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NOVEL METHOD OF EXTRACTION FOR RADIOCARBON MEASUREMENTS OF ATMOSPHERIC CARBON DIOXIDE

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ABSTRACT. In this paper, we present the first data from an alternative extraction method for
10 atmospheric ¹⁴CO₂ analysis, based on the direct trapping of whole air samples onto a molecular sieve
zeolite (13X) trap, incorporated into a commercially available automated graphitisation system. Results
are presented for both inter-laboratory comparison samples and an in-house reference standard. The in-
house reference was used to calculate the standard deviation of measurements (2.0 ‰). This newly
developed method will facilitate faster sample processing and therefore lower cost per analysis, critical
15 for scaling up such studies.

KEYWORDS: Sample pre-treatment, atmospheric ¹⁴CO₂ measurements, zeolite, CO₂ fossil fuel
emissions

INTRODUCTION

Carbon dioxide (CO₂) is the most important greenhouse gas contributing to climate change (IPCC
20 2014). The 2015 Paris climate agreement highlights the need to reduce anthropogenic emissions by
2050 (Rogelj *et al.* 2016). Understanding the relative contributions of global CO₂ sources is
fundamental to support mitigation policies. However, CO₂ source apportionment calculations currently
have large uncertainties (IPCC 2014).

Radiocarbon (¹⁴C), produced in the upper atmosphere by collisions of cosmic rays with nitrogen atoms,
25 is subsequently rapidly oxidised to ¹⁴CO₂ which is distributed throughout the terrestrial, oceanic and
atmospheric carbon reservoirs via the carbon cycle. There are also anthropogenic sources of ¹⁴C. A
large amount of ¹⁴C was produced and released as a result of nuclear weapons testing in the 1950s and
60s and subsequently distributed via the carbon cycle, doubling the atmospheric inventory of ¹⁴CO₂

(Nydal and Lovseth 1983; Levin *et al.* 1985; Manning *et al.* 1990). The nuclear power industry also
30 releases $^{14}\text{CO}_2$, which offsets the depletion caused by fossil fuel combustion (Kopal 1997; McNamara
et al. 1998; Fontugne *et al.* 2004; Yim & Caron 2006; Magnusson 2007; Molnar *et al.* 2007; Dias *et al.*
2009; Graven & Gruber 2011; Aulagnier *et al.* 2012; Svetlik *et al.* 2012; Vokal & Wang *et al.* 2012,
2013, 2014; Vogel *et al.* 2013; Metcalfe & Mills 2015; Tierney *et al.* 2016).

Initially, following the atomic weapons testing activities of the mid-20th century, the atmospheric levels
35 of ^{14}C were high, with $\Delta^{14}\text{C}$ values of several hundred ‰. This has now decreased to only several ‰
because of the uptake of CO_2 by the ocean and biosphere (Stuiver & Robinson 1974; Bozhinova *et al.*
2013; LaFranchi *et al.* 2016). Atmospheric concentration of CO_2 is rising because of fossil fuel burning
(Bronk *et al.* 2016). $^{14}\text{CO}_2$ measurements therefore provide a method of measuring fossil fuel CO_2
because the absence of ^{14}C in fossil fuels results in a dilution of atmospheric $^{14}\text{CO}_2$, a phenomenon
40 known as the Suess effect (Suess 1955), that can be quantified via atmospheric measurements. $\Delta^{14}\text{C}$
determinations of atmospheric CO_2 samples provide a method of quantifying fossil fuel emissions on
both regional and global scales but, the scale of such efforts is limited by measurement precision, time
and cost.

Historically, atmospheric $^{14}\text{CO}_2$ measurements were made using proportional counting methods, which
45 required large sample sizes; these were prepared by trapping CO_2 using NaOH over one or two weeks,
then extracting the CO_2 directly in the sampling device in a laboratory vacuum system by the addition
of H_2SO_4 , before cleaning with a charcoal column, to provide a time-integrated measurement (Levin
et al. 1980, 2003). Since the development of AMS, around 1000 times less CO_2 is required, making
whole air flask sampling, and therefore instantaneous sampling, possible (Graven *et al.* 2007). Samples
50 are collected into glass flasks (e.g. 0.7 L flasks) either instantaneously or integrated over hours (Turnbull
et al. 2012). In the laboratory, CO_2 is extracted from these whole air samples. Typically, cryogenic
methods use dry-ice to remove water, then CO_2 is isolated and purified using liquid nitrogen (Turnbull
et al. 2007, 2010; Hammer *et al.* 2017). This CO_2 is then transferred and graphitised using traditional
vacuum lines (Turnbull *et al.* 2007). These methods require multiple extraction steps and manual
55 intervention and are therefore, generally slow and costly. The development of automated graphitisation

