Supplement of 

Characterization of water-soluble brown carbon chromophores from wildfire plumes in the western USA using size-exclusion chromatography

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**Figure S11.** Absorption contribution of high (>500 Da), low (<500 Da), and unidentified molecular weight species of aqueous samples collected during (a) 24 Aug 2019, (b) 25 Aug 2019 L2, (c) 28 Aug 2019 L1, (d) ) 28 Aug 2019 L2, and (e) ) 28 Aug 2019 L3.

**Figure S12.** Relationship between BrC absorption as measured by SEC–UV offline and by PILS–LWCC online during the Southern Oxidant and Aerosol Study (SOAS) in the Talladega National Forest in Alabama, USA. Offline samples were collected on a filter assembly of a broadband cavity enhanced spectrophotometer and online particle absorption was measured using a PILS-LWCC (Di Lorenzo et al., 2017).

**Figure S13.** Absorption cross section of 4-nitrocatechol measured by injection onto the LWCC of the BrC-PILS and onto the diode array detector of the SEC-UV set-up compared to Hinrichs et al. 2016.

**Figure S14.** Size separation of a fresh SRHA solution which was re-run 20 months later.

**Table S1:** Summary of flights completed by the NOAA “Chemistry” Twin Otter aircraft during FIREX-AQ 2019.
**Figure S1.** A schematic showing the liquid flow within the BrC-PILS. MFM is liquid mass flow meter and LWCC is liquid waveguide capillary cell.

**Figure S2.** Selected map of second flight on 21 Aug 2019 of the Castle fire. The circular markers are coloured by absorption at 250 nm measured by SEC-UV and represent the average location where liquid flow was diverted into a single polypropylene sample tube.

**Text S1:** Mobile phase impact on SEC-UV elution profile
Analysis by SEC-UV allows us to examine the molecular size of BrC chromophores as a function of plume age. As the mobile phase carries the sample through the porous stationary phase of a SEC column, high MW molecules that exceed the size of the pores will flow directly through the column, passing the pores. This will result in an early elution time within the void volume. If absorbing species are small enough to penetrate the pores, they will spend more time in the column and elute later. All molecules that are smaller than the limit to fully penetrate the pores of the stationary phase will elute in the exclusion volume. Figure S3 displays single-wave size-exclusion chromatogram at 250 nm for aqueous samples collected on 21 Aug 2019. The absorption density of the aqueous samples listed in Table S1 had consistent size-resolved features with varying magnitude in absorption. The first peak between 8.5 to 9 minutes is characteristic of relatively higher MW compounds than the second peak between 9.5 to 11.

**Figure S3.** Example of absorption density plot at (a) 250 nm and (b) 300 nm of FIREX-AQ aqueous samples collected during the second flight 21 Aug 2019, of the Castle fire. The mobile phase is equal parts acetonitrile and 25 mM ammonium acetate solution at a flow rate of 1 mL min\(^{-1}\).
Figure S4 displays single-wavelength size-exclusion chromatograms of an aqueous separation on the SEC-UV with varying mobile phase compositions. With equal parts methanol and 25 mM ammonium acetate solution, the low MW fractions eluted after exclusion volume of the SEC column (<250 Da). With the addition of acetonitrile, it appears that the apparent higher MW fraction then co-eluted with the low MW fraction (represented by a single peak in the orange trace; Figure S4). Similar to the findings of Lyu et al. (2021) using equal parts acetonitrile and 25 mM ammonium acetate solution appears to be effective in suppressing interactions between the stationary phase of the SEC column and chromophores, as a narrow elution profile of a low MW fraction was observed.

Figure S4. Single-wavelength chromatogram at 250 nm of an aqueous sample run with equal parts 25 mM ammonium acetate solution and methanol (black) and equal parts 25 mM ammonium acetate solution and acetonitrile (orange).

The default mobile phase used was equal parts acetonitrile and DIW with 25 mM ammonium acetate. When ammonium acetate is dissolved in water, sub-stoichiometric acidification converts acetate to acetic acid producing conditions that can stabilize pH at 4.75 (Konermann, 2017). The alkalization of ammonium acetate solution generates NH₃ via the deprotonation of NH₄⁺, creating buffering capacity around the pka of ammonium (9.25). Therefore, ammonium acetate dissolved in water has an inherent buffering capacity in acidic (pH at 4.75±1) and basic ranges (9.25±1) (Konermann, 2017). The addition of acetonitrile to ammonium acetate dissolved in water reduces the buffer capacity and shifts the buffering ranges of ammonium acetate dissolved in water to approximately pH 5.5±1 and pH 9 ±1 (Subirats et al., 2009). The purpose of the addition of the ammonium acetate to the mobile phase was to minimize electrostatic interactions between the compounds and the stationary phase of the column. This has proven effective in previous SEC-UV analyses of biomass burning derived samples investigating MW properties of fresh and aged BrC (Di Lorenzo et al., 2017; Wong et al., 2017; Lyu et al., 2021). If the electrostatic interactions are negligible, SEC separation is based on hydrodynamic
volume, which is a function of MW and the density of the compounds (Pelekani et al., 1999). In Figure S6, there is a red shift when mobile phase conditions have a pH greater than the pKa of the single compound. However, Figure S7 shows that wavelength dependent absorption of the SRFA looks similar under all mobile phase conditions. This indicates that we do not anticipate pH impacting wavelength dependent absorption in the SEC-UV analysis.

