Laboratory studies of fresh and aged biomass burning aerosol emitted from east African biomass fuels - PART 2 - Chemical properties and characterization

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20 **Abstract:** There are many fuels used for domestic purposes in east Africa, producing a significant atmospheric burden of the resulting aerosols, which includes biomass burning particles. However, the aerosol physicochemical properties are poorly understood. Here, combustion of Eucalyptus, Acacia, and Olive fuels were performed at 500 and 800 °C in a tube furnace, followed by immediate filter collection for fresh samples or introduction into a photochemical chamber to simulate atmospheric photochemical aging under the influence of anthropogenic emissions. The aerosol
25 generated in the latter experiment was collected onto filters after 12 hours of photochemical aging. 500 and 800 °C were selected to simulate smoldering and flaming combustion, respectively, and to cover a range of combustion conditions. Methanol extracts from Teflon filters were analyzed by ultra-performance liquid chromatography interfaced to both a diode array detector and an electrospray ionization high-resolution quadrupole time-of-flight mass spectrometer (UPLC/DAD-ESI-HR-QTOFMS) to determine the light-absorption properties of biomass burning
30 organic aerosol constituents chemically characterized at the molecular level. Few chemical or UV/Visible differences were apparent between samples for either fuel when combusted at 800 °C. Differences in single scattering albedo (SSA) between fresh samples at this temperature were attributed to compounds not captured in this analysis, with eucalyptol being one suspected missing component. For fresh combustion at 500 °C, many species were present, where lignin pyrolysis and distillation products are more prevalent in Eucalyptus, while pyrolysis products of cellulose
35 and at least one nitroaromatic species were more prevalent in Acacia. SSA trends are consistent with this, particularly if the absorption of those chromophores extends to the 500 – 570 nm region. Upon aging, both show that resorcinol or catechol was removed to the highest degree, and both aerosol types were dominated by loss of pyrolysis and distillation products, though both differed in the specific compounds being consumed by the photochemical aging process.

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1 Introduction

This is the second part of a two-part laboratory measurement of optical and chemical properties of biomass burning (BB) aerosols emitted by east African fuels, focused on chemical properties. Biomass fuels were combusted at two different temperatures: Combustion at 500 °C corresponds to a smoldering-dominated fire, which leads to the formation of large quantities of molecular organic compounds and brown carbon (BrC) constituents in the resulting aerosol (Collier et al., 2016). These smoldering BB aerosols typically have modified combustion efficiency (MCE) values < 0.9. Combustion at 800 °C corresponds to a flaming-dominated fire that mainly forms black carbon (BC) aerosols (Reid et al., 2005), with MCE values > 0.9. Given the lack of MCE measurements in wildfires or in-cook stoves in east Africa, a range of combustion conditions were covered in this work. This also highlights the paucity of available measurements of combustion conditions in Africa and the need for such measurements, so that appropriate laboratory scale experiments can be performed.

A soot photometer aerosol mass spectrometer (SP-AMS) characterization of the chemical composition of BC-containing aerosol emitted by BB fuel sources was conducted during the Fire Lab at Missoula Experiments (FLAME) (McClure et al., 2019;May et al., 2014). The work focused on analyzing variations in the particle composition as a function of fuel and combustion conditions. Differences in the refractory black carbon (rBC) were observed with some fuel sources. Organic species were found to vary by orders of magnitude relative to rBC, depending on the fuel source. Chemical differences were also observed between the aged and fresh samples (Fortner et al., 2018).

A component of organic aerosol, BrC, has emerged as a significant type of BB aerosol. Also known as light-absorbing organic carbon, BrC in atmospheric particles has received much attention for its potential role in global radiative forcing (Qin et al., 2018). Primary organic aerosol (POA) contributes an estimated 10% of total solar absorption by BB aerosols, while aged OA contributes an estimated 30%, making secondary organic aerosol (SOA) production an important source of BrC (Kumar et al., 2018). As few as 20 −25 BrC chromophores are responsible for 40−50% of total UV absorbance in the 300 − 500 nm range (Laskin et al., 2018). While several field measurement campaigns have differentiated light absorption by BC and BrC, the chemical characteristics of BrC are not currently well understood (Qin et al., 2018).

BrC is often associated with emissions from incomplete combustion (Andreae and Gelencsér, 2006), but can also form via secondary reactions in the atmosphere (Laskin et al., 2015). Aerosol obtained from 500 °C burn classified as BrC aerosol are spherical in morphology as seen in the TEM images and are yellow-brown in color due to values of the imaginary portion of the refractive index (κ) that increase sharply toward shorter visible and ultraviolet wavelengths (Bond and Bergstrom, 2006). BrC is comprised of a wide range of poorly characterized compounds exhibiting highly variable light-absorption properties, with reported κ values spanning two orders of magnitude (Saleh et al., 2013;Chen and Bond, 2010;Kirchstetter et al., 2004;McMeeking et al., 2009). Its optical properties have been shown to change through atmospheric processing, such as oxidation, absorption of solar radiation leading to particle-phase reactions, and aqueous-phase processing within aerosol particles (Lambe et al., 2013;Lambe et al., 2011;Sareen et al., 2013;Lee et al., 2014;Zhao et al., 2015;Nguyen et al., 2012;Laskin et al., 2015).

These factors make the chemical composition of BrC dependent on location and source of fuel (Laskin et al., 2015;Moise et al., 2015). Many BrC chromophores vary significantly among biomass species, although some, such as sinapaldehyde and coniferaldehye are common among certain types of biomass fuels, such as angiosperms and gymnosperms, respectively. Other chromophores depend on combustion conditions, such as vanillic acid, which has only been observed as a product of smoldering combustion, when the MCE is low (Fleming et al., 2019). Further, an effect known as photo-bleaching can occur, where POA chromophores can lose their absorptivity or be destroyed entirely when irradiated by UV light for several hours (Laskin et al., 2018). Since the light-absorption spectra of organic compounds is dependent on their molecular structure, identifying atmospheric chromophores making up BrC is essential for understanding the changes in aerosol optical properties (Moise et al., 2015;Kitanovski et al., 2012a;Mohr et al., 2013;Teich et al., 2017;Zhang et al., 2013). Several recent studies have shown that BrC cannot be separated or easily distinguished from the rest of the organic aerosol (Saleh, 2020). BrC is classified into four optical regions as a continuum associated with a continuum of physicochemical properties, including molecular sizes, volatility, and solubility(Saleh et al., 2018;Saleh, 2020) . The language used in this work for BrC was mainly to contrast it with BC, based on the combustion temperature, which is decidedly a different component of aerosol, even if inextricably mixed in BBA.

In the companion paper, Part I, we described the laboratory measurements of a fuel specific study of the optical properties of BB aerosol emitted by three different fuels sourced from east Africa (Smith et al., 2020). This study was conducted under different aging and combustion conditions using a tube furnace and indoor smog chamber. Optical properties were measured for BB aerosols produced under smoldering and flaming conditions for each fuel type. For each combustion condition, we reported the measured optical properties. These include scattering and extinction cross sections, single scattering albedo (SSA), absorption and extinction for fresh emissions, emissions aged in the dark and emissions photochemically aged in the absence of added volatile organic compounds (VOCs) and with VOCs added to represent urban emissions from a representative African megacity. In this part of the manuscript, we report the chemical composition of fresh and BB aerosols photochemically aged in the presence of VOCs, which were combusted at two different temperatures and collected onto filters.

Identifying the light-absorbing chromophores and non-absorbing aerosol constituents by chemically characterizing BrC is a challenging task (Lin et al., 2015b). In this Part 2 manuscript, ultra-performance liquid chromatography (UPLC) was interfaced to both photodiode array spectrophotometry (or DAD) and high-resolution quadrupole time-of-flight mass spectrometry (HR-QTOFMS) to chemically characterize BrC chromophores at the molecular level for both fresh and aged emissions, and is compared with similar measurements for BC. This technique (i.e., UPLC/DAD-ESI-HR-QTOFMS) has previously been used for the molecular level characterization of BrC constituents in laboratory-generated SOA (Lin et al., 2015a;Lin et al., 2016;Lin et al., 2014) and in ambient aerosols and cloud water impacted by BB (Kitanovski et al., 2012b;Budisulistiorini et al., 2017;Zhang et al., 2013).

2 Experimental methods

115   In companion paper Part I, (Smith et al., 2020) we described the aerosol generation and combustion system, the smog chamber characteristics, and the aging conditions and process. A summary is provided here. For laboratory samples, BB aerosols were generated by combusting 0.5 g of biomass samples in a tube furnace (Carbolite Gero, HST120300-120SN). Smoke and gases produced from combustion were sent directly into the smog chamber via a heated (200 °C), ¼ inch corrugated stainless-steel transfer tube. The North Carolina A&T State University (NCAT)

120   indoor smog chamber has a volume of 9.01 m^3 and is lined by FEP Teflon. Two sides of the chamber each have a bank of 32 ultraviolet (UV) lights (Sylvania, F30T8/350BL/ECO, 36"), for a total of 64 lamps, to produce photochemical reactions. For the purposes of this experiment, a clean environment is maintained by flushing the smog chamber for at least 24 hours with clean air coming from a clean air generator until negligible particle concentrations are reached.

125   The particle size distribution was continuously measured during the experiments. In general, it took about 15−20 minutes for the size distribution to become a Gaussian distribution. In addition, we checked the total mass concentration reported by the aerosol instrument manager software and confirmed that after 15−20 minutes of combustion the mass loading become stable. We characterized this as a well-mixed condition in the chamber. An example of the size distribution for each fuel during an experiment is given in Figure S1.

130   For this study, authentic fuel plants were obtained from east Africa. These samples were weighed on a calibrated analytical balance so that it would approximately yield a total aerosol loading representative of a particular scenario (urban, wildfire, etc.). We utilized previously measured emission factors (Akagi et al., 2011;Simoneit, 2002;Yokelson et al., 2013;Andreae and Merlet, 2001), such as 18.5±4.1 g PM_{10} kg^{-1} wood (dry weight) for tropical forest fuels (Akagi et al., 2011). For instance, to achieve a mass loading of 1100 μg m^{-3}, which is the mean loading

135   found in urban/suburban residential locations (Oyem and Igbafe, 2010), 0.5 g of wood was burned in these experiments. As determined by SMPS spectra and assuming a particle density of 1 g cm^{-3}, typical peak particle concentrations were 800-900 μg m^{-3} for combustion at 500 °C and ~80 μg m^{-3} for combustion at 800 °C. Since aerosol densities are generally larger than 1 g cm^{-3}, our estimated mass represents the lower limit of the actual values.

        Optical measurements were performed after the chamber was mixed, since they could not be taken directly

140   from the furnace, given the long measurement time and high particle concentrations. While several types of aging were performed in Part 1, the work presented here focuses on photochemical aging with anthropogenic VOCs, specifically benzene, toluene, and xylene. No other atmospherically active species were added, such as HONO, NOx, O_3, etc., aside from those produced during combustion or resulting from photochemical aging. VOCs were injected into the chamber just before combustion while the UV lights are turned off, followed by 12 hours of irradiation. While

145   this is somewhat long for a chamber experiment, this length of time would represent a maximum change upon photochemical aging. Details of VOC addition is discussed in the first part of this paper (Smith et al., 2020). The purpose of adding the VOCs was to represent a polluted urban environment, where we used the emission inventory for urban environments from South Africa. This does not necessarily represent the east African emission inventory, but this serves as a baseline, since this is the only available data to us for the continent Anthropogenic VOC

150   concentrations were based on observations from several urban sites that were in the South African Air Quality Information System (SAAQIS). Average values for mid-July, the middle of the peak burning season for South Africa

for the year 2016, were ~1.25, 3.5, and 1.5 ppm for benzene, toluene, and xylene, respectively. While these ratios were maintained in this chamber work, amounts introduced into the chamber were ~12 times more concentrated (15, 42, and 18 ppm, respectively) due to sample preparation constraints, since the amounts needed for an exact match were too small to weigh accurately. In previous work, the total particle volume concentration (nm^3 cm^{-3}) was found to undergo a first order decay (Smith et al., 2019). The loss rate constant for this process was $(1.34 \pm 0.02) \times 10^{-3}$ min^{-1}, which corresponds to a lifetime of 12.4 hours, so these samples underwent ~1 lifetime of particle losses. The chamber relative humidity was 0% or under the detection limit of the measurement device. Chamber temperature was $20 - 23$ °C, depending on the temperature of the room. The temporal evolution of chamber temperature and other characteristics have been extensively investigated for the NCAT indoor smog chamber (Smith et al., 2019).

