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Journal Article**Author(s):**

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Publication date:

2016

Permanent link:

<https://doi.org/10.3929/ethz-b-000123414>

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Originally published in:

Atmospheric Chemistry and Physics 17, <https://doi.org/10.5194/acp-17-3233-2017>



Sources and formation mechanisms of carbonaceous aerosol at a regional background site in the Netherlands: insights from a year-long radiocarbon study

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Received: 14 July 2016 – Discussion started: 14 September 2016

Revised: 17 January 2017 – Accepted: 23 January 2017 – Published: 6 March 2017

Abstract. We measured the radioactive carbon isotope ¹⁴C (radiocarbon) in various fractions of the carbonaceous aerosol sampled between February 2011 and March 2012 at the Cesar Observatory in the Netherlands. Based on the radiocarbon content in total carbon (TC), organic carbon (OC), water-insoluble organic carbon (WIOC), and elemental carbon (EC), we estimated the contribution of major sources to the carbonaceous aerosol. The main source categories were fossil fuel combustion, biomass burning, and other contemporary carbon, which is mainly biogenic secondary organic aerosol material (SOA).

A clear seasonal variation is seen in EC from biomass burning (EC_{bb}), with lowest values in summer and highest values in winter, but EC_{bb} is a minor fraction of EC in all seasons. WIOC from contemporary sources is highly correlated with EC_{bb}, indicating that biomass burning is a dominant source of contemporary WIOC. This suggests that most biogenic SOA is water soluble and that water-insoluble carbon stems mainly from primary sources. Seasonal variations in other carbon fractions are less clear and hardly distinguishable from variations related to air mass history.

Air masses originating from the ocean sector presumably contain little carbonaceous aerosol from outside the Nether-

lands, and during these conditions measured carbon concentrations reflect regional sources. In these situations absolute TC concentrations are usually rather low, around 1.5 µg m⁻³, and EC_{bb} is always very low (~0.05 µg m⁻³), even in winter, indicating that biomass burning is not a strong source of carbonaceous aerosol in the Netherlands. In continental air masses, which usually arrive from the east or south and have spent several days over land, TC concentrations are on average by a factor of 3.5 higher. EC_{bb} increases more strongly than TC to 0.2 µg m⁻³. Fossil EC and fossil WIOC, which are indicative of primary emissions, show a more moderate increase by a factor of 2.5 on average.

An interesting case is fossil water-soluble organic carbon (WSOC, calculated as OC-WIOC), which can be regarded as a proxy for SOA from fossil precursors. Fossil WSOC has low concentrations when regional sources are sampled and increases by more than a factor of 5 in continental air masses. A longer residence time of air masses over land seems to result in increased SOA concentrations from fossil origin.

1 Introduction

Carbonaceous material constitutes a significant fraction of the atmospheric aerosol in almost all environments. In Europe this fraction is typically between 30 and 60 % of PM_{2.5} (e.g., Fuzzi et al., 2015; Pöschl, 2005; Putaud et al., 2004). With the continuing reduction of inorganic aerosol constituents in Europe (e.g., Chin et al., 2014), carbonaceous material will likely be the dominant aerosol component in the future. For further improvements in air quality, it is necessary to target the carbonaceous aerosol fraction. However, source apportionment of the organic aerosol fraction is notoriously difficult due to the large number of constituents and the complexity of chemical formation and transformation processes of the organic aerosol (Fuzzi et al., 2006).

Analysis of the radioactive carbon isotope ¹⁴C in various carbonaceous aerosol fractions has become an important tool for source apportionment in the recent years (e.g., Currie, 2000; Gelencsér et al., 2007; Heal et al., 2014, and references therein; Szidat et al., 2006). The success of this method lies in its clear-cut separation between fossil and contemporary sources of carbonaceous aerosol. ¹⁴C is continually produced in the upper troposphere and stratosphere. Thermal neutrons produced by cosmic rays react with ¹⁴N to ¹⁴C, which is oxidized via ¹⁴CO to ¹⁴CO₂. ¹⁴C decays with a half-life of 5730 years (Godwin, 1962). In the living biosphere it is continually replenished from atmospheric ¹⁴CO₂, so that a typical contemporary level is established. Aerosol carbon originating from the living biosphere, such as plant emissions or wood combustion, has therefore the contemporary ¹⁴C signature. Fossil fuels, however, have been buried for so long that ¹⁴C has completely decayed and as a consequence aerosol carbon from fossil fuels contains no ¹⁴C.

In aerosol source apportionment, the ¹⁴C content of a sample is usually reported as fraction modern (Mook and van der Plicht, 1999; Reimer et al., 2004):

$$F^{14}\text{C} = \frac{\left(\frac{^{14}\text{C}}{^{12}\text{C}}\right)_{\text{sample}}}{\left(\frac{^{14}\text{C}}{^{12}\text{C}}\right)_{1950}}, \quad (1)$$

which relates the ¹⁴C / ¹²C ratio of the sample to the ratio of the unperturbed atmosphere in the year 1950. Both ratios are normalized to a δ¹³C value of −25 ‰ to account for isotopic fractionation during sample pretreatment and measurement. Aerosol carbon derived from living biomass should therefore have F¹⁴C ~ 1 in an atmosphere unperturbed by human activities, whereas carbon from fossil sources has F¹⁴C = 0. Human activities, however, especially the atomic bomb tests which nearly doubled the natural ¹⁴CO₂ levels in the 1960s and 1970s, are perturbing the natural equilibrium. Currently, the atmospheric CO₂ has F¹⁴C of approximately 1.04 (Levin et al., 2010), which is decreasing every year, because the ¹⁴CO₂ produced by bomb testing is taken up by oceans and the biosphere. Moreover, fossil fuel consumption introduces

¹⁴C-free CO₂ into the atmosphere, leading to a further decrease in atmospheric F¹⁴C. This has consequences for the F¹⁴C of contemporary aerosol sources: biogenic primary and secondary organic aerosols, as well as aerosols from cooking emissions, have F¹⁴C close to the value of current atmospheric CO₂. F¹⁴C of aerosol from wood combustion is higher than that, because a significant fraction of carbon in the wood burned today was fixed during times when ¹⁴C levels in the atmosphere were high. Estimates based on tree growth models (e.g., Lewis et al., 2004; Mohn et al., 2008) range from 1.08 to 1.30 for biomass combusted in western Europe (Genberg et al., 2011; Gilardoni et al., 2011; El Hadad et al., 2011; Minguillón et al., 2011; Szidat et al., 2006, 2007, 2009).

If F¹⁴C is separately measured on different carbon sub-fractions, such as organic carbon (OC) and elemental carbon (EC), then the three major sources of carbonaceous aerosol can be separated, namely biogenic, biomass combustion, and fossil fuel combustion (Bernardoni et al., 2013; Minguillón et al., 2011; Szidat et al., 2006). For example, Szidat et al. (2007) showed that residential biomass combustion is the dominant source of carbonaceous aerosol in Alpine valleys in winter. Gustafsson et al. (2009) demonstrated that in the brown haze over South Asia the majority of the carbonaceous aerosol originates from biomass burning sources. Other studies showed that carbonaceous aerosol is dominated by biogenic sources in rural areas in the summer (Genberg et al., 2011; Yttri et al., 2011b).

Measurements of F¹⁴C in either water-insoluble OC (e.g., Szidat et al., 2006) or water-soluble OC (Kirillova et al., 2010) can help to distinguish secondary from primary fossil OC. Primary OC / EC ratios for fossil sources are relatively well characterized and tend to produce mainly water-insoluble organic carbon (WIOC). Therefore fossil WIOC can be taken as indicative of primary fossil OC material and the fossil water-soluble organic carbon (WSOC) as indicative of fossil SOA material. Using this modified tracer approach Zhang et al. (2014) showed that secondary fossil OC clearly exceeds primary fossil OC in southern China. Measurements downwind of urban centers in California (Zotter et al., 2014a) and Japan (Morino et al., 2010) have highlighted the rapid formation of fossil SOA, using ¹⁴C measurements together with positive matrix factorization of aerosol mass spectrometer data and a chemical mass balance model, respectively.

The radiocarbon content of different subfractions of the carbonaceous aerosol has therefore the potential to give information on not only sources but also formation pathways and processes of the organic aerosol. In this study we apply radiocarbon measurements to study the sources and formation processes of organic aerosol at a regional background site in the Netherlands, a heavily industrialized region with a high population density and generally high aerosol concentrations, where PM₁₀ values of larger than 50 μg m^{−3} were still observed 20–30 times in the year 2011 at many measurements stations. Carbonaceous material contributes on av-

erage around 30 % to the total PM_{2.5} at regional background locations in the Netherlands with somewhat higher concentrations in urban centers and at curbside locations (Weijers et al., 2011). Few radiocarbon measurements have been reported for the aerosol in the Netherlands. A fraction modern of organic carbon ($F^{14}C_{(OC)}$) of around 0.7 has been measured at urban background sites both in Amsterdam (Dusek et al., 2013) and in Rotterdam (Keuken et al., 2013) in spring, which is in the typical range of western European cities (Heal et al., 2011; Szidat et al., 2006, 2009). $F^{14}C_{OC}$ of total suspended particles at a coastal site in the Netherlands was considerably higher (0.86 on average) (Dusek et al., 2013).

These previous studies were limited in duration and only representative for a small geographical region. For the current study, we chose a regional background site surrounded by the biggest urban centers of the Netherlands and collected aerosol samples for more than 1 year. Due to the location of the site and the sampling approach, which accounted for transport of air masses, the results presented in this study reflect seasonal changes as well as differences between regional air masses and pollution due to long-range transport.

2 Methods

2.1 Measurement site

Measurements were taken at the Cesar Observatory (<http://www.cesar-observatory.nl/>), which is located in the western part of the Netherlands in an agricultural region 0.7 m below sea level. The immediate surroundings are dominated by open pastures with little variation in surface elevation and few wind breaks. The cities of Utrecht and Rotterdam lie 20–30 km to the northwest and southeast, respectively, and the cities of The Hague, Leiden, and Amsterdam at a distance of around 50 km to the west and north. The closest highways connecting these major cities are at a distance of 10–20 km from the site. This site therefore represents the regional background of a relatively polluted area in western Europe.

2.2 Sampling and filter handling

Samples of particulate matter with a diameter less than 2.5 μm were taken with a high-volume sampler (Digital DHA-80) from February 2011 to March 2012 at a flow rate of 500 L min⁻¹. The sampling duration varied between 2 and 7 days. Since the concentration and properties of organic aerosol can be significantly influenced by long-range transport, the start and end dates of each sampling were chosen based on air mass back-trajectory forecasts calculated every day by the NOAA HYSPLIT model (Stein et al., 2015). This way, individual filter samples generally contain aerosol from specific source regions.

Aerosol particles were collected on circular quartz fiber filters (Whatman QMA1851-150) with a diameter of 15 cm. The filters were pre-heated at 800 °C overnight to remove

adsorbed organic contaminants. Before and after sampling the filters were kept wrapped in pre-backed aluminum foil and stored at –20 °C, except during transport to and from the sampling site (approx. 1 h). After sampling, loaded filters were immediately removed from the filters stacks to avoid adsorption of volatile organic compounds to the quartz fibers. All instruments that came in contact with the filters (e.g., cutters, tweezers) were pre-cleaned with acetone followed by ethanol. Blank filters were treated exactly like the sample filters, except that they were kept in the sampler only for a few minutes without switching it on.

2.3 Combustion of carbon fractions and standards

The separation of OC and EC for all filter samples was performed on the aerosol combustion system (ACS) that was developed at the University of Utrecht and has been described in detail by Dusek et al. (2014). Briefly, the ACS consists of a combustion tube, where aerosol samples are combusted at different temperatures in pure O₂, and a purification line where the resulting CO₂ is isolated and separated from other gases, such as water vapor or NO_x. The mass of carbon in each sample is determined manometrically. The purified CO₂ is then stored in flame-sealed glass ampoules until ¹⁴C analysis.

We analyze radiocarbon in the following carbon fractions: total carbon (TC), OC, WIOC, and EC. For TC combustion an aliquot of the filter is heated at 650 °C for 15 min. The carbon fraction considered representative of OC is combusted by heating of a different filter piece at 360 °C for 15 min in O₂. For WIOC a water-extracted (see Dusek et al., 2014) filter piece is heated at 360 °C for 15 min in O₂. EC is combusted after OC has been completely burned off the filter. To achieve complete OC removal, WSOC is first removed from the filter by water extraction to prevent charring of organic material (Dusek et al., 2014). Subsequently, most WIOC is removed by heating the filter piece at 360 °C in O₂ for 15 min. Then the oven temperature is raised to 450 °C for 2 min and in this step a mixture of the most refractory OC and EC is burned off the filter in O₂. The remaining EC is then combusted by heating at 650 °C. Zhang et al. (2012) and Dusek et al. (2014) estimate that after water extraction and flash combustion in O₂, charred organic compounds can contribute roughly 5 % to the recovered EC. A mean bias for charring of 0.04 is therefore subtracted from the fraction modern of EC ($F^{14}C_{(EC)}$) before source apportionment, assuming 5 % of the measured EC fraction consists of charred OC and that $F^{14}C_{(OC)}$ is approximately 0.8. Moreover, it is possible that highly refractory OC could survive the thermal treatment and combusted together with EC.

As quality control two sets of standards with known ¹⁴C content are analyzed at regular intervals: the HOxII oxalic acid standard with a nominal $F^{14}C$ of 1.3408 and a graphite standard with $F^{14}C$ of 0. The standards are directly put on the filter holder and heated at 650 °C for 15 min. From

the deviation of the measured $F^{14}\text{C}$ from the nominal values, the contamination introduced by the combustion procedure can be estimated. This contamination is on average below $3 \mu\text{g C}/\text{combustion}$ (Dusek et al., 2014) and relatively small compared to typical sample amounts between 50 and $200 \mu\text{g C}$.

