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10	The age of	CO ₂ released	from soils i	n contrasting	ecosystems	during the	arctic winter
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35 Abstract

36

In arctic ecosystems, winter soil respiration can contribute substantially to annual CO_2 release, yet the source of this C is not clear. We analysed the ¹⁴C content of C released from plant-free plots in mountain birch forest and tundra-heath. Winter-respired CO_2 was found to be a similar age (tundra) or older (forest) than C released during the previous autumn. Overall, our study demonstrates that the decomposition of older C can continue during the winter, in these two contrasting arctic ecosystems.

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44 Key words: ¹⁴CO₂, passive sampling, mountain birch, radiocarbon, tree-line, tundra45 heath, winter respiration

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47 Arctic soils contain globally significant C stores (Post et al., 1982; Ping et al., 2008). As these areas are warming rapidly (AMAP, 2012), C may be lost if decomposition rates 48 49 increase in response (Davidson and Janssens, 2006). There is growing recognition that the CO₂ released during the long winters in high-latitude/altitude ecosystems can 50 51 represent a substantial proportion (up to 30%) of annual respiration (Elberling, 2007; 52 Williams et al., 2009), but, for practical reasons, flux measurements are biased towards 53 the growing season. Furthermore, debate continues as to whether the source of the CO₂ 54 released during the winter is similar to that released during the summer, or is derived 55 mainly from recently-fixed, labile C (Grogan et al., 2001; Grogan and Jonasson, 2005; 56 Nobrega and Grogan, 2007). Importantly, climate change has greater potential to affect 57 rates of winter respiration in the long term, either positively or negatively, if there is a 58 substantial contribution from the large reserves of older SOM, than if most of the 59 respired CO₂ is derived from small labile C pools (Jones et al., 2005; Hartley and Ineson, 2008). Here, by undertaking the first ¹⁴C analyses of CO₂ released from soils 60

61 during the arctic winter, we investigated whether the decomposition of older SOM62 continues during the winter.

elevation ~520 m) and tundra-heath (68°18'07"N, 18°51'16"E; elevation ~710 m), near
Abisko, northern Sweden. We collected samples of CO ₂ released over the 2007-2008
winter from three non-vegetated plots in each ecosystem. To allow only soil-respired
CO_2 to be collected, the plots were clipped and trenched in late summer 2006. In the
centre of each plot, 7-cm tall collars were sealed to the surface using putty, with
respiration rates and 14 CO ₂ contents being monitored during the 2007 growing season
(Hartley et al. 2012). To collect winter-respired CO ₂ , we developed a new technique
using molecular sieve cartridges (MSCs) to collect passively (by diffusion)
representative samples of CO ₂ over extended time periods (Garnett et al., 2009).
In order to minimise chamber height and potential effects on snow lie, lids were
directly placed on top of the collars. MSCs were then connected through the collar sides
via auto-shut-off Quick Couplings [™] (Colder Products Company, USA), with
hydrophobic filters (Accurel PP V8/2 HF, Membrana GmbH, Germany) attached inside
each collar to prevent liquid water passing into the MSCs (Fig. 1). For protection, the
MSCs were then placed inside foam insulation, PVC pipes and guttering. The lids were
closed on 16 th -17 th September 2007 and the MSCs connected on 20 th September. MSCs
were recovered on 23 rd -24 th May 2008. Air samples were collected at the tundra-heath
site on 20 th September 2007 and 24 th May 2008, by pumping air through MSCs for
approximately 60 minutes.
Soil temperatures at 5 cm were monitored throughout the winter (thermistor

84 probes and CR10x datalogger, Campbell Scientific, Leics, UK). After MSC collection,

soil temperatures at 2, 5 and 8 cm depth (digital thermometer, E.T.I. Ltd., West Sussex,

86 UK) and soil moisture at 6 cm (Theta probe: ML2, Delta-T Devices, Cambridge, UK)
87 were measured inside and outside the collars.

All ¹³C and ¹⁴C analyses were performed on CO₂ recovered from the MSCs 88 using established procedures (Hardie et al., 2005). Following convention, ¹⁴C results 89 were normalised to a δ^{13} C value of -25 ‰ and expressed as %modern (Stuiver and 90 91 Polach, 1977). Because collars were not inserted into the soil, it was not possible to 92 avoid some atmospheric contamination. Samples were corrected for atmospheric 93 contamination using the approach of Hartley et al. (2012), after accounting for the 4 ‰ ¹³C fractionation associated with passive sampling (see Garnett et al., 2009). 94 95 After MSC collection, on the tundra, both soil moisture and temperature were 96 near identical inside and outside the collars. In the forest, temperatures at 5 and 8 cm 97 were $0.6-0.7^{\circ}$ C higher within the collar, but there was no significant effect on soil 98 moisture. Therefore, the chambers appeared to have little effect on the soil physical 99 environment. During the winter, temperature at 5 cm was greater in the forest (mean: 100 0.07° C, range: -3.02° C to 6.94° C) than the tundra (mean: -1.73° C, range: -7.18° C to 101 4.66°C). Warmer winter temperatures can increase winter CO₂ production, and thus 102 influence annual carbon balances (Grogan and Jonasson, 2006; Nobrega and Grogan,

103 2007; Sullivan, 2010), potentially contributing to the lower C storage in forest than

104 tundra soils. However, it should be emphasised that previous research at the current

105 field sites (Hartley et al., 2012), as well as studies at lower latitudes (Mitchel et al.,

106 2007), have identified the important role plant-soil interactions and priming play in

107 controlling soil carbon storage in forest-heath transitions.