2.1 Fuel types and condition

Eucalyptus, Acacia, and Olive have many things in common besides their wide distribution of growth. All are Angiosperms and Eudicots. Acacia and Eucalyptus are more closely related, with both belonging to the Rosid clade, while Olive is an Astrid. This is where their evolutionary similarities end, with Acacia, Eucalyptus, and Olive in the orders Fabale, Myrtale, and Oleaceae, respectively. The common names of these species, followed by their botanical author citation are Eucalyptus (L'Hér), Acacia (Mart.), and Olive (L.). The fuel moisture content was 10%. As such, they had minimal moisture content and the fuels studied in this work are likely most relevant for situations where these fuels are combusted in home heating and cooking or as the litter component of wildfires.
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2.2 Collection of filter samples

BB aerosols were collected onto 47-mm Teflon filters (Tisch Environmental, SF18040) for chemical analysis and tunneling electron microscope (TEM) grids (Ted Pella, 01844, carbon film on 400 mesh copper) for image analysis. For fresh BB aerosol, these samples were collected inline immediately after the furnace, only allowing enough distance from the furnace for the sample to cool before reaching the filters. This allowed fresh samples to be collected as close to the initial combustion as possible. Flow of the aerosol through the Teflon filter was driven by the chamber instrumentation, specifically the CO and CO_2 analyzers, at ~2.5 L min^{-1}, while flow through the TEM grid was provided by a modified aquarium pump at 0.5 L min^{-1}. With the output from the furnace at 10 L min^{-1}, pressure was maintained through the system by diverting the excess flow (7 L min^{-1}) to the exhaust. Samples were collected for the duration of the burn, which was typically around 10 minutes. In this manner, approximately 25% and 5% of the total BB aerosol were collected onto the Teflon filter and TEM grid, respectively, after losses in the tube furnace and transfer tubing.

For aged BB aerosol, samples were taken directly from the chamber. Since the aerosol had already been diluted in the chamber, a larger sample volume was necessary for deposition onto the filters. For these samples, aerosol was taken from one of the chamber outputs, with a combined flow of 5 L min^{-1} through the Teflon filter and TEM grid setup produced by a separate pump. Samples were collected for 18 min, which corresponded to 90 L of chamber

air, or ~1% of the total chamber volume. With aged filter samples taken after approximately 24 hours in the chamber, particle concentration had been significantly reduced since initial combustion, due to particle loss and dilution. However, the resulting samples were still large enough to be seen by the naked eye on the filter paper. Unused filters underwent the same extraction process as the sample filters to serve as a blank sample.

We estimated the approximate mass loading in the chamber during each combustion case by converting the SMPS size distribution into the total mass by assuming a density of 1 g cm^{-3}. Our estimated mass represents the lower limit of the actual values. During the 500 °C combustion cases, the typical concentration of aerosol after being well mixed was about 800 µg m^{-3}, while that of the 800 °C cases was about 100 µg m^{-3}. After 12 to 15 hours of aging, the initial mass loading was reduced by a factor of two, with a resulting concentration of about 400 µg m^{-3} and 50 µg m^{-3} during the 500 °C and 800 °C combustion cases, respectively.

2.3 UPLC/DAD-ESI-HR-QTOFMS

An Agilent 1200 Series UPLC system interfaced to a 6520 Series Accurate Mass Q-TOFMS instrument (Agilent Technologies, Santa Clara, CA) and equipped with an ESI source (Lin et al., 2014) was operated in negative ion mode to chemically characterize BB organic aerosol (BBOA) constituents in filter samples collected from the tube furnace for fresh emissions and from the smog chamber for aged experiments. In brief, chromatographic separations were carried out using a Waters ACQUITY UPLC HSS T3 C$_{18}$ column (2.1×100 mm, 1.7 µm particle size, Waters Corporation, Milford, MA) at 45 °C. The mobile phases consisted of eluent (A) 0.1% acetic acid in ultrapure water (>18 MΩ cm), and eluent (B) 0.1% acetic acid in pure methanol (99.9%, Fisher Chemical). The gradient elution program was as follows: eluent (A) held at 100% from 0 − 2 min, decreased linearly to 10% from 2−10 min, held constant at 10% between 10 −11 min, increased linearly to 100% from 11−15 min, and held constant at 100% during a 5-min post run for column re-equilibration. The sample injection volume was 5 µL at a flow rate of 0.3 mL min^{-1}. A diode array detector (DAD) between the UPLC system and the mass spectrometer was operated to measure UV-visible (UV-vis) absorbance from 200 to 800 nm with a step of 2 nm. At the beginning of each analysis period, the mass axis of the Q-TOFMS was calibrated using a commercially available ESI-L low-mass tuning mixture (Agilent Technologies, Santa Clara, CA) containing seven masses ranging from 60 -1700 Da. For real-time mass correction, a solution containing 3 reference mass components was continuously infused for real-time mass axis correction (calibration). The resultant mass resolution of the ESI-HR-QTOFMS over *m/z* 60 −1700 ranged from ~11,000 at the low mass end to ~17,000 at the high mass end. Raw data were acquired and processed with MassHunter Software (Version B.06.00, Build 6.0.633.0, Agilent Technologies, Santa Clara, CA).

2.4 Sample preparation and BrC identification

Filter samples collected as described in the last section were extracted individually in 22 mL of pure methanol by sonication for 45 minutes, blown dry under a gentle nitrogen stream at room temperature (21−22°C), and reconstituted in 300 µL of 50:50 (v/v) methanol/water solvent mixture for the subsequent UPLC/DAD-ESI-HR-QTOFMS analysis. While the water-soluble fraction of BrC is less than 70%, almost 90% of BrC can be extracted by organic solvents, such as methanol. Furthermore, the water-insoluble fraction of BrC is more absorbing than the water-

soluble fraction (Lin et al., 2017;Bergstrom et al., 2007;Laskin et al., 2018). Immediately prior to analysis, each reconstituted extract was filtered through a PTFE syringe filter (Agilent, 0.2-μm pore size) to remove undissolved particles such as soot components. A similar procedure was used by other research studies (Kumar et al., 2018;Jiang et al., 2019).

Previous work went into quantifying BrC aerosol constituents identified by UPLC/DAD-ESI-HR-QTOFMS, seven external (2-nitrophenol, 4-nitro-1-naphthol, 4-methyl-5-nitrocatechol, vanillin, vanillic acid, coniferaldehyde, and benzoic acid) commercially-available authentic standards were prepared for 6-point calibration curves from 0.004 −10 μg mL^{-1}, with detection limits below 0.004 μg mL^{-1}. While several of these methods were done in this work, determining the absolute quantity of a BrC constituent was not pursued. From the previous tests of quality control, the recovery rate of ketopinic acid was ~90% among 20 BB aerosol samples with the same extraction process with methanol and the UPLC/DAD-ESI-HR-QTOFMS method. Two filter samples (with the collected PM$_{2.5}$ masses > 1 mg) were re-extracted using 70:30 (v/v) acetonitrile/toluene solvent mixture. Most of the identified BrC species were below detection limit from this second extraction, suggesting excellent extraction efficiency for the first time with methanol. Only a few BrC aerosol constituents (e.g., vanillic acid, coniferaldehyde) were detected from the second extraction, but their abundance was lower than 1.5% of that from the first extraction. The only exception was 2-nitrophenol, whose extraction efficiency using methanol was ~55% for the first time.

2.5. Non-targeted analysis

Non-targeted MS analyses were performed to compare the filter samples collected from the smog chamber experiments under different conditions, such as fuel type, combustion temperature, and aging, to reveal the major molecular secondary BBOA products, with a focus on BrC. The raw data files acquired from UPLC/ESI-HR-QTOFMS were first transformed into ".mzData" format by Agilent MassHunter and then processed by the online metabolomics platform XCMS (https://xcmsonline.scripps.edu/) (Tautenhahn et al., 2012). During this transformation, only peaks that had intensities above 0.6% of the base peak were selected. Workflow parameters were adopted from previous work with modifications (Tian et al., 2017). In brief, the "centWave" algorithm was used for peak detection, with a peak width range from 5 − 60 seconds, and the mass error tolerance was set at 20 ppm. Peak alignment required an *m/z* width (mzwid) at 0.025. Generally, around 30 major BBOA constituents were filtered out by XCMS, and were then examined in MassHunter to make sure they possessed good peak shapes and were not present in the filter blank samples. The resulting blank and sample chromatograms are presented in Figure S2.

The following data fields were used for each compound returned by the XCMS platform: median *m/z* ratio, median retention time (rt), and the intensities for each sample in the pairwise comparison. Compounds with a similar *m/z* ratio, regardless of observed retention times had their relative intensities combined, as they most likely returned the same chemical formula. Later, for relatively large peaks with slightly different median *m/z* values, when species were found to have the same molecular formula, as determined by MassHunter (see below), their ion intensities were also combined. The amount of aerosol varies significantly from sample to sample, especially between fresh filter samples collected directly from the furnace during combustion and aged filter samples collected from the chamber, with the former always being much greater. Due to the lack of authentic standards for quantifying individual BBOA,

these masses were not determined at the time of their collection, and their spectral intensities could not be normalized to a specific aerosol mass or tracer for aerosol mass. As such, no direct comparison between samples could be made without first normalizing spectra. We normalized the intensity data by generating a scaled intensity difference between the samples using the following equation:

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$$\frac{x_1}{\sum x_{1,i}} - \frac{x_2}{\sum x_{2,i}} \qquad (1)$$

where x_1 is the intensity from the first sample and x_2 is the intensity from the second sample, and each term represents the fractional integrated ion intensity of their respective samples. Positive values indicate that a compound was more abundant in the first sample, while negative values indicate that a compound was more abundant in the second sample.

275 The results of the XCMS platform were then sorted to exclude any absolute scaled intensity differences less than 0.5 %, which left around 30 compounds or fewer to identify for each pairwise comparison.

The MassHunter software was then used to confirm and identify the compounds found by the XCMS platform. For each pairwise comparison, extracted-ion chromatograms (EICs) were generated for both samples and

280 any relevant blank samples for each of the *m/z* ratios found by XCMS. MS results were then generated from any peaks found in the EICs. MassHunter's built-in formula generator was used to identify the most likely compounds associated with the *m/z* ratio and retention time identified by XCMS. These chemical formulas, along with their double bond equivalent (DBE), mass difference, and number of chromatographic peaks (at different retention times, most likely isomers of the compound) were cataloged, with updated *m/z* ratios and retention times, as identified by MassHunter.

285 Lastly, suggested identities were assigned to each chemical formula based on relevant literature. Any compounds that were also found in the blank sample(s) were removed from the catalog.

2.6 UV/Visible analysis

290 The UV/Visible spectra in this work were not done for specific molecular species, but for all molecular species that were extracted and chromatographically/spectroscopically measured. Spectra were derived by integrating across the entire chromatogram within MassHunter. Each was background subtracted from a blank run, though the resulting spectra could not be compared directly because of the reasons mentioned earlier. Since some negative absorption values were produced after background subtraction, baseline values were shifted such that the smallest

295 absorption intensity above 250 nm was made barely positive (by 0.00001 mAU) and all values were divided by the spectral maximum to produce a peak value of 1 in the 250 – 800 nm range. The reason all values were made slightly positive was that it enabled the determination of a normalized Angstrom absorption exponent (NAAE). Because the baseline was shifted to allow for a log plot, this is mathematically different than the AAE, and should only be used for comparison purposes. With a focus on the 500 − 570 nm region, the NAAE and the normalized slope vs.

