Application of the ECT9 protocol for radiocarbon-based source apportionment of carbonaceous aerosols

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Abstract: Carbonaceous aerosol is mainly composed of organic carbon (OC) and elemental carbon (EC). Both OC and EC originate from a variety of emission sources. Radiocarbon (^{14}C) analysis can be used to apportion bulk aerosol, OC, and EC into their sources. However, such analyses require the physical separation of OC and EC.

Here, we apply of ECT9 protocol to physically isolate OC and EC for ^{14}C analysis and evaluate its effectiveness. Several reference materials are selected, including: two pure OC (fossil "adipic acid", contemporary "sucrose"), two pure EC (fossil "regal black" and "C1150"), and three complex materials containing contemporary and/or fossil OC and EC ("rice char" and NIST urban dust standards "SRM1649a", i.e., bulk dust and "SRM8785", i.e., fine fraction of re-suspended SRM1649a on filter). The pure materials were measured for their OC, EC and total carbon (TC) mass fractions and corresponding carbon isotopes to evaluate the uncertainty of the procedure. The average accuracy of TC mass, determined via volumetric injection of a sucrose solution, was approximately 5%. Ratios of EC/TC and OC/TC were highly reproducible, with analytical precisions better than 2% for all reference materials, ranging in size from 20 to 100 μg C. Consensus values were reached for all pure reference materials for both δ^{13}C and FM^{14}C with an uncertainty of <0.3‰ and approximately 5%, respectively. The procedure introduced 1.3±0.6 μg of extraneous carbon, an amount compatible to that of the Swiss_4S protocol.

In addition, OC and EC were isolated from mixtures of pure contemporary OC (sucrose) with pure fossil EC (regal black) and fossil OC (adipic acid) with contemporary EC (rice char EC) to evaluate the effectiveness of OC and EC separation. Consensus FM^{14}C values were reached for all OC (~ 5-30 μg) and EC (~10-60 μg) fractions with an uncertainty of <5%. We found that the ECT9 protocol efficiently isolates OC or EC from complex mixtures. Based on δ^{13}C measurements, the average contribution of charred OC to EC is likely less than 3% when the OC loading amount is less than 30 μg C.

Charring was further assessed by evaluating thermograms of various materials, including aerosol samples collected in the Arctic and from tailpipes of gasoline or diesel engines. These data demonstrate that the ECT9 method effectively removes pyrolyzed OC. Thus, the ECT9 protocol, initially developed for concentration and stable isotope measurements of OC and EC, is suitable for ^{14}C-based apportionment studies for environment samples, including μg C-sized samples from Arctic environments.

1 Introduction

35　Carbonaceous aerosol is a major component (15-90%) of airborne particulate matter (PM) (Jimenez et al., 2009; Putaud et al., 2010; Yang et al., 2011; Hand et al., 2013; Ridley et al., 2017), and a complex mixture composed of mainly light-scattering organic carbon (OC) and highly-refractory, light-absorbing elemental carbon (EC, also referred to as black carbon) (Pöschl, 2005). The OC and EC fractions play important and often distinct roles in climate (Bond et al., 2013; Hallquist et al., 2009; Kanakidou et al., 2005; Laskin et al., 2015), air pollution and human health (Cohen et al., 2017; Grahame et al., 2014;

40　Janssen et al., 2012). Moreover, both OC and EC were identified as short lived climate forcers (SLCFs) by the IPCC expert meeting (https://www.ipcc-nggip.iges.or.jp/public/mtdocs/1805_Geneva.html) in 2018. To develop and monitor the efficiency of mitigation strategies for both climate change and air pollution, it is required to have a better understanding of the temporal and spatial dynamics of OC and EC emission sources.

　　The majority (>50%) of carbonaceous aerosol is OC, which has a wide size range. Coarse OC (in PM_{10}) consists of plant

45　debris, microorganisms, fungal spores, and pollen. Fine OC (in $PM_{2.5}$) is formed predominantly via the oxidation or nucleation/coagulation of volatile organic compounds, such as mono- and sesquiterpenes, from both biogenic and anthropogenic sources (Shrivastava et al., 2017), but can also be directly emitted from combustion sources (Hallquist et al., 2009; Fuzzi et al., 2015; Liggio et al., 2016). In contrast, EC is found primarily in fine particles, e.g., $PM_{1.0}$ or smaller (Chan et al., 2013; Bond et al., 2013). It is emitted through incomplete combustion of fossil fuels and biomass/biofuels (Bond et al.,

50　2013; Huang et al., 2010; Evangeliou et al., 2016; Winnie et al., 2016; 2017; 2019).

　　Measuring the isotopic signature and composition, i.e. radiocarbon (^{14}C) content and stable isotope ratio ($^{13}C/^{12}C$) of aerosol, offers a powerful tool for quantifying the sources of bulk aerosol and its OC and EC fractions. Aerosol ^{14}C content can be used to quantify the relative contributions from contemporary biomass and fossil sources (Heal, 2014). ^{14}C is a naturally occurring radioisotope (5,730-year half-life) produced in the atmosphere. After its oxidation to carbon dioxide ($^{14}CO_2$), ^{14}C

55　enters the food chain through photosynthesis so that all living organisms are labeled with a characteristic $^{14}C/^{12}C$ ratio and described as "modern" carbon. Materials containing carbon older than about 50,000 years ($^{14}C<<^{12}C$) are described as "fossil" carbon. Over the past centuries, the ^{14}C content of the atmosphere has undergone distinct changes (Graven, 2015; Levin et al., 2010): Anthropogenic combustion of fossil fuels emit ^{14}C-depleted carbon into the atmosphere (i.e. dilute the proportion of ^{14}C relative to ^{12}C). In contrast, nuclear weapons testing doubled the ^{14}C content of CO_2 in the Northern

60　Hemisphere in the mid-20th century, followed by mixing of this bomb-derived ^{14}C-enriched carbon into the ocean and biosphere. Similarly, aerosol stable isotope ratios provide insight to different types of anthropogenic sources (e.g. combustion of solid and liquid vs. gaseous fossil fuels). However, ^{13}C data cannot distinguish emissions from mixed fossil fuel combustion and live C3 plant biomass (Huang et al., 2006; Winiger et al., 2016). Thus, isotope-based source apportionment studies become particularly insightful when both ^{14}C and stable carbon isotopes are considered (Andersson et al., 2015;

65　Winiger et al., 2016, 2017) or when combined with analyses of specific source tracers, such as levoglucosan or potassium for wood burning emissions (Szidat et al., 2006; Zhang et al., 2008) and/or remote sensing data and modeling analysis (Barrett et al., 2015; Mouteva et al., 2015b; Wiggins et al., 2018).

The objective of this study is to evaluate the effectiveness of separating OC and EC via the ECT9 (EnCan-Total-900) protocol (Huang et al., 2006; Chan et al. 2010; Chan et al., 2019) for ^{14}C-based source apportionment studies of carbonaceous

70   aerosols. The ECT9 technique was originally developed to physically separate OC and EC mass fractions for concentration quantification and stable carbon isotope analysis. This protocol has been used since 2006 to monitor carbonaceous aerosol mass concentrations and stable isotope composition over Canada, including in the Arctic at Alert, as part of the Canadian Aerosol Baseline Measurements (CABM) Network by Environment & Climate Change Canada (Chan et al., 2010; 2019; Eckhardt et al., 2015; Sharma et al., 2017; Xu et al., 2017; Leaitch et al., 2017; 2018; Huang, 2018). It has also been used to

75   monitor carbonaceous aerosol over China (Yang et al., 2011). Furthermore, EC concentration measurements made with the ECT9 protocol correlate well with those derived from light absorption by an aethalometer as well as refractory black carbon (rBC) using a Single Particle Soot Photometer (SP2) (Sharma et al., 2017; Chan et al., 2019). It was demonstrated that the ECT9 protocol can be used to quantify OC/EC concentrations and provide source information at the same time.

The ECT9 protocol is a thermal evolution analysis (TEA) protocol which is different from commonly used thermal optical

80   analysis (TOA) methods for monitoring air quality, such as the Interagency Monitoring of Protected Visual Environments (IMPROVE) protocol (Chow et al., 2001; Watson et al., 2007), the National Institute for Occupational Safety and Health protocol (NIOSH method 5040, Birch, 2002), as well as the European Supersites for Atmospheric Aerosol Research (EUSAAR) protocol (Cavalli et al., 2010). In those protocols, the OC fraction is thermally desorbed from filter samples in an inert helium (He) atmosphere at relatively lower temperatures and the EC fraction is combusted at higher temperatures by

85   introducing oxygen (O_2) in He stream while the filter reflectance or transmittance for a laser signal is continuously monitored. During the analysis, a fraction of the OC may char (forming pyrolyzed OC or POC), causing the transmittance or reflectance to decrease. While TOA methods use the changes in laser signal to mathematically correct for POC within the measured EC fraction, the ECT9 protocol aims to minimize or remove POC, together with carbonate carbon (CC), during an intermediate temperature step of 870°C in pure He via high temperature evaporation (Chan et al., 2019). With much longer

90   retention times at each temperature step (see Methods) and without either reflectance or transmittance used, the ECT9 protocol effectively isolates OC, POC+CC, and EC.

It should be noted that other methods have been also developed mainly for ^{14}C analysis of OC and EC, such as the CTO-375 (Zencak et al., 2007), the Swiss_4S protocol (Mouteva et al., 2015a; Zhang et al., 2012), or hydropyrolysis (Meredith et al., 2012; Zhang et al., 2019), which use distinct temperature protocols, gas mixture and/or remove water-soluble OC or

95   inorganic carbon prior to EC analysis. In contrast to the ECT9 protocol, however, these approaches differ substantially from the protocols that are widely used for monitoring OC/EC mass concentrations in the field, which limits the relevance of this data for improving the representation of carbonaceous aerosols in chemical transport models.

Here we analyzed the ^{14}C content of OC and EC fractions (<100 µg C) isolated with the ECT9 protocol from four pure fossil and contemporary reference materials. These materials were analyzed on their own to quantify the amount and source

100   (modern or fossil) of extraneous carbon introduced by the procedure as well as its reproducibility. Mixtures of two reference materials were measured to elucidate how efficiently the ECT9 protocol isolates OC from EC. In addition, we investigated the laser signals of three reference materials and three aerosol samples (tailpipe emissions, ambient aerosol from Alert, and SRM8785) to assess how efficiently the ECT9 protocol removes POC. Our evaluation of the ECT9 protocol on its ability to

physically separate OC from EC for ^{14}C-based source apportionment studies significantly expands the existing opportunities

105 for characterizing and monitoring sources of carbonaceous aerosol at regional or global scales at the same time providing

solid base for EC and OC concentration measurements.

2 Methods

2.1 The ECT9 protocol for the physical separation of OC and EC

The ECT9 protocol was developed at the carbonaceous aerosol & isotope research (CAIR) lab of Environment and Climate

110 Change Canada (ECCC) to quantify the amount of OC and EC in carbonaceous aerosol and their $\delta^{13}C$ values (Huang et al.,

2006; Chan et al., 2010; 2019). Carbon fractions are isolated with an OC/EC analyzer (Sunset Laboratory Inc.) coupled to a

custom-made gas handling and cryogenic trapping system for CO_2 collection from OC and EC fractions (Fig. 1a). The

fractions are separated based on their thermal refractory. Specifically, carbon fractions are released by the ECT9 protocol in

three steps (Fig. 1b): (1) OC at 550°C for 600 seconds in pure He (99.9999% purity); (2) POC and CC at 870°C for 600

115 seconds in pure He; and (3) EC at 900°C for 420 seconds in a mixture of 2% O_2 with 98% He. All fractions are fully oxidized

to CO_2 by passing through a furnace containing MnO_2 maintained at 870°C. For concentration determination, the CO_2 passes

through a methanator at 500°C, is converted to CH_4, and quantified with a flame ionization detector. For isotope analysis, the

CO_2 is cryo-trapped with liquid N_2 (-196°C) in a U shaped glass trap, purified on a vacuum system (to remove He), sealed

into a Pyrex ampoule, and analyzed for its $\delta^{13}C$ ratio with an Isotopic Ratio Mass Spectrometer (IRMS), i.e., MAT253 or

120 $FM^{14}C$ with an Accelerated Mass Spectrometer (AMS).

2.2 Reference materials and their composition

To evaluate the ECT9 method for separating OC and EC for ^{14}C analysis, we isolated and measured the ^{13}C and ^{14}C content

of the OC or EC fraction or TC from 5-6 modern or fossil reference materials (Table 1), including two pure OC (adipic acid,

sucrose), two EC (C1150, regal black), and two natural OC/EC-mixtures (rice char and urban dust SRM1649a).

125 Some of the reference materials have previously been utilized to compare different protocols that quantify OC/EC fractions

(Hammes et al., 2007; Willis et al., 2016) as well as determine the mass of extraneous carbon introduced during OC/EC

isolation from carbonaceous aerosol (Mouteva et al., 2015a). Table 1 provides an overview of their chemical compositions,

i.e., total carbon contents and relative fraction of OC and EC, respectively (for individual measurements see Table S1).

Primary methods (i.e., gravimetric or volumetric) are used for mass loading of the materials, whereas the mass of TC, OC,

130 and EC are quantified via the ECT9 thermal protocol. Based on repeat injections of sucrose results (20-80 μg sucrose, n

=117), the accuracy of the TC mass is about 5%. The reproducibilities of both OC/TC and EC/TC percentages are 2% or

better. Although uncertainties of weighing pure EC mass (i.e., Regal black and C1150) via microbalances are relatively large

(due to static electricity and variable relative humidity), the EC/TC and OC/TC ratios for all reference materials are highly

reproducible (one s.d. <2%). The results show that the two EC materials (i.e., regal black and C1150) contain 97% and 98%

135 EC, with only 3% and 2% OC, respectively. The two OC materials (i.e., sucrose and adipic acid) are 99% and 100% OC, and

less than 1% EC (likely due to charred OC contribution), respectively. Thus, the materials are suitable for the purpose of this study.

We also analyzed the ^{13}C and ^{14}C isotopic composition of each reference material, using off-line combustions and ECT9 coupled with cryo-purification to convert them into CO_2. The results are summarized in Table 2 (for individual results see

140   Tables S2 & S3). The ^{14}C analysis of µg C-sized carbonaceous aerosol samples requires the assessment of extraneous carbon (Santos et al., 2010). This is achieved by measuring multiple smaller-sized materials with known ^{14}C content. Consequently, the results in Table 2 are critical, as those ^{14}C values provide the reference for quantifying the extraneous carbon introduced during the isotope analysis procedures.

2.3 Isolation of OC, EC or TC with the ECT9 protocol and purification of CO_2

145   The isotopic analysis of carbonaceous aerosol via the ECT9 system involves three steps (Fig. 1a): 1) OC and EC isolation/CO_2 collection and 2) CO_2 purification, followed by 3) isotope analysis for either ^{13}C/^{12}C by IRMS or ^{14}C by AMS (coupled measurements of ^{13}C/^{12}C and ^{14}C/^{12}C of µg C- sized graphite targets), as desired.

The initial masses of the pure reference materials ranged from 5 to 47 µg C (n=3-13; Table S6), whereas those for the mixed materials ranged from 5-30 µg C for OC and 5-60 µg C for EC (n=5-6; Table S7). The loaded mass of each material was

150   determined via a microbalance (MX5, Mettler Toledo or CCE6, Sartorius) with the lowest reading to 1 µg C or 0.1 µg C, respectively. OC materials were dissolved in MQ-water with known volumes and volumetrically loaded onto a pre-cleaned quartz filter punch (1.5 cm^2, Pall Canada Limited). EC (i.e., Regal black and C1150) and mixed materials (rice char or SRM 1649a), which cannot be completely dissolved in water, were directly weighed onto pre-cleaned quartz filter punches. These filters were pre-combusted at 900°C in a muffle furnace overnight and wrapped into aluminum foil before cooling below

155   200°C. A filter punch with the loaded mass was put into the Sunset analyzer and analyzed with the ECT9 protocol. OC and EC were separated and the combusted OC or EC fractions as CO_2 were cryo-collected in a U-shaped flask submerged in liquid N_2 (Fig. 1a, step 1). Then, this flask containing CO_2 and He was connected to a vacuum line with 4 cryo-traps and several open ports (Fig. 1a, step 2), where the CO_2 is purified by sequential distillation when passing cryo-traps 1 through 3. Finally the pure CO_2 is transferred and sealed into a 6 mm glass ampoule for ^{13}C or ^{14}C analysis. Pressure is read by a Pirani

160   gauge before sealing the ampoule for an estimation of the amount of gas, and consequently, sample size determination as µg C.

2.4 ^{14}C measurements

At the KCCAMS facility, the OC and EC fractions or TC (in form of CO_2) were reduced to graphite on iron powder via hydrogen (H_2) reduction using equipment and protocols specifically developed for smaller-sized (≤15 µg C) samples (Santos

165   et al., 2007b; 2007a). Briefly, sample-CO_2 was introduced into a vacuum line, cryogenically isolated from any water vapor, monometrically quantified, and then transferred to a heated reaction chamber, where it was mixed with H_2 and reduced to filamentous graphite. To characterize the graphitization, handling and AMS analysis, two relevant standards (Oxalic Acid II as modern carbon and Adipic acid as fossil carbon), with similar size ranges of the samples prepared via ECT9, were also

170 processed into graphite. The graphite was then pressed into aluminum holders and loaded into the AMS unit alongside measurement standards (Table S6) and blanks for ^{14}C measurement (Beverly et al., 2010). The data are reported in fraction modern carbon (FM[14]C), following the conventions established by Stuiver and Polach (1977) and also described elsewhere (Trumbore et al., 2016).