systems such as the Automated Graphitisation Equipment (AGE) (Wacker *et al.* 2010c) enables increased sample throughput in the preparation of samples for ^{14}C analysis with minimal manual interventions. The continuous-flow system and zeolite CO_2 trapping in the AGE3, the commercially available third generation AGE system, provides an alternative to traditional vacuum systems utilising cryogenic trapping of CO_2 . The CO_2 is absorbed onto a packed zeolite column and released into the reaction volume by heating the trap. We aimed to develop a simple, easily automated method for the extraction of air samples taken in simple glass flasks or tedlar bags.

Typically, current precision for most atmospheric ^{14}C laboratories is ca. 2-5 ‰ (Zhao *et al.* 1997; Meijer *et al.* 2006; Turnbull *et al.* 2007, 2010, 2015; Hammer *et al.* 2017;) a range which is now similar to the seasonal and spatial variability in some regions. (Graven *et al.* 2007) reported precisions of 1.7 ‰, expanding the usefulness of ^{14}C analysis in identifying and quantifying sources and fluxes of CO_2 .

The aim of the work reported in this paper was to develop and test an alternative method for CO_2 extraction from air samples using an existing AGE3 system, thus providing a simple and low-cost solution for users with such equipment.

70 METHODS

A prototype system for the extraction of CO_2 from whole air samples was designed and built in the Bristol Radiocarbon AMS laboratory (Figure 1) and is described here: Samples (in glass flasks (2L, 1.2 bar) or gas cylinders (50 L, 200 bar)) were transferred using a KNF pump (KNF N86KN.18, KNF Neuberger UK Ltd) via a phosphorus pentoxide water trap and mass flow controller (MFC, red-y smart series, GSC-B4SS-BB23, 0-600 mL/min, G1/4", Icentra, UK) to the sample inlet of the AGE3 system. The Fe catalyst was conditioned and the AGE3 system operated according to Wacker *et al.* (2010c). The samples were transferred to the AGE3 zeolite trap at a maximum flow rate of 180 mL/min, accurately controlled using the MFC. Atmospheric CO_2 was trapped on the zeolite trap of the AGE3 system at ambient temperature before being thermally desorbed and transferred into reaction tubes, CO_2 was quantified by measuring the pressure change in the reactors. A three-way valve was employed after the KNF pump to enable flushing and cleaning of the zeolite trap with Helium, to ensure the zeolite

trap is under an inert Helium environment before heating. The graphitization reaction was carried out at 580°C for 120 minutes, the graphite samples produced were pressed into aluminium cathodes using a pneumatic sample press.

85 A Luxfer cylinder was filled with dried ambient air for ^{14}C analysis (408.2 ppm), using a SA-6 pump (200 bar, 50 L) at the School of Chemistry, University of Bristol on the 29th April 2016. This cylinder had a similar CO_2 mole fraction to in situ ambient atmospheric samples. This in-house air reference cylinder will henceforth be referred to as the reference tank.

A number of samples ($n=38$) from our in-house reference tank were analysed using this new method. 90 In addition to these 38, seven samples extracted on the system in the initial testing phase were used. These have not been included in the subsequent analysis, all other samples extracted for 15 minutes at 180 $\text{mL}\cdot\text{min}^{-1}$ were included in the analysis. Radiocarbon ‘dead’ CO_2 gas ($^{14}\text{CO}_2$ -free 400 ppm in zero air, purchased from BOC) was used as a processing blank and inter laboratory comparison samples ($n=11$) were used as further quality control air standards. Normalisation and AMS quality control 95 standards, Oxalic acid II (NIST SRM 4990C) (OXII), IAEA-C7 oxalic acid and phthalic anhydride chemical blank (each at the equivalent of 500 $\mu\text{g C}$), were prepared by combustion in an elemental analyser interfaced to the AGE3.

Measurements were performed on a MICADAS AMS system (Synal *et al.* 2007; Wacker *et al.* 2010a). Data reduction was performed using BATS software (Wacker *et al.* 2010b). The $F^{14}\text{C}$ values generated 100 were converted to $\Delta^{14}\text{C}$ using Equation 1, correcting for mass dependent fractionation and age (Stuiver & Polach 1977) where x is the year of sample collection.

$$\Delta^{14}\text{C} = (F^{14}\text{C} e^{((1950-x)/8267)} - 1) \times 1000 \quad (1)$$

Samples were analysed until the OXII standards had achieved greater than 500,000 counts of ^{14}C .