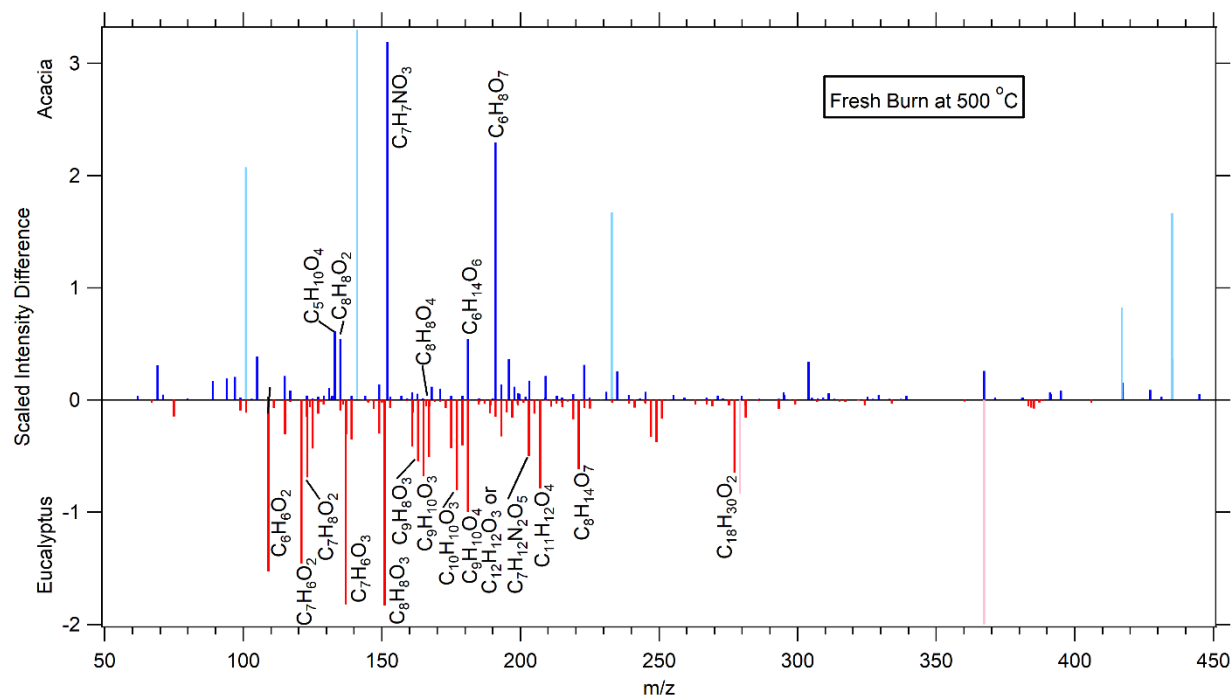
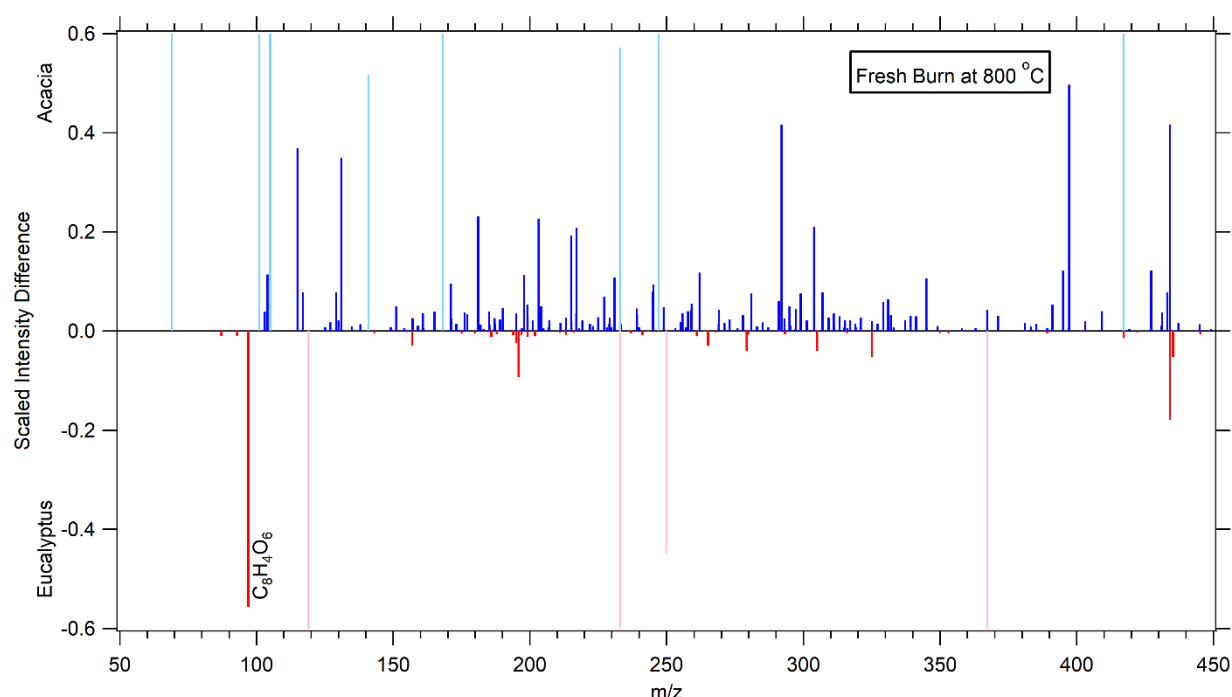
300 wavelength was determined. Heat map chromatograms of absorption vs wavelength and retention time are given in Figure S3.

3 Results and discussion

 3.1 MS analysis of fresh emissions

       Because the chemical measurements of the BB aerosols derived from Acacia and Eucalyptus fuels were the most robustly sampled in this work, a comparison of those results is presented here, though a few references to Olive will be made. Figure 1 shows the UPLC/ESI-HR-QTOFMS negative ion mode difference spectra produced for these

310   BB aerosol types. Peaks that are positive are more abundant in the Acacia-derived BB aerosol sample, while negative peaks are more present in the Eucalyptus-derived BB aerosol sample. Table 1 highlights the molecular differences between the BB aerosols derived from freshly combusted Acacia and Eucalyptus fuels, showing the scaled intensity difference for species with an absolute difference greater than 0.5 %, the mean peak m/z, the mass difference between the measured and the monoisotopic mass of the formula determined using MassHunter (listed here as a measure of

315   formula identification quality), the determined formula, the double bond equivalence (DBE) of the formula, the number of chromatographic peaks seen from an EIC, and the suggested identity based on the formula and relevant literature.

320 **Figure 1.** A comparison of fresh Acacia and Eucalyptus fuels combusted at 500 °C (above) and 800 °C (below) in terms of the scaled intensity difference (see section 2.5). Species more present in Acacia-derived BB aerosols have positive values (blue lines), while those more present in Eucalyptus-derived BB aerosols have negative values (red lines). Peaks in lighter colors were present in the XCMS analysis but were found to be present in significant amounts in the blank sample. Many of these peaks that were found in the blank extend past the scale of the plot but were cut off to focus on species associated with BB aerosol. Compounds

325 having an absolute difference greater than 0.5% have been labeled. Far more differences are revealed at the lower temperature burn and different classes of compounds are expressed in different relative amounts for Acacia and Eucalyptus combustion.

Table 1. Molecular differences between fresh Acacia- and Eucalyptus-derived BB aerosols emitted by combustion at 500 °C (upper section) and 800 °C (lower section), measured by negative ion mode UPLC/ESI-HR-QTOFMS

330 analysis. Species are ordered in terms of increasing scaled intensity difference, with positive values associated with Acacia-derived BB aerosols and negative values associated with Eucalyptus-derived aerosols. The mass difference is calculated as the difference between the observed median mass and the monoisotopic peak (typically singly deprotonated) of the formula determined using MassHunter. Peak counts labeled 'M' have multiple indistinct peaks, while those labeled 'W' elute over a very long retention time (1−4 minutes), both of which could easily consist of

335 several isomers. Suggested species marked with an asterisk can, and likely do, have multiple isomers present. Those marked with a 't' are tentative assignments. Identities marked with a dagger (†) have been confirmed against an authentic standard. See the supplemental information tables for associated retention times, wherein retention times in bold are associated with an authentic standard.

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	Scaled Intensity Difference	m/z	Mass Difference (ppm)	Formula	DBE	# of Peaks	Suggested Identity
500 °C	-1.831%	151.0403	-0.95	$C_8H_8O_3$	5	5	vanillin,[†] andmethoxybenzoic acid,* or hydroxyanisaldehyde*
	-1.822%	137.02442	0.72	$C_7H_6O_3$	5	4-5	salicylic acid* or dihydroxybenzaldehyde*
	-1.526%	109.02969	-3.37	$C_6H_6O_2$	4	1	dihydroxybenzene
	-1.459%	121.02951	-0.59	$C_7H_6O_2$	5	3	benzoic acid[†]and hydroxybenzaldehyde*
	-0.998%	181.05101	0.3	$C_9H_{10}O_4$	5	3	homovanillic acid*, dimethoxybenzoic acid*, or syringaldehyde*
	-0.801%	177.05542	1.16	$C_{10}H_{10}O_3$	6	1	coniferaldehyde[†]
	-0.790%	207.06602	1.5	$C_{11}H_{12}O_4$	6	1	sinapaldehyde
	-0.691%	123.04541	-0.81	$C_7H_8O_2$	4	2	guaiacol*
	-0.677%	165.05548	5	$C_9H_{10}O_3$	5	M	hydroxy-methoxyacetophenone* (apocynin, paeonol, etc), caffeyl alcohol*, veratraldehyde,* or phloretic acid*
	-0.647%	277.21747	-1.11	$C_{18}H_{30}O_2$	4	1	octadecatrienoic acid (likely linolenic acid)
	-0.616%	221.06665	0.61	$C_8H_{14}O_7$	2	1-2	dihydroxydimethoxyoxane-2-carboxylic acid*[t]
	-0.545%	163.0399	nc	$C_9H_8O_3$	6	8+	caffeic aldehyde,* coumaric acids*
	-0.510%	167.03508	-0.03	$C_8H_8O_4$	5	3+	vanillic acid*[†]
	-0.499%	203.07032	-10 - 10	$C_{12}H_{12}O_3$ or $C_7H_{12}N_2O_5$	7 or 2	W	dimethylmethoxycoumarin* or glycyl-glutamic acid[t]
	0.543%	135.04513	-2.25	$C_8H_8O_2$	5	2	anisaldehyde*, acetophenone, methylsalicylaldehyde*, or methylbenzoic acid*
	0.545%	181.07191	-1.1	$C_6H_{14}O_6$	0	4	mannitol, sorbitol, or galactitol
	0.612%	133.05005	3.92	$C_5H_{10}O_4$	1	1	deoxyribose or monoacetalglycerol[t]
	0.994%	666.05964	-	-	-	1	too many options
	2.293%	191.0198	-2.5	$C_6H_8O_7$	3	W	citric acid
	3.190%	152.03565	-4.2	$C_7H_7NO_3$	5	1	nitroanisole*, nitrocresol, or nitrobenzyl alcohol*
800 °C	-0.556%	96.960662	9	$C_8H_4O_6$	7	1	dihydroxyphthalic acid
	0.497%	397.27546	-	-	-	1	too many options
	0.560%	666.05967	-	-	-	1	too many options
	0.789%	982.993	-	-	-	W	no formula found

Except for dihydroxyphthalic acid and other tentatively assigned compounds, there is very little difference

345   between the two fuel types when they are combusted at 800 °C. This is consistent with previous work, where flaming combustion leads to formation of predominantly BC aerosol (Reid et al., 2005). The modified combustion efficiencies (MCE), discussed in companion paper (Part I), were 0.974 ± 0.015 when averaged from 12 different burns performed at 800 °C, which supports that these aerosols represent particles that were emitted during the flaming stage of combustion, which is dominated by BC aerosol. Since most of the chemical differences between samples are very

350   small, this suggests that either very few BrC species are produced for either fuel, or there are numerous species that are essentially the same between the samples. In Part I, Eucalyptus was shown to have a higher SSA, ~0.43 for 300 nm particles, than Acacia, which is more absorbing and has an SSA of ~0.3 at that size (see Figures 3 and 4 of Part I).

This would suggest that Eucalyptus has more non-absorbing OA, or at least less absorbing than BC. It is unlikely that dihydroxyphthalic acid would produce a significant, but wavelength dependent, absorption in the 500−570 nm region of the spectrum. Conversely, it is Acacia that appears to have many more low-abundant organic constituents. This suggests one or more of the following: 1) that Eucalyptus has significantly more complex combustion products than Acacia, such that their abundances fall below the lower threshold of XCMS analysis; 2) that Eucalyptus combustion products are either unresponsive in the negative ion mode of the UPLC/ESI-HR-QTOFMS analysis, such as having very high molecular weights (MWs), which prevents them from eluting off the UPLC column, or having no acidic functionalities; 3) that those products are not extractable by methanol; 4) that those products are semi-volatile and were removed during the offline sample processing for UPLC/DAD-ESI-HR-QTOFMS analyses; or 5) that differences between the observed SSA values are due to morphology differences. Very high MW compounds, such as PAHs, are potentially present, but it is unlikely that such large molecules would lack the ability to absorb light (see Section 3.3) to produce a more scattering aerosol for Eucalyptus, thus discounting part of reason 2. Negative ion mode ESI is known to bias detection in favor of acidic compounds, such as those with carboxylic acids, nitroaromatics, sulfated organics, and those that have sufficiently acidic -OH groups. Given that light-absorbing OA seems significantly soluble in methanol (Kumar et al., 2018), reason 3 is also unlikely. If one assumes that XCMS is capturing the complexity of molecular species, it is likely that Eucalyptus products are low MW species that are semi-volatile, that those low MW species are not captured by UPLC/DAD-ESI-HR-QTOFMS analyses, that the observed differences in SSA are due to morphology differences, or a combination of the three. One potential explanation would be a compound such as eucalyptol (1,8-cineole), which is a large fraction of Eucalyptus oil, that is a cyclic ether that lacks any basic functionality, has good solubility in alcohols, and does not absorb in the UV and visible. When extracting essential oil from Eucalyptus leaves by water-steam distillation, an oil yield of 0.3 – 2.0% (dry wt.) was produced of which 7 – 75% was eucalyptol, depending on the species (Masamba et al., 2001;Subramanian et al., 2012). The presence of such a compound should be apparent under positive ion mode ESI or atmospheric pressure chemical ionization MS analysis and possibly FT-IR if present in sufficient quantities.