To establish consensus values (Table 2), we also analyzed the ^{14}C content of the bulk reference materials ranging in size from 0.06 to 1 mg C, using our standard combustion and graphitization methods. Larger aliquots of material were weighed into

175 pre-combusted quartz tube with 80 mg CuO, evacuated, and combusted at 900°C for 3 hours. The resulting CO_2 was cryogenically purified on a vacuum line, reduced to graphite using a closed-tube zinc-reduction method (Xu et al., 2007), and analyzed as described above.

2.5 Quantification of extraneous carbon

Any type of sample processing and analysis introduces extraneous carbon (C_{ex}). Therefore, the measured mass of any sample

180 will include the mass of this sample and of any C_{ex} incorporated throughout the analysis [Eq. 1]:

$$m_{spl\_meas} = m_{spl} + m_{ex}$$ [Eq. 1],

where m_{spl_meas}, m_{spl}, and m_{ex} are the measured and theoretical mass of the sample and of C_{ex}, respectively. For small samples (with a mass of a few µg C), the mass of C_{ex} can compete with or overwhelm the sample mass and cause the measured FM[14]C value of a sample to deviate from its consensus value.

185 Here, we estimated the mass of C_{ex} introduced during the ECT9 protocol and the ^{14}C analysis following Santos et al. (2010), where C_{ex} is understood to consist of a modern and of fossil component [Eq. 2]:

$$m_{ex} = m_{mex} + m_{fex}$$ [Eq. 2],

where m_{mex} and m_{fex} is the mass of the modern and fossil C_{ex}, respectively.

Following an isotope mass balance approach, the measured isotopic ratio (^{14}C/^{12}C) of a sample (R_{spl_meas}) can be expressed

190 as [Eq. 3].

$$R_{spl\_meas} = \frac{m_{spl}R_{spl} + m_{mex}R_m + m_{fex}R_f}{m_{spl\_meas}}$$ [Eq. 3],

where R_{spl} is the theoretical isotopic ratio of the sample, and R_m and R_f are the consensus isotopic ratios of a modern and fossil standard, respectively. This equation can be further simplified because R_f is 0. R_m is determined by measuring regular-sized aliquots of this reference material. In addition, all ^{14}C/^{12}C ratios are corrected for isotope

195 fractionation using their δ^{13}C measured alongside ^{14}C on the AMS (Beverly et al., 2010).

The mass of modern C_{ex} can be quantified by analyzing fossil reference materials, which are highly sensitive to modern and insensitive to fossil pollutants. Based on [Eq. 3] the measured isotopic ratio of the fossil reference (R_{f_meas}) can be expressed as [Eq. 4]:

6

$$R_{f\_meas} = \frac{m_{mex}R_m}{m_{spl\_meas}} \qquad \text{[Eq. 4]}$$

200 The smaller the mass of the fossil reference material, the greater the effect of the constant mass of modern C_{ex} on the isotope ratio of the fossil reference material, i.e. R_{f_meas} deviates toward R_m.

Similarly, the mass of fossil C_{ex} can be quantified by analyzing modern reference materials. With decreasing mass, the measured isotopic ratio of the modern reference (R_{m_meas}) will deviate more strongly from R_m (toward R_f). Based on [Eq. 1-3] and assuming $m_{spl} \gg m_{mex}$, the R_{m_meas} can be expressed as [Eq. 5]:

205 $$R_{m\_meas} = \frac{m_{spl}R_m + m_{mex}R_m}{m_{spl\_meas}} \approx \frac{(m_{sp\ meas} - m_{fex})R_m}{m_{spl\_meas}} \qquad \text{[Eq. 5]}$$

Finally, we can calculate the C_{ex}-corrected isotope ratio of an unknown sample (FM_{spl_cor}). This value reported as the ratio between the theoretical isotopic ratio of this sample and the accepted value of a modern standard $\left(R/R_m\right)$ also known as Fraction Modern (FM; with all R corrected for stable isotope fractionation). This data is reported as [Eq. 6]:

210 $$FM_{spl\_cor} = \frac{R_{spl}}{R_m} \approx \frac{R_{spl\_meas} - R_{f\_meas}}{R_{m\_meas} - R_{f\_meas}} \approx FM_{m*} \times \frac{\left[\frac{R_{spl\_meas}}{R_m} - \frac{m_{mex}}{m_{spl\_meas}}\right]}{\left[1 - \frac{m_{mex}}{m_{spl\_meas}} - \frac{m_{fex}}{m_{spl\_meas}}\right]} \qquad \text{[Eq. 6]},$$

where FM_{m*} is determined from the direct measurement of the modern primary reference material (OX1) used to produce six time-bracketed graphite targets measured in a single batch, after isotopic fractionation correction and normalization (Santos et al., 2007a,b). The individual uncertainty of FM_{spl_cor} is determined from counting statitics and by propagating the quantified blanks using a mass balance approach. Long-term and continuous measurements

215 of various types of blanks indicate that the mass of C_{ex} within one analytical method or system can vary as much as 50% (see Santos et al., 2010; Fig. 1). Therefore, we applied a 50% error in m_{fex} and m_{mex} from long-term measurements of variance in m_{ex} of small samples (Santos et al., 2007a).

In this study, we used a multi-step approach to quantify m_{ex} introduced by the ECT9 protocol and ^{14}C analysis. First, we quantified m_{ex} introduced during ^{14}C analysis by analyzing different masses of our bulk reference

220 materials. Extraneous carbon is introduced during sealed tube combustion and graphitization followed by graphite target handling and measurement. Typically, ^{14}C analysis contributes a small portion to m_{ex} (Mouteva et al., 2015a; Santos et al., 2010). Second, we quantified the portion of m_{ex} added during the isolation of OC and EC with the ECT9 protocol. This portion of m_{ex} allows us to determine the practical minimum sample size limit for the entire method, including m_{ex} contributions from filter handling before OC/EC analysis, instrument separation, and transfer

225 to cryo-collection system and Pyrex ampoules. To isolate this portion, we quantified m_{ex} of the entire procedure (ECT9 protocol plus ^{14}C analysis) by analysing the ^{14}C signature of OC and EC from different masses of a large set of reference materials, and then subtracted the portion of m_{ex} introduced during ^{14}C analysis.

3. Results and Discussion

3.1. Recovery estimation

230 The reference materials used in this study, including the modern and fossil endmembers (i.e., the major carbon

sources) found in carbonaceous aerosol, and their TC, OC, and EC concentrations are shown in Table 1. Reference

materials were separated into OC, EC, or TC using the ECT9 method at ECCC's CAIR lab (Fig. 1) and analyzed for

their ^{14}C content at UC Irvine's KCCAMS facility.

Fig. 2 shows the cross-validation of carbon-mass between the mass determined at ECCC's CAIR lab and the mass

235 quantified at UC Irvine's KCCAMS lab indicating a very good positive correlation ($R^2 = 0.93$ for pure materials and

$R^2 = 0.95$ for two-material-mixtures in Fig. 2a and 2b, respectively). It is suggested that no major losses or gains of

carbon occurred during the entire analytical process and the overall recovery was close to 100%, with a 5%

uncertainty for samples ranging in size from about 5 to 60 μg C.

3.2 Quantification of extraneous carbon and its sources

240 All types of samples, regardless of size, show deviations in their measured $FM^{14}C$ value from their consensus values to

certain degree due to C_{ex} introduced during sample analysis. In μg C-sized samples (mass <15 μg C), significant bias from

any C_{ex} can be observed, because C_{ex} constitutes a large fraction of the total sample. Previous work (using solvent-free

analytical protocols) has shown that modern C_{ex} is introduced mostly through instrumentation and sample handling

techniques, while fossil C_{ex} originates from iron oxide used as a catalyst for the reduction of CO_2 to graphite prior to AMS

245 analysis (Santos et al., 2007a; 2007b).

The $FM^{14}C$ values of the pure modern or fossil reference materials generally agreed with their accepted $FM^{14}C$ values for

both OC and EC fractions (within approximately 5% uncertainty, Fig. 3 and Table S6) after applying a constant amount C_{ex}

correction in $FM^{14}C$ determination. This constant C_{ex} is a critical prerequisite for accurately correcting the $FM^{14}C$ value of

unknown samples. Hence, our data demonstrated that the ECT9 method (and subsequent ^{14}C analysis) introduces a small,

250 reproducible amount of C_{ex}.

According to equations [4]-[5] in section 2.5, C_{ex} can be quantified by measuring $FM^{14}C$ of pure modern or fossil materials

with different sizes. Fig. 3 demonstrates that regardless what ^{14}C content are in carbon fractions isolated from the reference

materials and what sizes they are, the corrected $FM^{14}C$ values match with consensus value within propagated uncertainty.

To evaluate the suitability of ECT9 for ^{14}C analysis of aerosol samples, a comparison is made between the results of a

255 published method (i.e., Swiss_4S) and those of ECT9. The two protocols are listed in Table 3 and their C_{ex} distribution is

shown in Table 4. The total amount of C_{ex} introduced by the complete procedure through ECT9, and determined based on all

reference materials, was 1.3±0.6 μg C, with 70% originating from contamination with modern carbon (Table 4). The isolation

of OC and EC with the ECT9 protocol introduced 65% of total C_{ex} (0.85 out of 1.35 μg C), with 65% derived from modern

carbon. Overall, the total amount of C_{ex} introduced during OC/EC isolation with the ECT9 protocol is comparable to that for

260 the Swiss_4S protocol established at UC Irvine within uncertainties (Table 3, Mouteva et al. (2015a)). Thus, it is

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demonstrated that the ECT9 protocol serves as a suitable alternative for the ^{14}C analysis of aerosol samples with masses >2 µg C.

3.3 Effectiveness of OC/EC separation

To investigate the effectiveness of the ECT9 to separate OC from EC in more complex mixtures with minimizing OC into the

265 EC fraction via pyrolysis, mixtures of the modern and fossil reference materials (Table 2) were used for measuring δ^{13}C (Table S4 - S5) and FM^{14}C (Table S7).

First, it was found that the FM^{14}C values of OC and EC fractions isolated from mixtures of pure sucrose (modern OC) and pure regal black (fossil EC) were within the measurement uncertainty of their accepted FM^{14}C values, after correction for a constant amount of C$_{ex}$ (Fig. 4) for samples with 5 – 34 µg OC carbon and 10 – 60 µg EC carbon, showing a good separation

270 of OC from EC. This amount of C$_{ex}$ was identical to that applied to the pure reference materials above, further corroborating the constant background introduced by the ECT9 protocol and ^{14}C analysis.

Next, the mixtures of fossil adipic acid (pure OC) and modern rice char (mixture of OC and EC) were isolated and analyzed. It was found that after correction for C$_{ex}$, the FM^{14}C values of the OC (from the mixture) were systematically greater than the consensus value of the pure adipic acid, i.e., a FM^{14}C of zero (Fig. 5a), indicating that there was certain level of modern

275 fraction contributed to the measured OC from the modern rice char. This was because rice char contains about 14% of modern OC (Table 1).

To confirm that ECT9 could remove OC contained in rice char, an additional step was taken before mixing modern rice char's EC with the fossil OC (adipic acid). Specifically, we stripped the OC fraction of rice char by running rice char (on a filter) through the ECT9 protocol. Adipic acid (fossil OC) was then injected onto the filter with the remaining rice char-EC.

280 The results show that the FM^{14}C of OC values of this mixture lie well within the expected range of the consensus value (Fig. 5b) after a C$_{ex}$ correction as described above, demonstrating an excellent remove of rice char OC.

In both mixtures (fossil adipic acid with modern bulk rice char or rice char-EC), the corrected FM^{14}C values of the isolated EC fractions were within the expected range for the rice char reference material (Fig. 5c, d). This provides further evidence that the ECT9 protocol isolates modern EC from fossil OC with no obvious evidence of transferring fossil OC into the EC

285 fraction. Together, the three sets of mixing experiments (Figs. 4 & 5) provide strong evidence for the effectiveness of separating OC from EC via ECT9 protocol.

In addition to FM^{14}C measurements, δ^{13}C measurements in mixtures of OC and EC can also provide quantitative information on the effectiveness of OC and EC separation via ECT9. Various amounts of sucrose (pure OC, 10 – 30 µg C) were first mixed with varying amounts of Regal black (pure EC, 20 – 66 µg C). The mixtures were then physically separated into OC

290 and EC fractions by ECT9 for δ^{13}C measurements. The measured δ^{13}C values of OC and EC from these mixing experiments are listed in Table S4. Based on the δ^{13}C values of individual pure reference materials (Table S3) and a two-end-member mixing mass balance, it is estimated that the average fraction contributed into each other in the mixtures (i.e., sucrose fraction into Regal black or vice versa) was likely less than 3% (Table S5).

3.4 Charring evaluation & POC removal using the ECT9 protocol

295   It is known that some of OC (e.g., oxygenated OC or water soluble OC) would char to form pyrolyzed organic carbon (POC) when heated in an inert He atmosphere, darkening the filter (Chow et al., 2004; Watson et al. 2005) and causing decreased laser signals due to light-absorption of charred OC. In most TOA protocols, this POC would combust and contribute to EC when O_2 is added. However, POC can be also be gasified and released as CO at high temperatures (>700°C) with limited O_2 supply, e.g., oxygenated OC at 870°C (Huang et al., 2006; Chan et al., 2010;

300   2019). Most TOA protocols estimate POC by quantifying the mass associated with reflectance/transmittance changes, i.e., the mass released between the time when O_2 is introduced and the OC/EC split point (where the reflectance/transmittance returns to the initial value). In contrast to other TOA protocols, ECT9 defines POC as the mass released at the temperature step of 870°C (during a period of 600 seconds). This includes charred OC, calcium carbonate ($CaCO_3$) that decomposes at 830°C, and any refractory OC not thermally released at 550°C (Huang et al.,

305   2006; Chan et al., 2010; 2019).

  Although ECT9 do not use laser signals to quantify POC, it is expected that the changes of laser signals during the stage of 870°C would provide useful information about POC. Thus, four sets of samples were selected, including those of pure reference materials and ambient aerosol samples from different sources with heavy or light mass loading (e.g., those Arctic sample filters from different seasons) to evaluate the possible charring via ECT9. Their

310   thermograms are shown in Figures 6 to 9.

  Figure 6 shows thermograms of pure or bulk references for Regal black, sucrose, and rice char, respectively. It is observed in all three that the laser transmittance signals first decrease and then increases again during the 870°C step, and that they return to their initial values just before EC is released at the next step of 900°C. This demonstrates that the ECT9 method minimizes POC-contributions to the EC fraction.

315   The thermograms of aerosol (on filters) collected directly from tailpipe exhaust of a diesel engine vehicle and a gasoline engine passage car, respectively are shown in Figure 7. These data suggest that the amount of POC generated during analysis are sample/matrix dependent. Specifically, the mass fraction during the 870°C temperature is larger for the gasoline than the diesel engine. This finding supports previous work showing that POC is proportional to the amount of oxygenated OC (Chan et al., 2010). It is noticed that the laser signal reaches the

320   initial value before the EC step, further demonstrating that the charring contribution to EC is minimized.

  Another set of thermograms of two total suspended particle filter samples collected during the summer (August) and winter (December) of 2015 at an Arctic site (i.e., Alert) are shown in Figure 8. More details about these samples can be found in Wex et al. (2019). The laser signal patterns are similar to those shown in Figures 6 & 7, yet more pronounced. During the 550°C step, the laser signals decrease. During the 870°C step, the signals further decrease,

325   then increase, and finally increase to their initial point before EC is released at 900°C. These thermograms further demonstrate ECT9 is able to minimize POC by gasification.

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Finally, the thermographs of NIST urban dust reference material SRM 8785 (the re-suspended SRM 1649a urban dust with a fine fraction <2.5 μm collected on quartz filter) analyzed with ECT9 and Swiss_4S are shown in Figure 9. Both thermograms obtained with the ECT9 method (Fig. 9 a&b) show the similar patterns as those in Figs. 6-8,

330 i.e., the laser signals reaching the initial value just before the EC release at 900°, suggesting that the charring contribution to EC is minimized during the stage of 870°C even though some POC might remain.

In the thermogram obtained with the Swiss-4S protocol (Fig. 9c), the laser signal increases from the beginning of the run while the first two stages (375°C and 475°C) are under the conditions of pure O_2 stream, inferring that light absorbing carbon is released during the first two OC stages. The laser signal continues to increase while the

335 temperature increases up to 650°C (the third stage) under the pure He gas stream, indicating that no charred OC is formed. However, when the temperature starts decreasing from 650°C, the laser signal decreases, indicating POC formation below that temperature. This signal decrease continues until the beginning of the next pure O_2 stage. It is important to note that to obtain EC fraction, the Swiss-4 (Table 3) method calls for filter sample pre-treatment, i.e., extraction with water before the thermal separation of OC/EC to minimize the contribution of charred OC from the

340 3rd stage to EC at the 4th stage. While it is difficult to make direct comparisons between OC and EC from b) and c) in Figure 9, laser profiles from those thermograms indicate that in both cases charred OC is negligible or minimum.