2.4 ^{14}C measurement and data correction

After purification the CO_2 collected from the combustion of various aerosol fractions on the ACS system is sealed in glass ampoules. Most of these CO_2 samples were sent to the Centre for Isotope Research (CIO) at the University of Groningen for graphitization and AMS measurements. The CO_2 extracted from all samples collected in spring 2011 and several blanks and standards were analyzed without graphitization at the AMS facility at the Laboratory of Ion Beam Physics of ETH Zurich.

At CIO CO_2 is reduced to graphite by reaction with molecular hydrogen at a molecular ratio H_2 / CO_2 of 2.5. A porous iron pellet (de Rooij et al., 2010) at a temperature of 600°C is used as a catalyst for the reaction. The resulting water vapor is cryogenically removed using Peltier cooling elements. The yield of graphite is usually $>95\%$ for samples larger than $30 \mu\text{g}$. After graphite has formed on the iron pellet, it is pressed into a 1.5 mm target holder, which is introduced into the AMS system for subsequent measurement. The AMS system (van der Plicht et al., 2000) is dedicated to ^{14}C analysis and simultaneously measures $^{13}\text{C} / ^{12}\text{C}$ and $^{14}\text{C} / ^{12}\text{C}$ ratios. Samples below $500 \mu\text{g}$ are analyzed together with varying amounts of reference materials ranging from 50 to $500 \mu\text{g C}$. Two reference materials with known ^{14}C content are used: the HOxII standard ($F^{14}\text{C} = 1.3407$) and a ^{14}C -free CO_2 gas ($F^{14}\text{C} = 0$). The differences between actually measured and nominal $F^{14}\text{C}$ values are used for correcting the sample values (de Rooij et al., 2010) for contamination during graphitization and AMS measurement. The contamination is typically below $2 \mu\text{g C}$ (Prokopiou, 2010).

At ETH Zurich, ^{14}C is measured directly without graphitization from the purified CO_2 using the AMS system MICADAS with a gas ion source (Ruff et al., 2007). In an automated gas interface, the CO_2 is released from ampoules with a cracker, mixed with helium and transferred into the ion source with a constant gas flow (Wacker et al., 2013). This system allows ^{14}C analysis of ultra-small samples down to $2 \mu\text{g C}$. The samples are corrected for blank and isotope fractionation as well as normalized to $F^{14}\text{C}$ values using HOxII and a ^{14}C -free material as gaseous standards. A constant contamination of $<0.1 \mu\text{g C}$ with an $F^{14}\text{C}$ value of 0.5 was determined for the ^{14}C gas analysis (Ruff et al., 2010), which was applied for the correction of the samples.

2.5 Blank correction

The amount of carbon on a blank filter was on average $0.34 \mu\text{g cm}^{-2}$ for OC, based on the analysis of three blank filters. The amount of OC on sample filters varied from roughly 10 to $100 \mu\text{g cm}^{-2}$, with an average of around $30 \mu\text{g cm}^{-2}$. Due to the small amount of carbon on the blank filters it was only possible to analyze ^{14}C for one of these blank filters. The $F^{14}\text{C}_{(\text{OC})}$ on this blank filter was 0.797 ± 0.019 . The amount of blank EC is mainly added during the combustion process and is therefore not necessarily proportional to the area of the filter used for analysis. Blank EC was found to be $0.3 \pm 0.1 \mu\text{g cm}^{-2}$ for a filter piece of 7 cm^2 (average of five samples), $0.17 \mu\text{g cm}^{-2}$ for a piece of 14 cm^2 (based on one sample), and $0.11 \pm 0.03 \mu\text{g cm}^{-2}$ for a piece of 21 cm^2 (average of three samples). ^{14}C was analyzed on the pooled carbon collected from the five single blank filter pieces (with a total area of 35 cm^2) and the $F^{14}\text{C}$ was 0.54 ± 0.03 . The concentration of EC on the sample filters varied from around 2 to $30 \mu\text{g cm}^{-2}$. In most cases two filter pieces were combusted at the same time, but for the smallest samples three filter pieces were used, so that in all cases but one the blank is 5 % or less of the sample amount. The concentration of TC on the blank filters was calculated by adding the carbon concentrations of OC and EC and was $0.68 \mu\text{g cm}^{-2}$. $F^{14}\text{C}_{(\text{TC})}$ of the blank filters was calculated as the weighted average of $F^{14}\text{C}_{(\text{OC})}$ and $F^{14}\text{C}_{(\text{EC})}$ and was 0.67.

The ^{14}C values of all carbon fractions were blank corrected according to the mass balance equation:

$$F^{14}\text{C}_S = \frac{F^{14}\text{C}_m \cdot M_m - F^{14}\text{C}_b \cdot M_b}{M_m - M_b}, \quad (2)$$

where $F^{14}\text{C}_S$ is the fraction modern of the aerosol carbon collected on the filter, $F^{14}\text{C}_m$ and M_m are the measured fraction modern and the measured concentration of the respective carbon fraction (TC, OC, WIOC, or EC), and $F^{14}\text{C}_b$ and M_b the fraction modern and the concentration of the respective carbon fraction on the blank filter.

2.6 Measurements of sugars

In addition to $F^{14}\text{C}$ we also measured atmospherically relevant sugars (e.g., levoglucosan, sucrose, glucose, mannosan). Levoglucosan can serve as an independent tracer of biomass burning and several other sugars can indicate primary biological material, which cannot be traced by ^{14}C measurements alone. The determination of carbohydrates is performed with an improved high-performance anion-exchange chromatography with pulsed amperometric detection (HPAEC-PAD). This method is recommended for atmospherically relevant sugars (sugar alcohols, monosaccharides and monosaccharide anhydrides) and based on the method described by Iinuma et al. (2009). Before the analysis a filter aliquot ($1 \times \varnothing 10 \text{ mm}$, 0.8 cm^2) was eluted with 3 mL of ultra-pure water (Milli-Q Plus, 185, Millipore), sonicated for 20 min

and subsequently centrifuged for 30 min. The extract was analyzed with an ion chromatography system (Dionex[®] ICS 3000), operated with a NaOH gradient (initial concentration: 480 mM NaOH, final concentration: 650 mM NaOH) run on a Carbo Pac MA1 column. The quantification is carried out with external standards prepared from pure substances (Merck[®], Fluka[®]). The relative standard deviation of the measurements, determined based on replicate analysis of standards and blanks, was below 10 %.

2.7 Estimation of OC and EC by thermal–optical analysis

Filter pieces of 1.5 cm² were analyzed for OC and EC with a dual-optical Sunset Analyzer (Sunset Laboratory, Inc.) using the QUARTZ temperature protocol (Birch and Cary, 1996). In this instrument, filter samples are heated stepwise in an He atmosphere to a maximum temperature of 840 °C to volatilize organic material, then cooled down to 550 °C and subsequently heated again stepwise to 870 °C in a, He–O₂ (10 % O₂) atmosphere to oxidize EC. All carbon gases are converted to CH₄ and detected with a flame ionization detector. As organic material partly pyrolyzes in the first part of the cycle, a laser beam is used to correct for charring by monitoring both transmission and reflection signals. The split between OC and EC is set when the transmission or reflection signals reach their initial values. The analytical uncertainty for OC and EC varied slightly with filter loading from 5 % at loadings above 20 μg cm⁻² to 7 % for loadings below 10 μg cm⁻² and 10 % for loadings around 2 μg cm⁻². EC values determined by both transmission and reflection method are compared later in the paper. EC, OC, and TC measured by the thermal optical method are used to evaluate the estimated values of OC and EC based on the data from the ACS system (see Sect. 2.8).

2.8 Estimation of OC, EC, and WIOC concentrations based on carbon fractions recovered by the ACS system

The OC combustion temperature of 360 °C in the ACS is likely not high enough to recover 100 % of OC. The concentrations of OC and EC extracted for ¹⁴C analysis ($M_{OC,e}$ and $M_{EC,e}$) are therefore lower than the actual concentrations of OC and EC in the aerosol sample (M_{OC} and M_{EC}). If $F^{14}C$ of TC is known, M_{OC} and M_{EC} can be estimated as follows:

$$M_{OC} = M_{TC} \frac{F^{14}C_{(TC)} - F^{14}C_{(EC)}}{F^{14}C_{(OC)} - F^{14}C_{(EC)}}, \quad (3)$$

$$M_{EC} = M_{TC} - M_{OC}, \quad (4)$$

where M_{OC} , M_{EC} , and M_{TC} are the mass concentrations of OC, EC, and TC, respectively, and $F^{14}C_{(OC)}$, $F^{14}C_{(EC)}$, and $F^{14}C_{(TC)}$ are the fraction modern of OC, EC and TC, respectively. The recovery of OC and EC can consequently be estimated as $M_{OC,e}/M_{OC}$ and $M_{EC,e}/M_{EC}$.

In addition, we measured ¹⁴C on WIOC. Estimating the concentration of WIOC is not straightforward, because the recovery of WIOC is unknown and cannot be derived in the same way as for OC. To estimate M_{WIOC} we therefore assume two extreme cases. (1) WIOC is completely recovered: In this case M_{WIOC} is equal to the mass of WIOC that was determined by the ACS system. This is most likely results in an underestimate of WIOC, because a fraction of WIOC might not combust at 360 °C. In this case the measured WIOC is less than the actual M_{WIOC} . (2) WIOC shows the same recovery as OC: in this case M_{WIOC} is calculated as the measured WIOC divided by the recovery of OC. This results probably in an overestimate of M_{WIOC} , since usually less WIOC than OC is lost to charring. We therefore consider a range of possible WIOC concentrations from a minimum of $M1_{WIOC}$ (complete recovery) to a maximum $M2_{WIOC}$ (recovery as OC).

2.9 Estimating source contributions to OC, EC, WIOC, and WSOC

The $F^{14}C$ values of the different carbon fractions can be used to estimate source contributions to the carbonaceous aerosol. For EC two main sources are considered: fossil-fuel-dominated sources (EC_f) and biomass burning (EC_{bb}), which in northwestern Europe is mainly from residential heating and fireplaces, since open wildfires are rare. EC_{bb} can be estimated as

$$EC_{bb} = M_{EC} \frac{F^{14}C_{(EC)} - F_f}{F_{bb} - F_f}, \quad (5)$$

where F_{bb} stands for $F^{14}C$ of biomass burning emissions (see Table 1), and F_f stands for the average $F^{14}C$ of EC from other (fossil-fuel-dominated) combustion sources, such as natural gas and coal burning and emissions from mobile sources. In Europe, these sources emit predominantly fossil carbon and usually F_f has been approximated as 0. However, in recent years an increasing fraction of biofuels has been added to gasoline and diesel used in road transport. This is expected to increase $F^{14}C$ in EC emitted from mobile emissions, although it is not clear how much biofuel addition affects $F^{14}C_{(EC)}$ (e.g., Bennett et al., 2008).

EC_f can be calculated as

$$EC_f = EC - EC_{bb}. \quad (6)$$

Once EC_{bb} is known, OC_{bb} is calculated using an estimate of the primary OC / EC ratio of wood combustion emissions (r_{bb}); (e.g., Bernardoni et al., 2013; Genberg et al., 2011; Szidat et al., 2006). This is a very crude estimate, since r_{bb} of domestic wood combustion in Europe is variable and not very well known. However, it is still useful to get an approximate idea of the relative contribution of primary biomass combustion vs. other modern carbon sources.

$$OC_{bb} = EC_{bb} \cdot r_{bb} \quad (7)$$

Table 1. The range of input values for crucial parameters used in the Monte Carlo simulations: the fraction modern of biomass burning aerosol (F_{bb}), the fraction modern of fossil-dominated sources (F_f), the fraction modern of biogenic aerosol (F_{bio}), and the OC/EC ratio of domestic biomass combustion (r_{bb}). These values are based on critical evaluation of the literature as discussed in Sect. 2.9.

	Min.	Central value	Max.
F_{bb}	1.1	1.15	1.2
F_f	0	0.07	0.1
F_{bio}	–	1.04	–
r_{bb}	3	5	7

OC_{bb} can be subtracted from the total OC and the remaining OC can be apportioned between OC from fossil-dominated combustion (OC_f) and other modern OC. The latter consists of primary and secondary biogenic OC but can also contain secondary OC from wood burning, especially in winter. This OC fraction is therefore named $OC_{c,o}$ (OC contemporary, other) and can be calculated as

$$OC_{c,o} = \frac{(F^{14}C_{(OC)} - F_f) \cdot M_{OC} + (F_f - F_{bb}) \cdot OC_{bb}}{F_{bio} - F_f}, \quad (8)$$

where the fraction modern of $OC_{c,o}$ is approximated by F_{bio} , the fraction modern of biogenic OC. $OC_{c,o}$ can also contain primary biogenic material, but in most cases the contributions to $PM_{2.5}$ are likely small. Other studies have used $F_f \approx F_{fossil} = 0$ for the fossil-fuel-dominated combustion sources, and OC emitted by biofuel combustion has been subsumed under $OC_{c,o}$. Currently in Europe, OC from biofuels is a very minor fraction of OC, but this might change in the future if biofuels are used more extensively. In urban areas OC could also include a contribution of modern carbon from cooking sources (e.g., Allan et al., 2010; Schauer et al., 1996), but at a regional background site this contribution is likely small.

Finally, OC from fossil-dominated combustion sources (OC_f) can be estimated as

$$OC_f = M_{OC} - OC_{c,o} - OC_{bb}. \quad (9)$$

To propagate experimental uncertainties and uncertainties in the parameters F_f , F_{bb} , and r_{bb} to the final results, we conducted a Monte Carlo simulation with 10 000 individual calculations of EC_{bb} , EC_f , OC_{bb} , OC_f , and $OC_{c,o}$ using Eqs. (2)–(7). For each calculation input values for OC and EC concentrations, as well as $F^{14}C_{(OC)}$ and $F^{14}C_{(EC)}$, are chosen randomly from a normal distribution centered around the measured value with the experimental uncertainties as standard deviation. Similarly, random values for F_f , F_{bb} , and r_{bb} are estimated as follows: for each of the parameters we estimated a central value and an upper and lower limit based on values reported in the literature. The random values are then chosen from a triangular frequency distribution, which

has its maximum at the central value and is 0 at the upper and lower limits. The central values, maxima, and minima for F_f , F_{bb} , and r_{bb} are reported in Table 1. Finally, the average and standard deviation of the 10 000 different estimates of EC_{bb} , EC_f , OC_{bb} , OC_f , and $OC_{c,o}$ are calculated. The average represents the best estimate of the concentration of each fraction and the standard deviation represents the uncertainty arising from uncertainties in the measurements and in F_f , F_{bb} , and r_{bb} .