RESULTS AND DISCUSSION

105 Full characterisation of the sample pre-treatment was carried out. This involved: (1) investigation of the trapping efficiency under different conditions, (2) multiple analyses of air from the in-house reference

tank to assess the precision of the method, and (3) comparison to other laboratories to examine the accuracy of the method.

Trapping Time

110 To determine the optimal length of time required to trap CO₂ from a 2 L whole air sample in a glass flask, the trapping time was varied under different conditions. Samples were extracted directly, both from glass flasks filled from the reference tank, and directly from the reference tank itself over a range of flow rates. The zeolite trap temperatures and the measured mass of CO₂ trapped were recorded (Figure 2). The AGE3 system is designed to isolate CO₂ from combusted samples from the Elemental
115 Analyser (EA) in a stream of helium carrier gas, at a flow rate of 180 mL/min. The samples extracted from flasks with a maximum flow rate set at 180 mL / min (this flow rate drops as the pressure of the flask is reduced). The samples extracted from the flasks (blue) demonstrated a plateau at 10 – 20 minutes, at 400 µg C, indicating complete isolation of the CO₂. The flask trapping experiment was continued after the plateau was observed to ensure all of the sample had been trapped. A slight increase
120 was seen from 20 to 25 minutes, there are two possible reasons for this: either the flasks used in these experiments were filled to slightly higher pressures than those tested at 10 - 20 minutes therefore the plateau would have been slightly higher, or at this point, the large pressure difference between the flask (lower than ambient pressure) and the laboratory a leak into the system via the pump or flask attachment may have occurred.

125 To improve counting statistics and therefore improve the analytical precision, larger samples could be prepared (i.e. 1 mg C instead of 0.5 mg) by filling and sampling 2L flasks in pairs, or filling flasks of larger volumes or at higher pressures. With larger samples however, there is the potential risk of CO₂ breakthrough in the zeolite trap due to its saturation. To determine the trapping capacity of the zeolite, samples from the in-house reference tank were extracted directly from the reference tank cylinder at a
130 flow rate of 180 mL / min (orange), ensuring that sample size was not a limiting factor. A linear increase in the quantity of CO₂ trapped was observed over the initial 20 minutes of trapping, however, after this period, at ~685 µg C a plateau was observed due to breakthrough of CO₂ since the zeolite trap had reached its capacity under these conditions. The flow rate was increased to 250 mL/min (green), and

again, a linear increase was observed initially, until the same plateau level was reached at 760 µg, but
135 after a shorter sampling time. The temperature of the zeolite trap was reduced to 10 °C from an ambient
trapping temperature (20 °C), and the capacity of the zeolite trap was observed to increase to 1200 µg
(red).

Sample Uncertainty

Multiple flasks filled from the reference tank were extracted for 15 minutes, at 180 mL/min, with the
140 zeolite trap at 20 °C. The extraction was performed on 38 samples. The weighted average of $\Delta^{14}\text{C}$ values
was determined as $-3.45 \pm 0.3 \text{ ‰}$. For each sample, the uncertainty of the measurement was plotted as
the uncertainty (σ_{tot}) for individual measurements (Figure 3). The uncertainty includes instrument error
(σ_{AMS}) that incorporates counting statistics, normalisation errors and blank uncertainties. Generally, a
sample scatter factor (σ_{ss}) (Equation 2), will be added using a sum-of-squares approach. This sample
145 scatter factor is determined using chi-squared tests on repeat measurements. This accounts for any
additional uncertainty resulting from sample preparation including graphitisation and extraction
(observed as scatter in the repeat analysis of a standard).

$$\sigma_{\text{tot}}^2 = \sigma_{\text{AMS}}^2 + \sigma_{\text{ss}}^2 \quad (2)$$

Assessment of Sample Uncertainty Contributions

150 The 38 samples extracted using our new direct zeolite trapping method were measured across 6
magazines from October 2017 to August 2018 (Table 1). We performed a Pearson's chi-squared test to
assess how well the observed distribution of data fits with the distribution that would be expected if the
variables are independent. The right-tailed p-value (α) calculated for this distribution was 0.87,
suggesting that the instrument uncertainty accounts for all observed scatter. If a sample scatter value
155 was to be added, the errors would be overestimated. It is likely that the uncertainty is currently
dominated by the scatter in oxalic acid II standards that are prepared by combustion in the EA. The
blanks from the EA method were also used in the calculation of the sample uncertainty. This is because
the standard deviation of the oxalic acid was 2.5 ‰, larger than that of the reference samples which was
1.7 ‰. It is likely that the EA combustion step introduced a larger degree of sample scatter to these

160 standards than was observed in air samples. In the future, we will investigate this further by using air standards containing CO₂ from bomb-combusted oxalic acids and blanks to determine a truer precision of the method.