Together, the thermograms (Figs. 6-9) elucidate that the ECT9 protocol can effectively remove or minimize charred OC (POC) to achieve good physical separation of OC and EC. Another great advantage of using ECT9 to separate OC from EC for isotope analysis (both ^{13}C & ^{14}C) is its consistency with the protocol used for OC and EC

345 concentration measurements. Moreover, the ECT9 method does not require filter samples to be pre-extracted with water before EC analysis (to reduce POC).

4. Conclusions

We demonstrate the effectiveness of the ECT9 protocol to physically isolate OC and EC from aerosol samples for ^{14}C and ^{13}C analysis by using OC and EC reference materials on their own and as mixtures. It was found that the

350 ECT9 protocol successfully separates OC and EC fractions with a low (but largely modern) total carbon blank of 1.3±0.6 μg C. The majority (65%) of this extraneous carbon originates from the isolation with the ECT9 protocol, with 35% contributed from graphitization and ^{14}C measurement of the samples at the KCCAMS facility. After mass balance background corrections, the FM^{14}C results from both bulk pure materials and mixtures (with sample size as small as 5 μgC) can reach the consensus values (Table 2) with an average uncertainty of about 5%.

355 In addition, we evaluated potential POC formation during ECT9 by investigating thermograms of a variety of reference materials and ambient filter samples. It is demonstrated that ECT9 provides a good alternative for carbonaceous aerosol source apportionment studies, including ultra small sized (5-15 μg C) samples obtained from Arctic regions. To increase the application of isotope data (^{14}C or ^{13}C) in atmospheric research, future efforts should be focused on the comparison on OC/EC separation via different methods/protocols using the same sets of reference

360 materials. At the same time, the isolation results should be also compared among those methods/protocols widely used in long-term national monitoring network for OC/EC contents, ensuring a consistency in measurements between OC/EC concentrations and their corresponding isotopic compositions.

Nomenclature

AMS	Accelerator Mass Apectrometry
365 ASTD	Atmospheric Science & Technology Directorate
BC	Black carbon
CABM	Canadian Aerosol Baseline Measurement
CAIR	Carbonaceous Aerosol & Isotope Research
CCMR	Climate Chemistry Measurements and Research
370 CC	Carbonate carbon
CRD	Climate Research Division
EC	Elemental carbon
ECCC	Environment and Climate Change Canada
ECT9	EnCan-Total-900 protocol
375 EUSAAR	European Supersites for Atmospheric Aerosol Research
FID	Flame ionization detector
FM^{14}C	Fraction Modern Carbon
ICP	Inter-comparison study
IRMS	Isotopic Ratio Mass Spectrometer
380 IMPROVE	Interagency Monitoring PROtected Visual Environments
KCCAMS	W.M. Keck Carbon Cycle Accelerator Mass Spectrometry Facility
MAC	Mass absorption coefficient
NIST	National Institute of Standard and Technology
OC	Organic carbon
385 PM	Particulate matter
POC	Pyrolyzed organic carbon
PSAP	Particle Soot Absorption Photometer
rBC	Refractory Black Carbon
SP2	Single Particle Soot Photometer
390 SRM	Standard Reference Material
TC	Total carbon
TEA	Thermal evolution analysis
TOA	Thermal optical analysis
UCI	University of California, Irvine

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395 **Data availability**

All data presented in this article are included in the supplement.

Supplement

The supplement related to this article is available online at: https://doi.org/10.5194/amt-2020-201-supplement. (to be finalized)

400 **Author contributions**

Conceptualizing and designing the study: LH, CIC, and GMS

Developing analytical methods and ensuring data quality: LH, GMS, WZ, CIC, BTR

Performing the experiments and data acquisition: WZ, GMS, SRH, VV, BTR

Data organizing /analysis and interpretation: LH, CIC, BTR, GMS, WZ

405 Writing the paper, including editing and preparing figures and tables: LH, CIC, BTR, GMS, WZ

 Competing interests

The authors declare that they have no conflicts of interest.

Acknowledgements

Tables

Table 1. Overview of the bulk reference materials analyzed with the ETC9 method for their total carbon (TC), organic carbon (OC), and elemental carbon (EC) content

Reference material	EC				OC				EC + OC mixture			
	Regal black		C1150		Sucrose		Adipic acid		Rice char		SRM-1649a	
	mean	s.d.	mean	s.d.	mean	s.d.	mean	s.d.	mean	s.d.	mean	s.d.
TC (%)	96	9	98	12	101[a]	4	43[b]	5	52[c]	1	17.9[d]	1.1
OC/TC (%)	**3**	1	**1**	2	**99**	1	**100**	0	**14**	1	**51.5**	0.8
EC/TC (%)	**97**	1	**99**	2	**1**	1	**0**	0	**86**	1	**48.5**	0.8
n	41		24		117		5		6		6	
Bulk material	fine powder				solution		fine powder					
Loading method	gravimetric (via a balance with 1 - 0.1 µg accuracy)				volumetric injection		gravimetric (1 - 0.1 µg accuracy)					
Loading range (µg)	16 - 134		4 - 104		20 - 80		30 - 250		70 - 210		440 - 1100	
Analysis period	2015 – 2017		2006, 2013, 2015		2013 - 2018		2015, 2019		2018		2004 - 2005	
Supplier	Aerodyne Research, MA, USA		McMaster Univ., ON, Canada		Sigma-Aldrich, MO, USA		Fisher-Scientific, NH, USA		Univ. of Zurich, Switzerland		NIST, MD, USA	

[a]101% is obtained from the ratio of TC measured to TC calculated from the injected solution of sucrose; [b]49% of TC to bulk material in adipic acid based on its molecular mass; [c]58.6% of TC to bulk material in Rice char obtained from Hammes et al. (2006); [d]17% of TC to bulk material in SRM 1649a obtained from a critical evaluation of inter-laboratory data by Currie et al. (2002)

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425 **Table 2.** Overview of the isotopic composition of the reference materials used in this study. Radiocarbon ($^{14}C/^{12}C$, reported as fraction modern (FM^{14}C)) was measure at the KCCAMS facility and $\delta^{13}C$ at the CAIR lab.

Reference material	EC				OC				EC + OC mixture			
	Regal black		C1150		Sucrose		Adipic acid		Rice char		SRM-1649a	
	mean	s.d.	mean	s.d.	mean	s.d.	mean	s.d.	mean	s.d.	mean	s.d.
^{14}C analysis												
FM^{14}C_TC	-0.0001	0.0006	0.0027	0.0008	1.0586	0.0016	0.0000	0.0002	1.0675	0.0007	0.5118	0.001
n	2		3		2		5		3		1	
Loading range (µg)	700 - 750		60 - 560		730 - 770		740 - 1050		900 - 960		760	
CO_2 isolation & $^{14}C/^{12}C$ analysis	Reference material is combusted in 6 mm O.D. quartz tubes with 80 mg CuO for 3 hours at 900°C. Sample-CO_2 is purified cryogenically & reduced to graphite (Xu et al., 2007).											
$\delta^{13}C$ analysis												
$\delta^{13}C_{VPDB}$ (‰)	-27.61	0.08	-23.06	0.08	-12.22	0.16	n/a		-26.74		-25.84	0.07
n	5		5		9		n/a		1		2	
Loading range (µg or µg C*)	15 - 70		20 – 50		20		n/a		160		600	
CO_2 isolation	Material is loaded on a quartz filter and combusted in a Sunset OCEC aerosol analyzer (http://www.sunlab.com) using the ECT9 method. Sample-CO_2 is collected in a U-shaped flask submerged in liquid N_2 at -196°C (Fig. 1b).						n/a		See description for Regal black, C1150, and sucrose.			
CO_2 extraction & $^{13}C/^{12}C$ analysis	Sample-CO_2 is cryogenically purified on a vacuum line and sealed into an ampoule for analysis with a MAT253 Isotopic Ratio Mass Spectrometer (Huang et al., 2013).						n/a					

*Sucrose was loaded as a solution (µg C), Regal Black, C1150, Adipic acid, Rice char, and SRM-1649a as a fine powder (µg dry mass); n/a = not applicable

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Table 3. Comparison of the OC and EC ECT9 and Swiss-4S isolation protocols.

Carrier gas	Carbon fraction	Temperature °C	Duration s	Comments
ETC9[a]				
He-purge		20 – 50	90	Purging of volatile and semi-volatile OC
He	OC	550	600	
He	POC + CC	870	600	Minimizing charred OC contribution to EC
O_2/He[b]	EC	900	420	
Swiss-4S[c]				
O_2-purge		20 – 50	90	Purging of volatile and semi-volatile OC
O_2	S1_OC	375	240	
O_2	S2_OC	475	120	
He	S3_OC	650	180	
O_2	S4_EC	760	160	Water-soluble OC is removed by water extraction prio thermal analysis.

[a]POC + CC = pyrolysis OC + carbonate carbon; [b]The flow of 10% O_2 + 90% He mixing with the flow of 100% He resulting in 2% O_2 + 98%He. in [c]The EC punch is flushed with Milli-Q water prior the analysis to remove the water-soluble OC and minimize charring (Zhang et al., 2012; Mouteva et al., 2015a).

Table 4. Comparison of the procedural contamination with extraneous carbon for aerosol reference materials partitioned into organic carbon (OC) and elemental carbon (EC) with the ECT9 or Swiss_4S protocols based on their ^{14}C contents. We assume a measurement uncertainty of 50% (see Methods).

Contamination Source	ECT9 µg C	Swiss_4S[a]
OC/EC isolation + trapping		
Modern	0.55	0.37
Fossil	0.30	0.13
Total	0.85	0.50
[14]C analysis[b]		
Modern	0.35	0.43
Fossil	0.10	0.53
Total	0.45	0.97
Full set-up		
Modern	0.90	0.80
Fossil	*0.40*	*0.67*
Total	*1.30*	*1.47*

[a]From Mouteva et al. (2015a), [b]Carbon introduced during sample combustion, CO_2 purification and graphitization, and measurement with 14C-AMS.

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Figure 1: Overview of the carbonaceous aerosol analysis system at Environment and Climate Change Canada.
(a) Schematic flow chart for ^{13}C & ^{14}C measurements of OC/EC via ECT9, including 1) OC/EC isolation/CO_2
collection via cryo-trapping, 2) CO_2 purification, and 3) isotope analysis with IRMS ($^{13}C/^{12}C$ of CO_2) or AMS
460   ($^{13}C/^{12}C$ and $^{14}C/^{12}C$ of graphite targets).

(b) Thermogram of the ECT9 protocol on a Sunset OC/EC Analyzer. First, organic carbon (OC) is thermally
desorbed at 550ºC for 600 seconds in 100% He, then any pyrolyzed OC (POC), refractory OC, and carbonate carbon
(CC) is released at 870ºC in 100% He for 600 seconds. Finally, elemental carbon (EC) is combusted at 900ºC for 420
seconds by introducing 2% O_2 in He. All carbon fractions are oxidized to CO_2 followed by reduction to CH_4 and
465   quantification via flame ionization detection (FID) for carbon content or purified and cryo-trapped in Pyrex ampoules
for isotope analysis. Example FID signals are shown for a pure OC reference material (sucrose) mixed with a pure
EC material (regal black) along with the internal standard (CH_4).

Figure 2: Cross-validation of carbon-mass prepared, isolated by the ECT9 protocol and collected via cryo-trapping at ECCC
and then, retrieved during the purification and graphitization on a KCCAMS vacuum line. Carbon fractions (organic carbon
470   (OC), elemental carbon (EC), or total carbon (TC)) were isolated from two reference materials for OC (sucrose, adipic acid),
EC (regal black, C1150), and one OC & EC mixture (rice char). Most of the points deviating from the 1:1 line are carbon-rich
reference materials, e.g., Regal black and C1150 (>90% TC), which usually there are greater uncertainties in initial mass
determination via weighing using microbalance, because their sample sizes aimed were very small.

Figure 3: Radiocarbon (^{14}C) compositions, expressed as Fraction Modern Carbon, of total carbon (TC, circles),
475   organic carbon (OC, triangles) and elemental carbon (EC, squares) fractions isolated with the ECT9 protocol from
modern or fossil individual reference materials. **a)** Sucrose and **b)** adipic acid are modern and fossil OC, respectively,
c) regal black and **d)** C1150 are fossil EC, and **e)** rice char is a mixture of modern OC and EC. Open and solid
symbols represent ^{14}C data before and after correction for extraneous carbon introduced during OC/EC isolation and
subsequent ^{14}C analysis, respectively. The dashed line indicates the consensus value determined from regular-sized
480   bulk samples of these materials undergoing off-line combustions (see Table 2).

Figure 4: Radiocarbon (^{14}C) composition, expressed as Fraction Modern Carbon, of **a)** organic (OC, triangles) or **b)**
elemental (EC, squares) carbon fractions isolated with the ECT9 protocol from mixtures of pure modern OC
(sucrose) with fossil EC (regal black). Open and solid symbols represent ^{14}C data before and after correction for
extraneous carbon introduced during OC/EC isolation via ECT9 and subsequent ^{14}C analysis via AMS, respectively
485   (see Table S7). The dashed line indicates the consensus value (see Table 2).

Figure 5: Radiocarbon (^{14}C) compositions, expressed in fraction modern carbon, of organic (OC, triangles) and
elemental (EC, squares) carbon fractions isolated with the ECT9 protocol from the mixtures of reference materials.
Fraction of modern carbon **a)** OC and **c)** EC isolated from mixtures of pure fossil OC (adipic acid) with modern bulk
rice char (made of 14% OC and 86 % EC), and of **b)** OC and **d)** EC isolated from mixtures of pure fossil OC (adipic
490   acid) with modern EC from rice char_EC (rice char _OC has been removed before mixing). Open and solid symbols

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represent data before and after correction for extraneous carbon introduced during OC/EC isolation via ECT9 and subsequent ^{14}C analysis via AMS respectively (Table S7). The dashed line indicates the consensus value (see Table 2).

**Figure 6:** Thermograms of pure or bulk references. **a)** Regal black and **b)** Sucrose and **c)** Rice char. Temperature (blue solid line) and FID signals (integrated yellow area with green line) on the left axes and laser (red solid line) on the right axis. It is observed that on the three thermograms during the temperature stage of 870°C, the laser transmittance signals decrease first and increases again before the next temperatures stage, minimizing POC fraction, i.e., possible charred OC contribution to EC.

Figure 7: Thermograms of the filters directly collected from tailpipe exhaust of a diesel engine vehicle in **a)** and a gasoline engine passage car in **b)**. The legends are the same as Fig 6. It is noticed that the mass fraction from the temperature stage of 870°C in b) is obvious larger than that in a). The latter is negligible indicating that the amount of POC fraction is sample-matrix dependent. The amount of POC from gasoline vehicle emissions is likely larger than that from diesel vehicle emissions. It was noticed that the laser signal reaches the initial value before the 900°C stage for EC releasing, demonstrating that the charring contribution to EC is minimized.

Figure 8: Thermograms of fine particles (PM1.0 μm) from the filter samples collected at an Arctic site, i.e., Alert, NU, Canada in summer **a)** and in winter **b)** of 2015. The legends are the same as Fig 6. It is clearly shown on both thermograms that during 550°C stage, the laser signal starts decreasing (implying charred OC formation) and begins increasing during 870°C and reaches the initial value before the EC stage (indicating the contribution to EC by charred OC is minimized or removed).

Figure 9: Thermograms of the SRM 8785 filters (the fine fraction ($PM_{2.5}$) of re-suspended urban dust particles from SRM 1649a and collected on quartz filters) with various amount of materials ranging from 614 mg to 1723 mg via two different thermal protocols. **a)** and **b)** were obtained by ECT9. The legends are the same as Fig 6. Both thermograms in a) and b) show the similar patterns as in Fig. 6, 7, 8. that the laser signals reaching the initial value are just before the temperature stage of EC, suggesting that the charred OC contribution to EC is minimized. The thermogram in **c)** is obtained from the same filter in b) but by Swiss-4 protocol for comparison. The legends are similar except for the integrated area with green line, which stands for CO_2 in ppm (by NDIR) instead of FID signals.

