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Pedobiologia

DOI:
[10.1016/j.pedobi.2016.07.001](https://doi.org/10.1016/j.pedobi.2016.07.001)

Published: 01/07/2016

Peer reviewed version

[Cyswllt i'r cyhoeddiad / Link to publication](#)

Dyfyniad o'r fersiwn a gyhoeddwyd / Citation for published version (APA):
Marella, V. S., Hill, P., Jones, D., & Roberts, P. (2016). Microbial turnover of above and belowground litter components in shrublands. *Pedobiologia*, 59(4), 229-232.
<https://doi.org/10.1016/j.pedobi.2016.07.001>

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1 **Title: Microbial turnover of above and belowground litter components in**
2 **shrublands**

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22 **Abstract**

23 Shrublands cover a large proportion of the world's land surface, yet they remain
24 poorly studied in comparison to other ecosystems. Within shrublands, soil organic
25 matter (SOM) is replenished from inputs of both above- and below-ground plant litter,
26 however, their relative importance depends on their respective turnover rates. To
27 critically address this, we measured the biodegradation rates of the soluble and
28 insoluble components of ^{14}C -labelled above- and below-ground plant litter in soil.
29 During the 150 day incubation, the amount of plant-derived soluble-C lost as $^{14}\text{CO}_2$ was
30 similar for the different plant parts being $64.7 \pm 2.3\%$ for roots, $72.1 \pm 7.4\%$ for stems,
31 and $72.4 \pm 1.8\%$ for leaves. In comparison, the turnover of the insoluble fraction was
32 much slower. However, again little difference in mineralisation was seen for the
33 different plant parts with the total losses being $21.1 \pm 0.9\%$ for roots, $19.5 \pm 1.6\%$ for
34 stems, and $19.6 \pm 1\%$ for leaves. A double exponential first order kinetic model fitted
35 well to the experimental data. It also allowed the partitioning of C between microbial
36 anabolic and catabolic processes for the soluble C component. Using this model, we
37 deduced that the soluble fraction turns over *ca.* 40 times annually, whereas it takes *ca.*
38 2.5 years to turnover the insoluble fraction. For the soluble plant component, the overall
39 microbial carbon use efficiency (CUE) was estimated to be greater for root-derived C
40 in comparison to that derived from aboveground (no difference was observed for the
41 insoluble component). From this, we tentatively suggest that C sourced from
42 belowground plant components may persist longer in soil than C derived from
43 aboveground plant components.

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45 Key words: belowground carbon storage, mineralisation, nutrient cycling, litter
46 decomposition, root turnover

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48 Soil organic matter (SOM) represents a major store of terrestrial carbon (C)
49 (Schlesinger, 1997) and its turnover and replenishment represents a critical component
50 of the global C cycle. SOM is primarily derived from the continual input of above- and
51 below-ground plant components, however, their relative importance, particularly in
52 shrubland ecosystems, remains poorly understood (Vogt et al., 1986). Earlier studies
53 have suggested that plant roots contribute a larger proportion of C to soil organic carbon
54 (SOC) than plant shoots, due to their greater chemical recalcitrance in relation to
55 microbial enzymatic breakdown (Broadbent and Nakashima, 1974; Jane et al., 2007).
56 In contrast, within some agroecosystems, significant contributions by crop shoots have
57 also been observed (Barber, 1979).

58 The input of organic matter to the soil can be broadly classified into two pools
59 (van Hees et al., 2005). The first pool is described as the dissolved organic C component
60 that includes low molecular weight, highly bioavailable compounds such as organic
61 acids, peptides, amino acids, mono- and oligo-saccharides, amino sugars, phenolics and
62 siderophores (McKeague et al., 1986). The second pool consists of plant polymers such
63 as cellulose, hemicellulose, lignin and some proteins, which are relatively resistant to
64 microbial attack (Kalbitz et al., 2000). These two pools can have vastly different C:N:P
65 ratios which may subsequently influence their rate of processing and also microbial
66 carbon use efficiency (CUE; Schmidt et al., 2011).