We assume that F_f lies between 0 (i.e., all combustion sources other than wood combustion are purely fossil) and 0.1 (emissions of modern carbon from the biofuel added to road fossil fuels contributes 10 % to carbon emission from such sources). The latter is a reasonable upper limit, since a contribution of biofuels to fuels in road transport of at least 5 % was required by law in the Netherlands in 2011, but usually the contribution of modern carbon in the emitted particles is not equal to the contribution of modern carbon in the fuels (e.g., Bennett et al., 2008, and references therein). The range for F_{bb} is estimated as 1.1 to 1.2, with the most likely value of 1.15 based on Genberg et al. (2011), Gilardoni et al. (2011), El Haddad et al. (2011), Minguillón et al. (2011), and Szidat et al. (2006, 2007, 2009). Szidat et al. (2006) derived a best estimate of 6.25 for r_{bb} from an extensive literature study of wood combustion emissions. A few more recent studies give slightly lower values for r_{bb} , for example 5.3 for cooking fires (Christian et al., 2010), 3 ± 2.4 for typical wood stoves used in Austria (Schmidl et al., 2008), and 2.8 for log combustion in wood stoves (Zhang et al., 2013). Yttri et al. (2009) derived r_{bb} of approximately 2.5 by ambient measurements in a location, where wood burning emissions dominate the concentrations of carbonaceous aerosol. Some larger values of r_{bb} of more than 20 can also be found in the literature, but these are often for special situations, such as strongly smoldering combustion (McMeeking et al., 2009) or burning of leaves and grasses, that are not representative for domestic wood combustion (Zhang et al., 2013). Moreover, there are strong indications that pellet and wood log burners have lower OC/BC ratios than traditional stoves (Heringa et al., 2012). Therefore, we use a somewhat lower best estimate for r_{bb} than Szidat et al. (2006) with a wider range from 3 to 7.

Since ratios of $WIOC_{bb}/EC_{bb}$ and $WSOC_{bb}/EC_{bb}$ are not often reported in the literature, it is not possible to separate biogenic and biomass burning carbon for these carbon fractions with any certainty. Therefore, the source apportionment for WIOC and WSOC considers only the fossil-dominated combustion source discussed above and a mixed biomass burning/biogenic “contemporary” source. First the concentration and $F^{14}C$ of WSOC are calculated as

$$M_{WSOC} = M_{OC} - M_{WIOC}, \quad (10)$$

$$F^{14}C_{(WSOC)} = \frac{M_{OC} \cdot F^{14}C_{(OC)} - M_{WIOC} \cdot F^{14}C_{(WIOC)}}{M_{WSOC}}. \quad (11)$$

Since there is a range of possible values for M_{WIOC} , this uncertainty, along with the experimental uncertainties in $F^{14}C$ and M_{OC} , is propagated to M_{WSOC} and $F^{14}C_{(WSOC)}$ as a part of the Monte Carlo simulations described above. M_{WIOC} is assumed to vary in the range of $M1_{WIOC}$ (complete recovery of WIOC, see above) to $M2_{WIOC}$ (recovery as OC), with a most likely value at $M1_{WIOC} + 2/3 \cdot (M2_{WIOC} - M1_{WIOC})$, since it is more likely that WIOC has a similar recovery as OC rather than 100 % recovery. However, since the recovery of OC is on average 75 %, the results are not strongly sensitive to how the most likely value is chosen. If we shift the central value closer to $M1_{WIOC}$, e.g., $M1_{WIOC} + 1/3 \cdot (M2_{WIOC} - M1_{WIOC})$, the average values of $WIOC_f$, $WIOC_c$ change by less than 5 %.

The concentration of WIOC from fossil-dominated combustion sources is calculated as

$$WIOC_f = WIOC \frac{F^{14}C_{(WIOC)} - F_c}{F_f - F_c}, \quad (12)$$

where F_c is defined as $F^{14}C$ of the contemporary organic carbon (i.e., the sum of OC_{bb} and $OC_{c,o}$) in the respective sample, estimated as

$$F_c = \frac{OC_{bb} \cdot F_{bb} + OC_{c,o} \cdot F_{bio}}{OC_{bb} + OC_{c,o}}. \quad (13)$$

Contemporary WIOC is calculated as $WIOC_c = WIOC - WIOC_f$. The concentrations of fossil fuel combustion WSOC are calculated as $WSOC_f = OC_f - WIOC_f$ and contemporary WSOC as $WSOC_c = WSOC - WSOC_f$.

3 Results

3.1 Method evaluation and quality control

Several aspects of the combustion method, mainly focused on the separation of carbon fractions for ^{14}C analysis, were thoroughly tested and evaluated by Dusek et al. (2014). For the current study, it was also necessary to evaluate the accuracy of the calculated TC, OC, and EC concentrations based on Eqs. (2) and (4), since these concentrations are the basis for further source apportionment of ambient aerosol. As basis for comparison we used OC and EC concentrations determined by thermal–optical analysis as described in Sect. 2.8. TC concentrations were determined by thermal–optical analysis on a subset of 18 filter samples. For eight of these filter samples the aerosol loadings were too high for reliable OC–EC analysis and therefore OC and EC concentrations are available only for 10 filter samples. For these 10 samples OC–EC concentrations estimated based on Eqs. (2) and (4) could be compared with thermal optical analysis.

Figure 1a shows that there is excellent agreement between TC determined by the ACS system and TC determined by the thermo-optical method (TC sunset). The slope is 1.07 ± 0.03 ,

thus slightly higher than 1, which can be partly due to traces of water vapor or other impurities that are not removed entirely by the ACS method and that increase the sample pressure during the manometric determination of TC amount. It might also reflect a bias in the calibrated volume used for TC determination on the ACS.

Figure 1b shows the ratio of calculated EC concentrations (M_{EC}) to EC determined by the thermo-optical method with transmission ($M_{EC,t}$) and with reflection correction ($M_{EC,r}$) and using the average of $M_{EC,t}$ and $M_{EC,r}$. The data points correspond to individual filter samples and the red triangles are the average ratios for all filter samples. M_{EC} is on average roughly 10 % higher than $M_{EC,t}$ and 10 % lower than $M_{EC,r}$. The standard deviation is approximately 25 % of the average ratio, which is comparable to the accuracy of different thermo-optical methods for EC (e.g., Schmid et al., 2001). We can therefore conclude that our method of estimating EC and OC concentrations gives comparable results to well-established methods for measuring EC and OC concentrations.

The recovery of OC and EC, estimated according to Sect. 2.8, is on average 75 % for OC and 82 % for EC. For one sample the recovery of OC was calculated as 178 %, likely due to an unrealistically low $^{14}C_{(TC)}$ value. This sample was excluded from further analysis and was also not considered in Fig. 1b. For the subset of samples displayed in Fig. 1b, the average recovery of EC was 83 %, compared to the recovery with respect to thermo-optical analysis $M_{EC,e}/M_{EC,av}$ of 79 %.

In ^{14}C source apportionment, biomass burning is usually considered the main source of contemporary EC. However, there is evidence that primary biogenic particles, such as pollen fragments, can be thermally very refractory and be combusted together with EC (Wittmaack, 2005). Whereas pollen themselves are usually not found in $PM_{2.5}$, the fragments of ruptured pollen are in the submicrometer size range (Taylor et al., 2004) and can potentially contribute modern carbon to the EC fraction. Therefore, in Fig. 2 $F^{14}C_{(EC)}$ was evaluated against levoglucosan, which is another tracer of biomass burning. The values of $F^{14}C_{(EC)}$ shown in Fig. 2 are the raw data, which have not yet been corrected for charring bias (see Sect. 2.3). For most samples there is a clear correlation between the fraction of levoglucosan in TC and $F^{14}C_{(EC)}$. However, there are three samples with strongly elevated $F^{14}C_{(EC)}$ but low levels of levoglucosan (shown in red). These samples were taken in spring 2014 during a time period with reported high pollen concentrations. In this period, many surfaces were covered with a fine yellow dust. This dust was strongly visible on the $PM_{2.5}$ impactor plate of the high-volume sampler and also present on the filter itself. For these three samples glucose and sucrose concentrations, which are tracers for primary biological material, were strongly elevated. Glucose/TC levels were $18 \text{ ng } \mu\text{g}^{-1}$ compared to an average of $1.7 \text{ ng } \mu\text{g}^{-1}$ for the rest of the year and sucrose/TC levels were $81 \text{ ng } \mu\text{g}^{-1}$ compared to an average

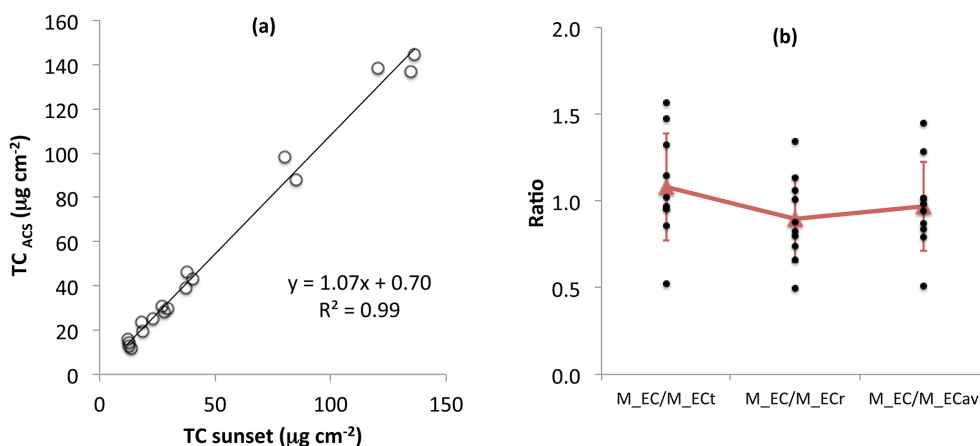


Figure 1. (a) Total carbon concentrations (TC) measured with a thermo-optical method (Sunset Analyzer) and the ACS system (extraction). (b) The ratio of EC concentrations estimated from the ACS system (M_{EC}), divided by EC from the Sunset Analyzer for three different cases: using transmission correction (M_{EC}/M_{ECt}), reflection correction (M_{EC}/M_{ECr}) and by the average of M_{ECr} and M_{ECt} (M_{EC}/M_{ECav}). The red triangles represent averages over all data points and the error bars 1 standard deviation.

of $11 \text{ ng } \mu\text{g}^{-1}$ for the rest of the year. These observations provide the first evidence that primary biological material can be responsible for elevated $F^{14}\text{C}_{(\text{EC})}$ during spring. Whether this contribution comes from strong charring or the fact that a part of the primary biological material survives the thermal treatment designed to purify EC is not entirely clear. For further calculations in the paper, the measured $F^{14}\text{C}_{(\text{EC})}$ of the three samples marked in red was replaced by $F^{14}\text{C}_{(\text{EC})}$ calculated based on measured levoglucosan concentration and the regression line in Fig. 2. The apparent modern EC that is assumed to be unburned OC is subtracted from the EC fraction and added to the $\text{OC}_{\text{c,o}}$ fraction. This can be seen as a crude correction for the highly refractory part of OC that was apparently incorrectly classified as EC from biomass burning. Without this correction we would overestimate the contribution of biomass burning to the carbonaceous aerosol in spring.

The offset of the regression line in Fig. 2 is not 0, which indicates that there is contemporary carbon in EC that is not related to biomass burning. One main reason is the addition of biofuel to diesel and gasoline, which in the Netherlands must currently contain at least 5 % biofuel. In addition, cooking could contribute to contemporary EC, although cooking usually emits much more OC than EC (e.g., Chow et al., 2004) and is probably not very relevant at a regional background site. Therefore any potential contribution from cooking is subsumed under EC_{bb} in this study. In addition there is evidence that levoglucosan is not a stable tracer but can be degraded by photochemical reactions during long-range transport (Hoffmann et al., 2010). This means that the offset in $F^{14}\text{C}_{(\text{EC})}$ could also partially be caused by underestimated levoglucosan concentrations.

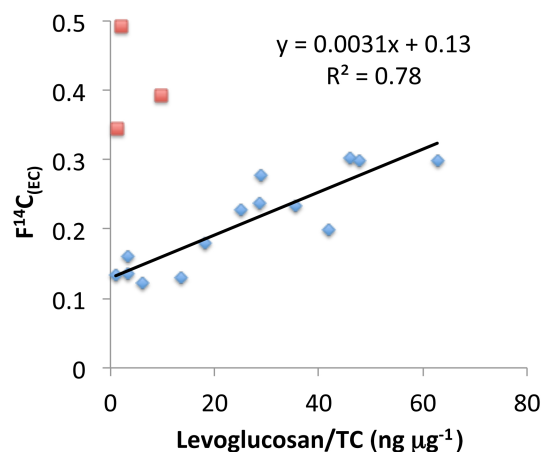


Figure 2. $F^{14}\text{C}_{(\text{EC})}$ vs. levoglucosan mass fraction in the carbonaceous aerosol. The data marked in red are from three periods in spring 2011, when high pollen concentrations were measured. The $F^{14}\text{C}$ data are not corrected for charring bias. The regression line and regression equation corresponds to a linear LS fit to the blue data points.