Figures

Figure 1a. Schematic procedures for ^{13}C & ^{14}C measurements of OC/EC via ECT9
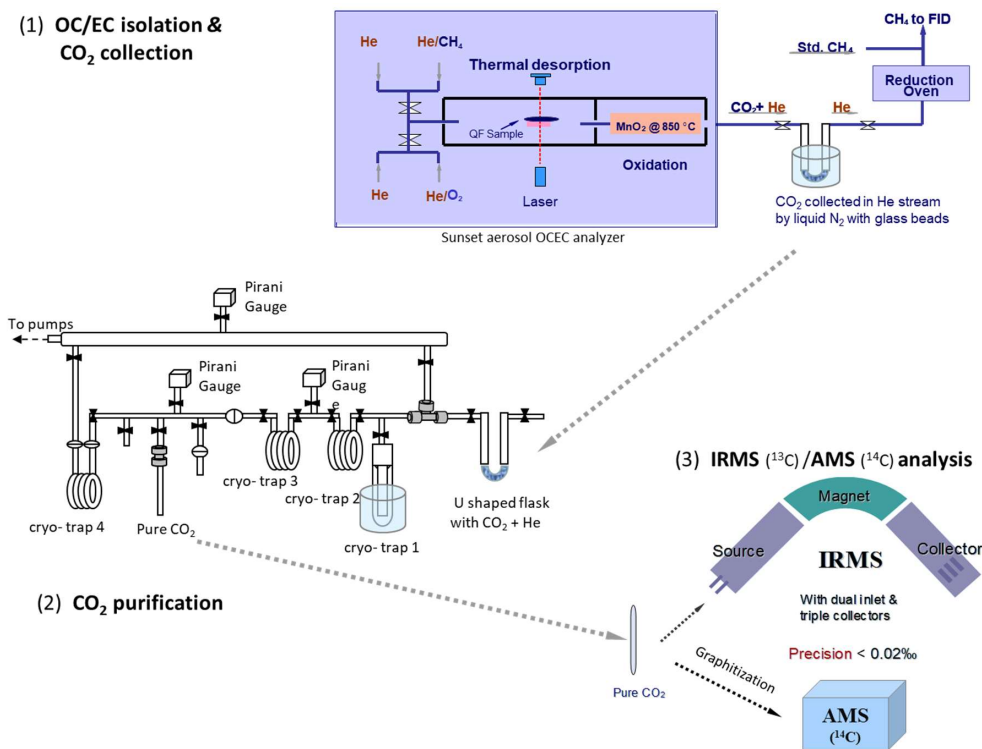
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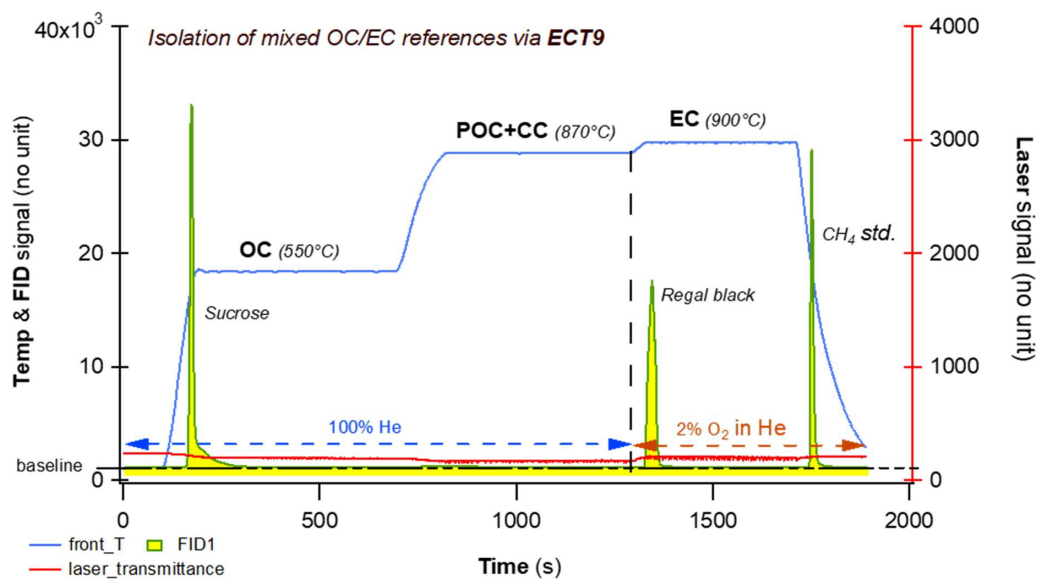
550   **Figure 1b**

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Figure 2

a)

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b)

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Figure 3

Figure 4

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590   **Figure 5**

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Figure 6

a) RegalBlack_82ug_012919_1
b) Sucrose_20ug_112212
c) Rice Char-050919
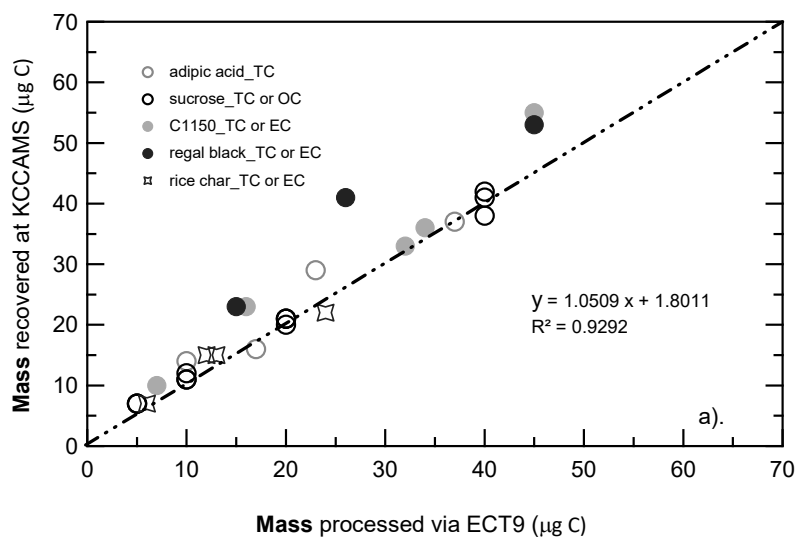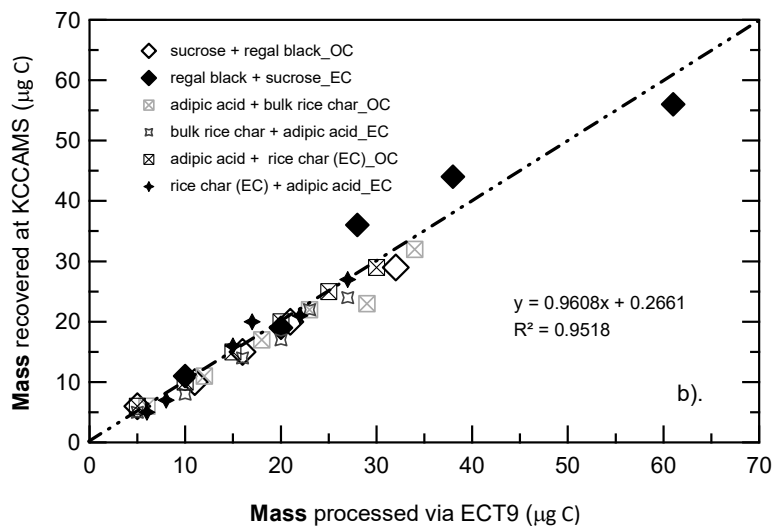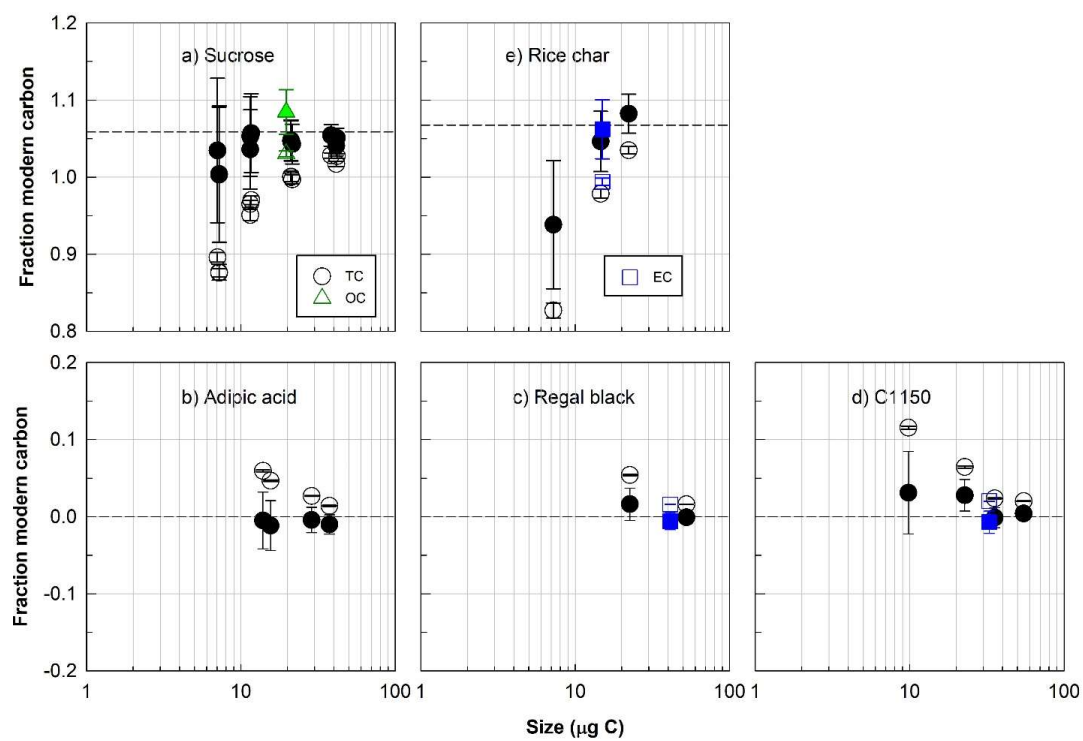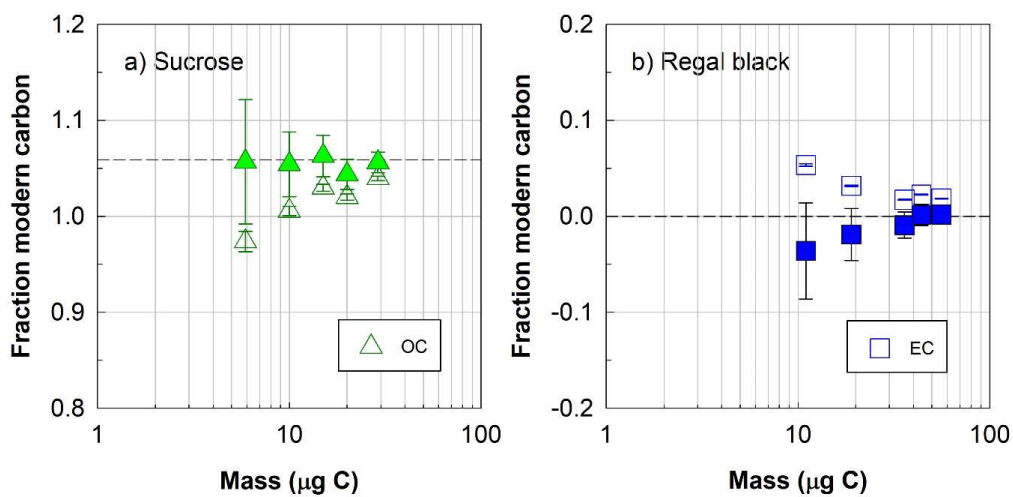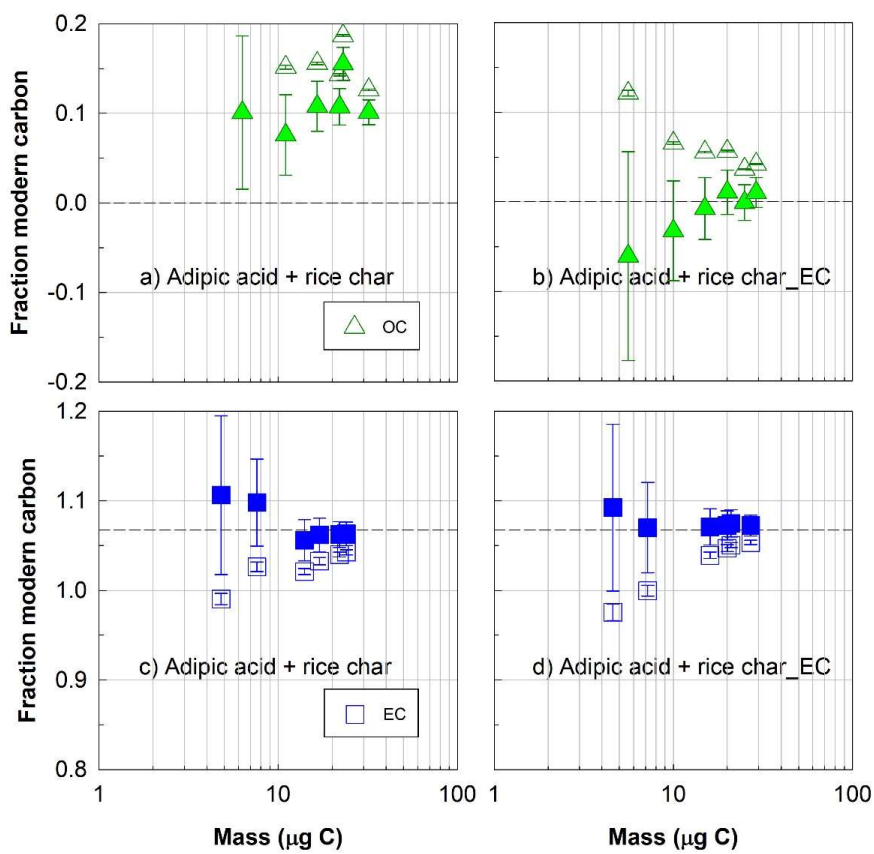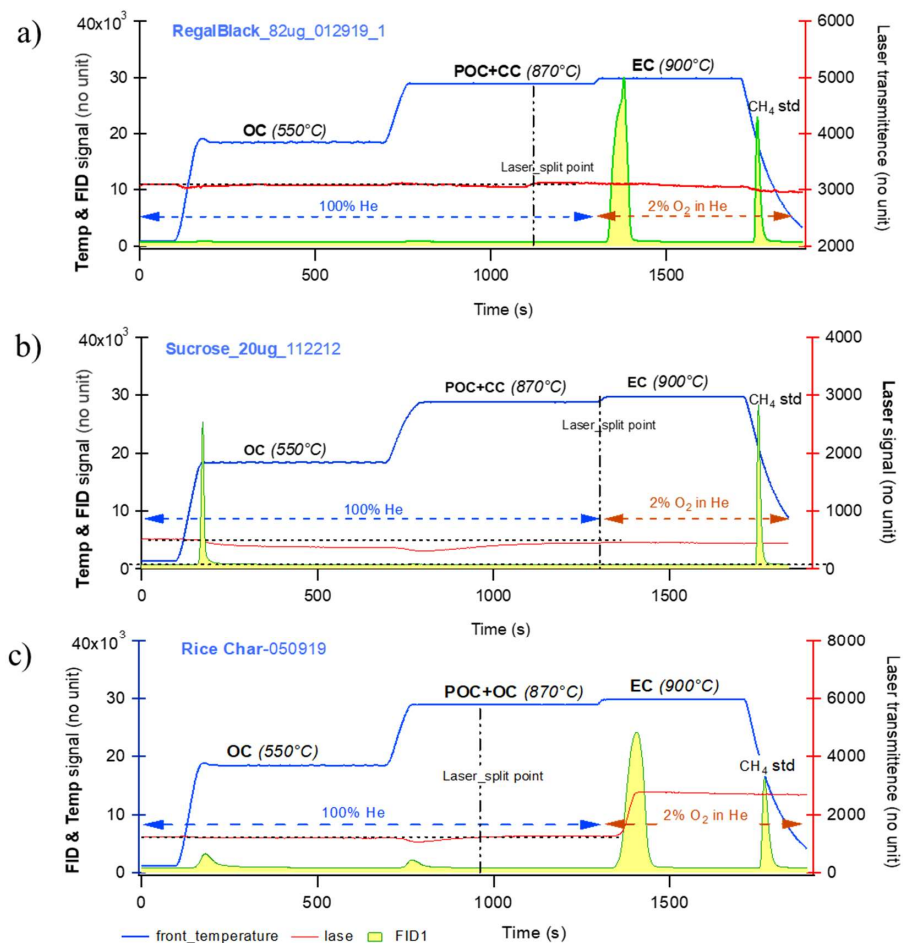
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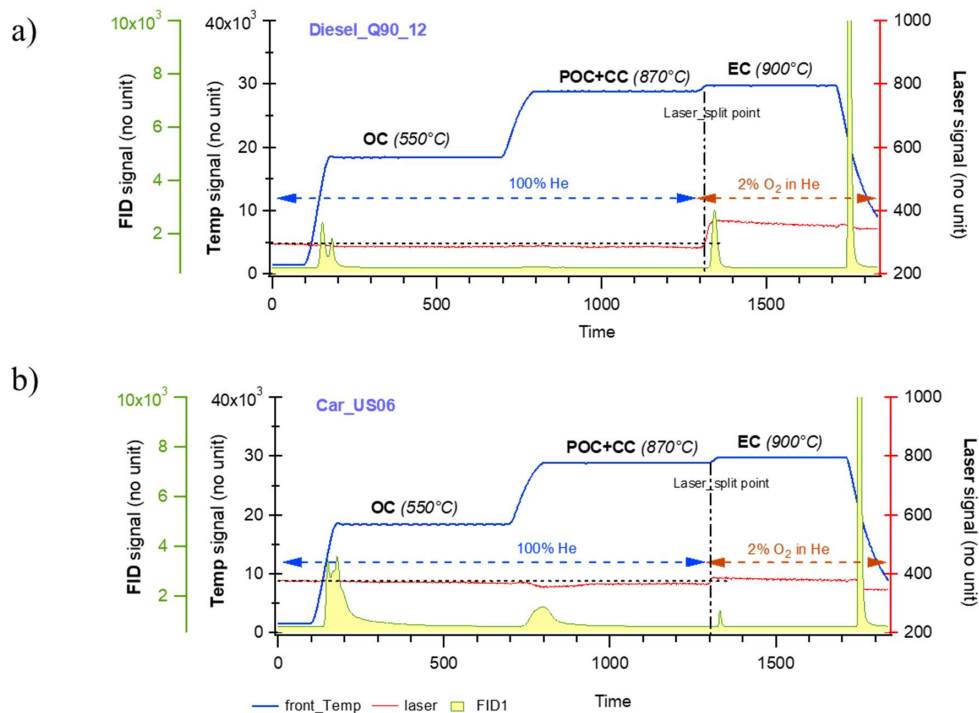
Figure 7