Combustion of each fuel at 500 °C presents a very different picture, not only because of the number of species differentiated by their mass, but also the number of isomers present for each mass. The number of chromatographic peaks in the EICs are listed for each mass in Table 1. In that table, all species are present in both samples to differing degrees, except for octadecatrienoic acid, which is likely linolenic acid, and is only present in Eucalyptus. Demonstrating this complexity, for instance, m/z 151.04 was vanillin (a hydroxy methoxy benzaldehyde confirmed with an authentic standard) (Budisulistiorini et al., 2017) and hydroxyanisaldehyde and/or methoxybenzoic acid (such as anisic acid), but consists of five isomers, likely differing in the placement of functional groups along the aromatic ring. These are previously observed lignin pyrolysis products (Simoneit, 2002) that are more prevalent in Eucalyptus. This includes only one isomer of dihydroxybenzene, which is likely catechol (Simoneit, 2002), though resorcinol is also possible, since it has been observed in gas-phase measurements of prescribed fires (Yokelson et al., 2013). At least four isomers of hydroxybenzoic acid, such as salicylic acid, and/or dihydroxybenzaldehyde, are present and more abundant in Eucalyptus. While these compounds have the same structural motifs as other lignin pyrolysis products, only salicylic acid (Fleming et al., 2019) and 4-hydroxybenzoic acid (Bi et al., 2008) have been observed previously

390   as BrC chromophores associated with BB, which suggests that there are at least two other unidentified isomers. Other lignin pyrolysis products are noticeably more abundant for Eucalyptus combustion at 500 °C and are likely BrC chromophores, such as benzoic acid (confirmed by an authentic standard) and two isomers of hydroxybenzaldehyde at m/z 121; homovanillic acid, dimethoxybenzoic acid, and/or syringaldhyde (3 isomers observed) at m/z 181 (Simoneit, 2002); coniferaldehyde at m/z 177 (confirmed); sinapaldehyde (and not another isomer) at m/z 207

395   (Fleming et al., 2019); guaiacol and some other isomer at m/z 123; and the ubiquitous vanillic acid at m/z 167 (confirmed) (Fleming et al., 2019;Simoneit, 2002). Distillation products more prevalent in Eucalyptus include an indistinct number of isomers of hydroxy-methoxyacetophenone (e.g. apocynin or paeonol), caffeyl alcohol, and/or phloretic acid at m/z 165. The last distillation product expressed more in Eucalyptus is at least one isomer of what is likely dimethyl methoxycoumarin. These species have been observed previously as a class of compounds (Fleming et

400   al., 2019;Lin et al., 2016), though not this one particularly to the author's knowledge. At least eight isomers of caffeic aldehyde, veratraldehyde, and/or coumaric acid are also present.

            Other species are present in greater abundance in Acacia at 500 °C. Saccharides such as mannitol and sorbitol have been observed as pyrolysis products of cellulose and in road dust (Simoneit et al., 2004), though two more isomers, one likely being galactitol ($C_6H_{14}O_6$), are also present. Similarly, citric acid has been observed as products

405   from saccharide combustion (Bi et al., 2008), though the authors have not been able to find references to deoxyribose or monoacetalglycerol in the literature. Nitroaromatic species have been shown to be produced during the pyrolysis process from fresh combustion in this work, where one isomer of nitroanisole, nitrocresol, or nitrobenxyl alcohol is present and produced in significantly higher quantities in Acacia than Eucalyptus.

            Work by Fleming et al. (2019) shows that sinapaldehyde and coniferaldehyde are two known lignin pyrolysis

410   products that are produced from their corresponding lignin monomeric units; specifically sinapyl and coniferyl alcohol. They also state that these species are anticorrelated, where sinapaldehyde is more prevalent in angiosperms and flowering fuels, while coniferaldehyde is more abundant in conifers and soft woods. We examined our BBOA mass spectra for sinapaldehyde ($C_{11}H_{12}O_4$, $[M-H]^- = 207.0657$) and coniferaldehyde ($C_{10}H_{10}O_3$, $[M-H]^- = 177.0552$) and found that both had a single peak with a coeluting retention time of ~9.2 minutes. Additionally, both species were

415   found in all three fuels – Eucalyptus, Acacia, and Olive. When integrating the chromatogram between 9.0 and 9.5 minutes and extracting the resulting mass spectrum, a comparison of relative ion intensities was made. The coniferaldehyde/sinapaldehyde ratio was found to be 1.6, 3.4, and 2.6 for Eucalyptus, Acacia, and Olive, respectively. In contrast to Fleming et al. (2019), we found that all of these species (all angiosperms) had a consistently higher amount of confieraldehyde than sinapaldehyde, though this ratio varied by about a factor of 2 between species studied

420   here. One possible reason for this was that Fleming et al. reported their results in terms of absorption intensity; not ion intensity. Since these species coeluted during our separation, it was not possible to distinguish them on that basis.

            According to the above analysis, there should be more, and more varied, species present in Eucalyptus than Acacia. The main species present in greater abundances in Acacia are derived from sugars and cellulose, and are not chromophores. In Part I of this work, we found that Acacia has higher SSA than Eucalyptus when combusted at 500

425   °C, by 0.1 − 0.2, depending on particle size. This suggests that either Acacia has larger absolute amounts of non-chromophore compounds or Eucalyptus has a greater quantity of chromophores whose absorptive properties extend

to the 500 − 570 nm region of the visible spectrum. This will be discussed in further depth when relating MS results to UV/Visible absorption results in Section 3.3.

430 **3.2. MS Analysis of Aging Effects**

Figure 2 is similar to Figure 1, but rather than contrast different species, a single species, Eucalyptus, is differentiated in terms of the aging process. Peaks with positive values indicate those that are more present in the fresh sample, while those with negative values are more present in the aged one. Another way to interpret this is that those
435 with positive values are no longer present in the aged sample, because they have been chemically changed due to the aging process. Those with negative values are either being produced by the aging process or are durable enough to be more significant, in a relative sense, in the aged sample. Again, we have attempted to identify compounds that change by more than 0.5 %, indicating the molecular formula for those in Figure 2, and listing them in Table 2 along with their suggested identities. Similarly, Figure 3 and Table 3 show the results for fresh versus aged Acacia.

440
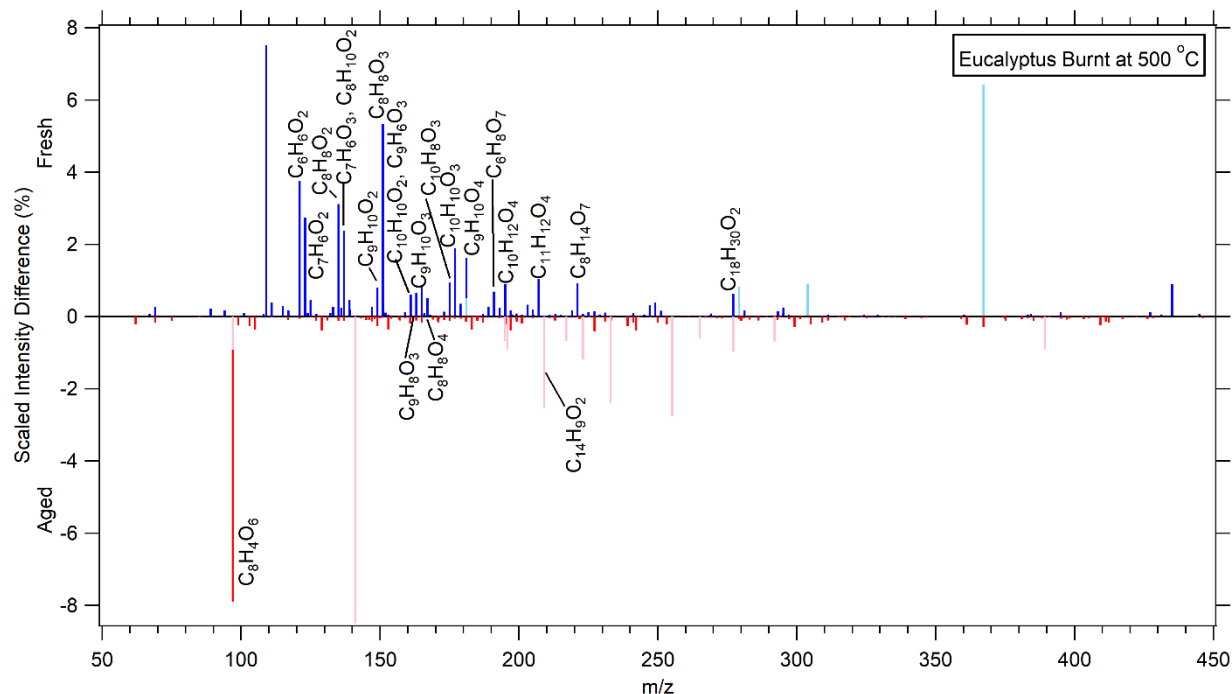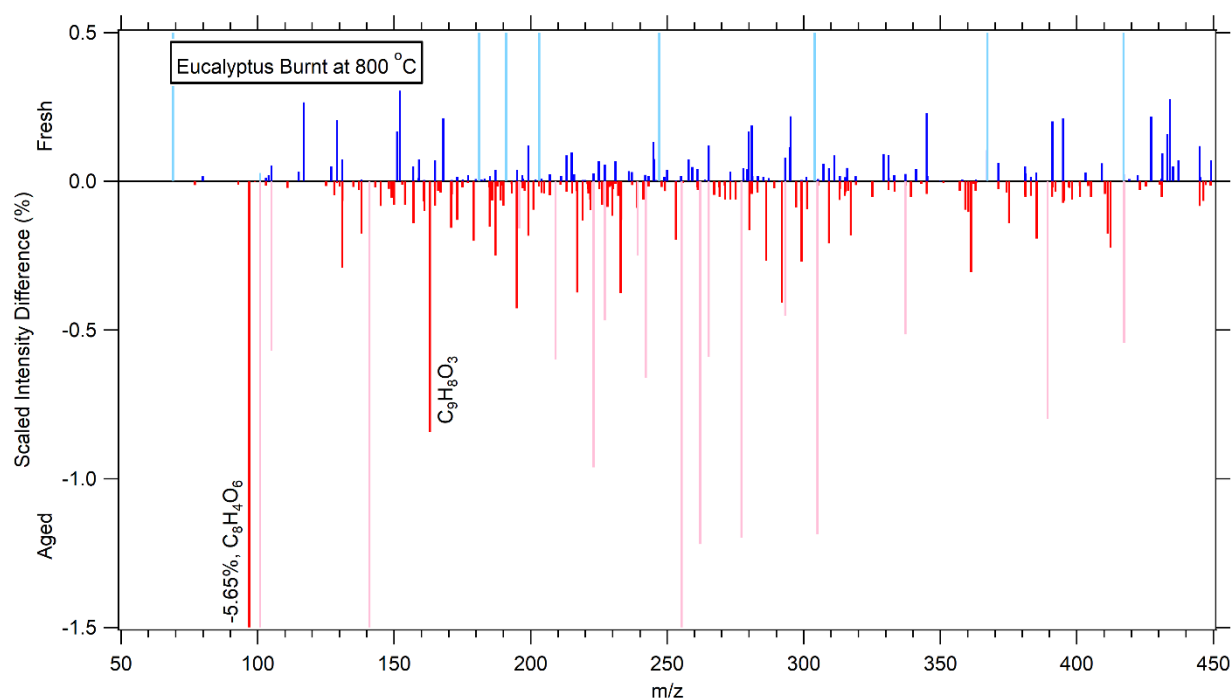
Figure 2. A comparison of Eucalyptus-derived BB aerosol combusted at 500 °C (above) and 800 °C (below) under fresh and photochemically aged conditions with VOCs in terms of the scaled intensity difference (see section 2.5). Species more present in the fresh sample have positive values (blue lines), while those more present in the aged sample have negative values (red lines) (i.e. species with positive values have been removed because of the aging process, while those with negative values are either produced or represent a greater relative contribution to the aged sample). Peaks in lighter colors were present in the XCMS analysis but were found to be present in significant amounts in the blank sample. Many of these peaks that were found in the blank extend past the scale of the plot but were cut off to focus on species associated with BB aerosol. Compounds having an absolute difference greater than 0.5% have been labeled. There is very little change for combustion at 800 °C, while 500 °C shows a net removal of many compounds.