3.2 Source apportionment in different seasons and air mass conditions

We analyzed $F^{14}\text{C}$ in TC, OC, WIOC, and EC fractions for 26 PM_{2.5} samples taken in 2011 and 2012. Four-day air mass back trajectories were calculated with the HYSPLIT model every 24 h for each sample from the sampling start time to the sample end time. These are not the forecast trajectories used for deciding the sampling times, but rather the actual back trajectories of the filter samples that were selected for analysis. Most filter samples can be associated with one of two main source regimes – continental or modified marine

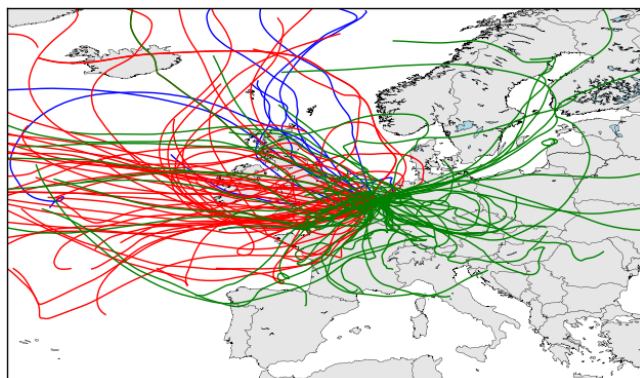


Figure 3. Ninety-six hour air mass back trajectories (NOAA HYSPLIT) ending at a height of 50 m for the filter samples classified as modified marine (red), marine (blue), and continental (green). For further analysis the blue trajectories were combined with the modified marine trajectories (red) to represent aerosol samples, where the total carbon is dominated by regional emissions. Only samples for which air mass back trajectories were relatively stable over several days have been selected for analysis.

– and are shown in Fig. 3. The air mass back trajectories for the filters classified as “continental” are shown as green lines. The trajectories originate mostly to the east or south of the Netherlands and spend considerable time over land. A few trajectories are also from the west, because occasionally a filter sample could not be stopped in time before a change of wind direction. However, for each continental filter sample such trajectories accounted only for a small fraction of the total sampling time. The air mass back trajectories for filter samples classified as “modified marine” originate over the Atlantic and spend only a brief time over land before reaching the Netherlands, mostly over Great Britain and Belgium. These trajectories are shown in red. The trajectories colored in blue correspond to the cleanest air mass conditions, when the air arrived from the north and did not cross any land masses before reaching the Netherlands. However, there were only two filter samples with these air mass conditions, and they were therefore included into the modified marine cases.

Since the concentration of carbonaceous aerosol over the ocean is low, the aerosol samples in the modified marine case are likely dominated by regional emissions from in and around the Netherlands. The majority of the particles in these samples had been emitted within the previous 24 h and the aerosol carbon is therefore characterized by a short aging time. These samples will be referred to as regional pollution (reg) samples hereafter. In the continental case, long-range transport contributes to the carbonaceous aerosol in addition to the regional emissions. The continental samples also contain particles that were emitted several days earlier, which results in a longer average atmospheric residence time of the collected particles. Due to the longer residence times the

aerosol carbon in the continental samples is on average more aged.

Table 2 shows $F^{14}C$ for the different seasons (winter: December, January, February; spring: March, April, May; summer: June, July, August; fall: September, October, November) and for the two main air mass types. The number of samples averaged for each case is given in the second column. Each of the samples was collected over several days and the total sampling days for each season and air mass condition are given in Table 2. The average $F^{14}C_{(EC)}$ in spring excludes the three EC samples that were influenced by primary biogenic material and therefore only three samples are considered. For all seasons EC is dominated by fossil sources and OC by modern sources. Water-insoluble OC has a considerably lower $F^{14}C$ than OC, which clearly indicates a higher contribution of fossil sources to WIOC. $F^{14}C$ of all carbon fractions is lowest in summer. The average values of $F^{14}C_{(OC)}$ varied by $\pm 4\%$ percent for all winter, summer, fall, and continental and regional air mass conditions, but the average $F^{14}C_{(OC)}$ in spring (0.89) was 20% higher than the average of the rest of the seasons (0.74). $F^{14}C_{(EC)}$ increased by almost 70% from summer to winter, likely caused by an increase in biomass combustion activities in winter. $F^{14}C_{(EC)}$ was on average 0.17 in air masses with regional pollution and around 0.3 in air masses with continental air mass origin.

The $F^{14}C$ values were used as input for source apportionment calculations to estimate OC and EC from biomass burning (OC_{bb} and EC_{bb}), OC and EC from fossil-dominated sources (OC_f and EC_f), and other contemporary OC ($OC_{c,o}$) with the Monte Carlo approach described in Sect. 2.9. Figure 4 shows a typical frequency distribution of OC_f , OC_{bb} , and $OC_{c,o}$ values, resulting from 10 000 individual calculations, each with randomly chosen input parameters from the ranges given in Sect. 2.9. These frequency distributions indicate the range and probability of possible values for each carbon fraction, taking into account all uncertainties in measurements and assumptions.

OC_f was constrained within 0.3 and $0.5 \mu\text{g m}^{-3}$, whereas estimates for OC_{bb} and $OC_{c,o}$ varied over a much wider range of roughly $0.5 \mu\text{g m}^{-3}$ each, reflecting the large uncertainty in r_{bb} . The individual estimates of OC_{bb} and $OC_{c,o}$ are not independent. Since the sum of OC_{bb} and $OC_{c,o}$ is narrowly constrained as $OC - OC_f$, large values of OC_{bb} correspond to small values of $OC_{c,o}$ and vice versa. Despite the considerable uncertainty in OC_{bb} and $OC_{c,o}$ some conclusions can be drawn from the distributions in Fig. 4, for example that $OC_{c,o}$ was the most abundant of the three components and that both OC_{bb} and $OC_{c,o}$ were clearly present in the sample.

The results of the source apportionment calculations are summarized in Table 3. The values represent averages for different seasons and air mass conditions and the corresponding standard deviations. Since Fig. 3 illustrates that the separation of OC_{bb} and $OC_{c,o}$ is rather uncertain, the sum of OC_{bb} and $OC_{c,o}$ is reported as contemporary OC

Table 2. Average fraction modern ($F^{14}C$) after blank correction \pm standard deviations of $F^{14}C$ determination for total carbon (TC), organic carbon (OC), elemental carbon (EC), and water-insoluble organic carbon (WIOC) in different seasons and for regional (reg) and continental (cont) air mass origin. The second column gives the number of samples averaged for each case.

	Samples	Days	$F^{14}C_{(TC)}$	$F^{14}C_{(OC)}$	$F^{14}C_{(WIOC)}$	$F^{14}C_{(EC)}$
Winter	$n = 5$	19	0.664 ± 0.068	0.742 ± 0.071	0.529 ± 0.093	0.262 ± 0.092
Spring	$n = 6$	23	0.760 ± 0.093	0.886 ± 0.052	0.660 ± 0.173	0.173 ± 0.095
Summer	$n = 6$	39	0.574 ± 0.073	0.713 ± 0.052	0.480 ± 0.127	0.157 ± 0.042
Fall	$n = 7$	31	0.627 ± 0.100	0.782 ± 0.058	0.595 ± 0.097	0.227 ± 0.079
Reg	$n = 13$	61	0.618 ± 0.065	0.766 ± 0.091	0.509 ± 0.088	0.170 ± 0.050
Cont	$n = 11$	51	0.698 ± 0.135	0.810 ± 0.082	0.637 ± 0.152	0.307 ± 0.107

Table 3. Average concentrations of OC, EC, WIOC, and WSOC and the respective contribution of contemporary (c) and of fossil-dominated (f) sources for different seasons and for regional (reg) and continental (cont) air mass conditions. The subscript c stands for all contemporary carbon, i.e., the sum of biomass burning and other contemporary carbon (e.g., $OC_c = OC_{bb} + OC_{c,o}$).

	Winter $C_{av} (\mu g m^{-3})$	Spring $C_{av} (\mu g m^{-3})$	Summer $C_{av} (\mu g m^{-3})$	Fall $C_{av} (\mu g m^{-3})$	Reg $C_{av} (\mu g m^{-3})$	Cont $C_{av} (\mu g m^{-3})$
TC	2.7 ± 3.0	3.6 ± 2.2	1.4 ± 0.5	4.1 ± 3.2	1.4 ± 0.7	4.9 ± 3.7
OC	2.1 ± 2.3	3.1 ± 2.0	1.0 ± 0.3	3.0 ± 2.3	1.0 ± 0.5	3.8 ± 2.9
OC_f	0.69 ± 0.78	0.46 ± 0.31	0.35 ± 0.11	0.88 ± 0.71	0.28 ± 0.10	1.0 ± 0.7
OC_c	1.4 ± 1.5	2.6 ± 1.6	0.68 ± 0.24	2.1 ± 1.6	0.76 ± 0.43	2.8 ± 2.1
EC	0.66 ± 0.72	0.51 ± 0.22	0.34 ± 0.16	1.2 ± 0.92	0.38 ± 0.18	1.1 ± 0.9
EC_f	0.51 ± 0.51	0.47 ± 0.17	0.33 ± 0.15	1.0 ± 0.78	0.35 ± 0.16	0.89 ± 0.69
EC_{bb}	0.15 ± 0.22	0.04 ± 0.05	0.02 ± 0.01	0.14 ± 0.15	0.02 ± 0.02	0.16 ± 0.18
WIOC	0.80 ± 0.88	0.66 ± 0.29	0.29 ± 0.12	1.2 ± 1.0	0.35 ± 0.15	1.3 ± 1.0
$WIOC_f$	0.44 ± 0.50	0.23 ± 0.07	0.17 ± 0.07	0.52 ± 0.36	0.19 ± 0.07	0.54 ± 0.43
$WIOC_c$	0.37 ± 0.38	0.43 ± 0.22	0.12 ± 0.05	0.70 ± 0.65	0.16 ± 0.07	0.74 ± 0.54
WSOC	1.3 ± 1.3	2.3 ± 1.5	0.73 ± 0.27	1.7 ± 1.3	0.70 ± 0.43	2.5 ± 1.4
$WSOC_f$	0.26 ± 0.29	0.23 ± 0.32	0.18 ± 0.07	0.36 ± 0.37	0.09 ± 0.11	0.45 ± 0.32
$WSOC_c$	1.0 ± 1.0	2.1 ± 1.2	0.55 ± 0.20	1.4 ± 1.0	0.60 ± 0.31	2.1 ± 1.1
OC / EC	4.0 ± 2.0	8.8 ± 3.7	3.2 ± 1.0	2.9 ± 1.1	3.2 ± 1.5	6.4 ± 3.5

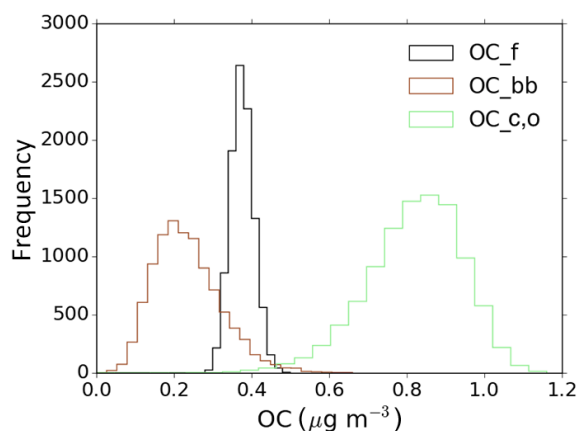


Figure 4. An example of the frequency distribution of OC concentrations from fossil-dominated, biomass burning, and biogenic sources for sample CA36, taken in June 2011. The concentrations of OC_{bb} and $OC_{c,o}$ are not independent, AND larger concentrations of OC_{bb} correspond to smaller concentrations in $OC_{c,o}$.

(OC_c). The seasonal variation is characterized by low concentrations in summer, with average TC concentrations of $1.4 \mu g m^{-3}$, which is less than half of the averages in other seasons. This can be attributed partially to rainy conditions and probably also to higher planetary boundary layers in summer. The main source of fossil elemental carbon (EC_f) in the Netherlands is traffic, which does not show a strong seasonal variation. Natural gas is mostly used for domestic heating, which produces very little aerosol. The concentrations of EC_f should therefore be relatively constant throughout the year. However, EC_f concentrations in the fall are 2 times higher than in spring and winter and 4 times higher than in summer. The concentrations OC_f are elevated in fall as well. This might be caused by easterly air masses that often contain higher levels of pollution than westerly ones from the sea and occur more frequently during fall. The contemporary OC fractions are elevated in spring: OC_c accounts for 70 % and $WSOC_c$ for 60 % of TC in spring, whereas OC_c accounts only for roughly 50 % and $WSOC_c$ for 35–40 % of TC in other seasons. EC_{bb} is more than 4 times higher in winter and fall than in spring and summer 2011. $WSOC_f$ is

lower than $WSOC_c$ in all seasons except for summer. The standard deviations reflect the variability of pollution levels, which in winter are higher than the mean value. In summary, there are some indications of a seasonal variation in carbon concentrations, but the variability within each season is high and strongly influenced by weather and air mass conditions.

Comparing the concentrations for different air mass conditions shows clearer differences. For regional pollution carbon concentrations are rather low and less variable than seasonal averages. If we assume that aerosol concentrations from long-range transport are superimposed on the regional background, then the increase in aerosol concentrations during continental air mass conditions gives an indication of the importance of long-range transport for the carbonaceous aerosol concentrations in the Netherlands. Carbon fractions that only show a small increase under continental air mass conditions are mostly of regional origin, whereas for carbon fractions that increase more strongly, regional sources are less prominent.

The concentrations of most carbon fractions increase by a factor of 3.4–3.6 in continental compared to regional air masses. Notable exceptions are fossil EC and fossil WIOC, which only increase by a factor of approximately 2.5. The biggest source of fossil EC and fossil WIOC is traffic emissions. Our data therefore indicate that the regional contribution to OC and EC from traffic sources is higher than for other carbon fractions and the influence of long-range transport less important. In contrast, EC_{bb} , $WIOC_c$, and $WSOC_f$ increase by more than a factor of 4 under continental air mass conditions. Especially concentrations of EC_{bb} are 8 times lower in regional than in continental pollution, suggesting that most of EC_{bb} originates from outside the Netherlands. Since primary OC from fossil sources is usually water insoluble, $WSOC_f$ is most likely of secondary origin. Its concentration increases considerably in continental air masses that contain emissions from more distant sources after a longer aging time in the atmosphere. $WIOC_c$ and its potential origin will be discussed in more detail below.