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Figure 8
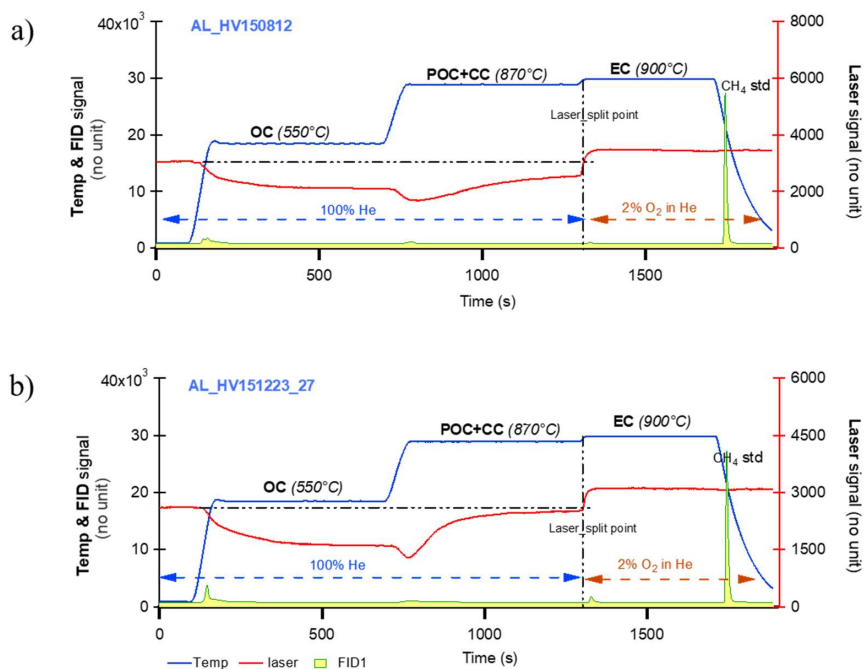
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Figure 9
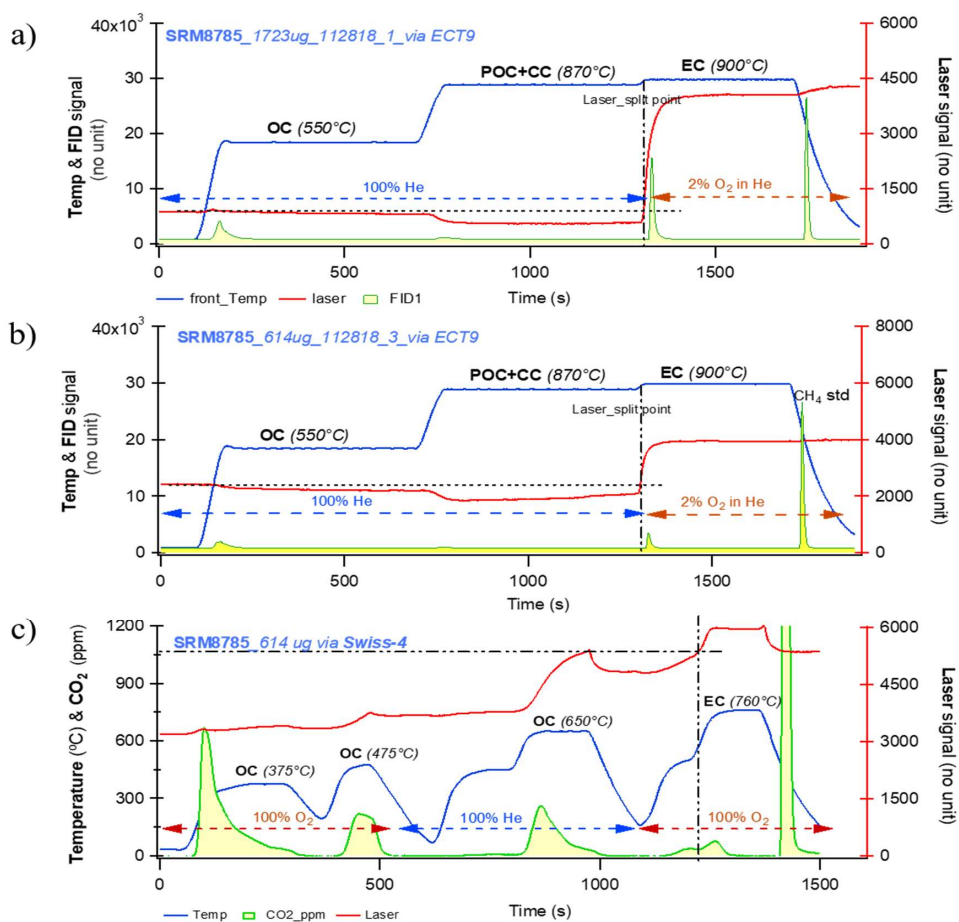
655 **References**

Andersson, A., Deng, J., Du, K., Zheng, M., Yan, C., Sköld, M. and Gustafsson, Ö.: Regionally-Varying Combustion
Sources of the January 2013 Severe Haze Events over Eastern China, Environ. Sci. Technol., 49(4), 2038–2043,
doi:10.1021/es503855e, 2015.

Barrett, T. E., Robinson, E. M., Usenko, S. and Sheesley, R. J.: Source Contributions to Wintertime Elemental and
660   Organic Carbon in the Western Arctic Based on Radiocarbon and Tracer Apportionment, Environ. Sci. Technol.,
49(19), 11631–11639, doi:10.1021/acs.est.5b03081, 2015.

Beverly, R. K., Beaumont, W., Tauz, D., Ormsby, K. M., Reden, K. F. Von, Santos, G. M. and Southon, J. R.: The
Keck Carbon Cycle AMS laboraoty, University of California, Irvine: Status report, Radiocarbon, 52(2), 301–309,
2010.

665 Birch, M. E.: Applied Occupational and Environmental Hygiene Occupational Monitoring of Particulate Diesel Exhaust
by NIOSH Method 5040, Appl. Occup. Environ. Hyg., 17(6), 400–405, doi:10.1080/10473220290035390 To, 2002.

Bond, T. C., Doherty, S. J., Fahey, D. W., Forster, P. M., Berntsen, T., DeAngelo, B. J., Flanner, M. G., Ghan, S.,
Kärcher, B., Koch, D., Kinne, S., Kondo, Y., Quinn, P. K., Sarofim, M. C., Schultz, M. G., Schulz, M.,
Venkataraman, C., Zhang, H., Zhang, S., Bellouin, N., Guttikunda, S. K., Hopke, P. K., Jacobson, M. Z., Kaiser, J.
670   W., Klimont, Z., Lohmann, U., Schwarz, J. P., Shindell, D., Storelvmo, T., Warren, S. G. and Zender, C. S.:
Bounding the role of black carbon in the climate system: A scientific assessment, J. Geophys. Res. Atmos., 118(11),
5380–5552, 2013.

Cavalli, F., Viana, M., Yttri, K. E., Genberg, J. and Putaud, J.: Toward a standardised thermal-optical protocol for
measuring atmospheric organic and elemental carbon : the EUSAAR protocol, , 79–89, 2010.

675 Chan, T. W., Huang, L., Leaitch, W. R., Sharma, S., Brook, J. R., Slowik, J. G., Abbatt, J. P. D., Brickell, P. C., Liggio,
J., Li, S. M. and Moosmüller, H.: Observations of OM/OC and specific attenuation coefficients (SAC) in ambient
fine PM at a rural site in central Ontario, Canada, Atmos. Chem. Phys., 10(5), 2393–2411, doi:10.5194/acp-10-2393-
2010, 2010.

Chan, T., Meloche, E., Kubsh, J., Brezny, R. et al.: Impact of Ambient Temperature on Gaseous and Particle Emissions
680   from a Direct Injection Gasoline Vehicle and its Implications on Particle Filtration, *SAE Int. J. Fuels Lubr.* 6(2):350-
371, doi.org/10.4271/2013-01-0527, 2013.

Chan, T. W., Huang, L., Banwait, K., Zhang, W., Ernst, D., Wang, X., John, G., Chow, J. C., Green, M., Czimczik, C.
I., Santos, G. M. and Sharma, S.: Inter-comparison of the Elemental and Organic Carbon Mass Measurements from
Three North American National Long-term Monitoring Networks, Atmos. Meas. Tech., 12, 4543–4560, 2019 ,
685   https://doi.org/10.5194/amt-12-4543-2019.

Chow, J. C., Watson, J. G., Crow, D., Lowenthal, D. H. and Merrifield, T.: Comparison of IMPROVE and NIOSH
Carbon Measurements, Aerosol Sci. Technol., 34(1), 23–34, 2001.

Chow, J. C., Watson, J. G., Chen, L. W. A., Arnott, W. P., Moosmüller, H. and Fung, K.: Equivalence of elemental
carbon by thermal/optical reflectance and transmittance with different temperature protocols, Environ. Sci. Technol.,
690   38(16), 4414–4422, doi:10.1021/es034936u, 2004.

Cohen, A. J., Brauer, M., Burnett, R., Anderson, H. R., Frostad, J., Estep, K., Balakrishnan, K., Brunekreef, B.,
Dandona, L., Dandona, R., Feigin, V., Freedman, G., Hubbell, B., Jobling, A., Kan, H., Knibbs, L., Liu, Y., Martin,
R., Morawska, L., Pope, C. A., Shin, H., Straif, K., Shaddick, G., Thomas, M., van Dingenen, R., van Donkelaar, A.,
Vos, T., Murray, C. J. L. and Forouzanfar, M. H.: Estimates and 25-year trends of the global burden of disease
695   attributable to ambient air pollution: an analysis of data from the Global Burden of Diseases Study 2015, Lancet,
389(10082), 1907–1918, doi:10.1016/S0140-6736(17)30505-6, 2017.

Currie, L. A., Benner, B. A. J., Kessler, J. D., Klinedinst, D. B., Klouda, G. A., Marolf, J. V., Slater, J. F., Wise, S. A.,
Cachier, H., Cary, R., Chow, J. C., Watson, J., Druffel, E. R. M., Masiello, C. A., Eglinton, T. I., Pearson, A., Reddy,
C. M., Gustafsson, Ö., Quinn, J. G., Hartmann, P. C., Hedges, J. I., Prentice, K. M., Kirchstetter, T. W., Novakow, T.,
700   Puxbaum, H. and Schmid, H.: A Critical Evaluation of Interlaboratory Data on Total, Elemental, and Isotopic Carbon
in the Carbonaceous Particle Reference Material, NIST SRM 1649a, J. Res. Natl. Inst. Stand. Technol., 107(3), 279–
298, 2002.

Després, V. R., Alex Huffman, J., Burrows, S. M., Hoose, C., Safatov, A. S., Buryak, G., Fröhlich-Nowoisky, J., Elbert,
W., Andreae, M. O., Pöschl, U. and Jaenicke, R.: Primary biological aerosol particles in the atmosphere: A review,
705   Tellus, Ser. B Chem. Phys. Meteorol., 64(1), 15598, doi:10.3402/tellusb.v64i0.15598, 2012.

Eckhardt, S., B. Quennehen,a, D. J. L. Olivié, T. K. Berntsen, R. Cherian, J. H. Christensen, W. Collins, S. Crepinsek,
N. Daskalakis, M. Flanner, A. Herber, C. Heyes, Ø. Hodnebrog, L. Huang, M. Kanakidou, Z. Klimont, J. Langner, K.
S. Law, M. T. Lund, R. Mahmood, A. Massling, S. Myriokefalitakis, I. E. Nielsen, J. K. Nøjgaard, J. Quaas, P. K.
Quinn, J.-C. Raut, S. T. Rumbold, M. Schulz, S. Sharma, R. B. Skeie, H. Skov, T. Uttal, K. von Salzen, and A. Stohl.
710   Current model capabilities for simulating black carbon and sulfate concentrations in the Arctic atmosphere: a multi-
model evaluation using a comprehensive measurement data set, Atmos. Chem. Phys., 15, 9413–9433, 2015
www.atmos-chem-phys.net/15/9413/2015/doi:10.5194/acp-15-9413-2015

Fuzzi, S., Baltensperger, U., Carslaw, K., Decesari, S., Denier Van Der Gon, H., Facchini, M. C., Fowler, D., Koren, I.,
Langford, B., Lohmann, U., Nemitz, E., Pandis, S., Riipinen, I., Rudich, Y., Schaap, M., Slowik, J. G., Spracklen, D.
715   V., Vignati, E., Wild, M., Williams, M. and Gilardoni, S.: Particulate matter, air quality and climate: Lessons learned
and future needs, Atmos. Chem. Phys., 15(14), 8217–8299, doi:10.5194/acp-15-8217-2015, 2015.

Grahame, T. J., Klemm, R., Schlesinger, R. B., Gwen Eklund, A., Chow, J. C., Greenbaum, D. S., Hidy, G. M.,
Kleinman, M. T., Watson, J. G., Wyzga, R. E., Grahame, T. J., Klemm, R. and Schlesinger, R. B.: Public health and
components of particulate matter: The changing assessment of black carbon, J. Air Waste Manag. Assoc., 64(11),
720   1221–1231, doi:10.1080/10962247.2014.960218, 2014.

Graven, H. D.: Impact of fossil fuel emissions on atmospheric radiocarbon and various applications of radiocarbon over
this century, Proc. Natl. Acad. Sci., 112(31), 9542–9545, doi:10.1073/pnas.1504467112, 2015.

Hallquist, M., Wenger, J. C., Baltensperger, U., Rudich, Y., Simpson, D., Claeys, M., Dommen, J., Donahue, N. M.,
George, C., Goldstein, A. H., Hamilton, J. F., Herrmann, H., Hoffmann, T., Iinuma, Y., Jang, M., Jenkin, M. E.,

725   Jimenez, J. L., Kiendler-Scharr, A., Maenhaut, W., McFiggans, G., Mentel, T. F., Monod, A., Prévôt, A. S. H.,
Seinfeld, J. H., Surratt, J. D., Szmigielski, R. and Wildt, J.: The formation, properties and impact of secondary
organic aerosol: current and emerging issues, Atmos. Chem. Phys., 9, 5155–5236, 2009.

Hand, J. L., B. A. Schichtel, W. C. Malm, and N. H. Frank, Spatial and Temporal Trends in PM2.5 Organic and
Elemental Carbon across the United States, Advances in Meteorology

730 Volume 2013, Article ID 367674, 13 pages http://dx.doi.org/10.1155/2013/367674.

Hammes, K., Smernik, R. J., Skjemstad, J. O., Herzog, A., Vogt, U. F. and Schmidt, M. W. I.: Synthesis and
characterisation of laboratory-charred grass straw (Oryza sativa) and chestnut wood (Castanea sativa) as reference
materials for black carbon quantification, Org. Geochem., 37(11), 1629–1633,
doi:10.1016/j.orggeochem.2006.07.003, 2006.

735 Hammes, K., Schmidt, M. W. I., Smernik, R. J., Currie, L. A., Ball, W. P., Nguyen, T. H., Louchouarn, P., Houel,
S.Elmquist, M., Cornelissen, G., Skjemstad, J. O., Dunn, J. C., Hatcher, P. G., Hockaday, W. C., Smith, D. M.,
Hartkopf-fro, C., Bo, A., Gschwend, P. M., Flores-cervantes, D. X., Largeau, C., Rumpel, C., Guggenberger, G.,
Kaiser, K., Rosa, M. De, Manning, D. A. C. and Lo, E.: Comparison of quantification methods to measure fire-
derived (black / elemental) carbon in soils and sediments using reference materials from soil , water , sediment and

740   the atmosphere, 21, doi:10.1029/2006GB002914, 2007.

Hammes, K., Smernik, R. J., Skjemstad, J. O. and Schmidt, M. W. I.: Characterisation and evaluation of reference
materials for black carbon analysis using elemental composition , colour , BET surface area and C NMR
spectroscopy, Appl. Geochemistry, 23, 2113–2122, doi:10.1016/j.apgeochem.2008.04.023, 2008.

Heal, M. R.: The application of carbon-14 analyses to the source apportionment of atmospheric carbonaceous

745   particulate matter: A review, Anal. Bioanal. Chem., 406(1), 81–98, doi:10.1007/s00216-013-7404-1, 2014.

Huang, L., The issue of harmonizing the methodologies for emission inventories of GHGs with those of SLCFs,
presentation at the IPCC Expert Meeting on SLCFs, Geneva, May, 2018 (https://www.ipcc-
nggip.iges.or.jp/public/mtdocs/1805_Geneva.html)

Huang, L., Brook, J. R., Zhang, W., Li, S. M., Graham, L., Ernst, D., Chivulescu, A. and Lu, G.: Stable isotope

750   measurements of carbon fractions (OC/EC) in airborne particulate: A new dimension for source characterization and
apportionment, Atmos. Environ., 40(15), 2690–2705, doi:10.1016/j.atmosenv.2005.11.062, 2006.