Table 2. Molecular difference between fresh and aged Eucalyptus revealed by negative mode MS analysis. See the caption of Table 1 for more details.

	Scaled Intensity Difference	*m/z*	Mass Difference (ppm)	Formula	DBE	# of Peaks	Suggested Identity
	-8.301%	666.05954	-	-	-	1	too many options
	-7.909%	96.960595	9	$C_8H_4O_6$	7	1	dihydroxyphthalic acid
	-1.292%	209.06166	-6.5	$C_{14}H_{10}O_2$	10	1	dihydroxyanthracene
	-0.546%	649.55411	0.06	$C_{39}H_{74}N_2O_5$	4	1	unknown, potential dinitro species
500 °C	0.501%	161.02415	-0.12	$C_9H_6O_3$	7	6+	umbelliferone or other hydroxycoumarin*
	0.504%	167.03476	-0.03	$C_8H_8O_4$	5	3+	vanillic acid[†]
	0.597%	161.06082	-0.94	$C_{10}H_{10}O_2$	6	2	methoxycannamaldehyde, methylcinnamic acid, methylhydrocoumarin
	0.618%	277.21747	-1.11	$C_{18}H_{30}O_2$	4	1	octadecatrienoic acid (likely linolenic acid)
	0.641%	163.04002	-0.14	$C_9H_8O_3$	6	8+	caffeic aldehyde,* coumaric acids*

0.686%	191.01981	-1.54 or -8.07	$C_7H_4N_4O_3$ or $C_6H_8O_7$	8 or 3	4-5	unknown; citric acid
0.796%	149.06075	-0.03	$C_9H_{10}O_2$	5	4-5	acetylanisole,* hydrocinnamic acid,* tolylacetic acid
0.814%	165.0556	-0.14	$C_9H_{10}O_3$	5	M	hydroxy-methoxyacetophenone* (apocynin, paeonol, etc), caffeyl alcohol,* veratraldehyde,* phloretic acid*
0.865%	137.06088	0.98	$C_8H_{10}O_2$	4	2	anisyl alcohol, creosol, dimethoxybenzene*
0.891%	195.06638	-0.15	$C_{10}H_{12}O_4$	5	2	acetosyringone*, homoveratic acid*
0.919%	221.06627	0.61	$C_8H_{14}O_7$	2	1-2	dihydroxydimethoxyoxane-2-carboxylic acid*[t]
0.942%	175.0402	0.03	$C_{10}H_8O_3$	7	7	methoxycoumarin,* methylhydroxycoumarin*
1.030%	207.06603	1.5	$C_{11}H_{12}O_4$	6	1	sinapaldehyde
1.621%	181.05061	0.3	$C_9H_{10}O_4$	5	3	homovanillic acid*, dimethoxybenzoic acid,* or syringaldehyde*
1.888%	177.05541	1.16	$C_{10}H_{10}O_3$	6	1	coniferaldehyde[†]
2.371%	137.02443	0.72	$C_7H_6O_3$	5	4-5	salicylic acid* or dihydroxybenzaldehyde*
2.737%	123.04531	-0.81	$C_7H_8O_2$	4	2	guaiacol
3.107%	135.0451	-2.25 - 0.05	$C_8H_8O_2$	5	5	anisaldehyde,*methylbenzoic acid*, acetophenone, or methylsalicylaldehyde*
3.746%	121.02951	-0.59	$C_7H_6O_2$	5	3	benzoic acid[†]and hydroxybenzaldehyde*
5.335%	151.04027	-0.95	$C_8H_8O_3$	5	5	vanillin,[†] and methoxybenzoic acid,* or hydroxyanisaldehyde*
7.514%	109.02957	-3.37	$C_6H_6O_2$	4	1	dihydroybenzene
800 °C	-5.654%	96.961	9	$C_8H_4O_6$	7	1	dihydroxyphthalic acid

-5.654%	96.961	9	$C_8H_4O_6$	7	1	dihydroxyphthalic acid
-1.247%	649.55378	-2.61	$C_{39}H_{74}N_2O_5$	4	W	unknown, potential dinitro species
-0.844%	163.040	-0.14	$C_9H_8O_3$	6	8+	caffeic aldehyde,* coumaric acids*
0.856%	666.060	-	-	-	1	too many options
1.237%	982.99221	-	-	-	W	no formula found

(The leftmost label "800 °C" spans the last five rows.)

At 800 °C, there appears to be very little occurring due to the aging process in the presence of UV light and VOCs. This suggests that there were very few BrC species to begin with and that the presence of anthropogenic VOCs did not significantly impact the chemical makeup of these aerosols. It is also possible that semivolatile species produced during oxidation of these VOCs were lost in significant quantities to the chamber walls rather than the particle phase. The only significant products of oxidation were dihydroxyphthalic acid, which is likely a product of xylene oxidation, and a mixture of caffeic aldehyde, veratraldehyde, and/or hydroxy cinnamic acid isomers, such as coumaric acid. Examining the SSA changes due to aging under either dark or UV illuminated conditions was not possible for the experimental reasons stated in Part I.

Eucalyptus combusted at 500 °C is far more complex. Since many of these compounds have been discussed in more depth earlier, this discussion will only focus on compounds that are distinctly different as a function of age or those not previously mentioned. Another distillation product, benzoic acid and the lignin pyrolysis product hydroxybenzaldehyde isomer (such as salicylaldehyde), likely underwent similar processing. Derivatives of the aromatic lactone coumarin are also known distillation products present in biomass burning (Fleming et al., 2019;Laskin et al., 2015). While not as prevalent as lignin pyrolysis products, a significant number do seem to be consumed due to UV irradiation and/or photochemistry. This includes species that were not distinctly different from

Acacia, as discussed earlier, such as seven isomers of methoxy- and/or methylhydroxycoumarin ($C_{10}H_8O_3$, [M-H]$^-$ = 175.0402), neither of which have been previously observed to the author's knowledge. Others include all six possible isomers of hydroxycoumarin ($C_9H_6O_3$, [M-H]$^-$ = 161.0241), of which umbelliferone has been previously observed (Fleming et al., 2019). There are also multiple indistinct peaks that can correspond to one of several hydroxy methoxyacetophenones (apocynin (acetovanillone), paeonol, etc.), caffeyl alchohol, and/or phloretic acid ($C_9H_{10}O_3$, [M-H]$^-$ = 165.0556), many of which have been extracted from various plants but have not been observed as constituents of BBOA. There are 4 − 5 isomers of acetylanisole or, less likely, hydrocinnamic acid or tolyacetic acid. Octadecatrienoic acid has also been removed, likely by a reaction of one of its double bonds with ozone or OH radical.

In comparison, the compounds being removed from Eucalyptus combusted at 500 °C is dominated by lignin pyrolysis products. Most of these have already been discussed earlier. One isomer of dihydroxybenzene, being either resorcinol or catechol, was a pyrolysis product, and seems sensitive to the aging process, such as oxidation by OH or by bleaching, since it absorbs in the UV at 290 nm (Dewar et al., 1958). Both proper isomers of acetosyringone and/or homoveratic acid ($C_{10}H_{12}O_4$, [M-H]$^-$ = 195.0664), have previously been observed (Simoneit, 2002) and may be present here, though it cannot be definitively stated that these specific isomers are present. Highlighting the smoldering conditions present during combustion at this temperature, there are two isomers of either anisyl alcohol, creosol, or some dimethoxybenzene compound ($C_8H_{10}O_2$, [M-H]$^-$ = 137.0609). These are thought to be precursors of other, more highly oxygenated pyrolysis products of lignin, such as anisic acid, vanillin, or vanillic acid (Simoneit, 2002). These have similar structural motifs as lignin products and they may be regarded as distillation products. There are only a few compounds produced because of aging, such as dihydroxyphthalic acid and a very high MW compound identified as $C_{39}H_{74}N_2O_5$, which may be a dinitro compound. As mentioned previously, dihydroxyphthalic acid could be a product of xylene oxidation, but no enhancement of OA was observed in Part I of this work when comparing aging under lights to lights with anthropogenic VOCs. While there is a slight increase in SSA due to the presence of VOCs, it would be difficult to point to this one compound as the sole difference. Lastly, there is dihydroxyanthracene, which is likely a secondary oxidation product of the PAH anthracene. This compound should absorb in the UV and visible, given the absorption of 1-hydroxyanthracene from the UV to ~410 nm (Tichy, 1967).

Figure 3 and Table 3 show the results from aged Acacia. As before, very little has changed for the Acacia fuel combusted at 800 °C. Combustion at 500 °C is far richer and consists almost exclusively of compounds that have been consumed (i.e. have a positive scaled intensity difference). The species undergoing the most significant loss is either resorcinol or catechol. The second most consumed compound, which was not seen previously, is one isomer that is either nitroanisole, nitrocresol, or possibly nitrobenzyl alcohol ($C_7H_7NO_3$, [M-H]$^-$ = 152.0356). Given that it is being consumed over the course of the aging process, it was likely formed as a primary species during combustion of lignin in the presence of NO$_x$. While this species has not been observed previously, related compounds such as nitrocatechol and methyl nitrocatechol have (Fleming et al., 2019). Given that these compounds have absorption spectra that extend to ~500 nm, they were likely decomposed due to photo-bleaching, though heterogeneous reactions with OH cannot be ruled out. It is likely that the products of this photo-bleaching and/or oxidation were numerous, and each had an abundance that was too small to be captured in this analysis.
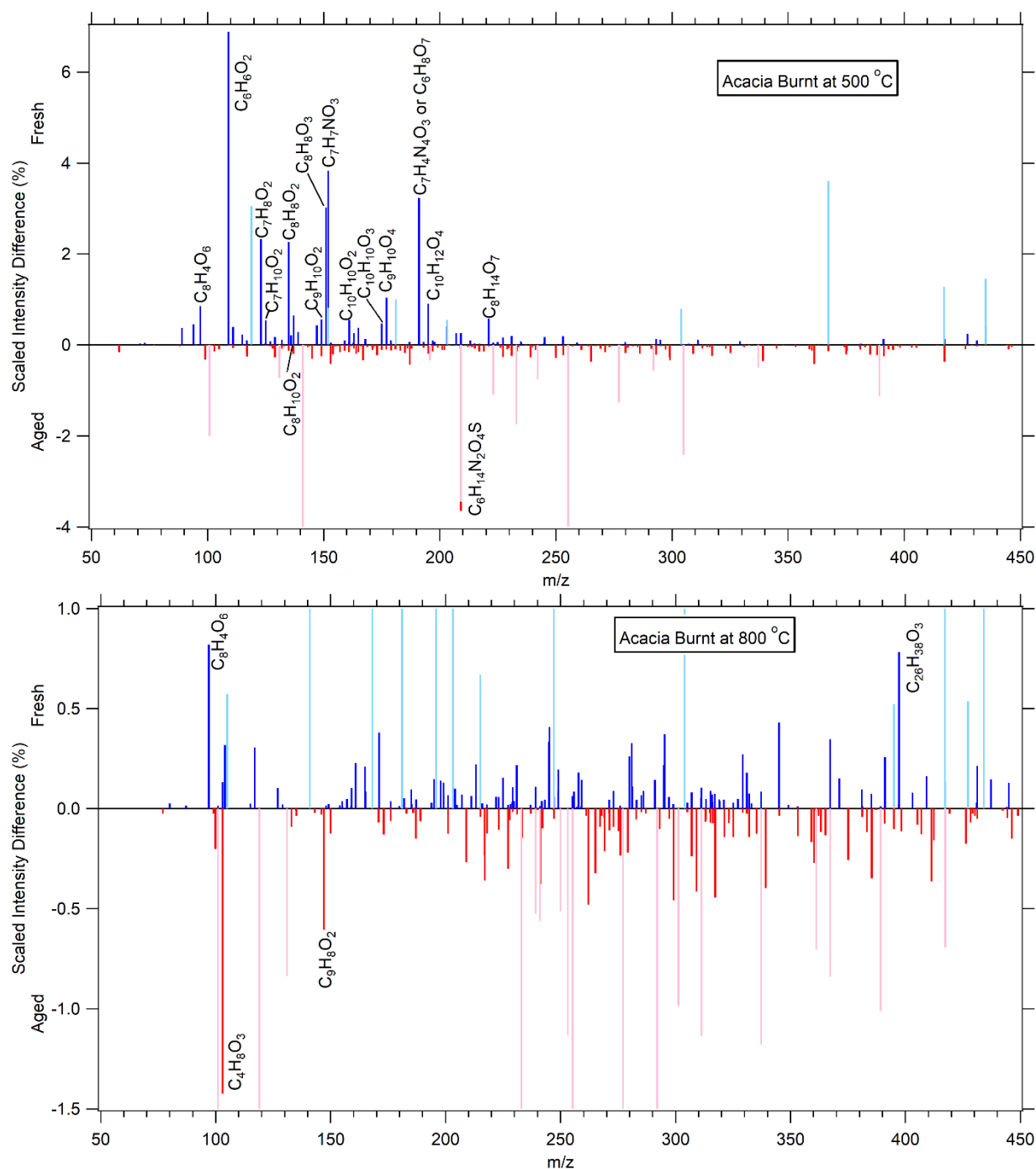
Figure 3. A comparison of Acacia-derived BB aerosol combusted at 500 °C (above) and 800 °C (below) under fresh and photochemically aged conditions with VOCs in terms of the scaled intensity difference (see section 2.5). Species more present in the fresh sample have positive values (blue lines), while those more present in the aged sample have negative values (red lines) (i.e. species with positive values have been removed because of the aging process, while those with negative values are either produced or represent a greater relative contribution to the aged sample). Peaks in lighter colors were present in the XCMS analysis but were found to be present in significant amounts in the blank sample. Many of these peaks that were found in the blank extend past the scale of the plot but were cut off to focus on species associated with BB aerosol. Compounds having an absolute difference

greater than 0.5% have been labeled. There is very little change for combustion at 800 °C, while 500 °C shows a net removal of many compounds.