Differences in precipitation are a first-order effect that could explain part of the difference between carbon concentrations during regional and continental air masses conditions. Usually precipitation is higher under westerly wind directions (i.e., marine modified) than under the easterly wind directions that are associated with continental air mass conditions. Wet removal of aerosol strongly lowers aerosol concentrations during precipitation events (e.g., Ohata et al., 2016). Therefore we compared the average duration and amount of precipitation measured at the Cesar Observatory during the sampling times of filters representative for regional and continental air mass conditions. The highest TC concentrations above $5 \mu\text{g m}^{-3}$ occur only on dry days, but on dry days concentrations below $2 \mu\text{g m}^{-3}$ are also measured frequently. The rainfall duration was on average 1 h day^{-1} for continental conditions and 2 h day^{-1} for regional conditions and the amount was 1.2 vs. 3.4 mm day^{-1} .

Even though there is on average more precipitation under regional air mass conditions, one additional hour of rain per day is unlikely to reduce aerosol concentrations by a factor of 3. Furthermore, the continental/regional ratios are different for EC_{bb} and EC_f and it is not likely that these compounds would be scavenged differently. Therefore, the influence of precipitation is unlikely to be the main reason for the difference in carbonaceous aerosol concentrations between air mass conditions. Also the wind speed is not much lower during continental air mass conditions. The wind speeds averaged around 3.5 m s^{-1} under continental air mass conditions and 4.5 m s^{-1} under regional air mass conditions. Despite the fact that higher wind speeds are usually associated with a higher boundary layer and more diluted pollutants, there was not a very clear relationship between TC concentrations and wind speed. For example, the highest TC concentrations of $9.5 \mu\text{g m}^{-3}$ were measured in a period with relatively low wind average speed (2.9 m s^{-1}), but almost equally high concentrations of $8.2 \mu\text{g m}^{-3}$ were measured in a period with above average wind speeds of 4.4 m s^{-1} . The three lowest TC concentrations (below $1 \mu\text{g m}^{-3}$) were measured during time periods with average wind speeds (3.7 m s^{-1}), high wind speeds (8.3 m s^{-1}), and slightly elevated wind speeds (4.8 m s^{-1}). Therefore we think it is unlikely that meteorological conditions are mainly responsible for the differences in carbon concentrations under regional and continental air mass conditions.

Figure 5 shows the concentrations of OC and EC apportioned to the various sources averaged over different seasons as colored bars and the concentrations corresponding to the individual filter samples as black dots. The red lines show the medians for each season. The error bars show uncertainties of the average calculated by propagating the uncertainties of the individual data points, obtained by the Monte Carlo simulation. These error bars therefore reflect the methodological uncertainties rather than variability of the data. These methodological uncertainties are smaller than the standard deviation of the individual measurement points that were reported in Table 3. The strong variability in the concentrations of EC and OC fractions therefore primarily reflects the actual differences in atmospheric concentrations and not the uncertainties of our measurements and assumptions.

Figure 5a shows that the high average concentration of EC_f in fall is mainly due to three pollution events. A similar pollution event also occurred in winter. This example shows the importance of occasional pollution events for aerosol concentrations in the Netherlands. Constructing seasonal averages therefore requires a large number of samples that also capture typical frequencies of pollution events. This would require sampling over several years and careful selection of samples to represent average meteorological conditions and air mass origin for the respective season. This was not possible for this study and therefore the seasonal variation should not be overinterpreted.

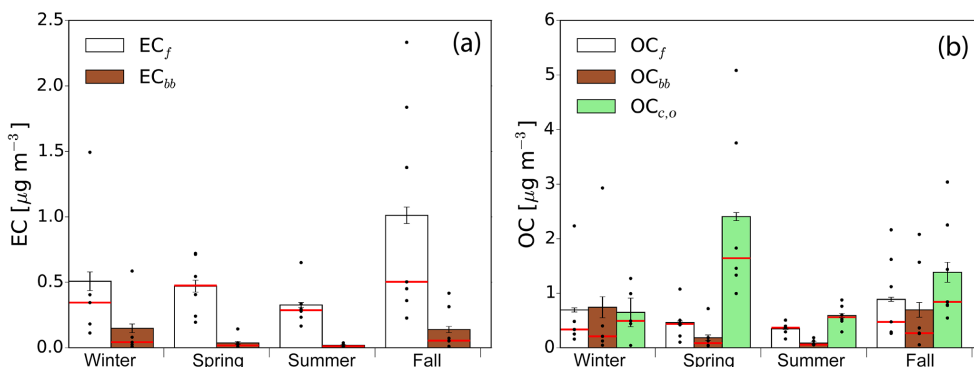


Figure 5. Concentrations of different subfractions of OC and EC from various sources averaged for winter, spring, summer, and fall in 2011/12. The bars show the average concentrations and the error bars the propagated uncertainties from the Monte Carlo simulations. The red line shows the median concentration. The individual data points are shown as black dots. (a) EC from fossil-dominated sources (EC_f) and from biomass combustion (EC_{bb}); (b) OC from fossil-dominated sources (OC_f), biomass combustion (OC_{bb}), and biogenic sources ($OC_{c,o}$).

The medians are less influenced by outliers and most carbon fractions in Fig. 5a and b show a much weaker seasonal variation in the median than in the mean. However, the median $OC_{c,o}$ is still clearly elevated in spring. The two samples with very high $OC_{c,o}$ concentrations correspond to two of the three pollen events. EC_{bb} and OC_{bb} show also clear seasonal variation with low median concentrations in spring and summer, when domestic biomass combustion is not a major source of aerosol carbon in western Europe.

Figure 6 shows the concentrations of OC and EC during regional and continental air mass conditions. Pollution episodes characterized by high carbon concentrations occur only during continental air mass conditions. When pollutants originate in and around the Netherlands, the concentrations of EC and OC are below $0.5 \mu\text{g m}^{-3}$ for EC and below $1 \mu\text{g m}^{-3}$ for OC_f , OC_{bb} , and $OC_{c,o}$, respectively, with average values less than half of the average concentrations encountered during continental air mass conditions. OC and EC concentrations are also less variable in recent pollution with relative standard deviations roughly 50% of the mean value of most carbon fractions, whereas standard deviations nearly approach the mean value for many carbon fractions continental air mass conditions. Carbon concentrations comparable to regional air mass conditions occur for EC_f , EC_{bb} , and OC_{bb} in 50–60% of the continental samples and for OC_f and $OC_{c,o}$ in 30–40% of the continental samples. This shows that despite higher average concentrations, continental air mass conditions do not always carry high concentrations of carbonaceous aerosol concentrations to the Netherlands. The concentrations under continental air mass conditions are not correlated with precipitation or wind speed, and therefore likely depend on the air mass origin.

EC_{bb} and OC_{bb} are low in regional pollution, even in winter, suggesting low biomass combustion emission from within the Netherlands. Under continental air mass conditions, carbon concentrations from biomass combustion are

elevated during fall and winter. Very low concentrations of EC_{bb} and OC_{bb} occur in summer and spring, when residential biomass combustion is low throughout Europe.

The higher average concentrations of fossil carbon in continental air masses are mainly caused by pollution events and the median concentrations are comparable. However, the median concentrations of $OC_{c,o}$ are clearly higher under continental than under regional air mass conditions.

3.3 Source apportionment of water-soluble and water-insoluble OC

Figure 7a shows the contribution of $WSOC_f$, $WIOC_f$, and $WIOC_c$ to the total organic carbon in different seasons during 2011/12. The fraction of $WSOC$ is the sum of the blue areas, which accounts for about three-quarters of the total OC in spring and summer and roughly two-thirds in fall and winter. Throughout the year, OC is dominated by contemporary $WSOC$, which reflects that the two main sources of contemporary OC, namely biomass combustion and biogenic SOA, are largely water soluble. $WIOC$ consists of roughly equal parts of fossil and contemporary carbon.

The contributions of fossil and contemporary carbon fractions to OC (Fig. 7b) stay within 5% (absolute) for different air mass origins, even though the average concentrations of OC increased from $1 \mu\text{g m}^{-3}$ in regional to almost $4 \mu\text{g m}^{-3}$ in continental air masses.

The sources of $WIOC_c$, which contributes up to a quarter of the total OC in this study, are currently not very well known. Figure 8 shows a positive correlation between the concentrations of $WIOC_c$ and EC_{bb} . The three points marked in red correspond to the pollen events, where high $F^{14}C_{(EC)}$ values were caused by refractory primary biogenic aerosol. EC_{bb} concentrations were corrected for this effect as described in Sect. 3.1. $WIOC_c$ is high for the three samples collected during pollen events, which shows that primary bi-

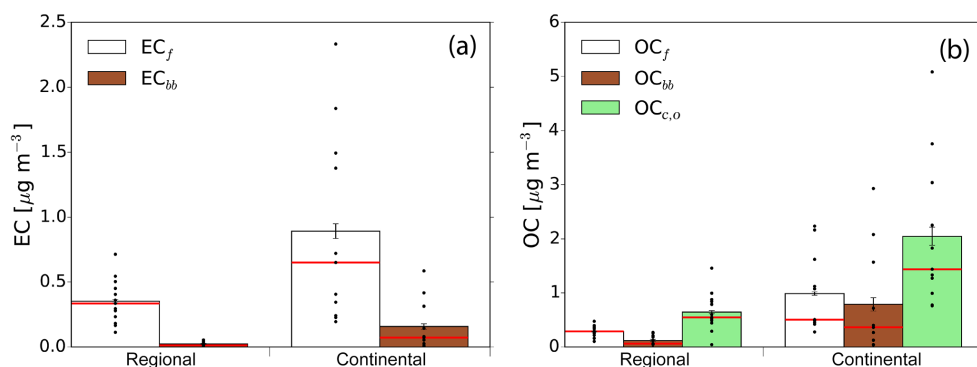


Figure 6. Concentrations of different subfractions of OC and EC from various sources in the time period from February 2011 to March 2012 averaged for regional and continental air mass origin. The bars show the average concentrations and the error bars the propagated uncertainties from the Monte Carlo simulations. The red line shows the median concentration. The individual data points are shown as black dots. **(a)** EC from fossil-dominated sources (EC_f) and from biomass combustion (EC_{bb}); **(b)** OC from fossil-dominated sources (OC_f), biomass combustion (OC_{bb}), and biogenic sources ($OC_{c,o}$).

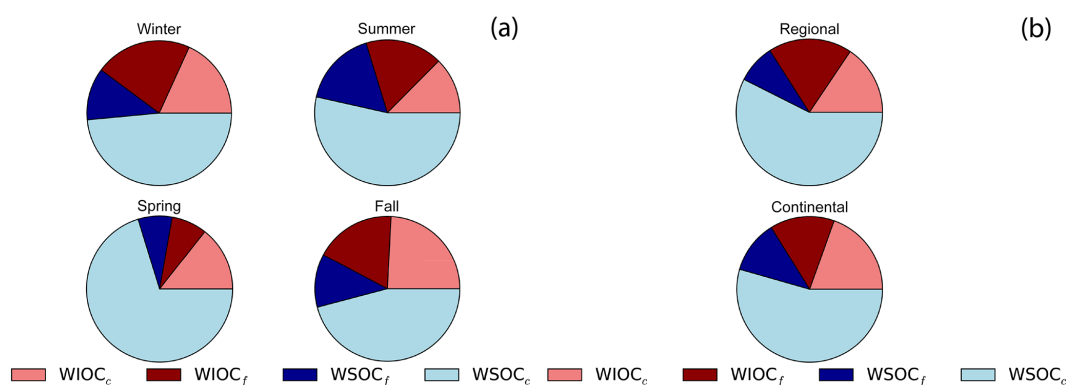


Figure 7. The contribution of water-soluble and water-insoluble organic carbon from fossil-dominated sources ($WSOC_f$ and $WIOC_f$) as well as water-soluble and water-insoluble organic carbon from contemporary sources ($WSOC_c$ and $WIOC_c$) to the total organic carbon. **(a)** Contribution of different carbon fractions during winter, summer, spring, and fall. **(b)** Contribution of different carbon fractions for modified marine and continental air mass origin.

ological material can contribute significantly to the water-insoluble carbon.

A linear least squares fit to the data, excluding the three highest data points and the pollen events, yields an intercept of 0.10 ± 0.02 and an R^2 of 0.74. The offset is small, corresponding to approximately $0.1 \mu\text{g m}^{-3}$ of $WIOC_c$ that is independent of EC_{bb} . In other words, a large part of the variability in contemporary $WIOC$ in the Netherlands seems to be associated with biomass combustion. There is no strong evidence of a major contribution of biogenic SOA to $WIOC$, which should be highest in spring and summer, when EC_{bb} and $WIOC_c$ concentrations are both small.

Figure 9 shows a scatter plot of $WIOC_f$ and EC_f concentrations. EC_f is emitted by the combustion of fossil fuels and a positive correlation of $WIOC_f$ with EC_f implies that primary emissions by fossil fuel combustion are an important source for $WIOC_f$ as well. There is one outlier data point, for which it is possible that the EC concentration calculated by

Eqs. (1) and (2) is overestimated. A linear regression excluding the four data points with highest EC_f concentrations has a slope of 0.32, an intercept of 0.1, and an R^2 of 0.68. In contrast, a similar linear regression of $WSOC_f$ against EC_f yields an R^2 of 0.01, indicating that fossil water-soluble $WSOC$ does not have a common source with EC_f . If the four highest data points are included the R^2 is generally higher (0.92 for $WIOC_f$ and 0.48 for $WSOC_f$), but this is mainly due to the fact that for these four samples TC concentrations are much higher than average (ranging from 5 to $10 \mu\text{g m}^{-3}$) and this leads to both higher OC and EC concentrations in general. It is not good practice to fit such bimodal data with a linear regression. The low intercept indicates that primary emissions from fossil fuel combustion is the predominant source of fossil $WIOC$. The slope of such a regression line can give real world constraints on OC / EC emissions ratios of fossil sources; in this case we can suggest that OC / EC ratios of an integrated fossil source should be below 0.8.