Huang, L., Gong, S. L., Sharma, S., Lavoué, D. and Jia, C. Q.: A trajectory analysis of atmospheric transport of black
carbon aerosols to Canadian high Arctic in winter and spring (1990–2005), Atmos. Chem. Phys., 10(11), 5065–5073,
doi:10.5194/acp-10-5065-2010, 2010.

755 Irei, S., Laboratory study of stable carbon isotope ratio of secondary particulate organic matter in the gas-phase, PhD
thesis, York University, Sept. 2008.

Janssen, N. A., Gerlofs-Nijland, M. E., Lanki, T., Salonen, R. O., Cassee, F., Hoek, G., Fischer, P., Brunekreef, B. and Krzyzanowski, M.: Health effects of black carbon, WHO Regional Office for Europe, Copenhagen, Denmark., 2012.

Jimenez, J. L., Canagaratna, M. R., Donahue, N. M., Prevot, a. S. H., Zhang, Q., Kroll, J. H., DeCarlo, P. F., Allan, J.
760   D., Coe, H., Ng, N. L., Aiken, a. C., Docherty, K. S., Ulbrich, I. M., Grieshop, A. P., Robinson, a. L., Duplissy, J., Smith, J. D., Wilson, K. R., Lanz, V. a., Hueglin, C., Sun, Y. L., Tian, J., Laaksonen, A., Raatikainen, T., Rautiainen, J., Vaattovaara, P., Ehn, M., Kulmala, M., Tomlinson, J. M., Collins, D. R., Cubison, M. J., Dunlea, E. J., Huffman, J. A., Onasch, T. B., Alfarra, M. R., Williams, P. I., Bower, K., Kondo, Y., Schneider, J., Drewnick, F., Borrmann, S., Weimer, S., Demerjian, K., Salcedo, D., Cottrell, L., Griffin, R., Takami, A., Miyoshi, T., Hatakeyama, S., Shimono,
765   A., Sun, J. Y., Zhang, Y. M., Dzepina, K., Kimmel, J. R., Sueper, D., Jayne, J. T., Herndon, S. C., Trimborn, a. M., Williams, L. R., Wood, E. C., Middlebrook, A. M., Kolb, C. E., Baltensperger, U., Worsnop, D. R., Dunlea, J., Huffman, J. A., Onasch, T. B., Alfarra, M. R., Williams, P. I., Bower, K., Kondo, Y., Schneider, J., Drewnick, F., Borrmann, S., Weimer, S., Demerjian, K., Salcedo, D., Cottrell, L., Griffin, R., Takami, A., Miyoshi, T., Hatakeyama, S., Shimono, A., Sun, J. Y., Zhang, Y. M., Dzepina, K., Kimmel, J. R., Sueper, D., Jayne, J. T.,
770   Herndon, S. C., Trimborn, a. M., Williams, L. R., Wood, E. C., Middlebrook, A. M., Kolb, C. E., Baltensperger, U., Worsnop, D. R., Dunlea, E. J., Huffman, J. A., et al.: Evolution of organic aerosols in the atmosphere, Science (80-.)., 326(5959), 1525–1529, doi:10.1126/science.1180353, 2009.

Kanakidou, M., Seinfeld, J. H., Pandis, S. N., Barnes, I., Dentener, F. J., Facchini, M. C., Van Dingenen, R., Ervens, B., Nenes, A., Nielsen, C. J., Swietlicki, E., Putaud, J. P., Balkanski, Y., Fuzzi, S., Horth, J., Moortgat, G. K.,
775   Winterhalter, R., Myhre, C. E. L., Tsigaridis, K., Vignati, E., Stephanou, E. G. and Wilson, J.: Organic aerosol and global climate modelling: a review, Atmos. Chem. Phys., 5(4), 1053–1123, doi:10.5194/acp-5-1053-2005, 2005.

Klouda, G., Filliben, J., Parish, H., Chow, J., Watson, J. and Cary, R.: Reference Material 8785: air particulate matter on filter media, Aerosol Sci. Technol., 39(2), 173–183, 2005.

Laskin, A., Laskin, J. and Nizkorodov, S. A.: Chemistry of Atmospheric Brown Carbon, Chem. Rev., 115(10), 4335–
780   4382, doi:10.1021/cr5006167, 2015.

Levin, I., Naegler, T., Kromer, B., Diehl, M., Francey, R. J., Gomez-Pelaez, A. J., Steele, L. P., Wagenbach, D., Weller, R., Worthy, D. E. and Deihl, M.: Observations and modelling of the global distribution and long-term trend of atmospheric 14CO2, Tellus B, 62, 26–46, 2010.

Meredith, W., P. L. Ascough, M. I. Bird, D. J. Large, C.E. Snape, Y. Sun, E. L. Tilston, Assessment of hydrprolysis as
785   a method for the qualification of black carbon using standard reference materials, Geochem. Cosmochim. Acta 97, 131-147, 2012.

Mouteva, G. O., Fahrni, S. M., Santos, G. M., Randerson, J. T., Zhang, Y.-L., Szidat, S. and Czimczik, C. I.: Accuracy and precision of ^{14}C-based source apportionment of organic and elemental carbon in aerosols using the Swiss_4S protocol, Atmos. Meas. Tech., 8(9), 3729–3743, doi:10.5194/amt-8-3729-2015, 2015a.

790 Mouteva, G. O., Czimczik, C. I., Fahrni, S. M., Wiggins, E. B., Rogers, B. M., Veraverbeke, S., Xu, X., Santos, G. M., Henderson, J., Miller, C. E. and Randerson, J. T.: Black carbon aerosol dynamics and isotopic composition in Alaska

Atmospheric
Measurement
Techniques
Discussions

linked with boreal fire emissions and depth of burn in organic soils, Global Biogeochem. Cycles, 29(11), 1977–2000, doi:10.1002/2015GB005247, 2015b.

Pöschl, U.: Atmospheric aerosols: Composition, transformation, climate and health effects, Angew. Chemie - Int. Ed., 795   44(46), 7520–7540, doi:10.1002/anie.200501122, 2005.

Putaud, J. P., Van Dingenen, R., Alastuey, A., Bauer, H., Birmili, W., Cyrys, J., Flentje, H., Fuzzi, S., Gehrig, R., Hansson, H. C., Harrison, R. M., Herrmann, H., Hitzenberger, R., Hüglin, C., Jones, A. M., Kasper-Giebl, A., Kiss, G., Kousa, A., Kuhlbusch, T. A. J., Löschau, G., Maenhaut, W., Molnar, A., Moreno, T., Pekkanen, J., Perrino, C., Pitz, M., Puxbaum, H., Querol, X., Rodriguez, S., Salma, I., Schwarz, J., Smolik, J., Schneider, J., Spindler, G., ten 800   Brink, H., Tursic, J., Viana, M., Wiedensohler, A. and Raes, F.: A European aerosol phenomenology - 3: Physical and chemical characteristics of particulate matter from 60 rural, urban, and kerbside sites across Europe, Atmos. Environ., 44(10), 1308–1320, doi:10.1016/j.atmosenv.2009.12.011, 2010.

Ridley, D.A., C.L. Heald, K.J. Ridley and J. H. Kroll, Cause and consequences of decreasing atmeopheric organic aerosol in the United States, Proc Natl Acad Sci USA, January 9, 2018 115 (2) 290-295, 805   https://doi.org/10.1073/pnas.1700387115

Leaitch,W.R., L. M. Russell, J. Liu, F. Kolonjari, D. Toom, L. Huang, S. Sharma, . Chivulescu, D. Veber, W. Zhang, Organic Functional Groups in the Submicron Aerosol at 82.5°N from 2012 to 2014, Atmos. Chem. Phys., 18, 3269–3287, 2018

Leaitch,W.R., S. Sharma, L. Huang, D. Toom-Sauntry, A. Chivulescu, A. Marie Macdonald, K. von Salzen, J. R. 810   Pierce, A. K. Bertram, J. C. Schroder, N. C. Shantz, R. Y.-W. Chang, A.-L. Norman, 2013. Dimethyl sulfide control of the clean summertime Arctic aerosol and cloud, Elementa: Science of the Anthropocene • 1: 000017 • doi: 10.12952/journal.elementa.000017.

Sharma, S., W. R. Leaitch, L. Huang, D. Veber, F. Kolonjari, W. Zhang, S. J. Hanna, A. K. Bertram, and J. A. Ogren, 2017. An evaluation of three methods for measuring black carbon in Alert, Canada, Atmos. Chem. Phys., 17, 15225– 815   15243, 2017 https://doi.org/10.5194/acp-17-15225-2017.

Santos, G. M., Moore, R. B., Southon, J. R., Griffin, S., Hinger, E. and Zhang, D.: AMS 14C Sample Preparation at the KCCAMS/UCI Facility: Status Report and Performance of Small Samples, Radiocarbon, 49(2), 255–269, doi:10.2458/azu_js_rc.49.2925, 2007a.

Santos, G. M., Southon, J. R., Griffin, S., Beaupre, S. R. and Druffel, E. R. M.: Ultra small-mass AMS 14C sample 820   preparation and analyses at KCCAMS/UCI Facility, Nucl. Instruments Methods Phys. Res. Sect. B Beam Interact. with Mater. Atoms, 259, 293–302, 2007b.

Santos, G. M., Southon, J. R., Drenzek, N. J., Ziolkowski, L. A., Druffel, E. R. M., Xu, X., Zhang, D., Trumbore, S. E., Eglinton, T. I. and Hughen, K. A.: Blank assessment for ultra-small radiocarbon samples, Radiocarbon, 52, 1322– 1335, 2010.

825 Sharma, S., Richard Leaitch, W., Huang, L., Veber, D., Kolonjari, F., Zhang, W., Hanna, S. J., Bertram, A. K. and

Ogren, J. A.: An evaluation of three methods for measuring black carbon in Alert, Canada, Atmos. Chem. Phys., 17(24), 15225–15243, doi:10.5194/acp-17-15225-2017, 2017.

Shrivastava, M., Cappa, C.D., Fan, J., Goldstein, A.H., Guenther, A.B., Jimenez, J.L., Kuang, C., Laskin, A., Martin, S.T., Ng, N.L. and Petaja, T.: Recent advances in understanding secondary organic aerosol: Implications for global climate forcing. Reviews of Geophysics, 55(2), 509-559, 2017.

Stuiver, M. and Polach, H. A.: Discussion Reporting of 14C Data, Radiocarbon, 19(03), 355–363, doi:10.1017/S0033822200003672, 1977.

Szidat, S., Jenk, T. M., G¨aggeler, H. W., Synal, H.-A., Hajdas, I., Bonani, G., and Saurer, M.: THEODORE, a two-step heating system for the EC/OC determination of radiocarbon (^{14}C) in the environment, Nucl. Instrum. Methods B, 223–224, 829–836, 2004

Szidat, S., Jenk, T. M., Synal, H.-A., Kalberer, M., Wacker, L., Hajdas, I., Kasper-Giebl, A. and Baltensperger, U.: Contributions of fossil fuel, biomass-burning, and biogenic emissions to carbonaceous aerosols in Zurich as traced by 14C, J. Geophys. Res., 111(D7), D07206, 2006.

Trumbore, S. E., Sierra, C. A. and Hicks Pries, C. E.: Radiocarbon Nomenclature, Theory, Models, and Interpretation: Measuring Age, Determining Cycling Rates, and Tracing Source Pools, in Radiocarbon and Climate Change: Mechanisms, Applications and Laboratory Techniques, edited by E. A. G. Schuur, E. Druffel, and S. E. Trumbore, pp. 45–82, Springer International Publishing, Cham., 2016.

Watson, J. G., Chen, L. A., Chang, O., Chow, J. C., Watson, J. G., Chen, L. A., Chang, M. C. O., Robinson, N. F., Trimble, D. and Kohl, S.: The IMPROVE _ A Temperature Protocol for Thermal / Optical Carbon Analysis : Maintaining Consistency with a Long-Term Database, Air Waste Manag., 57, 1014–1023, doi:10.3155/1047-3289.57.9.1014, 2007.

Wex, H., Huang, L., Zhang, W., Huang, H., Traversi, R., Becagli, S., Sheesley, R.J., Moffett, C. E., Barrett, T.E., Bossi, R., Skov, H., Hunerbern, A., Lubitz, J., Loffler, M., Linke, O., Hartmann, M., Herenz, P., and Stratmann, F.: Annual variability of ice nucleating particle concentrations at different Arctic locations , Atmos. Chem. Phys., 19, 5293-5311, 2019.

Wiggins, E. B., Czimczik, C. I., Santos, G. M., Chen, Y., Xu, X., Holden, S. R., Randerson, J. T., Harvey, C. F., Kai, F. M. and Yu, L. E.: Smoke radiocarbon measurements from Indonesian fires provide evidence for burning of millennia-aged peat, Proc. Natl. Acad. Sci., 115(49), 12419–12424, doi:10.1073/pnas.1806003115, 2018.

Willis, M. D., Healy, R. M., Riemer, N., West, M., Wang, J. M., Jeong, C., Wenger, J. C., Evans, G. J., Abbatt, J. P. D. and Lee, A. K. Y.: Quantification of black carbon mixing state from traffic : implications for aerosol optical properties, , 4693–4706, doi:10.5194/acp-16-4693-2016, 2016.

Winiger, P., Andersson, A., Eckhardt, S., Stohl, A. and Gustafsson, O.: The sources of atmospheric black carbon at a European gateway to the Arctic, Nat. Commun., 7, doi:10.1038/ncomms12776, 2016.

Winiger, P., Andersson, A., Eckhardt, S., Stohl, A., Semiletov, I. P., Dudarev, O. V., Charkin, A., Shakhova, N.,

860   Klimont, Z., Heyes, C. and Gustafsson, Ö.: Siberian Arctic black carbon sources constrained by model and
       observation, Proc. Natl. Acad. Sci., 114(7), E1054–E1061, doi:10.1073/pnas.1613401114, 2017.

       Winiger, P., T. E. Barrett, R. J. Sheesley, L. Huang, S. Sharma, L. A. Barrie, K. E. Yttri, N. Evangeliou, S. Eckhardt, A.
       Stohl, Z. Klimont, C. Heyes, I. P. Semiletov, O. V. Dudarev, A. Charkin, N. Shakhova, H. Holmstrand, A.
       Andersson, Ö. Gustafsson, Source apportionment of circum-Arctic atmospheric black carbon from isotopes and
865   modeling. Sci. Adv. 2019;5: eaau8052

       Xu, J., Martin, R. V, Morrow, A., Sharma, S., Huang, L., Leaitch, W. R., Burkart, J., Schulz, H., Zanatta, M., Willis, M.
       D., Henze, D. K., Lee, C. J., Herber, A. B. and Abbatt, J. P. D.: Source attribution of Arctic black carbon constrained
       by aircraft and surface measurements, , 11971–11989, 2017.

       Xu, X., Trumbore, S. E., Zheng, S., Southon, J. R., McDuffee, K. E., Luttgen, M. and Liu, J. C.: Modifying a sealed
870   tube zinc reduction method for preparation of AMS graphite targets: Reducing background and attaining high
       precision, Nucl. Instruments Methods Phys. Res. Sect. B Beam Interact. with Mater. Atoms, 259(1), 320–329,
       doi:10.1016/j.nimb.2007.01.175, 2007.

       Yang, F., Huang, L., Duan, F., Zhang, W., He, K., Ma, Y., Brook, J. R., Tan, J., Zhao, Q. and Cheng, Y.: Carbonaceous
       species in PM2.5 at a pair of rural/urban sites in Beijing, 2005-2008, Atmos. Chem. Phys., 11(15), 7893–7903,
875   doi:10.5194/acp-11-7893-2011, 2011.

       Yang,F., J. Tan, Q. Zhao, Z. Du, K. He, Y. Ma, F. Duan, G. Chen, and Q. Zhao, Characteristics of PM2.5 speciation in
       representative megacities and across China, Atmos. Chem. Phys., Atmos. Chem. Phys., 11, 5207–5219, 2011
       www.atmos-chem-phys.net/11/5207/2011/ doi:10.5194/acp-11-5207-2011.

       Zencak, Z., Elmquist, M. and Gustafsson, Ö.: Quantification and radiocarbon source apportionment of black carbon in
880   atmospheric aerosols using the CTO-375 method, Atmos. Environ., 41, 7895–7906, 2007.

       Zhang, X., Li, J., Mo, Y., Shen, C., Ding, P., Wang, N., Zhu, S., Cheng, Z., He, J., Tian, Y., Gao, S., Zhou, Q., Tian, C.,
       Chen, Y. and Zhang, G.: Isolation and radiocarbon analysis of elemental carbon in atmospheric aerosols using
       hydropyrolysis, Atmos. Environ., 198(August 2018), 381–386, doi:10.1016/j.atmosenv.2018.11.005, 2019.

       Zhang, X. Y., Wang, Y. Q., Zhang, X. C., Guo, W. and Gong, S. L.: Carbonaceous aerosol composition over various
885   regions of China during 2006, J. Geophys. Res., 113, D14111, 2008.

       Zhang, Y. L., Perron, N., Ciobanu, V. G., Zotter, P., Minguillón, M. C., Wacker, L., Prévôt, A. S. H., Baltensperger, U.
       and Szidat, S.: On the isolation of OC and EC and the optimal strategy of radiocarbon-based source apportionment of
       carbonaceous aerosols, Atmos. Chem. Phys., 12, 10841–10856, 2012.