520 **Table 3.** Molecular difference between fresh and aged Acacia revealed by negative mode MS analysis. See the caption of Table 1 for more details.

	Scaled Intensity Difference	m/z	Mass Difference (ppm)	Formula	DBE	# of Peaks	Suggested Identity
	-3.652%	209.06134	-7.41	$C_6H_{14}N_2O_4S$	1	1	unknown
	0.521%	736.13744	-0.53	$C_{44}H_{23}N_3O_9$	35	1	unknown
	0.539%	125.06093	-3.35	$C_7H_{10}O_2$	3	1	cyclohexene carboxylic or heptadienoic acid
	0.549%	161.0607	-0.94	$C_{10}H_{10}O_2$	6	2	methoxycannamaldehyde, methylcinnamic acid, or methylhydrocoumarin
	0.557%	149.06125	-0.03	$C_9H_{10}O_2$	5	4-5	acetylanisole,* hydrocinnamic acid,* tolylacetic acid
	0.575%	221.06668	0.61	$C_8H_{14}O_7$	2	1-2	dihydroxydimethoxyoxane-2-carboxylic acid*[t]
	0.641%	137.06111	-4.02	$C_8H_{10}O_2$	4	2	anisyl alcohol, creosol, dimethoxybenzene*
500 °C	0.763%	181.05141	0.3	$C_9H_{10}O_4$	5	3	homovanillic acid*, dimethoxybenzoic acid*, or syringaldehyde*
	0.852%	96.960701	9	$C_8H_4O_6$	7	1	dihydroxyphthalic acid*
	0.901%	195.06627	-0.15	$C_{10}H_{12}O_4$	5	2	acetosyringone* or homoveratic acid*
	1.038%	177.05542	1.16	$C_{10}H_{10}O_3$	6	1	coniferaldehyde[†]
	1.306%	666.05965	-	-	-	1	too many options
	2.259%	135.04538	-2.25	$C_8H_8O_2$	5	2	anisaldehyde*, acetophenone, methylsalicylaldehyde*, or methylbenzoic acid*
	2.327%	123.04542	-0.81	$C_7H_8O_2$	4	2	guaiacol
	3.016%	151.04032	-0.95	$C_8H_8O_3$	5	5	vanillin[†] and methoxybenzoic acid* or hydroxyanisaldehyde*
	3.233%	191.01941	-1.54 or -8.07	$C_7H_4N_4O_3$ or $C_6H_8O_7$	8 or 3	4-5	unknown; citric acid
	3.821%	152.03561	4.2	$C_7H_7NO_3$	5	1	nitroanisole*, nitrocresol*, or nitrobenzyl alcohol*
	6.886%	109.02984	-3.37	$C_6H_6O_2$	4	1	dihydroxybenzene
	-1.423%	103.02421	-4.1	$C_4H_8O_3$	1		hydroxybutanoic acid, methoxypropanoic acid
	-0.604%	147.04672	-8.72	$C_9H_8O_2$	6	W	cinnamic acid
800 °C	0.783%	397.26686	-0.08	$C_{26}H_{38}O_3$	8	1	uncertain
	0.819%	96.960676	nc	$C_8H_4O_6$	7		dihydroxyphthalic acid*
	3.467%	666.05938	-	-	-	1	too many options

Given that one isomer of dihydroxybenzene undergoes the greatest change upon aging, it is worth examining potential chemical transformations further. While resorcinol is a potential identity, catechol has gained significant
525 attention due to its involvement as a precursor to a strong chromatophore. Catechol can react in the atmosphere to

form 4-nitrocatechol ($C_6H_5NO_4$) through photochemical and dark processes in the gas and aqueous phase through a number of mechanisms (Finewax et al., 2018;Kroflič et al., 2018;Vidovic et al., 2018;Wang et al., 2019). Given the dryness of our chamber and the presence of UV irradiation, this mechanism would proceed through hydrogen abstraction from the hydroxyl by OH to produce a *β*-hydroxyphenoxy/*o*-semiquinone radical, followed by radical-radical combination with NO_2 (Finewax et al., 2018). Both fresh and aged spectra for 500 °C combustion were examined for nitrocatechol ((M-H)⁻ at *m/z* 154.0140), and it was found at a rt of 8.39-8.55 min in both fresh and aged Acacia and Eucalyptus. There was a greater abundance in the aged samples, despite the mass loading on the filter being lower, suggesting a small amount from primary emission, but mainly a product of secondary formation for both fuels.

Smoldering-dominated burns were also examined for other nitroaromatic species. Methyl nitrocatechol, which has been identified as a tracer for BB SOA (Iinuma et al., 2010), and nitroguaiacol both have the formula of $C_7H_7NO_4$ and would have an (M-H)⁻ peak at *m/z* 168.0297. Fresh Eucalyptus had two peaks at retention times of 7.71 and 9.31 min, while Acacia only had a single peak at 9.30 min. Using the exact mass and the rt, this peak at 9.30 min has been identified as 4-methyl-5-nitrocatechol, so the 7.71 min peak must be another isomer produced in fresh combustion. Since this compound must be more polar to elute earlier, it may be the 4-, 5-, or 6- isomer of nitroguaiacol (Bluvshtein et al., 2017;Kitanovski et al., 2012a). Upon aging, both fuels had two peaks with smaller absolute abundances at 9.5-9.6 and 10.0 min. The 9.5-9.6 min peak is likely a slightly shifted 4-methyl-5-nitrocatechol, suggesting both a primary and secondary source of the compound. Based on previous observations of BB aerosol, the 10.0 min peak is either 3-methyl-6-nitrocatechol, 3-methyl-5-nitrocatechol, or 4-nitroguaiacol, whose sensitivity can be altered for it to elute either before or after 4-methyl-5-nitrocatechol (Bluvshtein et al., 2017;Kitanovski et al., 2012a). Nitrophenol ($C_6H_5NO_3$, (M-H)⁻ at *m/z* 138.0191) and nitrocresol ($C_7H_7NO_3$, (M-H)⁻ at *m/z* 152.0348) are secondary products of phenol and cresol, respectively. Neither was observed for fresh Eucalyptus and Acacia. There are at least two forms of methyl nitrophenol in aged Acacia (rt of 10.15 and 10.46 min) and only one (rt 10.47 min) in aged Eucalyptus. Several isomers have been observed in BB aerosol in previous studies, including 3-methyl-4-, 2-methyl-4-, and 2-methyl-6-nitrophenol (Bluvshtein et al., 2017;Kitanovski et al., 2012a). The peak with a rt of ~10.46 was likely a slightly shifted 4-nitro-*o*-cresol, which was a standard compound run during the analysis of fresh BB aerosol and had a rt of 10.30 min. One nitrophenol peak at 9.37 min was observed in aged Acacia, while aged Eucalyptus did not exhibit any nitrophenol. This had a different rt than the 2-nitrophenol standard, which had a rt of 10.05 min, suggesting it was some other isomer. Nitronaphthol ($C_{10}H_7NO_3$, (M-H)⁻ at *m/z* 188.0348) was not observed in any spectra.

3.3 UV/Visible analysis of fresh versus aged emissions

In the companion paper Part I of this work, we found, as expected, that the SSA of Eucalyptus and Acacia increased after photochemical aging for 12 hours. For instance, 300 nm particles from Eucalyptus combustion at 500 °C went from an SSA of ~0.67 to almost unity, with only a slight increase due to the presence of VOCs. Acacia increased from ~0.78 to 0.90 − 0.95 after aging, with no apparent change due to VOCs. Since these measurements

were size selected and this increase was observed for all selected sizes, the SSA change could not be attributed to bulk changes within the aerosol, such as particle coagulation. Such an increase in SSA resulting from morphology changes alone seems implausible, so the particles themselves must become less absorbing with age. This would be because of a loss of absorbing compounds in the $500 - 570$ nm region, the formation of non-absorbing SOA that alters the real portion of the refractive index, or both. In this section, we discuss how these changes in molecular composition affect the integrated UV/Visible absorption spectrum and how these relate to the SSA of *in situ* aerosol studied in Part I of this work.

Schuster et al. (2016) found a strong correlation between AAE and the spectral variability of the refractive index of BB aerosols ($\kappa_{440}/\kappa_{700}$), with a linear correlation of $R \geq 0.939$. BrC is spectrally dependent, with a greater imaginary part of the refractive index at 440 nm (κ_{440}) than the red region of the spectrum (κ_{700}). Figure 4a shows that UV absorption dominates for fresh BB aerosol combusted at 500 °C, with a negative slope in the $500 - 570$ nm range, indicating the presence of BrC can alter the absorption in the mid-visible region. This is corroborated by large *in situ* values of AAE ($\gtrsim 1.5$) for these samples as seen in Table 4. BC, on the other hand, is spectrally independent ($\kappa_{440}/\kappa_{700} \approx 1$), and has AAE values between 0.5 and 1.5, as seen for the CRDS measurements in Table 4. This is corroborated by Figure 4b and Table 4, which shows that absorption in this range has a slope of zero within the measurement uncertainty for fresh BB aerosol combusted at 800 °C; all within 1σ, except Olive, which is within 2σ. The NAAE values based on UV/Vis measurements of molecular chromophores do not conform to this trend at 800 °C; however, this is likely because they have been normalized. Lastly, AAE can reach negative values when absorption is higher at longer wavelengths (near 700 nm). Table 4 suggests that as BB aerosols are aged in the presence of UV light, they become more absorbing at longer wavelengths (except for Eucalyptus combusted at 800 °C, as measured by CRDS). This could be caused by formation of species that absorb at such long wavelengths, such as nitro-aromatic species and functionalized PAHs, or lensing effects from hygroscopic growth (Fleming et al., 2019;Liu et al., 2017).
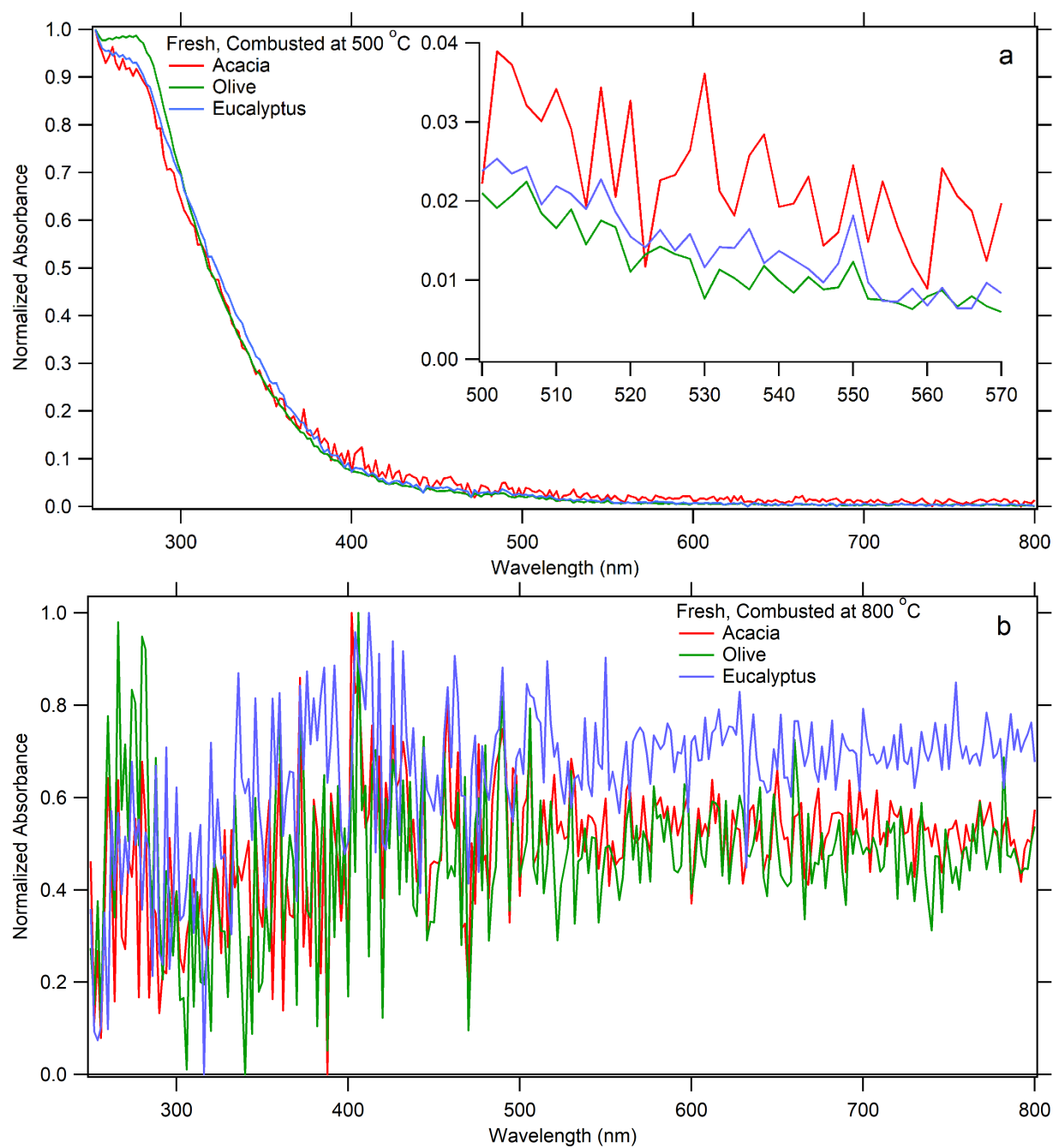
Figure 4. The normalized UV/Visible spectra for fresh Acacia-, Olive-, and Eucalyptus-derived BB aerosol combusted at 500 °C (a) and 800 °C (b). These spectra have been averaged over the entire chromatogram, background subtracted, and normalized, as described in section 2.6.

