

promoting access to White Rose research papers



Universities of Leeds, Sheffield and York
<http://eprints.whiterose.ac.uk/>

This is an author produced version of a paper published in **Ecological Engineering**

White Rose Research Online URL for this paper:

<http://eprints.whiterose.ac.uk/id/eprint/77447>

Paper:

Armstrong, A, Holden, J, Luxton, K and Quinton, JN (2012) *Multi-scale relationship between peatland vegetation type and dissolved organic carbon concentration*. Ecological Engineering, 47. 182 - 188. ISSN 0925-8574

<http://dx.doi.org/10.1016/j.ecoleng.2012.06.027>

1 **Multi-scale relationship between peatland vegetation type and dissolved organic carbon**
2 **concentration**

3 **Armstrong, A.*¹, Holden, J.², Luxton, K.³, and Quinton, J. N.⁴**

4 ¹School of Geographical and Earth Sciences, University of Glasgow, Glasgow, G12 8QQ,
5 UK

6 ²School of Geography, University of Leeds, Leeds, LS2 9JT, UK

7 ³United Utilities, Haweswater House, Lingley Mere Business Park, Great Sankey,
8 Warrington, Cheshire, WA5 3LP, UK

9 ⁴Lancaster Environment Centre, Lancaster University, Lancaster, LA1 4YQ, UK

10

11 *Corresponding author: Alona Armstrong; alona.armstrong@ges.gla.ac.uk

12

13 **Abstract**

14 Dissolved organic carbon (DOC) is a key component of the carbon cycle and has significant
15 impacts on aquatic ecosystems and potable water treatment. Upward trends in river and
16 lacustrine DOC concentrations have been observed and a number of key drivers have been
17 proposed. Here, we present DOC concentration data from plot scale pore waters at one site
18 and surface water from artificial drains sampled within a national survey which demonstrate a
19 significant correlation between peatland vegetation type and DOC concentration. *Calluna*
20 dominance was associated with the highest DOC concentration, *Molinia* and *Sphagnum*
21 dominance with lower concentrations, and sedge dominance with intermediate
22 concentrations. Water sampled from drains dominated by *Calluna* had greater DOC
23 concentrations than water sampled from pore waters in plots dominated by *Calluna*. In
24 contrast DOC concentrations from plots dominated by sedges were greater than those
25 sampled from drains dominated by sedges. We discuss these findings in relation to plant

26 functional traits and their influence on the physical and biotic conditions that regulate DOC
27 concentrations. Given the known effects of management activities and climate change on
28 peatland vegetation composition there is potential to manage plant community composition to
29 ameliorate the observed rising DOC concentration.

30

31 **Key words**

32 Moorland; vegetation; plant functional type; dissolved organic carbon; peatland; water
33 quality

34

35 **1. Introduction**

36 Peatlands are important terrestrial carbon stores, containing 20 to 30% of the global soil
37 carbon stock (Gorham 1991). Aquatic carbon loss is an important component of the peatland
38 carbon cycle, accounting for 30 to 50% of net ecosystem exchange fixed carbon (Dinsmore
39 and others 2010; Nilsson and others 2008; Roulet and others 2007). Peatlands release carbon
40 through sub-surface pathways as DOC and there is mounting evidence in the fluvial and
41 lacustrine literature that DOC concentrations, [DOC], in waters flowing from peatland
42 catchments are increasing in many parts of the northern hemisphere (Evans and others 2005;
43 Skjelkvåle and others 2005; Stoddard and others 2003; Worrall and others 2004), prompting
44 concern that these important terrestrial carbon stores are becoming carbon sources (Freeman
45 and others 2001a; Freeman and others 2004). The increase in [DOC] is a major problem in
46 water supply areas as DOC is strongly linked to water discolouration which has small legal
47 limits and also disinfection of the water results in reactions with the DOC to produce harmful
48 bi-products. Thus [DOC] is very important in water supply areas, and in the UK uplands
49 many water companies receive a large proportion of their supply from peatlands. It is not
50 possible to turn away these peat water supplies as they are critical to the volume of water

51 needed to fulfil water demand requirements and so instead water companies have to deal with
52 the increasing [DOC] and costly problem.