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Supplementary Information

Table S1. Individual measurements of OC and EC via ECT9 at ECCC for the references listed in Table 1.

Lab ID	Date	[a]Loaded mass on filter	Carbon fraction				OC$_{total}$/TC	EC/TC	TC/loaded mass
			OC	POC+CC	EC	TC			
		μg	μg/cm^2				%		
Regal Black	(n = 41)								
16-084-04	24-Mar-16	24	0.02	0.69	28.34	29.05	2	98	121
16-098-03	7-Apr-16	22	-0.05	0.48	18.63	19.06	2	98	87
16-098-04	7-Apr-16	23	0.43	0.91	23.89	25.23	5	95	110
16-097-04	6-Apr-16	19	0.44	0.48	20.02	20.94	4	96	110
16-098-06	7-Apr-16	18	0.15	0.49	19.05	19.69	3	97	109
17-052-07	21-Feb-17	21	0.17	0.50	18.42	19.09	4	96	91
17-053-03	22-Feb-17	16	0.14	0.76	13.24	14.14	6	94	88
17-240-06	28-Aug-17	18	0.27	0.59	15.12	15.98	5	95	90
17-243-03	31-Aug-17	20	0.00	0.42	20.22	20.64	2	98	104
17-243-04	31-Aug-17	24	0.14	0.20	18.79	19.13	2	98	79
15-117-07	27-Apr-15	30	0.22	0.95	27.46	28.63	4	96	95
16-094-06	3-Apr-16	32	0.80	0.76	38.23	39.79	4	96	124
16-095-04	4-Apr-16	27	0.39	0.57	26.11	27.07	4	96	100
16-099-06	8-Apr-16	27	0.03	0.87	24.68	25.58	4	96	95
16-099-07	8-Apr-16	26	0.14	0.95	25.37	26.46	4	96	102
17-052-07	21-Feb-17	25	0.12	0.92	23.47	24.51	4	96	98
15-104-08	14-Apr-15	52	0.00	0.85	47.21	48.06	2	98	92
16-095-07	4-Apr-16	47	0.30	1.18	48.19	49.67	3	97	106
16-097-05	6-Apr-16	43	0.32	1.03	39.78	41.13	3	97	96
16-098-08	7-Apr-16	50	0.12	0.67	47.38	48.17	2	98	96
17-052-05	21-Feb-17	53	0.90	1.74	44.31	46.95	6	94	89
17-052-06	21-Feb-17	42	0.22	1.37	35.51	37.10	4	96	88
17-241-07	29-Aug-17	44	0.52	1.51	38.78	40.81	5	95	93

17-241-08	29-Aug-17	49	0.80	0.89	40.80	42.49	4	96	87
17-243-06	31-Aug-17	43	0.00	0.53	38.07	38.60	1	99	91
15-117-10	27-Apr-15	71	0.50	1.59	65.55	67.64	3	97	95
16-098-05	7-Apr-16	61	0.18	1.17	64.91	66.26	2	98	109
16-099-03	8-Apr-16	71	0.00	0.56	64.60	65.16	1	99	92
16-099-04	8-Apr-16	63	0.00	1.36	54.53	55.89	2	98	89
17-052-09	21-Feb-17	83	0.83	2.08	76.60	79.51	4	96	96
17-243-05	21-Feb-17	74	0.67	1.99	63.36	66.02	4	96	89
17-243-07	31-Aug-17	68	0.00	1.14	57.82	58.96	2	98	87
17-243-09	31-Aug-17	71	0.24	1.49	60.34	62.07	3	97	88
15-117-04	27-Apr-15	134	0.00	0.61	123.52	124.13	0	100	93
16-098-07	7-Apr-16	107	0.64	0.42	99.88	100.94	1	99	94
17-240-03	28-Aug-17	95	0.85	2.30	85.17	88.32	4	96	93
17-241-02	29-Aug-17	101	0.83	2.23	88.23	91.29	3	97	90
17-241-06	29-Aug-17	93	0.43	1.24	82.44	84.11	2	98	91
17-240-05	28-Aug-17	116	0.86	2.85	103.57	107.28	3	97	92
17-243-10	31-Aug-17	123	0.11	2.06	109.73	111.90	2	98	91
17-244-02	1-Sep-17	122	0.63	2.11	108.41	111.15	2	98	91
						mean	3	97	96
						s.d.	1	1	9
C1150	(n = 24)								
06-195-07	14-Jul-06	4	0.05	0.05	3.17	3.26	3	97	81
06-195-09	14-Jul-06	7	0.23	0.00	6.35	6.57	3	97	94
06-195-10	14-Jul-06	10	0.48	0.18	8.91	9.57	7	93	96
06-198-03	17-Jul-06	18	0.12	0.09	18.18	18.39	1	99	102
06-198-04	17-Jul-06	25	0.32	0.42	23.22	23.96	3	97	96
06-198-05	17-Jul-06	42	0.02	0.32	39.75	40.08	1	99	95
06-198-06	17-Jul-06	34	0.48	0.42	32.94	33.84	3	97	100
06-198-07	17-Jul-06	15	0.26	0.41	14.45	15.11	4	96	101
13-225-03	13-Aug-13	25	0.00	0.00	20.29	20.29	0	100	80
13-225-04	13-Aug-13	89	0.28	0.00	91.34	91.62	0	100	102
13-225-05	13-Aug-13	30	0.00	0.00	27.50	27.50	0	100	93
13-225-06	13-Aug-13	46	0.00	0.00	38.35	38.35	0	100	84
13-226-03	14-Aug-13	10	0.05	0.01	7.33	7.39	1	99	78

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13-226-04	14-Aug-13	79	0.06	0.00	68.51	68.57	0	100	87
13-226-05	14-Aug-13	14	0.05	0.00	13.73	13.78	0	100	98
13-226-06	14-Aug-13	17	0.11	0.00	20.16	20.27	1	99	116
13-226-07	14-Aug-13	49	0.16	0.03	49.37	49.56	0	100	101
15-122-09	2-May-15	72	0.05	0.00	69.62	69.67	0	100	97
15-122-07	2-May-15	71	0.19	0.00	70.79	70.98	0	100	100
15-122-08	2-May-15	104	0.00	0.00	97.78	97.78	0	100	94
15-123-03	3-May-15	22	0.13	0.00	29.38	29.51	0	100	134
15-123-04	3-May-15	71	0.00	0.00	76.06	76.06	0	100	107
15-123-05	3-May-15	27	0.32	0.00	25.96	26.28	1	99	97
15-123-06	3-May-15	59	0.11	0.15	66.40	66.66	0	99	113
						mean	1	99	98
						s.d.	2	2	12
Sucrose	(n = 117)								
13-332-02	28-Nov-13	20	19.76	0.35	0.00	20.11	100	0	101
13-332-03	28-Nov-13	20	19.77	0.48	0.02	20.27	100	0	101
13-333-02	28-Nov-13	20	19.46	0.44	0.00	19.90	100	0	100
13-332-08	28-Nov-13	40	37.50	1.00	0.00	38.50	100	0	96
13-332-10	28-Nov-13	40	38.77	0.98	0.00	39.75	100	0	99
13-333-03	29-Nov-13	40	39.51	1.11	0.01	40.63	100	0	102
13-333-05	29-Nov-13	80	75.63	1.73	0.22	77.58	100	0	97
13-333-08	29-Nov-13	80	74.25	2.14	0.07	76.46	100	0	96
13-333-07	29-Nov-13	80	76.43	2.05	0.07	78.55	100	0	98
14-129-02	9-May-14	20	19.39	0.29	0.06	19.74	100	0	99
14-129-03	9-May-14	20	19.33	0.16	0.05	19.54	100	0	98
14-132-02	12-May-14	20	19.71	0.00	0.00	19.71	100	0	99
14-133-03	13-May-14	40	39.16	0.66	0.60	40.42	99	1	101
14-133-04	13-May-14	40	39.67	0.53	0.10	40.30	100	0	101
14-134-02	14-May-14	40	39.44	0.31	0.11	39.86	100	0	100
14-134-03	14-May-14	80	80.11	0.80	0.10	81.01	100	0	101
14-134-04	14-May-14	80	79.39	1.01	0.36	80.76	100	0	101
14-134-05	14-May-14	80	78.49	1.86	1.46	81.81	98	2	102
14-231-02	19-Aug-14	20	19.03	0.28	0.12	19.43	99	1	97
14-234-02	22-Aug-14	20	19.20	0.50	0.13	19.83	99	1	99

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14-235-02	23-Aug-14	20	19.06	0.55	0.00	19.61	100	0	98
14-233-05	21-Aug-14	40	38.76	0.99	0.20	39.95	99	1	100
14-233-06	21-Aug-14	40	38.22	0.00	0.00	38.22	100	0	96
14-233-07	21-Aug-14	40	38.32	0.04	0.00	38.36	100	0	96
14-235-08	23-Aug-14	80	78.25	1.44	0.18	79.87	100	0	100
14-235-09	23-Aug-14	80	79.46	0.27	0.00	79.73	100	0	100
14-238-04	26-Aug-14	80	76.15	1.47	0.38	78.00	100	0	98
15-015-03	15-Jan-15	20	18.67	1.22	0.10	19.99	99	1	100
15-015-04	15-Jan-15	20	18.65	1.51	0.18	20.34	99	1	102
15-019-02	19-Jan-15	20	18.95	1.01	0.01	19.97	100	0	100
15-019-03	19-Jan-15	40	35.12	2.62	1.07	38.81	97	3	97
15-020-02	20-Jan-15	40	36.63	1.84	0.17	38.64	100	0	97
15-020-05	20-Jan-15	40	37.43	2.43	0.29	40.15	99	1	100
15-020-06	20-Jan-15	80	75.34	3.27	0.87	79.48	99	1	99
15-020-07	20-Jan-15	80	76.30	3.42	0.92	80.64	99	1	101
15-020-08	20-Jan-15	80	76.65	2.85	0.72	80.22	99	1	100
15-097-03	10-Apr-15	20	19.79	0.41	0.00	20.20	100	0	101
15-114-02	27-Apr-15	20	17.15	2.41	0.12	19.68	99	1	98
15-108-02	21-Apr-15	20	18.62	1.28	0.00	19.90	100	0	100
15-097-04	10-Apr-15	40	39.35	0.85	0.02	40.22	100	0	101
15-097-05	10-Apr-15	40	38.90	1.80	1.02	41.72	98	2	104
15-097-06	10-Apr-15	40	38.59	1.75	0.88	41.22	98	2	103
15-108-04	21-Apr-15	80	76.10	4.20	0.23	80.53	100	0	101
15-108-03	21-Apr-15	80	76.47	4.13	0.31	80.91	100	0	101
15-108-06	21-Apr-15	80	74.94	4.89	0.70	80.53	99	1	101
15-280-03	8-Oct-15	20	17.56	2.64	0.04	20.24	100	0	101
15-280-04	8-Oct-15	20	17.34	2.95	0.05	20.34	100	0	102
15-280-05	8-Oct-15	20	16.99	3.00	0.00	19.99	100	0	100
15-287-02	14-Oct-15	40	34.13	4.64	0.13	38.90	100	0	97
15-287-04	14-Oct-15	40	34.72	4.81	0.15	39.68	100	0	99
15-288-03	15-Oct-15	40	33.67	4.98	0.17	38.82	100	0	97
15-292-03	19-Oct-15	80	70.58	6.94	1.31	78.83	98	2	99
15-292-04	19-Oct-15	80	69.29	7.36	1.53	78.18	98	2	98
15-292-05	19-Oct-15	80	69.29	7.23	1.47	77.99	98	2	97

16-026-03	26-Jan-16	20	17.74	2.70	0.02	20.46	100	0	102
16-026-05	26-Jan-16	20	16.85	3.37	0.12	20.34	99	1	102
16-027-05	27-Jan-16	20	16.68	3.24	0.10	20.02	100	0	100
16-026-06	26-Jan-16	40	34.15	4.79	0.18	39.12	100	0	98
16-027-04	27-Jan-16	40	33.69	4.98	0.51	39.18	99	1	98
16-027-06	27-Jan-16	40	33.14	5.39	0.75	39.28	98	2	98
16-027-07	27-Jan-16	80	69.99	7.15	2.28	79.42	97	3	99
16-028-03	28-Jan-16	80	71.40	7.34	1.98	80.72	98	2	101
16-028-04	28-Jan-16	80	71.87	7.06	1.91	80.84	98	2	101
16-243-03	30-Aug-16	20	16.69	3.24	0.65	20.58	97	3	103
16-243-04	30-Aug-16	20	17.35	3.35	0.07	20.77	100	0	104
16-244-02	31-Aug-16	20	16.80	2.92	0.85	20.57	96	4	103
16-244-05	31-Aug-16	40	35.61	3.87	1.26	40.74	97	3	102
16-244-06	31-Aug-16	40	35.76	3.87	1.29	40.92	97	3	102
16-244-07	31-Aug-16	40	35.81	4.20	1.85	41.86	96	4	105
16-250-02	6-Sep-16	80	77.54	3.94	1.34	82.82	98	2	104
16-250-03	6-Sep-16	80	77.77	3.81	1.26	82.84	98	2	104
16-250-04	6-Sep-16	80	77.95	3.81	1.25	83.01	98	2	104
17-038-04	7-Feb-17	20	14.57	4.14	0.93	19.64	95	5	98
17-039-02	8-Feb-17	20	14.99	3.88	0.84	19.71	96	4	99
17-039-03	8-Feb-17	20	14.74	4.31	0.79	19.84	96	4	99
17-039-04	8-Feb-17	40	32.68	5.47	1.20	39.35	97	3	98
17-039-05	8-Feb-17	40	34.09	5.70	1.00	40.79	98	2	102
17-039-06	8-Feb-17	40	33.22	5.89	2.47	41.58	94	6	104
17-041-02	10-Feb-17	80	74.47	7.17	1.59	83.23	98	2	104
17-041-03	10-Feb-17	80	73.71	5.02	1.61	80.34	98	2	100
17-041-05	10-Feb-17	80	70.96	8.04	2.31	81.31	97	3	102
18-037-03	6-Feb-18	20	20.31	0.00	0.21	20.52	99	1	103
18-032-04	1-Feb-18	20	20.06	0.00	0.11	20.17	99	1	101
18-036-03	5-Feb-18	20	20.01	0.00	0.16	20.17	99	1	101
18-033-06	2-Feb-18	40	37.87	1.66	1.71	41.24	96	4	103
18-037-04	6-Feb-18	40	39.36	1.17	1.39	41.92	97	3	105
18-037-08	6-Feb-18	40	39.02	1.30	1.58	41.90	96	4	105
18-037-09	6-Feb-18	80	73.37	2.92	2.15	78.44	97	3	98

18-037-10	6-Feb-18	80	74.15	3.26	2.33	79.74	97	3	100
18-037-11	6-Feb-18	80	73.89	2.90	2.07	78.86	97	3	99
18-129-02	9-May-18	20	19.45	0.37	0.03	19.85	100	0	99
18-129-03	9-May-18	20	19.81	0.35	0.07	20.23	100	0	101
18-129-04	9-May-18	20	20.06	0.59	0.14	20.79	99	1	104
18-129-05	9-May-18	40	38.26	1.72	1.19	41.17	97	3	103
18-129-06	9-May-18	40	40.03	1.37	0.79	42.19	98	2	105
18-129-07	9-May-18	40	38.42	1.79	1.21	41.42	97	3	104
18-130-02	10-May-18	80	80.93	0.95	0.58	82.46	99	1	103
18-130-03	10-May-18	80	81.34	1.02	0.28	82.64	100	0	103
18-131-08	11-May-18	80	81.52	1.86	0.72	84.10	99	1	105
18-297-02	24-Oct-18	20	19.44	0.45	0.03	19.92	100	0	100
18-302-03	29-Oct-18	20	19.09	0.84	0.36	20.29	98	2	101
18-298-06	25-Oct-18	20	19.17	0.71	0.13	20.01	99	1	100
18-302-06	29-Oct-18	40	39.37	0.89	0.17	40.43	100	0	101
18-298-07	25-Oct-18	40	39.68	0.42	0.29	40.39	99	1	101
18-309-08	5-Nov-18	40	41.22	0.25	0.00	41.47	100	0	104
18-309-11	5-Nov-18	80	78.46	0.81	0.20	79.47	100	0	99
18-309-14	5-Nov-18	80	78.26	1.56	0.15	79.97	100	0	100
18-310-03	6-Nov-18	80	82.01	1.75	0.50	84.26	99	1	105
18-355-02	21-Dec-18	20	22.04	0.99	0.24	23.27	99	1	116
18-355-03	21-Dec-18	20	21.48	1.11	0.15	22.74	99	1	114
18-355-04	21-Dec-18	20	21.17	1.31	0.26	22.74	99	1	114
18-361-04	27-Dec-18	40	42.02	1.29	0.13	43.44	100	0	109
18-361-05	27-Dec-18	40	41.56	0.93	0.00	42.49	100	0	106
18-361-06	27-Dec-18	40	41.06	1.85	0.23	43.14	99	1	108
18-361-07	27-Dec-18	80	85.76	2.27	0.75	88.78	99	1	111
18-361-08	27-Dec-18	80	86.49	2.48	0.78	89.75	99	1	112
18-361-09	27-Dec-18	80	85.98	2.63	0.61	89.22	99	1	112
						mean	99	1	101
						s.d.	1	1	4
Adipic Acid	(n = 5)								
15-062-06	3-Mar-15	34	13.67	0.09	0.00	13.76	100	0	40
15-062-05	3-Mar-15	102	47.47	0.00	0.00	47.47	100	0	47

15-100-02	13-Apr-15	n/a	5.25	0.00	0.05	5.30	99	1	n/a
19-137-05	17-May-19	253	120.68	1.05	0.07	121.80	100	0	48
19-137-06	17-May-19	28	10.62	0.00	0.00	10.62	100	0	38
						mean	100	0	43
						s.d.	0	0	5
Rice Char	(n = 6)								
18-158-05	7-Jun-18	112	4.27	3.25	49.54	57.06	13	87	51
18-164-05	13-Jun-18	212	8.73	6.22	96.87	111.82	13	87	53
18-165-06	14-Jun-18	79	2.96	2.86	35.46	41.28	14	86	52
18-169-04	18-Jun-18	71	2.76	2.70	30.18	35.64	15	85	51
18-172-05	21-Jun-18	150	5.74	4.28	70.40	80.42	12	88	54
18-176-06	25-Jun-18	121	4.83	4.67	56.43	65.93	14	86	54
						mean	14	86	52
						s.d.	1	1	1
SRM-1649a	(n = 6)								
04-271-04	27-Sep-04	690	29.94	9.65	36.46	76.05	52.1	47.9	16.5
04-322-10	17-Nov-04	490	25.82	7.18	30.41	63.41	52.0	48.0	19.4
04-322-12	17-Nov-04	880	40.28	11.25	47.71	99.24	51.9	48.1	16.9
05-046-02	15-Feb-05	1101	51.66	16.59	67.16	135.41	50.4	49.6	18.5
05-046-03	15-Feb-05	441	21.06	6.41	25.35	52.82	52.0	48.0	18.0
05-046-04	15-Feb-05	855	40.33	12.37	51.22	103.92	50.7	49.3	18.2
						mean	51.5	48.5	17.9
						s.d.	0.8	0.8	1.1

895 [a]Loaded mass are the weighed mass (for Regal black, C1150, Adipic acid, Rice char and SRM-1649a) or injected mass (sucrose) on the filter.