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Table 4. The slope of absorption vs. wavelength and the Ångstrom absorption exponent for each sample, as measured by the UV/Vis spectrum during chemical analysis (normalized), and during CRDS/Nephelometer analysis (*in situ*). AAE was determined for the wavelength range 500 nm $\leq \lambda \leq$ 570 nm.

			AAE (normalized for UV/Vis)			Slope ($\times10^{-5}$ nm^{-1} for UV/Vis, $\times10^{-16}$ m$^2\bullet$particle$^{-1}\bullet$nm^{-1} for CRDS/Neph)		
			Acacia	Eucalyptus	Olive	Acacia	Eucalyptus	Olive
UV/Vis	500 °C	Fresh	5.55±1.21	9.53±0.72	9.10±0.64	-23.7±4.8	-24.6±1.9	-20.6±4.3
		Aged	-0.12±1.25	-1.87±1.25	-	-2.15±9.52	10.3±8.8	-
	800 °C	Fresh	0.40±0.60	0.91±0.51	0.22±0.84	-48.5±60.3	-120±69	-34.5±76.6
		Aged	-0.13±0.49	-1.88±1.01	-	4.33±70.5	145±91	-
CRDS/Neph (300 nm)	500 °C	Fresh	2.52±0.27	1.71±0.18	1.32±0.33	-1.98±0.21	-2.02±0.21	-1.63±0.412
		Aged	-3.25±0.98	-4.20±3.12	-	1.29±0.4	0.46±0.42	-
	800 °C	Fresh	0.57±0.25	0.66±0.22	0.25±0.15	-0.57±0.25	-1.64±0.53	-0.620±0.367
		Aged	-5.54±0.53	0.77±0.14	-	1.81±0.16	-2.98±0.55	-

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For both Eucalyptus and Acacia combusted at 500 °C, the greatest loss of UV absorptivity, as seen in Figure 5a, happens between 260 − 350 nm, peaking at ~290 nm. Many of the lignin pyrolysis products are consumed upon aging, due to either photo-bleaching or oxidation. Previous work has shown a reduction in BrC absorptivity due to photochemical aging (Sumlin et al., 2017) and OH oxidation (Dasari et al., 2019;Schnitzler and Abbatt, 2018). Most

605   of these compounds absorb in the UV between 200 and 350 nm, with the notable exception of coniferaldehyde and sinapaldehyde, which both peak near 340 − 345 nm and absorb past 400 nm (Fleming et al., 2019).
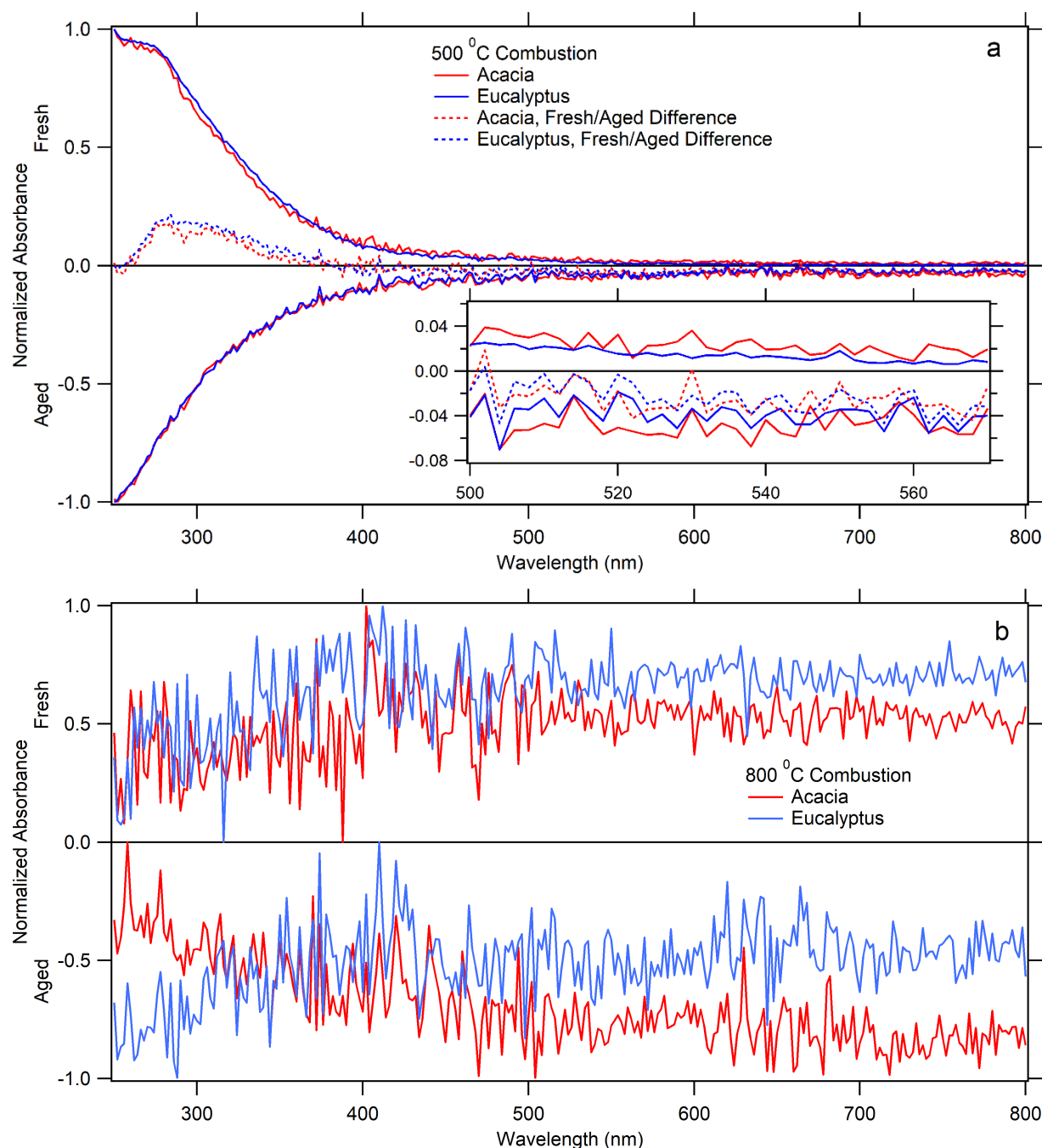
Figure 5. The UV/Visible spectrum for each fuel combusted at 500 °C (a) and 800 °C (b), showing the result from fresh combustion with positive values and photochemically aged with negative values. These spectra have been averaged over the entire chromatogram, background subtracted, and normalized, as described in section 2.6. The differences between them are highlighted with the dashed line and a blowup of the 500 − 570 nm region is presented in the inset.

3.4 Chromatographic analysis of potential chromophores in the visible

In addition to examining the integrated spectrum, we can also focus on potential chromophores that absorb in the 500 – 570 nm region of the spectrum. Considering that flaming-dominated combustion shows little in the way

of absorbing species and the abundance of these species decreases upon aging, only fresh emissions from combustion at 500 °C will be examined. Figure 6 shows the extracted wavelength chromatogram for the 500 – 510 nm region, where absorption would be most intense over this range, for Eucalyptus and Acacia combusted at 500 °C. This chromatogram has been background subtracted by the filter blank and, given that the amount of aerosol deposited on these filters was not measured, no scaling was applied. Remarkably, only a few peaks dominate the chromatogram, though there are likely several low abundance and indistinct peaks 11 – 13 minutes. The UV/visible absorption spectra for each designated peak in Figure 6 are given in Figure S4. As can be seen in Figure 6, several of these species have similar retention times. A comparison was made between these peaks, with respect to elution time. Peak A and H, despite coming from different fuels, show similar retention times and UV spectra, as can be seen in Figure S5. The same can be said for peaks B, C, I, and J, which have well separated peaks at ~340 and ~485 nm with a gradual decrease to 600 nm. Peaks D, E, and K also have the same general appearance, and may be related to B, C, I and J, but are somewhat broader and peak at slightly shorter wavelengths (332 and ~480 nm).
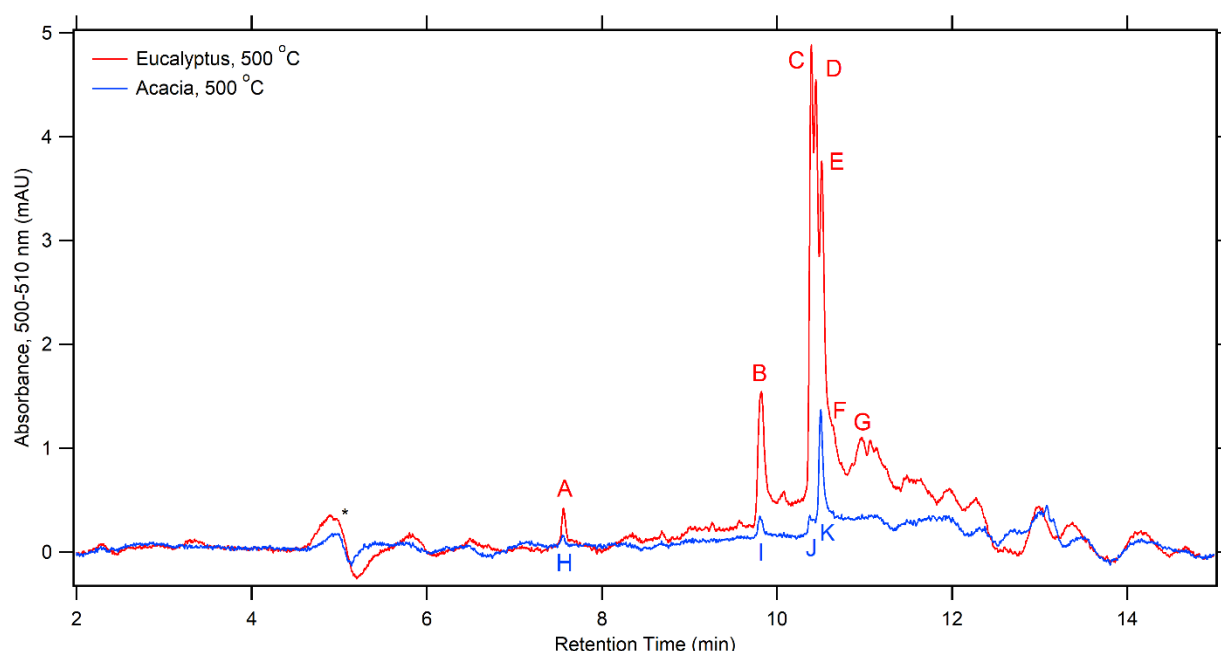
Figure 6. The extracted wavelength chromatogram for the 500 – 510 nm region for each fuel freshly combusted at 500 °C. The peak with the asterisk is an artifact from background subtraction.