The instrument uncertainty (σ_{AMS}) for each sample was therefore, used as the total analytical uncertainty for this study. The average uncertainty across the 38 replicate analyses was 2.0 ‰ (on the $\Delta^{14}\text{C}$ scale).

165 **Blank Analysis**

Extraction of air ‘blanks’, consisting of zero-air mixed with radiocarbon dead CO₂, using the direct zeolite trapping method produced an average value of 0.76 ± 0.13 pMC, calculated based on measurements of blank samples independent from contamination from ambient samples (n=15). The purpose of these blank analyses was to enable identical sample pre-treatment of all standards. The data 170 for these were higher than the Phthalic Anhydride blank prepared using the EA (0.34 ± 0.07 pMC). This is lower than observed from the isolation of our radiocarbon ‘dead’ air standard, suggesting some contamination; the source of this is yet to be confirmed. All samples in this study contained ca. 400 μg C, resulting in a slightly higher blank than would be observed for typical full-sized samples (1 mg C).

Cross Contamination

175 ‘Modern’ in-house reference samples were extracted and graphitised immediately preceding radiocarbon ‘dead’ samples to examine the effect of cross contamination between samples. Four sequences consisting of three blanks following a sample from the in-house reference tank ($\Delta^{14}\text{C} = -3.45$ ‰) were extracted, and the data are presented in Figure 4. The first blank of each sequence was observed to be higher than the second and third.

180 Cross contamination can be described by a simple mixing model. A measured pMC value of the blank sample (X , 0.942) depends linearly on the previous measured sample (s_0 , 100.45) and the ‘true’ pMC value of the blank (s_1 , 0.76), the cross-contamination level (c), is the coefficient for the previous sample.

$$c = \frac{X - s_1}{s_0 - s_1} \quad (3)$$

185 A cross-contamination level of 1.83 ± 1.52 ‰ (or 0.183 ± 0.152 ‰) was determined using Equation 3, which is significantly higher than that reported by Wacker *et al.* (2010c) (0.6 ± 0.1 ‰). Therefore, as with the EA-AGE3 system, when analysing samples with very different levels of ^{14}C , the zeolite should be pre-conditioned with a sample of similar ^{14}C content. The ‰ cross contamination is in parts per thousand. Therefore, 0.183 ‰ of the C in one samples comes from the sample before. For example, if two samples have a difference in $\Delta^{14}\text{C}$ of 10 ‰, the second sample will be shifted by 0.0183 ‰ on the
190 $\Delta^{14}\text{C}$ scale. To establish the amount of C from the sample transferred to the 2nd blank processed subsequently, the “effective c” value of this blank was calculated to be 0.43 ‰. The “real c”, calculated from the first blank cannot be changed regardless of the approach taken, whereas the effective c can change depending on the approach used (e.g. 1st + 2nd blank). This agrees with the findings of Wacker *et al.*, (2010c) and demonstrates the efficacy of a sacrificial sample (of similar ^{14}C content to subsequent
195 samples) between samples of very different ^{14}C content when using this system.

Inter-laboratory Comparisons

Of great importance to atmospheric ^{14}C laboratories, and ^{14}C laboratories in general is inter laboratory comparisons to confirm that laboratories are reporting with the same accuracy and give realistic values for precision. These exercises, though of great importance for global monitoring, rarely happen (Miller
200 *et al.* 2013; Hammer *et al.* 2017).

The results of our determination on the in-house reference standard were compared to seven flasks of the same reference tank analysed at an independent AMS laboratory, the Institute of Arctic and Alpine Research (INSTAAR), which has a long history of making atmospheric $^{14}\text{CO}_2$ measurements. These data are shown in Table 2. The average $\Delta^{14}\text{C}$ values are comparable within 1σ .