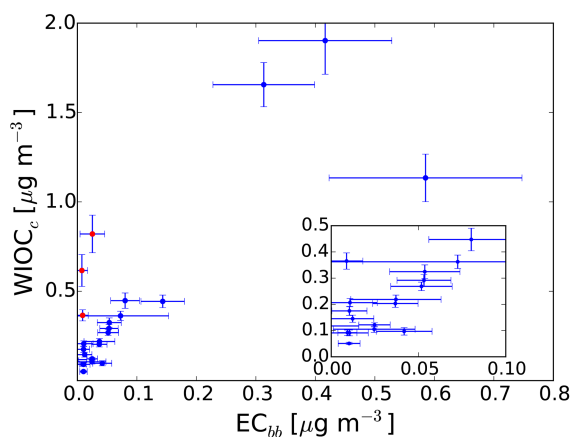


Figure 8. Scatter plot of contemporary water-insoluble carbon concentrations ($WIOC_c$) against EC derived from biomass burning (EC_{bb}). EC_{bb} has been corrected for the contribution of highly refractory primary OC during three pollen events (red points).

In the lower concentration range there is considerable scatter in the data points and individual $WIOC_f/EC_f$ is in the range of 0.4 to 1. However, compared to the typical variability of ambient OC/EC ratios, which can range from 1 to 10 or higher in remote locations (e.g., Khan et al., 2016; Sandrini et al., 2014, and references therein), or even of OC_f/EC_f ratios, this range of variability is relatively low. The values of $WIOC_f/EC_f$ measured in this study are comparable to OC/EC ratios of primary fossil emissions, such as vehicular traffic in tunnel studies (e.g., Chirico et al., 2011). OC/EC emission ratios of coal combustion can be in the same range or higher (e.g., Chen et al., 2015). Coal combustion is a negligible source in and around the Netherlands, but it might be important during long-range transport events of pollution from eastern Europe. $WIOC_f/EC_f$ values measured during this study were overall higher in winter than in the other seasons.

Average $WIOC_f/EC_f$ ratios are similar under regional pollution (0.6 ± 0.2) and under continental air mass conditions (0.6 ± 0.3). This indicates that $WIOC_f/EC_f$ ratios might stem from similar sources in both cases. The $WIOC_f/EC_f$ ratio is 0.6 ± 0.3 averaged over all samples, which might be indicative for an integrated primary OC/EC emission ratio for fossil sources in western Europe.

$WSOC_f$ is usually thought to result mainly from secondary formation. The ratio of $WSOC_f/EC_f$ is 0.44 ± 0.46 under regional air mass conditions, which sample relatively fresh emissions. It increases to 0.6 ± 0.4 under continental air mass conditions, where older and more processed aerosol is sampled. The difference is significant at the 90% confidence level but not at the 95% confidence level ($p = 0.06$). In general the large variability of the $WSOC_f/EC_f$ ratio indicates that $WSOC_f$ and EC_f do not originate from a common source.

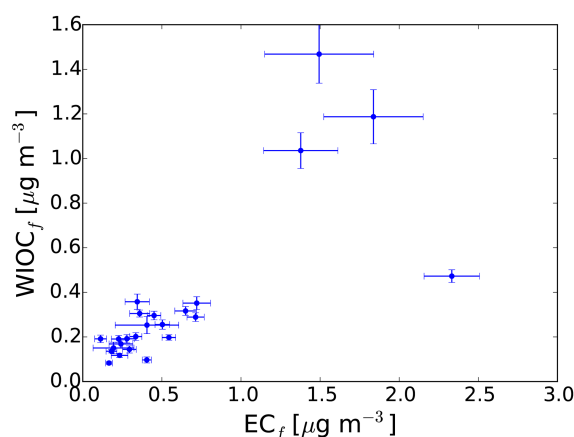


Figure 9. A scatter plot of EC concentrations from fossil-dominated sources (EC_f) versus concentrations of water-insoluble fossil carbon ($WIOC_f$).

4 Discussion: aerosol in the Netherlands compared to other regions

The average $F^{14}C_{(TC)}$ measured in the Netherlands over the course of a year is approximately 0.65. This is in the range of $F^{14}C_{(TC)}$ measured at different locations throughout Europe, roughly 0.5 to 0.85, depending on location and season (Heal, 2014, and references therein). $F^{14}C_{(TC)}$ values are usually at the higher end of the range in rural locations and at the lower end in urban areas. $F^{14}C_{(TC)}$ at our polluted regional background site lies roughly in the middle of this range, corresponding to expectations. $F^{14}C_{(TC)}$ values in this study were slightly lower in conditions dominated by regional pollution (0.6) than in conditions influenced by long-range transport (0.7). This indicates that aerosol carbon in the Netherlands is more strongly influenced by fossil sources than aerosol carbon transported from southern and central Europe. In the United States, $F^{14}C_{(TC)}$ values for suburban and rural sites are usually higher (0.7–1; Lewis and Stiles, 2006; Lewis et al., 2004; Schichtel et al., 2008). In Japan, $F^{14}C_{(TC)}$ is usually lower and ranged from 0.3 to 0.5 in studies conducted mostly in urban or suburban areas (Fushimi et al., 2011; Heal, 2014, and references therein; Morino et al., 2010). In China, $F^{14}C_{(TC)}$ can vary strongly by location from predominantly fossil in large urban centers (Zhang et al., 2015) to mainly biogenic in some rural areas (Zhang et al., 2014).

In this study $F^{14}C_{(TC)}$ was highest in spring (0.76) and lowest in summer (0.57). There are not many studies in which data were compared for the same site in different seasons, and most of them analyzed only a few samples per season, making them vulnerable for sampling bias, as discussed above for this study. However, a general survey seems to indicate that for sites situated in cleaner regions, such as Scandinavia (Genberg et al., 2011; Szidat et al., 2009; Yttri et al., 2011a), rural Spain (Minguillón et al., 2011), the rural US (Bench, 2004; Schichtel et al., 2008), and several other rural

background sites throughout Europe (Gelencsér et al., 2007), $F^{14}C_{(TC)}$ was lower in winter than in summer. In these locations, biogenic SOA presumably contributed significantly to the total aerosol concentrations in summer. In more polluted locations, such as the Po Valley (Gilardoni et al., 2011), Barcelona (Minguillón et al., 2011), and Tokyo (Minoura et al., 2012), $F^{14}C_{(TC)}$ tended to be higher in winter than in summer. In Birmingham (Heal et al., 2011), there was no significant seasonal variation in $F^{14}C_{(TC)}$. The results of our study are more in line with the observations at polluted sites, which seems reasonable given the high population and traffic density of the surrounding urban areas. However, the results of our study might also be influenced by the unusually high amount of precipitation and low solar radiation during the summer of 2011.

$F^{14}C_{(OC)}$ was around 0.8, which is on the higher end of several studies conducted in Europe, where $F^{14}C_{(OC)}$ varied between 0.55 and 0.8 (Ceburnis et al., 2011; Dusek et al., 2013; Heal et al., 2011; Minguillón et al., 2011; Szidat et al., 2006). However, most of these values were measured in urban areas and it is likely that $F^{14}C_{(OC)}$ is higher than that range at rural or regional sites. $F^{14}C_{(OC)}$ did not vary strongly between air mass conditions and seasons in our study, and other measurements in Europe have shown no or weak variation between the seasons (Heal, 2014, and references therein).

If the samples influenced by primary biogenic material are disregarded, the average $F^{14}C_{(EC)}$ measured at the Cesar site is 0.2, with on average 0.15 in summer and 0.25 in winter. This is in the range of or slightly higher than $F^{14}C_{(EC)}$ values reported for several European cities, such as Zurich (Szidat et al., 2006), Birmingham (Heal et al., 2011), Barcelona (Minguillón et al., 2011), or Göteborg (Szidat et al., 2009), but lower than $F^{14}C_{(EC)}$ measured in several background regions, especially in winter, such as ~ 0.4 in rural Sweden (Szidat et al., 2009), ~ 0.3 in rural Spain (Minguillón et al., 2011), and 0.3 to 0.8 during wintertime smog episodes in Switzerland (Zotter et al., 2014b). In addition to the limited number of studies that directly measured $F^{14}C_{(EC)}$ for $PM_{2.5}$ in Europe, several studies estimated the contribution of biomass burning to EC based on $F^{14}C_{(TC)}$ and other tracers, such as levoglucosan. In comparison with these studies we can conclude that the wintertime contribution of biomass burning to EC in the Netherlands is lower than in many other regions in Europe (e.g., Genberg et al., 2011; Gilardoni et al., 2011; Yttri et al., 2009). Additionally, our study shows that $F^{14}C_{(EC)}$ increases from 0.2 under regional pollution to 0.3 under continental air mass conditions. This is consistent with long-range transport of carbonaceous aerosol from other regions in Europe with higher $F^{14}C_{(EC)}$.

Relatively few studies determined $F^{14}C_{(WIOC)}$. In several locations in Europe WIOC contained roughly equal fractions of contemporary and fossil carbon (Szidat et al., 2007, 2009), comparable to our study. This is, however, not necessarily the case in other regions of the world. In Mexico city, where OC

is dominated by fossil emissions (Aiken et al., 2010), WIOC is also mainly fossil. At a regional background site in south-eastern China, where biomass burning is a strong source, WIOC is mainly modern (Zhang et al., 2014).

$WIOC_f / EC_f$ ratios determined in Switzerland or Sweden were comparable to our values ranging from roughly 0.5 to 1 (Szidat et al., 2004, 2009); the ratios were not directly reported but estimated from graphs in these publications. Two studies done in China report higher $WIOC_f / EC_f$ ratios of around 1 (Zhang et al., 2014) and in the range from 1 to 2.5 in southeastern China (Liu et al., 2013). These high values could be explained by less efficient combustion in an older vehicle fleet and also higher OC / EC emission ratios from coal burning that is much more common in China than in western Europe.

In contrast, WSOC is dominated by modern sources in all regions of the globe with usually only 0–20 % contributions from fossil sources (e.g., Kirillova et al., 2010, 2013, 2014; Szidat et al., 2006, 2009; Wozniak et al., 2012). The data from the Cesar site fall in this range with a fossil fraction of WSOC below 0.2 (Fig. 7).

We also determined the ratio of water-soluble to water-insoluble fossil carbon $(WSOC / WIOC)_f$. This ratio is higher in spring and summer than in winter (1.2 vs. 0.6). It is also higher in aged continental air mass conditions than in regional pollution (1.2 vs. 0.7). Szidat et al. (2009) also noticed that the fractional contribution of WSOC to fossil OC increases in summer in Scandinavia. This is consistent with a secondary origin of $WSOC_f$. SOA formation is usually enhanced in spring and summer due to higher solar radiation. Longer atmospheric aging times of primary emissions usually lead to higher SOA concentrations. If $WSOC_f$ is regarded as a proxy for SOA_f , then the above $(WSOC / WIOC)_f$ ratios indicate that secondary formation contributes roughly 50 % to fossil OC in spring and summer versus 40 % in winter and 50 % in continental air masses versus approximately 30 % in regional pollution. Much higher contributions of SOA_f (around 70 %) to fossil OC have been found in southern China (Zhang et al., 2014). Zotter et al. (2014b) observed that SOA_f concentrations are larger in the Los Angeles area than in European cities, in contrast to EC_f concentrations, which are higher in Europe. They attributed this to the higher fraction of diesel cars in Europe, which emit a larger fraction of EC and less SOA precursors than gasoline cars. Fushimi et al. (2010) concluded that in suburban Tokyo, fossil sources make a dominant contribution to the very high SOA concentrations observed in daytime. SOA concentrations in that study were up to 10 times higher than primary OC concentrations, so that fossil OA was predominantly of secondary origin. Even though secondary formation contributes significantly to fossil OA in the Netherlands, compared to other, more photochemically active, regions the contribution is relatively moderate.

5 Conclusions

This study presents source apportionment of carbonaceous aerosol at a regional site in the Netherlands over the course of a whole year. Radiocarbon was measured in several carbon fractions of $PM_{2.5}$, namely TC, OC, WIOC, and EC. $F^{14}C_{(EC)}$ is strongly correlated with the fraction of levoglucosan in TC, except for three samples in spring time, which were collected under very high pollen concentrations. Our results suggest that refractory primary biogenic material can survive the OC removal steps and bias $F^{14}C_{(EC)}$ to higher values.

$F^{14}C$ values of all carbon fractions are on average lower for regional pollution, which indicates that aerosol carbon in the Netherlands contains a higher contribution from fossil sources than the aerosol transported to the Netherlands. Overall, the highest $F^{14}C$ values were measured in spring and the lowest $F^{14}C$ values in summer. However, there are only a few data points per season, so specific meteorological conditions, such as the very wet summer of 2011, may have strongly affected the differences between seasons, which may not be representative for a multiyear mean.

The concentrations of all carbonaceous fractions were on average higher in continental air masses influenced by long-range transport than for regional pollution. The difference in TC concentrations was on average a factor of 3. However, the average concentrations in continental air masses were strongly influenced by a few pollution episodes with concentrations much higher than for regional pollution. This highlights the importance of long-range transport that episodically caused high TC concentrations in the Netherlands. Fossil EC and fossil WIOC concentrations increased less than TC in continental air masses. We conclude that these carbon fractions have a larger relative contribution from regional sources than TC. Carbonaceous aerosol from biomass burning and fossil WSOC increased strongly in continental air masses, suggesting major sources outside the Netherlands. Fossil WSOC can be attributed mainly to secondary formation and the longer aging times during long-range transport lead to higher concentrations of secondary carbon.

Fossil WIOC/EC was on average 0.6, well within the range of primary OC/EC ratios from vehicular emissions. This ratio did not change between regional and continental air masses. One of the most interesting results of our study is that, even though a large fraction of carbon emitted by biomass burning is water soluble, long-range transport of biomass smoke acts to significantly increase the rather low background concentrations of WIOC_c (around $0.1 \mu\text{g m}^{-3}$) in the Netherlands. This can be concluded from the strong correlation of WIOC_c with EC_{bb} and that a strong increase in EC_{bb} and WIOC_c only happens during continental air mass conditions. However, it is not clear if this holds in other regions of the world and especially for other particle size ranges, such as PM_{10} or total suspended particles. In these

size ranges primary biogenic material can also contribute to WIOC_c.