Table S2. Radiocarbon content of bulk reference materials, expressed as fraction modern carbon (FM) with and without background correction. CO_2 isolation and $^{14}C/^{12}C$ analysis were carried out at KCCAMS, UCI (the method is described in Table 2).

UCI AMS #	Size	Corrected FM		Uncorrected FM	
	µg C		±		±
Sucrose					
150230	735	1.0597	0.0021	1.0597	0.0021
150231	769	1.0575	0.0017	1.0574	0.0017
AdipicAcid					
123428	876	0.0002	0.0005	0.0020	0.0001
123430	851	0.0001	0.0005	0.0019	0.0001
123431	934	-0.0001	0.0005	0.0016	0.0001
123432	1053	-0.0003	0.0005	0.0015	0.0001
123433	740	-0.0001	0.0005	0.0016	0.0001
Regal Black					
150228	717	0.0004	0.0005	0.0019	0.0001
150229	752	-0.0005	0.0005	0.0011	0.0000
C1150					
150232	88	0.0026	0.0005	0.0042	0.0001
150233	64	0.0035	0.0005	0.0050	0.0002
150234	560	0.0019	0.0005	0.0035	0.0001
RiceChar					
123434	924	1.0683	0.0023	1.0683	0.0023
123435	913	1.0670	0.0018	1.0670	0.0018
123436	961	1.0673	0.0019	1.0672	0.0019

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Table S3. Stable isotopic composition ($^{13}C/^{12}C$) of OC and EC fractions or bulk materials. CO_2 isolation and $^{13}C/^{12}C$ analysis were carried out at the CA
925   lab, CRD, ASTD/ECCC (the method is described in Table 2).

Reference m	Lab ID	Date	Fraction	Loaded mass on filter	$\delta^{13}C_{VPDB}$
				µg or µg C[a]	‰
Regal Black	16-036-04	5-Feb-16	EC	16	-27.67
(n = 5)	16-036-05	5-Feb-16	EC	27	-27.49
	16-036-06	5-Feb-16	EC	22	-27.67
	16-036-08	5-Feb-16	EC	59	-27.62
	16-036-09	5-Feb-16	EC	68	-27.57
				mean	**-27.61**
				s.d.	**0.08**
C1150	13-013-05	13-Jan-13	EC	50	-23.01
(n = 5)	13-013-07	13-Jan-13	EC	22	-23.16
	13-013-08	13-Jan-13	EC	48	-22.96
	16-036-06	5-Feb-16	EC	30	-23.14
	16-036-07	5-Feb-16	EC	46	-23.05
				mean	**-23.06**
				s.d.	**0.08**
Sucrose[b]	15-146-07	26-May-15	OC	20	-12.08
(n = 9)	15-148-03	27-May-15	OC	20	-12.40
	15-148-04	27-May-15	OC	20	-12.31
		5-Oct-17	OC	20	-12.44
		18-Apr-18	OC	20	-12.04
		18-Apr-18	OC	20	-12.30
		26-Feb-19	OC	20	-12.21
		26-Feb-19	OC	20	-12.16
		26-Feb-19	OC	20	-12.04
				mean	**-12.22**
				s.d.	**0.15**
Rice Char	04-328-06	23-Nov-04	OC	n/m	-24.42
(n = 1)	04-328-07	23-Nov-04	POC	n/m	-26.67
	04-328-05	23-Nov-04	EC	n/m	-26.94

		fraction weighted	TC	160	**-26.74**
SRM-1649a					
(n = 2)	04-330-03	25-Nov-04	OC	n/m	-26.38
	04-338-08	3-Dec-04	OC	n/m	-26.29
	04-330-05	25-Nov-04	POC	n/m	-25.51
	04-338-07	3-Dec-04	POC	n/m	-25.66
	04-330-06	25-Nov-04	EC	n/m	-25.56
	04-338-09	3-Dec-04	EC	n/m	-25.43
		fraction[c] weighted	TC	~ 600	**-25.84 ± 0.07**

[a]Sucrose was loaded as a solution (μg C), Regal Black, C1150, Rice char, and SRM-1649a as a powder (μg dry mass); [b]$\delta^{13}C_{VPDB}$ of bulk material (sucrose) via off-line method: -12.0 ± 0.2‰ (Satoshi, 2008); [c]Mean fraction (of two measurements) weighted isotopic composition of TC; n/m = not measured.

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Table S4. Stable isotopic compositions of $^{13}C/^{12}C$ in OC and EC fractions from mixtures of reference materials. OC and EC fractions were isolated with the ECT9 protocol (Huang et al., 2006), purified in a vacuum system and analyzed on a MAT253 (Huang et al., 2013) at the CAIR lab, CRD, ASTD/ECCC.

Reference mate	Lab ID	ate	Initial mass		Measured fraction	$\delta^{13}C_{VPDB}$
			Sucrose	Regal Black		
			µg C	µg		(‰)
Regal Black	15-148-08	28-May-15	10	22	EC	-27.49
n = 9	15-148-05	28-May-15	15	26	EC	-27.73
	15-149-07	29-May-15	20	50.4	EC	-27.34
	15-148-09	28-May-15	30	66	EC	-27.32
	16-224-04	11-Aug-16	20	57	EC	-27.31
	16-224-07	11-Aug-16	20	53	EC	-27.27
	16-224-08	11-Aug-16	20	58	EC	-27.37
	16-225-07	12-Aug-16	10	20	EC	-27.57
	17-248-08	30-Aug-17	20	53	EC	-27.47
					mean	**-27.43**
					s.d.	**0.15**
Sucrose	15-149-04	29-May-15	10	22	OC	-12.82
n = 9	15-148-06	28-May-15	15	26	OC	-12.54
	15-149-05	29-May-15	20	50.4	OC	-12.54
	15-149-06	29-May-15	30	66	OC	-12.29
	16-224-05	11-Aug-16	20	57	OC	-13.04
	16-224-06	11-Aug-16	20	53	OC	-12.36
	16-225-03	12-Aug-16	20	58	OC	-12.72
	16-225-04	12-Aug-16	10	20	OC	-12.86
	17-242-06	30-Aug-17	20	53	OC	-12.34
					mean	**-12.61**
					s.d.	**0.26**

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Table S5. Calculated stable isotopic composition ($^{13}C/^{12}C$) in a two-end-member-mixing system with endmember #1 being Sucrose ($\delta^{13}CV_{PDB}$ =-12.22‰) and end member #2 being Regal black ($\delta^{13}CV_{PDB}$=-27.61‰) and where endmember #1 is mixed into endmember#2.

$\delta^{13}C_{VPDB}$ of pure endmember		fraction of sucrose in mixture (Sucrose + Regal black)	$\delta^{13}C_{VPDB}$ of the mixture calculated
Sucrose	Regal black		
‰	‰	%	‰
		0	-27.610
		1	-27.456
		2	-27.302
		3	-27.148
		4	-26.994
		5	-26.841
		10	-26.071
		20	-24.532
		30	-22.993
		40	-21.454
		50	-19.915
		60	-18.376
-12.22	-27.61	70	-16.837
		80	-15.298
		90	-13.759
		91	-13.605
		92	-13.451
		93	-13.297
		94	-13.143
		95	-12.990
		96	-12.836
		97	-12.682
		98	-12.528
		99	-12.374
		100	-12.220

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Table S6. Radiocarbon content, expressed as fraction modern carbon (FM), of total (TC), organic (OC), and elemental (EC) carbon fractions with and without background correction following Santos et al. (2010). OC and EC fractions were isolated with the ECT9 protocol (Huang et al., 2006) from pure reference materials (into the form of CO_2), then purified cryogenically and sealed in ampoules at the CAIR lab, ECCC. CO_2 is reduced to graphite (Santos et al., 2007b, 2007a) and analyzed at the KCCAMS facility.

UCIAMS#	Fraction	Mass after ECT9	Mass atKCCAMS	Corrected FM		Uncorrected FM	
		µgC	µgC		±		±
Adipicacid							
153279	TC	10	14	-0.0050	0.0367	0.0593	0.0010
153280	TC	17	16	-0.0116	0.0325	0.0465	0.0009
153281	TC	23	29	-0.0043	0.0165	0.0268	0.0005
153282	TC	37	37	-0.0102	0.0125	0.0140	0.0006
mean				-0.0078			
s.d.				0.0037			
Sucrose							
153283	TC	5	7	1.0041	0.0885	0.8766	0.0101
153284	TC	5	7	1.0031	0.0878	0.8759	0.0051
153285	TC	5	7	1.0346	0.0938	0.8960	0.0064
153286	TC	10	11	1.0529	0.0516	0.9652	0.0045
153287	TC	10	11	1.0360	0.0511	0.9510	0.0070
153288	TC	10	12	1.0571	0.0510	0.9702	0.0056
153289	TC	20	21	1.0477	0.0265	1.0006	0.0069
153290	TC	20	21	1.0429	0.0257	0.9971	0.0058
153291	TC	20	21	1.0470	0.0262	1.0000	0.0056
153292	TC	40	41	1.0405	0.0127	1.0170	0.0034
153293	TC	40	38	1.0543	0.0139	1.0282	0.0034
153294	TC	40	42	1.0509	0.0125	1.0272	0.0026
153295	OC	20	20	1.0844	0.0290	1.0305	0.0041
mean				1.0427			
s.d.				0.0213			
C1150							
153303	TC	7	10	0.0310	0.0535	0.1154	0.0020
153304	TC	16	23	0.0278	0.0205	0.0644	0.0012
153305	TC	34	36	-0.0012	0.0131	0.0237	0.0006
153306	TC	45	55	0.0041	0.0083	0.0201	0.0003

153307	EC	32	33	-0.0072	0.0144	0.0202	0.0004
mean				0.0109			
s.d.				0.0174			
RegalBlack							
153308	TC	16	23	0.0161	0.0209	0.0540	0.0008
153309	TC	47	53	-0.0008	0.0087	0.0160	0.0004
153310	EC	28	41	-0.0057	0.0112	0.0159	0.0004
mean				0.0032			
s.d.				0.0114			
Ricechar							
153299	TC	6	7	0.9383	0.0830	0.8272	0.0097
153300	TC	12	15	1.0463	0.0390	0.9784	0.0057
153301	TC	24	22	1.0823	0.0254	1.0348	0.0046
153302	EC	13	15	1.0621	0.0383	0.9940	0.0046
mean				1.0323			
s.d.				0.0643			
OxalicacidII[a]							
153316	TC	n/a	7	1.3141	0.0398	1.2411	0.0203
153315	TC	n/a	17	1.3365	0.0137	1.3080	0.0063
153314	TC	n/a	45	1.3342	0.0051	1.3235	0.0027
mean				1.3283			
s.d.				0.0123			
Adipicacid[a]							
153318	TC	n/a	6	-0.0020	0.0313	0.0544	0.0031
153317	TC	n/a	16	-0.0016	0.0115	0.0205	0.0011
153278	TC	n/a	56	-0.0014	0.0033	0.0051	0.0003
mean				-0.0017			
s.d.				0.0003			

[a]Reference standards that underwent combustion and graphitization process only for blank determination at KCCAMS (without ECT9); n/a. = not applicable

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Table S7. Radiocarbon content, expressed as fraction modern carbon (FM), of total (TC), organic (OC), and elemental (EC) carbon fractions with and without background correction following Santos et al. (2010). OC and EC fractions were isolated with the ECT9 protocol (Huang et al., 2006) from mixtures of reference materials (into the form of CO_2), then purified cryogenically and sealed in ampoules at ECCC. CO_2 is reduced to graphite (Santos al., 2007b, 2007a) and analyzed at KCCAMS facility.

UCI AMS #	Fraction measured	Initial loaded mass		Mass after ECT9	Mass at KCCAMS	Corrected FM		Uncorrected FM	
		µg C	µg	µg C			±		±
Sucrose + Regal black		Sucrose	Regal black						
159800	OC	5	10	5	6	1.0568	0.0648	0.9738	0.0107
159802	OC	10	21	11	10	1.0542	0.0337	1.0057	0.0049
159804	OC	15	29	16	15	1.0629	0.0216	1.0298	0.0037
159806	OC	20	39	21	20	1.0436	0.0156	1.0201	0.0034
159808	OC	30	63	32	29	1.0563	0.0107	1.0395	0.0025
159801	EC	5	10	10	11	-0.0361	-0.0502	0.0535	0.0014
159803	EC	10	21	20	19	-0.0189	-0.0270	0.0317	0.0007
159805	EC	15	29	28	36	-0.0091	-0.0136	0.0172	0.0005
159807	EC	20	39	38	44	0.0014	0.0110	0.0226	0.0004
159809	EC	30	63	61	56	0.0019	0.0085	0.0186	0.0003
Adipic acid + Bulk rice char		Adipic acid	Bulk rice char[a]						
159822	OC	5	11	6	6	0.1009	0.0856	0.2279	0.0027
159824	OC	10	22	12	11	0.0759	0.0450	0.1516	0.0021
159826	OC	15	35	18	17	0.1078	0.0278	0.1558	0.0013
159828	OC	20	44	23	22	0.1072	0.0204	0.1432	0.0014
159830	OC	25	51	29	23	0.1552	0.0185	0.1868	0.0011
159832	OC	30	60	34	32	0.1013	0.0138	0.1263	0.0009
159823	EC	5	11	5	5	1.1063	0.0887	0.9903	0.0063
159825	EC	10	22	10	8	1.0981	0.0486	1.0263	0.0052
159827	EC	15	35	16	14	1.0559	0.0231	1.0211	0.0034
159829	EC	20	44	20	17	1.0619	0.0190	1.0328	0.0040
159831	EC	25	51	23	22	1.0625	0.0143	1.0400	0.0027
159833	EC	30	60	27	24	1.0633	0.0131	1.0426	0.0028
Adipic acid + Rice char_EC[b]		Adipic acid	Rice char_EC						
159810	OC	5	13	5	6	-0.0605	-0.1166	0.1212	0.0032
159812	OC	10	19	10	10	-0.0324	-0.0558	0.0655	0.0015

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159814	OC	15	34	15	15	-0.0075	-0.0345	0.0556	0.0008
159816	OC	20	38	20	20	0.0107	0.0248	0.0568	0.0011
159818	OC	25	49	25	25	-0.0009	-0.0198	0.0366	0.0005
159820	OC	30	60	30	29	0.0103	0.0168	0.0421	0.0006
159811	EC	5	13	6	5	1.0926	0.0931	0.9755	0.0094
159813	EC	10	19	8	7	1.0702	0.0506	0.9997	0.0058
159815	EC	15	34	15	16	1.0709	0.0203	1.0392	0.0037
159817	EC	20	38	17	20	1.0726	0.0162	1.0471	0.0038
159819	EC	25	49	22	21	1.0749	0.0152	1.0505	0.0029
159821	EC	30	60	27	27	1.0723	0.0116	1.0535	0.0024

[a]The bulk rice char contains 52% of TC, on which 14% is OC and 86% EC, respectively; [b]Adipic acid was injected after the OC of rice char is removed through combustion at 870°C via ECT9. Thus, adipic acid was mixed only with rice char-EC, and the OC of the mixture is only from Adipic acid and EC of the mixture is only from Rice char.