205 Samples ($n = 4$) that previously were analysed at the Integrated Carbon Observation System Central Radiocarbon Laboratory (ICOS-CAL), in Heidelberg, Germany were also analysed on our new direct trapping extraction system. These samples were part of a previous inter laboratory comparison study (Hammer *et al.* 2017), therefore have been measured at multiple laboratories. The results of the analyses from this study were compared to the consensus values from the inter-comparison study, presented in

210 Table 3. Samples were split into two aliquots to enable replicate measurements. Unfortunately, for
samples 30864 and 30874 analysis of only one aliquot was possible. For samples 30996 and 31061 both
aliquots were analysed. The results presented in Table 3 are a weighted mean of these. Although all
measurements were within 2σ (and three of the four samples were within 1σ), all of the measurements
from this study were found to be slightly lower than the consensus value. The limited size of the dataset
215 makes interpretation of this difficult, however if there is a systematic offset it could point to a small
amount of contamination with atmospheric CO_2 during a leak to the system or may be because of the
lower blank values. Further analysis will be required to establish this. The uncertainties reported for the
consensus values are lower than any of the measurements by individual labs (including those reported
here) as they were calculated based on multiple measurements at several laboratories. The uncertainties
220 reported in this paper are similar in magnitude to the standard deviation of the measurements of the
inter-comparison study at the individual laboratories, meaning this is dependent on the factors outlined
above due to the combustion of the Oxalic Acid standard. The right tailed chi-squared p values for each
measurement shows that there is no significant difference in the measurements in this study and the
consensus values. A significance level of 5 % was used ($p = 0.05$) and all p values were higher than
225 this, meaning the null hypothesis was true and there is agreement between the measurements in this
study and the consensus values within errors. This analysis was performed on a small number of samples
and further comparisons will be required to ensure measurements between laboratories are comparable.
The World Metrological Organisation (WMO) Guidelines state that current compatibility between
laboratories is 2 – 4 ‰, short of the goal of 0.5 ‰ (Tans and Zellweger 2016). The comparisons made
230 in this paper are comfortably within the compatibility reported in the WMO guidelines (2 – 4 ‰) as are
comparable to a level of 2.7 ‰ which is the largest difference in the inter comparison experiments.
Overall, our new set up has shown comparable compatibility to other AMS laboratories making
atmospheric $^{14}\text{CO}_2$ measurements from other studies (Hammer *et al.* 2017).

CONCLUSION

235 We have developed and reported an alternative method for the extraction of atmospheric $^{14}\text{CO}_2$ samples.
It is anticipated that this method, with the graphitisation and analysis for ^{14}C using AMS, will be used

for the analysis of multiple samples for quantifying fossil fuels CO₂ emissions in the UK. Our initial results are promising for the future of these measurements, demonstrating agreement with other ¹⁴C laboratories. We have reported a range of tests that have characterised the direct trapping system. A trapping capacity of 400 µg C in 15 minutes at 180 mL·min⁻¹ from a 2 L flask has been achieved. The trapping capacity at room temperature achieved was 685 µg. The trapping capacity at 10 °C was 1200 µg. In this work, the maximum flow rate investigated for trapping on the zeolite trap was 250 mL / min. This method has been used for pre-treatment of 38 replicate samples to achieve standard deviation on long term measurements of 2.0 ‰ over 6 magazines. Assessment of the uncertainty suggests the use of an air sample containing OXII-derived CO₂ rather than OXII prepared by EA-AGE3 could be advantageous as a normalization standard. Blank analysis shows that the cross-contamination level is 1.83 ± 1.52 ‰, meaning if analysing samples with very different levels of ¹⁴C, the zeolite should be pre-conditioned with a sample of similar ¹⁴C content. Analysis of inter-comparison samples showed this method is comparable to two other global radiocarbon laboratories to a level of 2.7 ‰, within the WMO guidelines (2 – 4 ‰). In the future, we will fully characterise the system to further investigate why the air blank is higher than the combustion blank, and why the memory effect of our line and its uncertainty are much higher than those reported in Wacker *et al.* (2010c). We then aim to automate the whole extraction process, integrating this to the AGE3 system to increase sample throughput and precision, vital for atmospheric ¹⁴CO₂ measurements.

