>
<td>0.051</td>
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</table>
Table 2. Two product yield curve fitting parameters for one, two ( ortha, meta and para) and three alkyl substitutes

<table>
<thead>
<tr>
<th>Yield Curve</th>
<th>α_1</th>
<th>K_{om,1} (m³·µg⁻¹)</th>
<th>α_2</th>
<th>K_{om,2} (m³·µg⁻¹)</th>
<th>MSRE</th>
</tr>
</thead>
<tbody>
<tr>
<td>One Substitutes</td>
<td>0.144</td>
<td>0.039</td>
<td>0.137</td>
<td>0.005</td>
<td>5.38</td>
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<td>Two Substitutes-ortho</td>
<td>0.158</td>
<td>0.249</td>
<td>0.024</td>
<td>0.005</td>
<td>2.03</td>
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<tr>
<td>Two Substitutes-meta</td>
<td>0.156</td>
<td>0.040</td>
<td>0.080</td>
<td>0.005</td>
<td>2.51</td>
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<td>Two Substitutes-para</td>
<td>0.154</td>
<td>0.025</td>
<td>0.036</td>
<td>0.005</td>
<td>1.21</td>
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<tr>
<td>Three Substitutes</td>
<td>0.180</td>
<td>0.025</td>
<td>0.052</td>
<td>0.005</td>
<td>0.84</td>
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Table 3. Correlation among SOA density, volatility (VFR) and SOA chemical composition

<table>
<thead>
<tr>
<th></th>
<th>f_44</th>
<th>f_57</th>
<th>f_71</th>
<th>O/C</th>
<th>H/C</th>
<th>O/S</th>
<th>k_{OH}</th>
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<tbody>
<tr>
<td>Density</td>
<td>0.324</td>
<td>-0.056</td>
<td>-0.38</td>
<td>0.551</td>
<td>-0.301</td>
<td>0.540</td>
<td>-0.249</td>
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<td>VFR_{end}</td>
<td>0.537</td>
<td>0.862</td>
<td>0.223</td>
<td>0.063</td>
<td>0.341</td>
<td>0.070</td>
<td>0.435</td>
</tr>
<tr>
<td>p-value</td>
<td>0.304</td>
<td>0.089</td>
<td>0.073</td>
<td>0.224</td>
<td>0.144</td>
<td>0.058</td>
<td>0.000</td>
</tr>
</tbody>
</table>

Note: a) Mean squared error (MSRE)= [(Fitted Yield - Measured Yield)/ Measured Yield]²/(Number of Data Points)

Note: Song, et al, 2005; Song, et al, 2007; Li, et al., 2015a-b; 2016 data are also included; 123TMB- 1, 2, 3-Trimethylbenzene; 135TMB- 1, 3, 5-Trimethylbenzene; 124TMB- 1, 2, 4-Trimethylbenzene.
Fig. 1. Aromatic SOA yields as a function of $M_2$.

Fig. 2. $f_{44}$ vs. $f'_{43+57+71}$ in SOA formed from different aromatic hydrocarbon photooxidation under low NOx colored by aromatic isomer type and marked with individual aromatic hydrocarbon species: Ethylbenzene 2084A; Propylbenzene 1245A; Isopropylbenzene 1247A;
*Error bar stands for standard deviation when significant particles are formed (>5 μg/m³).*
Fig. 3. H/C vs. O/C in SOA formed from different aromatic hydrocarbon photooxidation under low NOx colored by aromatic isomer type and marked with individual aromatic hydrocarbon species (C8 and C9 on the lower left indicate the location of initial aromatic hydrocarbon precursor): Ethylbenzene 2084A; Propylbenzene 1245A; Isopropylbenzene 1247A; m-Xylene 1191A; m-Ethyltoluene 1199A; o-Xylene 1320A; o-Ethyltoluene 1179A; p-Xylene 1308A; p-Ethyltoluene 1194A; 1, 2, 3-Trimethylbenzene (123TMB) 1162A; 1, 2, 4-Trimethylbenzene (124TMB) 1119A; 1, 3, 5-Trimethylbenzene (135TMB) 1156A. Alkyl number trend is the linear fitting in (Li., et al., 2015a). Solid black cycles are SOA elemental ratio from C8 and C9 aromatic hydrocarbon predicted by SOA elemental ratio formed from benzene. *Error bar stands for H/C and O/C standard deviation when significant particles are formed (>5μg/m³).
Fig. 4. Oxidation state ($OS_c$) of SOA formed from different aromatic hydrocarbon photooxidation under low NOx: Ethylbenzene 2084A; Propylbenzene 1245A; Isopropylbenzene 1247A; m-Xylene 1191A; o-Xylene 1320A; p-Xylene 1308A; m-Ethyltoluene 1199A; o-Ethyltoluene 1179A; p-Ethyltoluene 1194A; 1,2,3-Trimethylbenzene (123TMB) 1162A; 1,2,4-Trimethylbenzene(124TMB) 1119A; 1,3,5-Trimethylbenzene(135TMB) 1156A.
Fig. 5. Measured and predicted SOA density from different aromatic hydrocarbon photooxidation under low NOx (Colored with substitute number and length, one substitute-red, xylenes-green, ethyltoluenes-blue and trimethylbenzene-purple; black line is predicted density according to Kuwata, et al., 2011); 123TMB- 1, 2, 3-Trimethylbenzene; 135TMB- 1, 3, 5-Trimethylbenzene; 124TMB- 1, 2, 4-Trimethylbenzene.
Fig. 6. SOA Volume fraction remaining (VFR_{end}) at the end of aromatic hydrocarbon photooxidation under low NO_x (Colored with substitute number and length, one substitute-red, xylenes-green, ethyltoluences-blue and trimethylbenzene-purple); 123TMB- 1, 2, 3-Trimethylbenzene; 135TMB- 1, 3, 5-Trimethylbenzene; 124TMB- 1, 2, 4-Trimethylbenzene.
Fig. 7. Comparison of measured and predicted elemental ratio (a) and oxidation state (b) of SOA formed from longer alkyl substitute (-C$_{2}$H$_{2n+1}$, n>1). Ethyltoluences are predicted by corresponding xylenes and one substitute aromatic hydrocarbons are predicted by toluene.*Predicted elemental ratio of isopropylbenzene is same as propylbenzene (not showed in Fig. a)