6 Data availability

The data are available at <http://www.rug.nl/research/isotope-research/projects/public-data-set> (Dusek, 2017).

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

Acknowledgements. This work was funded by the Dutch Science Foundation (NWO, grant no. 820.01.001). We would like to thank Dicky van Zonneveld, Henk Been, and Anita Aerts-Bijma for their ^{14}C analysis work. We gratefully acknowledge the work of Mattia Monaco and Arthur Kappetijn, who contributed to sampling and analysis as part of their masters' projects. The authors gratefully acknowledge the NOAA Air Resources Laboratory (ARL) for the provision of the HYSPLIT transport and dispersion model and/or READY website (<http://www.ready.noaa.gov>) used in this publication.

Edited by: E. Weingartner

Reviewed by: two anonymous referees

References

- Aiken, A. C., de Foy, B., Wiedinmyer, C., DeCarlo, P. F., Ulbrich, I. M., Wehrli, M. N., Szidat, S., Prévôt, A. S. H., Noda, J., Wacker, L., Volkamer, R., Fortner, E., Wang, J., Laskin, A., Shutthanandan, V., Zheng, J., Zhang, R., Paredes-Miranda, G., Arnott, W. P., Molina, L. T., Sosa, G., Querol, X., and Jimenez, J. L.: Mexico city aerosol analysis during MILAGRO using high resolution aerosol mass spectrometry at the urban supersite (T0) – Part 2: Analysis of the biomass burning contribution and the non-fossil carbon fraction, *Atmos. Chem. Phys.*, 10, 5315–5341, doi:10.5194/acp-10-5315-2010, 2010.
- Allan, J. D., Williams, P. I., Morgan, W. T., Martin, C. L., Flynn, M. J., Lee, J., Nemitz, E., Phillips, G. J., Gallagher, M. W., and Coe, H.: Contributions from transport, solid fuel burning and cooking to primary organic aerosols in two UK cities, *Atmos. Chem. Phys.*, 10, 647–668, doi:10.5194/acp-10-647-2010, 2010.
- Bench, G.: Measurement of Contemporary and Fossil Carbon Contents of PM Aerosols?: Results from Turtleback Dome, Yosemite National Park, *Environ. Sci. Technol.*, 38, 2424–2427, 2004.
- Bennett, M., Volckens, J., Stanglmaier, R., McNichol, A. P., Ellenson, W. D., and Lewis, C. W.: Biodiesel effects on particulate radiocarbon (^{14}C) emissions from a diesel engine, *J. Aerosol Sci.*, 39, 667–678, 2008.
- Bernardoni, V., Calzolari, G., Chiari, M., Fedi, M., Lucarelli, F., Nava, S., Piazzalunga, A., Riccobono, F., Taccetti, F., Valli, G., and Vecchi, R.: Radiocarbon analysis on organic and elemental carbon in aerosol samples and source apportionment at an urban site in Northern Italy, *J. Aerosol Sci.*, 56, 88–99, 2013.

- Birch, M. E. and Cary, R. A.: Elemental carbon-based method for monitoring occupational exposures to particulate diesel exhaust, *AS&T*, 25, 221–241, 1996.
- Ceburnis, D., Garbaras, A., Szidat, S., Rinaldi, M., Fahrni, S., Perron, N., Wacker, L., Leinert, S., Remeikis, V., Facchini, M. C., Prévôt, A. S. H., Jennings, S. G., Ramonet, M., and O'Dowd, C. D.: Quantification of the carbonaceous matter origin in sub-micron marine aerosol by ^{13}C and ^{14}C isotope analysis, *Atmos. Chem. Phys.*, 11, 8593–8606, doi:10.5194/acp-11-8593-2011, 2011.
- Chen, Y., Tian, C., Feng, Y., Zhi, G., Li, J., and Zhang, G.: Measurements of emission factors of $\text{PM}_{2.5}$, OC, EC, and BC for household stoves of coal combustion in China, *Atmos. Environ.*, 109, 190–196, 2015.
- Chin, M., Diehl, T., Tan, Q., Prospero, J. M., Kahn, R. A., Remer, L. A., Yu, H., Sayer, A. M., Bian, H., Geogdzhayev, I. V., Holben, B. N., Howell, S. G., Huebert, B. J., Hsu, N. C., Kim, D., Kucsera, T. L., Levy, R. C., Mishchenko, M. I., Pan, X., Quinn, P. K., Schuster, G. L., Streets, D. G., Strode, S. A., Torres, O., and Zhao, X.-P.: Multi-decadal aerosol variations from 1980 to 2009: a perspective from observations and a global model, *Atmos. Chem. Phys.*, 14, 3657–3690, doi:10.5194/acp-14-3657-2014, 2014.
- Chirico, R., Prévôt, A. S. H., DeCarlo, P. F., Heringa, M. F., Richter, R., Weingartner, E., and Baltensperger, U.: Aerosol and trace gas vehicle emission factors measured in a tunnel using an Aerosol Mass Spectrometer and other on-line instrumentation, *Atmos. Environ.*, 45, 2182–2192, 2011.
- Chow, J. C., Watson, J. G., Kuhns, H., Etyemezian, V., Lowenthal, D. H., Crow, D., Kohl, S. D., Engelbrecht, J. P., and Green, M. C.: Source profiles for industrial, mobile, and area sources in the Big Bend Regional Aerosol Visibility and Observational study, *Chemosphere*, 54, 185–208, 2004.
- Christian, T. J., Yokelson, R. J., Cárdenas, B., Molina, L. T., Engling, G., and Hsu, S.-C.: Trace gas and particle emissions from domestic and industrial biofuel use and garbage burning in central Mexico, *Atmos. Chem. Phys.*, 10, 565–584, doi:10.5194/acp-10-565-2010, 2010.
- Currie, L. A.: Evolution and multidisciplinary frontiers of ^{14}C aerosol science, *Radiocarbon*, 42, 115–126, 2000.
- de Rooij, M., van der Plicht, J., and Meijer, H. A. J.: Porous iron pellets for AMS ^{14}C analysis of small samples down to ultramicroscale size (10–25 μgC), *Nucl. Instruments Methods Phys. Res. B*, 268, 947–951, 2010.
- Dusek, U.: ^{14}C Public data set, University of Groningen, available at: <http://www.rug.nl/research/isotope-research/projects/public-data-set>, 2017.
- Dusek, U., ten Brink, H. M., Meijer, H. A. J., Kos, G., Mrozek, D., Röckmann, T., Holzinger, R., and Weijers, E. P.: The contribution of fossil sources to the organic aerosol in the Netherlands, *Atmos. Environ.*, 74, 169–176, 2013.
- Dusek, U., Monaco, M., Prokopiou, M., Gongriep, F., Hitzemberger, R., Meijer, H. A. J., and Röckmann, T.: Evaluation of a two-step thermal method for separating organic and elemental carbon for radiocarbon analysis, *Atmos. Meas. Tech.*, 7, 1943–1955, doi:10.5194/amt-7-1943-2014, 2014.
- Fushimi, A., Tanabe, K., Ohara, T., Hasegawa, S., Uchida, M., Takami, A., Yokouchi, Y., and Kobayashi, S.: Contrasting Diurnal Variations in Fossil and Nonfossil Secondary Organic Aerosol in Urban Outflow, Japan, *Environ. Sci. Technol.*, 44, 8581–8586, 2010.
- Fushimi, A., Wagai, R., Uchida, M., Hasegawa, S., Takahashi, K., Kondo, M., Hirabayashi, M., Morino, Y., Shibata, Y., Ohara, T., Kobayashi, S., and Tanabe, K.: Radiocarbon (^{14}C) diurnal variations in fine particles at sites downwind from Tokyo, Japan in summer, *Environ. Sci. Technol.*, 45, 6784–6792, 2011.
- Fuzzi, S., Andreae, M. O., Huebert, B. J., Kulmala, M., Bond, T. C., Boy, M., Doherty, S. J., Guenther, A., Kanakidou, M., Kawamura, K., Kerminen, V.-M., Lohmann, U., Russell, L. M., and Pöschl, U.: Critical assessment of the current state of scientific knowledge, terminology, and research needs concerning the role of organic aerosols in the atmosphere, climate, and global change, *Atmos. Chem. Phys.*, 6, 2017–2038, doi:10.5194/acp-6-2017-2006, 2006.
- 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. 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, 8217–8299, doi:10.5194/acp-15-8217-2015, 2015.
- Gelencsér, A., May, B., Simpson, D., Sánchez-Ochoa, A., Kasper-Giebl, A., Puxbaum, H., Caseiro, A., Pio, C., and Legrand, M.: Source apportionment of $\text{PM}_{2.5}$ organic aerosol over Europe: Primary/secondary, natural/anthropogenic, and fossil/biogenic origin, *J. Geophys. Res.*, 112, D23S04, doi:10.1029/2006JD008094, 2007.
- Genberg, J., Hyder, M., Stenström, K., Bergström, R., Simpson, D., Fors, E. O., Jönsson, J. Å., and Swietlicki, E.: Source apportionment of carbonaceous aerosol in southern Sweden, *Atmos. Chem. Phys.*, 11, 11387–11400, doi:10.5194/acp-11-11387-2011, 2011.
- Gilardoni, S., Vignati, E., Cavalli, F., Putaud, J. P., Larsen, B. R., Karl, M., Stenström, K., Genberg, J., Henne, S., and Dentener, F.: Better constraints on sources of carbonaceous aerosols using a combined ^{14}C – macro tracer analysis in a European rural background site, *Atmos. Chem. Phys.*, 11, 5685–5700, doi:10.5194/acp-11-5685-2011, 2011.
- Godwin, H.: Half-life of Radiocarbon, *Nature*, 195, 984, 1962.
- Gustafsson, Ö., Kruså, M., Zencak, Z., Sheesley, R. J., Granat, L., Engström, E., Praveen, P. S., Rao, P. S. P., Leck, C., Rodhe, H., and Asia, S.: Brown Clouds over South Asia?: Biomass or Fossil Fuel Combustion??, *Science*, 323, 495–498, 2009.
- El Haddad, I., Marchand, N., Temime-Roussel, B., Wortham, H., Piot, C., Besombes, J.-L., Baduel, C., Voisin, D., Armengaud, A., and Jaffrezo, J.-L.: Insights into the secondary fraction of the organic aerosol in a Mediterranean urban area: Marseille, *Atmos. Chem. Phys.*, 11, 2059–2079, doi:10.5194/acp-11-2059-2011, 2011.
- Heal, M. R.: The application of carbon-14 analyses to the source apportionment of atmospheric carbonaceous particulate matter: A review, *Anal. Bioanal. Chem.*, 406, 81–98, 2014.
- Heal, M. R., Naysmith, P., Cook, G. T., Xu, S., Duran, T. R., and Harrison, R. M.: Application of ^{14}C analyses to source apportionment of carbonaceous $\text{PM}_{2.5}$ in the UK, *Atmos. Environ.*, 45, 2341–2348, 2011.
- Heringa, M. F., DeCarlo, P. F., Chirico, R., Lauber, A., Doberer, A., Good, J., Nussbaumer, T., Keller, A., Burtscher, H., Richard,