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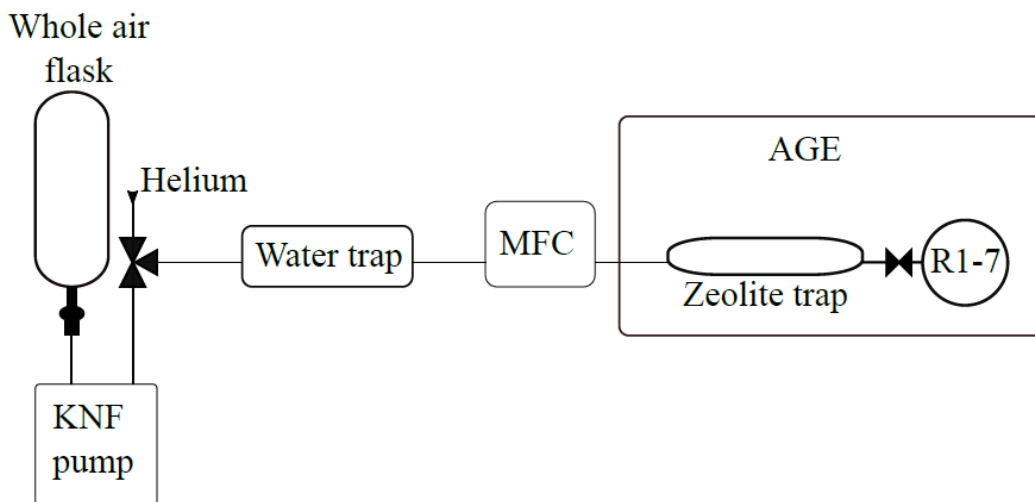
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Figure 1. Schematic of the direct trapping system - The flask (NOAA design, 2L, Normag, Germany)

is attached ($\frac{1}{2}$ " ultratorr) to the pump (KNF N86KN.18, KNF Neuberger UK Ltd), the sample is extracted via a phosphorus pentoxide water trap via a mass flow controller (MFC, red-y smart series, GSC-B4SS-BB23, 0-600 mL/min, G1/4", Icentra, UK) directly to the zeolite trap (13X) of the AGE3

365 system. The AGE3 is showed simplified here (see (Wacker et al. 2010c) for full details of this system)

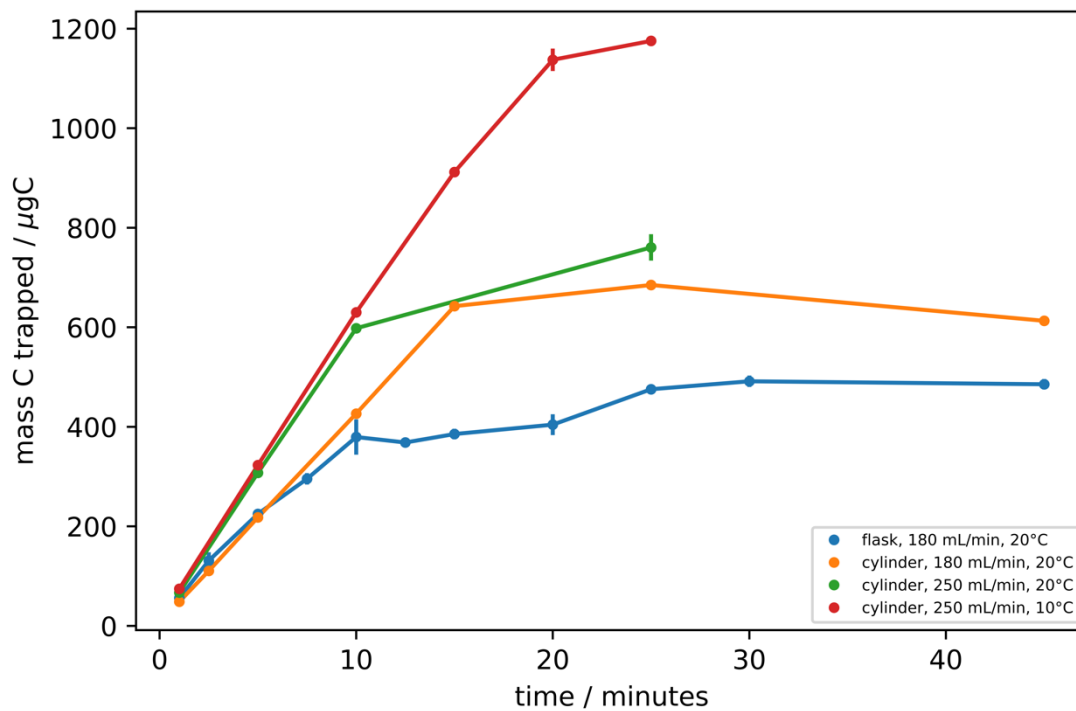


Figure 2. Time varying trapping recording the masses of C trapped on the zeolite, direct from flask at 180 mL·min⁻¹ (blue), direct from cylinder at 180 mL·min⁻¹ (orange), 250 mL·min⁻¹ (green), 10 °C (red). The uncertainty of each data point is represented as the standard deviation of repeat measurements (n=2).