67 Numerous studies have described the mineralisation of individual low molecular
68 weight compounds (Glanville et al., 2012), plant material (Simfukwe et al., 2011) and
69 have measured the subsequent rates of $^{14}\text{CO}_2$ evolution and/or microbial incorporation.
70 These studies have enhanced our understanding of the ^{14}C mineralisation process of
71 single or occasionally combinations of simple C compounds by the microbial

72 community. However, plant material consists of vast range of compounds
73 (Buckingham, 1993) and the mineralisation capacity of microorganisms to act upon
74 more complex suite of substrates provides a more representative estimate of the
75 potential for C storage in soil. Therefore, the aim of this study was to assess the
76 microbial turnover of the soluble and insoluble fractions of above- and below-ground
77 plant components (root, stem, leaf) from a common shrubland plant to assess their
78 persistence in the soil under laboratory conditions.

79 Soil was obtained from the Henfaes experimental station located in
80 Abergwyngregyn, Gwynedd, North Wales (53°14'N, 4°01'W) UK. The sandy clay loam
81 textured soil is classified as a Eutric Cambisol (FAO) or Dystric Eutrudepts (US Soil
82 Taxonomy) (see SM₁ and Table S₁). *Cistus monspeliensis* L. plants were grown in a
83 hydroponic system consisting of 50% strength Long Ashton nutrient solution under
84 laboratory conditions. Plants were labelled with ¹⁴C twice, 3 days apart for 5 h each
85 time to get sufficient translocation of ¹⁴C to all plant components (see SM₂).
86 Immediately after the second labelling, the plant components were separated into
87 leaves, stem, and roots and air-dried. The dried plant parts were finely ground using a
88 ball mill and stored in 50 ml polypropylene tubes at 20°C for further analysis. The
89 distribution of ¹⁴C label among soluble and structural fractions of plant material was
90 determined by performing a sequential chemical extraction. These results were tested
91 in parallel with unlabelled plants, using an automated fibre analyser (see SM₃). The
92 soluble and insoluble fraction from each of the three plant components were separated
93 using a hot water extract (see SM₄) and amended to field-moist soil contained in 50 cm³
94 polypropylene tubes. The mineralisation of the ¹⁴C-labelled components was studied
95 for 150 days and values were expressed as a percentage of the initial amount of ¹⁴C
96 applied to the soil (see SM₅). Similar extraction process was conducted with unlabelled

97 plant components and the soluble fraction from each component was analysed for
98 distribution of low molecular weight (≤ 300 Da) compounds using MALDI-TOF mass
99 spectrometry (Bruker Reflex IV) with TiO_2 as a matrix. At the end of the incubation
100 period, the amount of soluble ^{14}C remaining in the soil either as unaltered plant material
101 or fixed in the microbial biomass was determined by extracting the soil in $0.5\text{ M K}_2\text{SO}_4$
102 (see SM₆). A double exponential first order decay model was then fitted to the
103 experimental data (Glanville et al., 2016). Substrate-C pool distribution within the
104 microbial community, decay constants, CUE and half-lives (Newton-Raphson iteration
105 method) (Oburger and Jones, 2009) were calculated (see SM₇). The data was analysed
106 by one-way ANOVA with Post-Hoc least significant difference test using SPSSv20.0
107 (SPSS Inc., Chicago, IL) using $P < 0.05$ as an indication of statistical significance.

108 Following the labelling process, the distribution of ^{14}C into soluble and
109 structural fractions of the different plant components was broadly similar to the total
110 amount of unlabelled ^{12}C in each chemical fraction, although the data for stems is not
111 available (Table S₂). This indicates a fairly uniform dilution of the ^{14}C isotope within
112 the plant. The addition of ^{14}C -labelled soluble and insoluble fractions to soil caused an
113 initial rapid phase of $^{14}\text{CO}_2$ evolution followed by a secondary slower phase,
114 irrespective of plant tissue type (Fig. 1). The overall amount of ^{14}C mineralisation in
115 soils amended with soluble fractions was substantially higher compared to the values
116 obtained for the insoluble fractions ($P < 0.001$). This was presumably due to the
117 presence of more labile low molecular weight compounds in the soluble fractions.
118 Conversely, insoluble fractions broadly consist of structural polymers which require
119 enzymatic depolymerisation to promote solubilisation prior to uptake and assimilation
120 by the microbial community (van Hees et al., 2005). Among the soluble fractions, root-
121 derived ^{14}C showed the fastest mineralisation rate followed by stem and leaf ^{14}C during