53

54 Several global change drivers have been suggested to explain the increases in [DOC],
55 including increased temperatures (Clark and others 2009; Evans and others 2005; Freeman
56 and others 2001a), changes in hydrology (Clark and others 2009; Mitchell and McDonald
57 1992; Tranvik and Jansson 2002), increased atmospheric carbon dioxide concentrations
58 (Fenner and others 2007a; Freeman and others 2004), and changes in atmospheric deposition
59 (Clark and others 2008a; Evans and others 2008; Evans and others 2006a; Hruska and others
60 2009; Monteith and others 2007; Stoddard and others 2003; Tipping and Hurley 1988;
61 Vuorenmaa and others 2006). These drivers have the potential to act independently and
62 interactively to influence processes responsible for DOC production and transport in
63 peatlands.

64

65 The type of vegetation present within a peatland has been shown to influence peatland
66 conditions and resources that could contribute to the production and release of DOC
67 including effects on (1) the geochemistry of soil water (Clymo 1987; Kuhry and others 1993);
68 (2) the physical properties of the peat (including temperature and water table) (McNamara
69 and others 2008); (3) the biological agents which live within the peat (Artz and others 2007;
70 Artz and others 2008); and (4) the quality and quantity of plant and litter inputs (Moore and
71 Dalva 2001; Wickland and others 2007). Plant species can be characterised by functional
72 traits, which are measureable physiological and morphological characteristics including plant
73 structure, litter quantity and quality and root architecture. These functional traits dictate how
74 they assimilate and process carbon and influence soil properties and processes and provide a
75 useful mechanistic framework to understand the relationship between vegetation type and

76 carbon cycling (De Deyn and others 2008; Dorrepaal 2007). In peatland ecosystems ericoid
77 dwarf-shrubs, graminoids and bryophytes are the three dominant plant functional types and
78 have been shown to correlate with carbon dioxide and methane fluxes from UK ombrotrophic
79 peatlands (McNamara and others 2008; Ward and others 2009).

80

81 Both the historic vegetation type, which controls the physical and geochemical characteristics
82 of the peat mass (Ringqvist and Öborn 2002), and the current vegetation type, which controls
83 short-term carbon cycling, could exert control over pore water [DOC]. Evidence from a ¹⁴C
84 carbon dating study has shown that between 96% and 100% of DOC originated from surface
85 peat in three different peatland catchments (Tipping 2010). It has also been proven that in
86 blanket peatlands 80% of runoff originates from the top 5 cm of peat (Holden and Burt 2003),
87 litter and roots from the most recent vegetation, and that stream [DOC] is more strongly
88 correlated to soil water at 1 cm and 5 cm depth compared to 20 cm and 50 cm depth (Clark
89 and others 2008b). Also, experiments examining the impact of elevated CO₂ found significant
90 short-term increases in DOC and attributed this to increased root exudation and increased
91 vascular plant cover (Fenner and others 2007b).

92

93 In this study we aim to examine the value and variation in [DOC] at the plot and artificial
94 drain scale and the relationship with vegetation type. We hypothesised that vegetation type
95 would correlate with peatland [DOC] at the plot and drain catchment scales. To do this we
96 collected DOC samples (1) from soil water within plots of different vegetation types at a
97 single site, Bingley Moor, and (2) from surface water drains from sites throughout England
98 and Scotland.

99

100 **2. Materials & methods**

101 2.1 Field sites

102 *a) Plot soil waters*

103 The plot-scale analysis was undertaken at Bingley Moor, Yorkshire, northern England
104 (53°52'N, 1°8'W). The site consists of blanket peat of approximately 1 m in depth and is at
105 an altitude of 300 m. The site is used for sheep grazing and as a grouse moor, and therefore
106 there is a mosaic of prescribed heather patch burning over some of the site. Five vegetation
107 types - burnt *Calluna* (burnt <2 years ago), *Calluna*, sedges (predominantly *Eriophorum*
108 *angustifolium*), *Sphagnum* and *Molinia* - were selected. Nine one-metre long 22 mm
109 diameter dipwells with perforations along their entire length were installed in each vegetation
110 type on comparable slopes in terms of their angle, aspect, drainage, and peat depth. The exact
111 peat area the dipwells sampled from would vary depending on the peat hydraulic properties
112 and antecedent conditions but we anticipate the sample would be dominated by pore water
113 from the surrounding 1 m of peat. Samples were taken once per week for three weeks in June
114 and July 2008 (24/06/08, 29/06/08 and 06/07/08) using a piece of plastic tubing attached to a
115 syringe, yielding 135 samples for [DOC] analysis.