**Figure S5.** Absorption density as a function of molecular weight of aqueous samples collected during FIREX-AQ water samples and Suwannee River humic acid. The mobile phase is equal parts acetonitrile and 25 mM ammonium acetate solution at a flow rate of 1 mL min$^{-1}$. 
Figure S6. Absorption as a function of wavelength measured in various mobile phases of (a) 4-nitrocatechol (b) 4-hydroxy-3-methoxycinnamaldehyde (c) vanillin (d) 7-hydroxycoumarin and (e) a mixture of the four compounds by a diode array detector. The default mobile phase consists of 1:1 mixture of acetonitrile and DIW with 25 mM ammonium acetate. The purple and black traces represent pH controlled mobile phases, where the pH of the ammonium acetate solution was adjusted to pH 5 and pH 9 prior to the addition of the acetonitrile.
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Figure S8. Absorption as a function of wavelength of SRFA measured in a 25 mM ammonium acetate solution and in a mobile phase controlled to pH 5.
**Figure S9.** Average and standard deviation of CO measured during each flight. Blue markers denote the coefficient of variation, which represents the standard deviation as a percent of the mean.

**Text S2: Conversion of SEC-UV signal to ambient absorption units of Mm$^{-1}$**

To compare absorption of the water soluble BrC measured online by the BrC-PILS, the absorption data measured by SEC-UV was converted to inverse megameters, Mm$^{-1}$. The absorption of different MW fractions at a selected wavelength is determined by integrating the peak area. To apply the Beer-Lambert Law to the absorption profiles by the SEC-UV method, the dilution of the sample during this analysis must be considered. The absorbance of the sample, $A_\lambda$, is a function of injection volume, $V_i$, and the absorbance across a peak, $A_{\lambda p}$, is dependent on the volume across a peak, $V_p$. The value of $V_p$ can be determined by multiplying the flow rate by the peak width. The peak area, $a_p$, is the product of the absorbance intensity (mAU) multiplied by the peak width (minutes), $w_p$. 

<table>
<thead>
<tr>
<th>Absorbance across a peak</th>
<th>Absorbance of sample collected</th>
</tr>
</thead>
<tbody>
<tr>
<td>$A_{\lambda p} = \varepsilon \lambda c_p$</td>
<td>$A_\lambda = \varepsilon \lambda c$</td>
</tr>
<tr>
<td>$A_{\lambda p} = \varepsilon \lambda n/V_p$</td>
<td>$A_\lambda = \varepsilon \lambda n/V_i$</td>
</tr>
<tr>
<td>$\varepsilon = A_{\lambda p} \cdot V_p / l n$</td>
<td>$\varepsilon = A_\lambda \cdot V_i / l n$</td>
</tr>
</tbody>
</table>

$$A_{\lambda p} = A_\lambda \cdot V_i / V_p \quad \text{Eq (1)}$$

Volume across peak

$$a_p = A_{\lambda p} \cdot w_p$$

$$w_p = V_p / F$$

$$A_{\lambda p} = a_p \cdot F / V_p \quad \text{Eq (2)}$$

**Combine Eq 1 & 2**

$$a_p \cdot F / V_p = A_\lambda \cdot V_i / V_p$$

$$a_p \cdot F = A_\lambda \cdot V_i$$

$$A_\lambda = a_p \cdot F / V_i \quad \text{Eq (3)}$$

For this analysis, the conversion of the absorbance (mAU) of a water sample ($A_\lambda$) to Mm$^{-1}$ considers the volume collected by the PILS and the volume of air sampled by the inlet system as described in Washenfelder et al. (2022) and the PILS collection efficiency:
Absorption \( (Mm^{-1}) = \frac{\text{peak area} \times \text{SEC Flow rate}}{\text{Injection Volume}} \times \frac{\text{PILS Flow}}{\text{Gas flow} \times \text{Optical path}} \times \ln(10) \times \text{PILSCollectionEff} \)

**Figure S10.** Absorption at 365 nm to CO as a function of plume age of water extracts for smoke sampled of glass microfiber filters in Vancouver (Di Lorenzo et al., 2017) The green and blue markers represent absorption at 365 nm to CO of water extracts sampled on Teflon filters aboard the NASA DC-8 aircraft during SEAC4RS (Forrister et al., 2015).

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<table>
<thead>
<tr>
<th>US State</th>
<th>Fire Name</th>
<th>Fuel</th>
<th>Date and Flight leg</th>
<th>Number of water samples collected</th>
<th>Number of Transects</th>
<th>Plume age range (hr)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Arizona</td>
<td>Castle</td>
<td>Grassland, timber, mixed conifer</td>
<td>20190821-L2</td>
<td>10</td>
<td>8</td>
<td>0.2 – 4.1</td>
</tr>
<tr>
<td></td>
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<td></td>
<td>20190824-L2</td>
<td>10</td>
<td>8</td>
<td>0.5 – 2.4</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>20190825-L2</td>
<td>9</td>
<td>10</td>
<td>0.5 – 2.9</td>
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<tr>
<td></td>
<td>204 Cow</td>
<td>Sub-alpine fir, timber, grass</td>
<td>20190828-L1</td>
<td>6</td>
<td>6</td>
<td>0.7 – 1.8</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>20190828-L2</td>
<td>6</td>
<td>4</td>
<td>1.3 – 2.9</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>20190828-L3</td>
<td>6</td>
<td>4</td>
<td>2.2 – 4.2</td>
</tr>
</tbody>
</table>
References