122 the first hour, presumably because of relatively higher quantities of low molecular
123 weight compounds which exist in roots (Figs. S₁ and S₂). After 24 h, the amount of ¹⁴C
124 mineralisation of the root soluble fraction ($19.7 \pm 0.4\%$) was substantially higher than
125 for the stems ($8.7 \pm 0.3\%$) and leaves ($5.7 \pm 0.3\%$). Similarly, among the insoluble
126 fractions, the root-derived ¹⁴C fraction had the highest initial mineralization rate (0.62
127 $\pm 0.2\%$) within 24 h, followed by the stems ($0.43 \pm 0.02\%$) and leaves ($0.26 \pm 0.01\%$).
128 However, at the end of 150 days, the pattern had changed with $64.7\% \pm 2.3$, $72.1 \pm$
129 7.4% , and $72.4 \pm 1.8\%$ of the soluble fraction lost for the root, stem and leaf-derived
130 ¹⁴C, respectively. In contrast, for the three insoluble fractions the amount recovered as
131 ¹⁴CO₂ after 150 d was very similar, being $21.1 \pm 0.9\%$, $19.5 \pm 1.6\%$, and $19.6 \pm 1\%$ of
132 the total ¹⁴C added for the root, stem and leaves respectively.

133 The amount of ¹⁴C allocated to the rapid mineralisation pool (a_1) and
134 corresponding decay constant values (k_1) were much higher for soluble fractions than
135 insoluble fractions (Table 1), presumably due to their rapid assimilation by microbial
136 biomass (Boddy et al., 2007). This is supported by the lack of soluble-¹⁴C recovered
137 from the soil after 150 d (Fig. 2). The half-life periods calculated from k_1 for the
138 insoluble fractions were 3-5 fold longer than that of the soluble fraction. However, the
139 k_2 values were very low (100-200 times lower than the k_1 values) for both soluble and
140 insoluble fractions and were significantly different. Using the Newton-Raphson
141 iteration method, the combined half-life period for both pools together (a_1+a_2) was *ca.*
142 9 and 930 d for the soluble and insoluble fractions respectively (Oburger and Jones,
143 2009). Thus, soluble fractions turnover *ca.* 40 times annually, whereas insoluble
144 fractions take *ca.* 2.5 years to turnover.

145 It was interesting to note that approximately 20% more soluble C derived from
146 the aboveground plant components (leaf and stem) was allocated to microbial catabolic

147 C pools (pool a_1) than soluble C derived from the belowground component (despite
148 having an initial slower ^{14}C mineralisation rate). Conversely, more root-derived soluble
149 ^{14}C was allocated to anabolic microbial processes (pool a_2) thus resulting in a higher
150 CUE for the below-ground soluble component (Glanville et al., 2016). Hence, microbes
151 have shown more efficient usage of root soluble ^{14}C compared to leaf and stem which
152 could be major driver for ecosystem C storage potential (Sinsabaugh et al., 2013). Thus,
153 we tentatively suggest that C sourced from belowground plant components persists
154 longer than the above ground plant components in soil. However, overall contributions
155 can only be calculated once the total flux of each component into the ecosystem is
156 known. In addition, the amount of C associated with mycorrhizal turnover and root
157 exudation would be needed to complete the budget. Nevertheless, the results obtained
158 here highlight the importance of roots in soil C storage especially as plants in most
159 shrublands heavily invest in belowground biomass in the form of a deeper root system
160 (Meyer, 2011). Results also support suggestions that increased allocation of C to roots
161 under elevated atmospheric CO_2 may partially mitigate atmospheric CO_2 rise by
162 increasing soil C storage (Madhu and Hatfield, 2013).

163 In conclusion, this study has clearly demonstrated the faster mineralisation of
164 soluble fractions compared to the insoluble fractions. Additionally, modelling of the C
165 pools tentatively suggests the longer persistence of belowground components in soil
166 relative to shoots and leaves.

167

168 **Acknowledgements**

169 We would like to thank Barry Grail for his assistance in MALDI-TOF MS
170 analysis, Helen Glanville for her helpful advice ^{14}C modelling and Jonathan Roberts
171 for technical support.

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