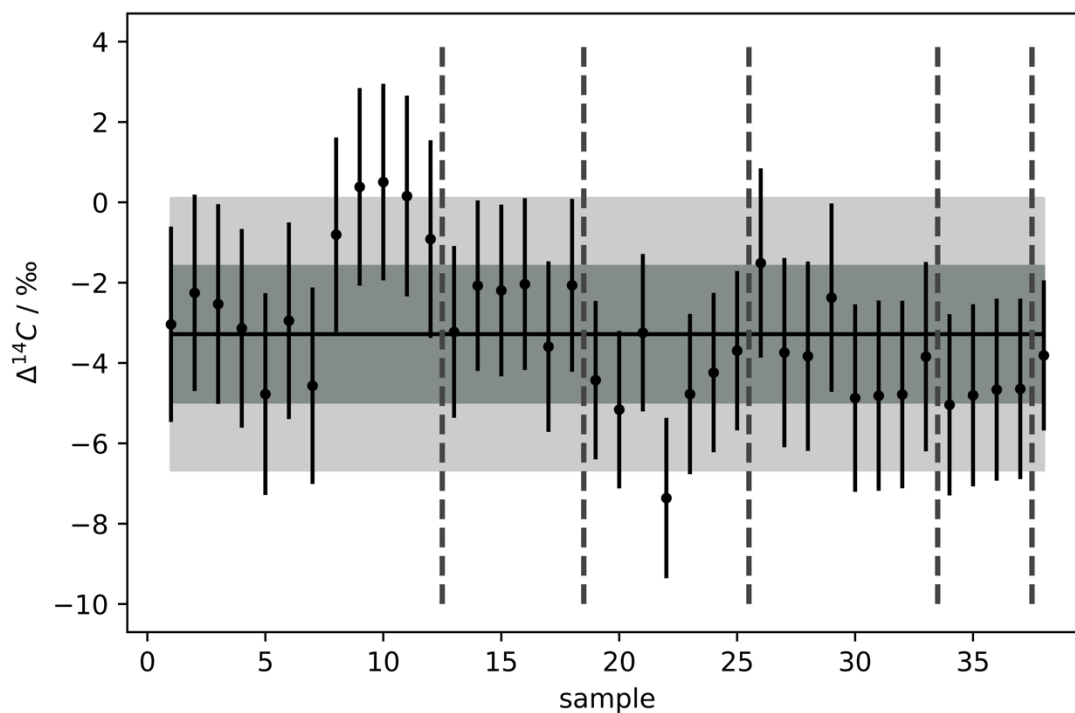


Figure 3. $\Delta^{14}\text{C}$ values determined for the in-house reference standard extracted for 15 minutes at a
 375 max. flow rate of $180 \text{ mL}\cdot\text{min}^{-1}$ for 38 samples, 1σ (grey - 1.97 ‰) and 2σ (pale grey - 3.94 ‰). Mean
 represented by solid black line ($\Delta^{14}\text{C} = -3.45 \text{ ‰}$). All measurements were within 2σ . Vertical grey
 dashed lines separate measurements from different AMS magazines.