To determine which species produced the absorption features shown in Figure 6, mass spectra at the absorption maxima were examined and assessed based on the peak height in the mass spectra, how well the retention times of extracted ion chromatograms matched the extracted visible wavelength chromatogram, and if a compound can be reasonably thought to absorb in the 500-510 nm region. Assignments using this method are tenuous, given that species detected by the DAD may not be seen by negative-mode ESI-MS. Additionally, many of the species described below have additional peaks outside those described, suggesting that not all isomers corresponding to a given formula absorb in this region. The extracted ion and visible chromatograms (shown in Figure 6) are in Figure S6a for

Eucalyptus. Peak A is either $C_9H_{10}O_3$ (m/z 165.0559), tentatively assigned earlier, and/or $C_9H_{12}O_4$ (m/z 183.0658), which could be an isomer of trimethoxyphenol or dimethoxy methyl catechol. Several isomers of $C_9H_{10}O_3$ are known to only absorb below 240 nm, though veratraldehyde and caffeyl alcohol UV/visible spectra have not been measured, nor has trimethoxyphenol or dimethoxy methyl catechol for $C_9H_{12}O_4$. A $C_9H_{10}O_3$ isomer may also contribute to peak B and, while $C_8H_{10}O_2$ (m/z 137.0612) is more abundant at peak B and previously identified as several isomers in this work, none would absorb in this region. Likewise, $C_{10}H_{10}O_2$ (m/z 161.0604) elutes with peak B, which may be methoxycinnamaldehyde, methylcinnamic acid, or methylhydrocoumarin. While the spectra of those compounds have not been previously measured, it is unlikely that they would absorb in this region either. One compound that could weakly absorb in the visible and elutes at peak B is hydroxycoumarin ($C_9H_6O_3$, m/z 161.0244). One such compound, umbelliferone, absorbs to ~420 nm (Abu-Eittah and El-Tawil, 1985), though a number of isomers are present which may have somewhat different spectra.

Several species show peaks in the mass chromatogram that correspond to peaks B-F, including $C_{10}H_8O_3$ (m/z 175.0405), $C_{11}H_8O_3$ (m/z 187.0401), and $C_{11}H_{10}O_3$ (m/z 189.0565). The species corresponding to $C_{10}H_8O_3$ could be methoxycoumarin or methylhydroxycoumarin, though these species mainly absorb below ~450 nm (Abu-Eittah and El-Tawil, 1985) and may only absorb very weakly above 500 nm if at all. Naphthalenetriol is a much more likely candidate for $C_{10}H_8O_3$, given its extended and functionalized aromatic system, though a spectrum for any of its isomers has not been measured. The less functionalized 1,4-naphthalenediol absorbs to ~680 nm (Linstrom and Mallard, 2020). The structures attributable to $C_{11}H_8O_3$ could be hydroxynaphthoic acid, acetylcoumarin, or methylcoumarinaldehyde, but none of these compounds absorb above 410 nm (Donovalová et al., 2012). Still, several of these compounds are yellow solids in their pure forms according to their SDS sheets, suggesting they have some interaction with visible light. Similarly, an indistinct number of peaks associated with $C_{11}H_{10}O_3$ are found within these retention times, which may be isomers of methylmethoxycoumarin or dimethylhydroxycoumarin; none of which have previously measured spectra. Peaks in the vicinity of if peaks F and G correspond to $C_{12}H_{10}O_3$ (m/z 201.0554) methoxynaphthoic acid or methoxyhydroxynaphthaldehyde. No UV/visible spectra are available for these compound and, while some spectra are available for related compounds such as 1- and 2-naphthoic acid (Linstrom and Mallard, 2020), these spectra seem truncated at ~350 nm. More telling is the SDS descriptions for these substituted naphthaldehydes and naphthoic acids, which are light yellow, yellow, and light brown. This suggest that their absorption extends into the visible and possibly into the 500 nm region.

One set of isomers in the vicinity of peaks C through F correspond to the formula of $C_{11}H_6O_3$ (m/z 185.0256). Given the high degree of unsaturation (DBE = 9), these can only be furanocoumarins, which have a fused furan and benzene ring. Two common isomers of these species are psoralen and angelicin. While these species do not absorb in this region of this visible, they are strong absorbers below 370 nm (Rutan et al., 2018) and have not been previously identified as BrC species in BB aerosol. The peaks for nitroguaiacol/methylnitrocatechol, discussed earlier in this work, do not align well with peaks in Figure 6.

Fresh Acacia combusted at 500 °C underwent a similar analysis, as shown in Figure S6b. All the previously mentioned extracted ion chromatograms appear very similar for Acacia, with the main difference being that the ion

abundance for Acacia was smaller by a factor or 2-4. Most ions had similar temporal profiles as Eucalyptus with the exception of $C_{11}H_{10}O_3$ (*m/z* 189.0565), which only matched peak K and onwards. Given this, it is very likely that peaks with similar DAD retention times are produced by the same species, and peak A corresponds with peak H, B with I, C with J, and E with K.

4 Conclusions

When both fuels in this work were efficiently combusted at 800 °C, the chemical characterization of fresh BB aerosol revealed very little apparent difference between the two species – Eucalyptus and Acacia. The higher SSA observed for Eucalyptus was not attributed to it containing more low abundance species, but some set of compounds that were not being captured in this analysis. This could be due to a prevalence of low abundance species, semi-volatile compounds lost during sample processing (Chen et al., 2017), or would be insensitive to ionization in the negative ion mode of our UPLC/DAD-ESI-HR-QTOFMS technique. A compound such as eucalyptol, which does not absorb in the UV or visible, could significantly alter the real portion of the refractive index of aerosols, but would not be apparent using negative ion mode ESI-MS, given its lack of basic functionality.

Comparing these species combusted at 500 °C revealed many species, differentiated by their mass and chromatographic retention time (i.e. different isomers possessing the same formula). This work focused on compounds that were significantly different between fuel types, which had scaled intensity differences greater than 0.5%. Both samples had a variety of compounds in common, such as lignin pyrolysis products, distillation products, and cellulose breakdown products. Only one isomer of dihydroxybenzene was observed for both fuels, which is either catechol (Simoneit, 2002) or resorcinol (Yokelson et al., 2013). Several lignin pyrolysis products and distillation products are more prevalent in Eucalyptus than Acacia, while pyrolysis products of cellulose and at least one nitro-aromatic species (nitroanisole, nitrocresol, or nitrobenzyl alcohol) were more prevalent in Acacia. When both fuels and Olive were examined for their sinapaldehyde and coniferaldehyde content, it was found that coniferaldehyde was more abundant by a factor of 1.6 – 3.4, depending on the species. Given that all three are angiosperms, this observation is counter to observations made by Fleming et al. (2019), though it is worth noting that their results were in terms of absorption intensity instead of ion intensity. Given that chromophores, such as lignin pyrolysis and distillation products are more prevalent in Eucalyptus than Acacia, which has a higher abundance of non-chromophores derived from sugars and cellulose, one would assume that Eucalyptus would be more absorbing in the visible (i.e. have a lower SSA) than Acacia. Despite this analysis not capturing absolute amounts of OA, Acacia was found to have an SSA that is higher than Eucalyptus by 0.1 to 0.2, which is consistent with these observations. This suggests that Acacia has either larger absolute amounts of non-chromophore compounds or Eucalyptus has a greater quantity of chromophores whose absorptive properties extend to the 500 – 570 nm region of the visible spectrum.

In examining the effects of aging in the presence of UV light and anthropogenic VOCs, it is clear from Figures 2 (Eucalyptus) and 3 (Acacia) that the amount of BrC decreases as the sample ages for combustion at 500 °C, and very few changes are occurring for particles produced from combustion at 800 °C. For both Eucalyptus and Acacia, an isomer of dihydroxybenzene, such as resorcinol or catechol, was removed to the highest degree from the

fresh BB aerosol. Both of these species have relatively broad UV absorption spectra, extending from the vacuum UV to ~292 nm, with a peak at 275 nm and another at shorter wavelengths (Dewar et al., 1958). This is where most of the reduction of UV absorption is occurring. While this could suggest that these two compounds are responsible for a significant portion of the bleaching effect, there are a great number of compounds that absorb in a similar wavelength range, so that could not be conclusively stated. The removal of dihydroxybenzene was investigated by examining spectra for a known reaction product, dihydroxynitrobenzene (e.g. nitrocatechol). Results for both fuels suggest a small amount was produced by primary emissions, but it was mainly a product of secondary formation. The second most removed compound for Eucalyptus was the distillation product benzoic acid and the lignin pyrolysis product hydroxybenzaldehyde. Several Eucalyptus distillation products are also sensitive to aging that have not been previously observed, such as seven isomers of methoxy- and/or methylhydroxycoumarin and all 6 isomers of hydroxycoumarin, of which only one has been previously observed. The second most consumed compound for Acacia was one isomer that is likely nitroanisole, nitrobenzyl alcohol, or nitrocresol, which was not seen previously and is a primary product of in-flame NO_x chemistry. Generally, both fuels were dominated by loss of chromatophoric lignin pyrolysis and distillation products. Focusing on nitroaromatic species, one isomer of nitrophenol was produced during the aging of Acacia combusted at 500 °C, while none was produced for Eucalyptus. Despite being thought of as a tracer for BB SOA (Iinuma et al., 2010), there is evidence for both primary emission and secondary production of 4-methyl-5-nitrocatechol.

Not surprisingly, the associated absorbance from these chromophores, mostly from 200 – 350 nm, also attenuates with respect to age, as seen in Figure 5. This may be caused in part by the photo-bleaching effect created by irradiation of UV light for 12 hours, heterogeneous OH oxidation, and SOA formation of non-chromophores. As stated in Part I of this companion paper, ignition temperature is the leading factor for differences in the SSA of the resulting BB aerosol, which is supported by the associated MCE values at each temperature. At lower temperatures, combustion is less efficient and releases more emissions in the form of the lignin pyrolysis and distillation products, which are primarily responsible for the change in absorption seen between fresh and aged samples, as they absorb strongly in the UV. Some larger compounds were discovered, but only one functionalized PAH was successfully identified. The identification of other functionalized PAHs would be beneficial in determining any changes to SSA because of photochemical aging with anthropogenic VOCs present, as these compounds exhibit wide absorption spectra extending into the visible.

An attempt was also made to identify which species produce absorption at wavelengths longer than 500 nm. An extracted wavelength chromatogram from 500 – 510 nm shows only a few peaks dominate the absorption from fresh emissions of combustion at 500 °C. Both Acacia and Eucalyptus have common absorbing species, as shown by peaks with common retention times, similar UV/Visible spectra, and extracted ion chromatograms. By comparing the retention times of extracted ion and wavelength chromatograms, several species were tentatively suggested for contributing to absorption in this wavelength range. This includes functionalized courmarin species (hydroxy-, methoxy- methylhydroxy-, methylmethoxy-, etc.), naphthalenetriol, and methoxyhydroxynaphthaldehyde or methoxynaphthoic acid. UV spectra available from the literature for several of these species and/or related compounds seem truncated and are very likely to extend into this visible region of the spectrum, which highlights the need for

broadband UV/Visible spectral measurements of these compounds. While not important for absorption in this visible region, a novel set of compounds were found that could be strong BrC species – furanocoumarins.

As shown in companion paper Part 1, there are changes in the SSA that occur in response to dark aging. Future work will examine those processes from a molecular standpoint, and how they are different from photochemical aging. Given that there was no SOA enhancement or significant difference in SSA observed for the addition of anthropogenic VOCs, it is unlikely that even these elevated concentrations would alter the molecular constituents either. However, this should be confirmed with at least one example.

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Author contribution: Damon Smith and Tianqu Cui conducted the experiments, Marc Fiddler, Tianqu Cui, and Damon Smith analyzed the data. Marc Fiddler, Jason Surratt and Solomon Bililign designed the experiments and contributed to writing and editing. Rudra Pokhrel contributed to the data analysis and interpretation.

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Competing interests: The authors declare that they have no conflict of interest.

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