- A., Miljevic, B., Prévôt, A. S. H., and Baltensperger, U.: Time-resolved characterization of primary emissions from residential wood combustion appliances, *Environ. Sci. Technol.*, 46, 11418–11425, 2012.
- Hoffmann, D., Tilgner, A., Iinuma, Y., and Herrmann, H.: Atmospheric stability of levoglucosan: A detailed laboratory and modeling study, *Environ. Sci. Technol.*, 44, 694–699, 2010.
- Iinuma, Y., Engling, G., Puxbaum, H., and Herrmann, H.: A highly resolved anion-exchange chromatographic method for determination of saccharidic tracers for biomass combustion and primary bio-particles in atmospheric aerosol, *Atmos. Environ.*, 43, 1367–1371, 2009.
- Keuken, M. P., Moerman, M., Voogt, M., Blom, M., Weijers, E. P., Röckmann, T., and Dusek, U.: Source contributions to PM_{2.5} and PM₁₀ at an urban background and a street location, *Atmos. Environ.*, 71, 26–35, 2013.
- Khan, M. B., Masiol, M., Formenton, G., Di Gilio, A., de Gennaro, G., Agostinelli, C., and Pavoni, B.: Carbonaceous PM_{2.5} and secondary organic aerosol across the Veneto region (NE Italy), *Sci. Total Environ.*, 542, 172–181, 2016.
- Kirillova, E. N., Sheesley, R. J., and Andersson, A.: Natural Abundance ¹³C and ¹⁴C Analysis of Water-Soluble Organic Carbon in Atmospheric Aerosols, *Anal. Chem.*, 82, 7973–7978, 2010.
- Kirillova, E. N., Andersson, A., Sheesley, R. J., Kruså, M., Praveen, P. S., Budhavant, K., Safai, P. D., Rao, P. S. P., and Gustafsson, Ö.: ¹³C- and ¹⁴C-based study of sources and atmospheric processing of water-soluble organic carbon (WSOC) in South Asian aerosols, *J. Geophys. Res.-Atmos.*, 118, 614–626, 2013.
- Kirillova, E. N., Andersson, A., Tiwari, S., Srivastava, A. K., Bisht, D. S., and Gustafsson, Ö.: Water-soluble organic carbon aerosols during a full New Delhi winter: Isotope-based source apportionment and optical properties, *J. Geophys. Res.-Atmos.*, 119, 3476–3485, 2014.
- Levin, I., Naegler, T., Kromer, B., Diehl, M., Francey, R. J., Gomez-Pelaez, A. J., Steele, L. P., Wagenbach, D., Weller, R., and Worthy, D. E.: Observations and modelling of the global distribution and long-term trend of atmospheric ¹⁴CO₂, *Tellus B*, 62, 26–46, 2010.
- Lewis, C. W. and Stiles, D. C.: Radiocarbon Content of PM_{2.5} Ambient Aerosol in Tampa, FL, *Aerosol Sci. Technol.*, 40, 189–196, 2006.
- Lewis, C., Klouda, G., and Ellenson, W.: Radiocarbon measurement of the biogenic contribution to summertime PM-2.5 ambient aerosol in Nashville, TN, *Atmos. Environ.*, 38, 6053–6061, 2004.
- Liu, D., Li, J., Zhang, Y., Xu, Y., Liu, X., Ding, P., Shen, C., Chen, Y., Tian, C., and Zhang, G.: The Use of Levoglucosan and Radiocarbon for Source Apportionment of PM_{2.5} Carbonaceous Aerosols at a Background Site in East China, *Environ. Sci. Technol.*, 47, 10454–10461, 2013.
- McMeeking, G. R., Kreidenweis, S. M., Baker, S., Carrico, C. M., Chow, J. C., Collett, J. L., Hao, W. M., Holden, A. S., Kirchstetter, T. W., Malm, W. C., Moosmüller, H., Sullivan, A. P., and Wold, C. E.: Emissions of trace gases and aerosols during the open combustion of biomass in the laboratory, *J. Geophys. Res.*, 114, D19210, doi:10.1029/2009JD011836, 2009.
- Minguillón, M. C., Perron, N., Querol, X., Szidat, S., Fahrni, S. M., Alastuey, A., Jimenez, J. L., Mohr, C., Ortega, A. M., Day, D. A., Lanz, V. A., Wacker, L., Reche, C., Cusack, M., Amato, F., Kiss, G., Hoffer, A., Decesari, S., Moretti, F., Hillamo, R., Teinilä, K., Seco, R., Peñuelas, J., Metzger, A., Schallhart, S., Müller, M., Hansel, A., Burkhardt, J. F., Baltensperger, U., and Prévôt, A. S. H.: Fossil versus contemporary sources of fine elemental and organic carbonaceous particulate matter during the DAURE campaign in Northeast Spain, *Atmos. Chem. Phys.*, 11, 12067–12084, doi:10.5194/acp-11-12067-2011, 2011.
- Minoura, H., Morikawa, T., Mizohata, A., and Sakamoto, K.: Carbonaceous aerosol and its characteristics observed in Tokyo and south Kanto region, *Atmos. Environ.*, 61, 605–613, 2012.
- Mohn, J., Szidat, S., Fellner, J., Rechberger, H., Quartier, R., Buchmann, B., and Emmenegger, L.: Determination of biogenic and fossil CO(2) emitted by waste incineration based on (14)CO(2) and mass balances, *Bioresour. Technol.*, 99, 6471–6479, 2008.
- Mook, W. G. and van der Plicht, J.: Reporting ¹⁴C activities and concentrations, *Radiocarbon*, 41, 227–239, 1999.
- Morino, Y., Takahashi, K., Fushimi, A., Tanabe, K., Ohara, T., Hasegawa, S., Uchida, M., Takami, A., Yokouchi, Y., and Kobayashi, S.: Contrasting diurnal variations in fossil and non-fossil secondary organic aerosol in urban outflow, Japan, *Environ. Sci. Technol.*, 44, 8581–8586, 2010.
- Ohata, S., Moteki, N., Mori, T., Koike, M., and Kondo, Y.: A key process controlling the wet removal of aerosols: new observational evidence, *Sci. Rep.*, 6, 34113, doi:10.1038/srep34113, 2016.
- Pöschl, U.: Atmospheric aerosols: composition, transformation, climate and health effects, *Angew. Chem. Int. Ed. Engl.*, 44, 7520–7540, 2005.
- Prokopiou, M.: Characterization of a thermal method for separating organic and elemental carbon from aerosol samples using 14-C analysis, MS thesis, University of Groningen, the Netherlands, 2010.
- Putaud, J.-P., Raes, F., Van Dingenen, R., Brüggemann, E., Facchini, M.-C., Decesari, S., Fuzzi, S., Gehrig, R., Hüglin, C., Laj, P., Lorbeer, G., Maenhaut, W., Mihalopoulos, N., Müller, K., Querol, X., Rodriguez, S., Schneider, J., Spindler, G., Brink, H. Ten, Tørseth, K., and Wiedensohler, A.: A European aerosol phenomenology – 2: chemical characteristics of particulate matter at kerbside, urban, rural and background sites in Europe, *Atmos. Environ.*, 38, 2579–2595, 2004.
- Reimer, P. J., Brown, T. A., and Reimer, R. W.: Discussion: Reporting and calibration of post-bomb ¹⁴C data, *Radiocarbon*, 46, 1299–1304, 2004.
- Ruff, M., Wacker, L., Gäggeler, H. W., Suter, M., Synal, H.-A., and Szidat, S.: A gas ion source for radiocarbon measurements at 200 kV, *Radiocarbon*, 49, 307–314, 2007.
- Ruff, M., Szidat, S., Gäggeler, H. W., Suter, M., Synal, H.-A., and Wacker, L.: Gaseous radiocarbon measurements of small samples, *Nucl. Instrum. Meth. B*, 268, 790–794, 2010.
- Sandrini, S., Fuzzi, S., Piazzalunga, A., Prati, P., Bonasoni, P., Cavalli, F., Bove, M. C., Calvello, M., Cappelletti, D., Colombi, C., Contini, D., de Gennaro, G., Di Gilio, A., Fermo, P., Ferrero, L., Gianelle, V., Giugliano, M., Ielpo, P., Lonati, G., Marinoni, A., Massabò, D., Molteni, U., Moroni, B., Pavese, G., Perrino, C., Perrone, M. G., Perrone, M. R., Putaud, J. P., Sargolini, T., Vecchi, R., and Gilardoni, S.: Spatial and seasonal variability of carbonaceous aerosol across Italy, *Atmos. Environ.*, 99, 587–598, 2014.

- Schauer, J. J., Rogge, W. F., Hildemann, L. M., Mazurek, M. A., and Cass, G. R.: Source apportionment of airborne particulate matter using organic compounds as tracers, *Atmos. Environ.*, 30, 3837–3855, 1996.
- Schichtel, B. A., Malm, W. C., Bench, G., Fallon, S., McDade, C. E., Chow, J. C., and Watson, J. G.: Fossil and contemporary fine particulate carbon fractions at 12 rural and urban sites in the United States, *J. Geophys. Res.*, 113, D02311, doi:10.1029/2007JD008605, 2008.
- Schmid, H., Laskus, L., Ju, H., Abraham, K., Baltensperger, U., Lavanchy, V., Bizjak, M., Burba, P., Cachier, H., Crow, D., Chow, J., Gnauk, T., Even, A., Brink, H. M., Giesen, K., Hitzenberger, R., Hueglin, C., Maenhaut, W., Pio, C., Carvalho, A., Putaud, J., Toom-saunty, D., and Puxbaum, H.: Results of the “carbon conference” international aerosol carbon round robin test stage I, *Atmos. Environ.*, 35, 2111–2121, 2001.
- Schmidl, C., Marr, I. L., Caseiro, A., Kotianová, P., Berner, A., Bauer, H., Kasper-Giebl, A., and Puxbaum, H.: Chemical characterisation of fine particle emissions from wood stove combustion of common woods growing in mid-European Alpine regions, *Atmos. Environ.*, 42, 126–141, 2008.
- Stein, A. F., Draxler, R. R., Rolph, G. D., Stunder, B. J. B., Cohen, M. D., and Ngan, F.: NOAA’s HYSPLIT atmospheric transport and dispersion modeling system, *B. Am. Meteorol. Soc.*, 96, 2059–2077, 2015.
- Szidat, S., Jenk, T. M., Gäggeler, H. W., Synal, H.-A., Fisseha, R., Baltensperger, U., Kalberer, M., Samburova, V., Reimann, S., Kasper-Giebl, A., and Hajdas, I.: Radiocarbon (^{14}C)-deduced biogenic and anthropogenic contributions to organic carbon (OC) of urban aerosols from Zürich, Switzerland, *Atmos. Environ.*, 38, 4035–4044, 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 ^{14}C , *J. Geophys. Res.*, 111, D07206, doi:10.1029/2005JD006590, 2006.
- Szidat, S., Prévôt, A. S. H., Sandradewi, J., Alfarra, M. R., Synal, H.-A., Wacker, L., and Baltensperger, U.: Dominant impact of residential wood burning on particulate matter in Alpine valleys during winter, *Geophys. Res. Lett.*, 34, L05820, doi:10.1029/2006GL028325, 2007.
- Szidat, S., Ruff, M., Perron, N., Wacker, L., Synal, H.-A., Hallquist, M., Shannigrahi, A. S., Yttri, K. E., Dye, C., and Simpson, D.: Fossil and non-fossil sources of organic carbon (OC) and elemental carbon (EC) in Göteborg, Sweden, *Atmos. Chem. Phys.*, 9, 1521–1535, doi:10.5194/acp-9-1521-2009, 2009.
- Taylor, P. E., Flagan, R. C., Miguel, A. G., Valenta, R., and Glovsky, M. M.: Birch pollen rupture and the release of aerosols of respirable allergens, *Clin. Exp. Allergy*, 34, 1591–1596, 2004.
- van der Plicht, J., Wijma, S., Aerts, A., Pertuisot, M., and Meijer, H. A.: Status report: The Groningen AMS facility, *Nucl. Instruments Methods Phys. Res. B*, 172, 58–65, 2000.
- Wacker, L., Fahrni, S. M., Hajdas, I., Molnar, M., Synal, H.-A., Szidat, S., and Zhang, Y. L.: A versatile gas interface for routine radiocarbon analysis with a gas ion source, *Nucl. Instrum. Meth. B*, 294, 315–319, 2013.
- Weijers, E. P., Schaap, M., Nguyen, L., Matthijsen, J., Denier van der Gon, H. A. C., ten Brink, H. M., and Hoogerbrugge, R.: Anthropogenic and natural constituents in particulate matter in the Netherlands, *Atmos. Chem. Phys.*, 11, 2281–2294, doi:10.5194/acp-11-2281-2011, 2011.
- Wittmaack, K.: Combustion characteristics of water-insoluble elemental and organic carbon in size selected ambient aerosol particles, *Atmos. Chem. Phys.*, 5, 1905–1913, doi:10.5194/acp-5-1905-2005, 2005.
- Wozniak, A. S., Bauer, J. E., Dickhut, R. M., Xu, L., and McNichol, A. P.: Isotopic characterization of aerosol organic carbon components over the eastern United States, *J. Geophys. Res.*, 117, D13303, doi:10.1029/2011JD017153, 2012.
- Yttri, K. E., Dye, C., Braathen, O.-A., Simpson, D., and Steinnes, E.: Carbonaceous aerosols in Norwegian urban areas, *Atmos. Chem. Phys.*, 9, 2007–2020, doi:10.5194/acp-9-2007-2009, 2009.
- Yttri, K. E., Simpson, D., Stenström, K., Puxbaum, H., and Svendby, T.: Source apportionment of the carbonaceous aerosol in Norway – quantitative estimates based on ^{14}C , thermal-optical and organic tracer analysis, *Atmos. Chem. Phys.*, 11, 9375–9394, doi:10.5194/acp-11-9375-2011, 2011a.
- Yttri, K. E., Simpson, D., Nøjgaard, J. K., Kristensen, K., Genberg, J., Stenström, K., Swietlicki, E., Hillamo, R., Aurela, M., Bauer, H., Offenberg, J. H., Jaoui, M., Dye, C., Eckhardt, S., Burkhardt, J. F., Stohl, A., and Glasius, M.: Source apportionment of the summer time carbonaceous aerosol at Nordic rural background sites, *Atmos. Chem. Phys.*, 11, 13339–13357, doi:10.5194/acp-11-13339-2011, 2011b.
- Zhang, Y. L., Perron, N., Ciobanu, V. G., Zotter, P., Minguilón, 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, doi:10.5194/acp-12-10841-2012, 2012.
- Zhang, Y., Obrist, D., Zielinska, B., and Gertler, A.: Particulate emissions from different types of biomass burning, *Atmos. Environ.*, 72, 27–35, 2013.
- Zhang, Y.-L., Li, J., Zhang, G., Zotter, P., Huang, R.-J., Tang, J.-H., Wacker, L., Prévôt, A. S. H., and Szidat, S.: Radiocarbon-based source apportionment of carbonaceous aerosols at a regional background site on Hainan Island, South China, *Environ. Sci. Technol.*, 48, 2651–2659, 2014.
- Zhang, Y. L., Schnelle-Kreis, J., Abbaszade, G., Zimmermann, R., Zotter, P., Shen, R. R., Schaefer, K., Shao, L., Prévôt, A. S. H., and Szidat, S.: Source Apportionment of Elemental Carbon in Beijing, China: Insights from Radiocarbon and Organic Marker Measurements, *Environ. Sci. Technol.*, 49, 8408–8415, 2015.
- Zotter, P., El-Haddad, I., Zhang, Y.-L., Hayes, P. L., Zhang, X., Lin, Y.-H., Wacker, L., Schnelle-Kreis, J., Abbaszade, G., Zimmermann, R., Surratt, J. D., Weber, R., Jimenez, J. L., Szidat, S., Baltensperger, U., and Prévôt, A. S. H.: Diurnal cycle of fossil and nonfossil carbon using radiocarbon analyses during CalNex, *J. Geophys. Res.*, 119, 6818–6835, 2014a.
- Zotter, P., Ciobanu, V. G., Zhang, Y. L., El-Haddad, I., Macchia, M., Daellenbach, K. R., Salazar, G. A., Huang, R.-J., Wacker, L., Hueglin, C., Piazzalunga, A., Fermo, P., Schwikowski, M., Baltensperger, U., Szidat, S., and Prévôt, A. S. H.: Radiocarbon analysis of elemental and organic carbon in Switzerland during winter-smog episodes from 2008 to 2012 – Part 1: Source apportionment and spatial variability, *Atmos. Chem. Phys.*, 14, 13551–13570, doi:10.5194/acp-14-13551-2014, 2014b.