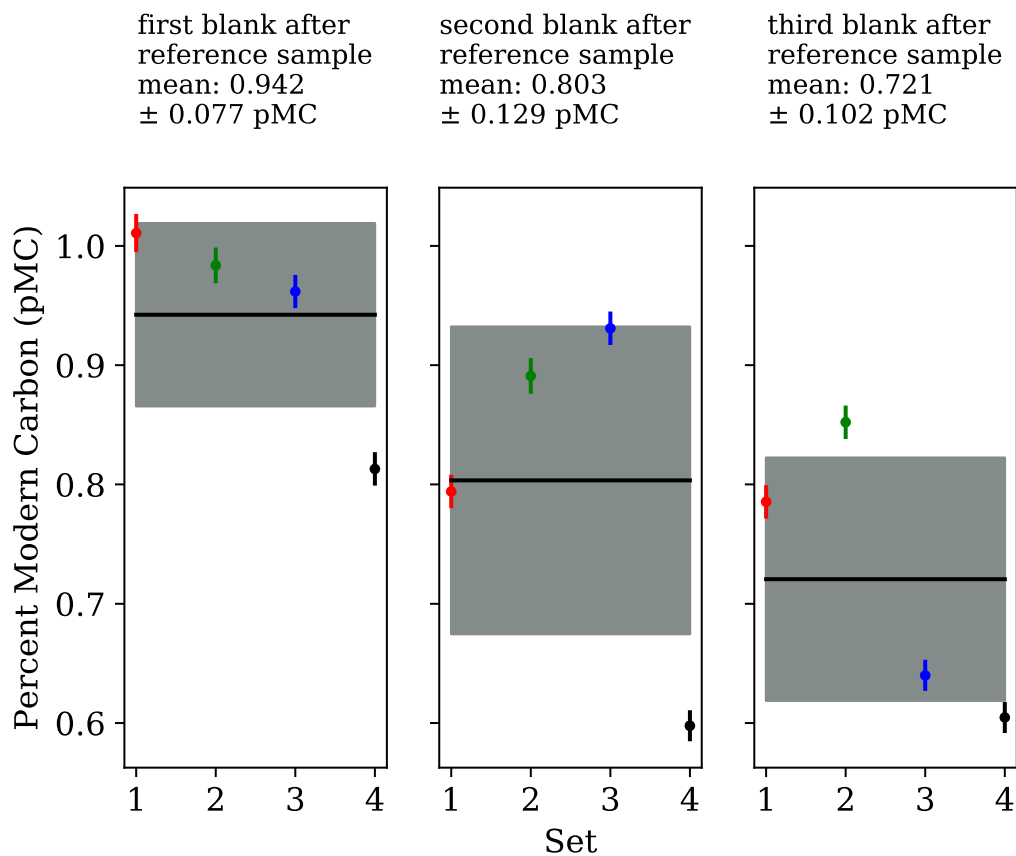


Figure 4. Cross contamination tests. Four sets of three consecutive radiocarbon blanks isolated and graphitized after a sample of our (modern) in-house reference gas. A cross contamination level of 1.83 ± 1.52 % was calculated using a simple mixing model. The mean value for the first blank after the reference was 0.942 ± 0.077 pMC, second blank was 0.803 ± 0.129 pMC and third blank was 0.721 ± 0.102 pMC.

| Magazine code | Number of in-house reference samples | Number of OX II standards | Number of Phthalic Anhydride Blanks | Average high energy ^{12}C Ion Current / μA | Measurement time per target / h |
|---------------|--------------------------------------|---------------------------|-------------------------------------|---|---------------------------------|
| 20171018 | 12 | 3 | 2 | 18.4 | 1.28 |
| 20171123 | 6 | 4 | 3 | 21.1 | 1.39 |
| 20171213 | 7 | 3 | 2 | 24.6 | 1.17 |
| 20180130 | 8 | 3 | 2 | 18.6 | 1.33 |
| 20180511 | 4 | 3 | 2 | 19.9 | 1.06 |
| 20180802 | 1 | 3 | 2 | 20.5 | 1.33 |

390 **Table 1.** Details regarding the AMS magazines containing samples measured as part of this study.

| Analysis | Number of samples analysed | Mean $\Delta^{14}\text{C}$ / ‰ | $\Delta^{14}\text{C}$ standard deviation / ‰ |
|------------------|----------------------------|--------------------------------|--|
| BRAMS, this work | 38 | -3.44 | 2.0 |
| INSTAAR | 7 | -3.80 | 1.80 |

Table 2. Summary of our in-house reference standard extracted and measured at two different

395 laboratories, Bristol Radiocarbon Accelerator Mass Spectrometer facility (BRAMS) and Institute of Arctic and Alpine Research (INSTAAR).

| Sample | Number of samples analysed | Measurement from this study $\Delta^{14}\text{C}$ / ‰ | Consensus value $\Delta^{14}\text{C}$ / ‰ | Right-tailed P-values (α values) |
|--------------|----------------------------|---|---|--|
| 30864 | 1 | 24.92 ± 1.83 | 25.2 ± 0.7 | 0.886 |
| 30874 | 1 | 37.75 ± 1.60 | 40.4 ± 0.7 | 0.129 |
| 30996 | 2 | 9.97 ± 1.60 | 10.9 ± 0.7 | 0.481 |
| 31061 | 2 | 21.20 ± 1.60 | 22.7 ± 0.7 | 0.259 |

Table 3. Four of the samples used in the inter comparison (Hammer et al. 2017) measured during this study and the intercomparison consensus values. The chi squared right-tailed P value for each sample

400 is also reported.