108 Consistent with warmer temperatures increasing winter respiraton, our 109 molecular sieves collected more CO_2 in the forest than tundra (means of 102.3 ml and 110 71.8 ml, respectively). However, the amount of CO_2 collected on our MSCs depends on 111 the average CO_2 concentration within the chamber (Garnett et al., 2009). This is

112 controlled not only by respiration rates, but also by rates of exchange between the 113 atmosphere and the headspace, which may have been greater on the tundra due to higher 114 wind speeds and reduced snow cover; atmospheric contamination was greater in tundra 115 samples (Table 1). Therefore, volumes collected cannot be translated directly into 116 respiration rates.

On the tundra, the ¹⁴C content of the CO₂ respired during the winter was similar 117 to that collected the previous September (Fig. 2), while in the birch forest, the ${}^{14}C$ 118 119 content of winter-respired CO₂ was significantly greater than at any point during the growing season. Mean residence time modelling, based on soil ¹⁴C measurements, 120 121 indicated that C fixed before the 1950s should contribute only a small proportion to total CO₂ release (Hartley et al., 2012). Therefore, the increase in the ¹⁴C content of the 122 winter-respired CO₂ in the birch forest indicates that more 'older' C, enriched in 14 C 123 from 20th century nuclear weapons testing, was being released (Fig. 2). This was 124 125 possibly caused by the gradual loss of recently-fixed, labile C which would have been relatively ¹⁴C-depleted (atmospheric $CO_2 = 105.15$ % modern). This process may have 126 127 been more pronounced in the forest due to both the smaller soil C stocks and the greater 128 inputs of contemporary C associated with the higher plant productivity (Hartley et al. 129 2012). Overall, our results indicate that the decomposition of decade-old SOM can 130 continue during the protracted arctic winter in both forest and tundra ecosystems 131 (Table 1). The fact that such C can be released during the Arctic winter makes it 132 possible for changes in winter conditions to affect substantially the C balance of arctic ecosystems. Finally, in the future, comparative analyses of the CO₂ released from both 133 134 plant-free and vegetated plots, would help further identify the sources of winter-respired 135 CO₂ in intact Arctic ecosystems.

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Table 1

The ¹⁴C content and δ^{13} C values of samples collected from the tundra-heath and birch forest, with associated measurement uncertainty (±1 σ) and radiocarbon laboratory codes. The δ^{13} C values have been corrected for fractionation during passive sampling (Garnett et al., 2009). The ¹⁴C content of the respired CO₂ was calculated after correction for contamination with atmospheric CO₂ based on the average δ^{13} C value of the atmospheric CO₂ at the site, and the δ^{13} C of pure samples of respired CO₂ collected from monoliths incubated on site in closed containers (see Hartley et al., 2012). The average age of the respired CO₂ (relative to the sampling date) was calculated from its bomb-¹⁴C concentration by reference to records of direct atmospheric ¹⁴CO₂ measurements (Levin et al., 2008).

Site	¹⁴ C (%modern)	Collected CO ₂ Measured δ^{13} C	Corrected δ^{13} C	Lab code	Monolith δ^{13} C ratio	Atmospheri ¹⁴ C (%modern)	$c CO_2 \delta^{13}C$ ratio	Atmospheric Fraction	Respired CO ₂ ¹⁴ C (%modern)	Age (years)
Tundra	108.30±0.51	-24.4±0.1	-20.4	SUERC-19528	-26.60	105.16	-8.8	0.350	109.98	9
Tundra	108.19±0.48	-25.3±0.1	-21.3	SUERC-19529	-26.60	105.16	-8.8	0.300	109.49	8
Tundra	108.61±0.51	-24.4±0.1	-20.4	SUERC-19532	-26.60	105.16	-8.8	0.349	110.46	10
Forest	108.70±0.51	-24.0±0.1	-20.0	SUERC-19533	-26.83	105.16	-8.8	0.380	110.86	11
Forest	109.67±0.49	-26.5±0.1	-22.5	SUERC-19534	-26.83	105.16	-8.8	0.243	111.12	11
Forest	110.25±0.52	-26.2±0.1	-22.2	SUERC-19535	-26.83	105.16	-8.8	0.259	112.04	13

Figure legends

Fig. 1. Photographs showing the installation of one of the systems for passively sampling soil respiration during the arctic winter. Panel (a) shows the hydrophobic filter inside the collar cover, prior to lid being attached. Panel (b) shows the MSC cartridge being attached, before the clips were removed, while panel (c) shows the final arrangement after the cartridge has been covered in insulating foam, protected inside pipe and plastic guttering, pegged in place, taped up and surrounded by stones. Although the sampling system was only 7 cm tall, the stones were arranged to smooth out the vertical profile and minimise any impact on snow drifting patterns.

Fig. 2. The bars indicate the ¹⁴C content of the CO₂ respired from two sites during the winter. Mean values ±1SE are shown (n = 3). The ¹⁴C contents of the CO₂ released during the previous growing season are also indicated (May/June, light grey line; July, dark grey line; September, black line; see also Hartley et al., 2012). The dashed line indicates the ¹⁴C content of the CO₂ in the contemporary atmosphere (~105.16% modern. On the tundra-heath, the ¹⁴C content of the winter respired CO₂ did not significantly differ from the growing season measurements made in July and September [All statistical tests were carried out using the SPSS version 16 (SPSS Science, Birmingham, UK)]. In contrast, in the birch forest, the '*' indicates that the winter respired CO₂ was significantly enriched in ¹⁴C compared with all growing season measurements (P < 0.05, repeated measures ANOVA). The winter respired CO₂ was also significantly more enriched in ¹⁴C in the birch forest compared with the tundra-heath (P < 0.05, *t*-test).