116

117 *b) National survey drain waters*

118 A national survey was undertaken to examine [DOC] in peatland drains (Armstrong and
119 others 2010). Thirty-two blanket bog sites were visited across England and Scotland (Figure
120 1) between 14/02/06 and 05/12/06 with 180 grab water samples taken from artificial drains
121 and analysed for [DOC]. The drain catchment areas were generally <1 km². The drains had a
122 maximum slope of 9°, ranged from 10 cm to 100 cm deep and from 20 to 280 cm wide. Peat
123 depth was greater than 180 cm at 107 sites and at the remaining 73 the minimum depth was
124 30 cm, the maximum 179 cm with a mean depth of 95 cm. Out of the 180 drains 146 were
125 blocked and 26 were unblocked, 44 had evidence of burning and 107 had evidence of

126 grazing. The vegetation type of each drain catchment was described as either *Calluna*-
127 dominated ($n = 22$), sedge-dominated ($n = 92$), or mixed (varying proportions and types of
128 sedges, *Sphagnum*, and dwarf shrubs, $n = 66$).

129

130 2.2 Laboratory analysis

131 The water samples were filtered through 0.45 μm filters. [DOC] was measured using a
132 Thermalox Total Carbon analyser, which has a precision of $\pm 0.1 \text{ mg C l}^{-1}$ and a minimum
133 detection limit of 1 mg C l^{-1} . Prior to analysis, the DOC samples were acidified and sparged
134 with oxygen in order to stabilise the sample and to remove any inorganic carbon and
135 subsequently analysed in duplicate (or triplicate if $\text{CV} > 1 \%$), with [DOC] determined from a
136 seven-point calibration determined using potassium hydrogen phthalate (KHP). In addition,
137 regular analysis of KHP standards and a certified reference material (VKI QC WW4a)
138 ensured that the level of error was kept to a minimum. All samples were stored in the dark at
139 4°C , filtered and analysed within one week of collection.

140

141 2.3 Data analysis

142 The [DOC] data were logarithmically transformed and statistical significance of vegetation
143 type was tested using a one way analysis of variance (ANOVA) followed by a Bonferroni
144 multiple-comparison test for each of the data sets. The statistical differences between the plot
145 and drain scale data were tested using a t-test with unequal variances and the equality of
146 variances assessed using a F-test. All statistical analysis was undertaken using Stata10
147 (StataCorp 2007) and P values are reported as <0.01 , <0.05 , and < 0.10 .

148

149 **3. Results**

150 3.1 Plot soil water

151 [DOC] ranged from 2.2 to 120.9 mg l⁻¹ with a mean of 24.2 and a standard deviation of 18.6
152 mg l⁻¹. The highest mean and median [DOC] were associated with *Calluna*, then sedges >
153 burnt > *Sphagnum* > *Molinia* (Table 1 & Figure 2). However the highest minimum [DOC]
154 were associated with sedges > burnt > *Sphagnum* > *Molinia* > *Calluna* and the highest
155 maximum [DOC] with *Calluna* > *Sphagnum* > sedges > burnt > *Molinia* (Table 1). The
156 differences in the [DOC] between vegetation types were statistically significantly different
157 ($p < 0.05$) except between burnt *Calluna* and *Sphagnum*, and sedges and *Sphagnum*.
158 Variability in [DOC] as determined by the CV was greatest for *Sphagnum* > *Molinia* >
159 *Calluna* > burnt > sedges whereas the IQR indicates that the greatest range in [DOC] were
160 associated with *Calluna* > sedges > *Sphagnum* > burnt > *Molinia* (Table 1 & Figure 2). There
161 was a weak significant difference ($p < 0.10$) in [DOC] between the first and second sampling
162 days. There was an intense storm prior to the first sampling day: the weather preceding the
163 second and third sample days was dry, although a rainfall event occurred during the third
164 sampling day.

165

166 3.2 National survey drain waters

167 The [DOC] of water sampled from artificial drains within the national survey varied from 4.7
168 to 114.0 mg l⁻¹ with a mean of 33.6 and a standard deviation of 19.7 mg l⁻¹. The highest
169 mean, median, minimum [DOC] were sampled from *Calluna*-dominated catchments, then
170 mixed > sedge-dominated catchments (Table 1 & Figure 3), with statistically significant
171 differences between sedge and *Calluna* ($p < 0.01$) and *Calluna* and mixed ($p < 0.01$). The
172 highest maximum [DOC] were sampled from sedge < *Calluna* < mixed dominated
173 catchments (Table 1). The variability and range, as defined by the CV and IQR, in [DOC]
174 were greatest for sedges > mixed > *Calluna* (Table 1 & Figure 3).

175

176 3.3 Comparison of the *Calluna* and sedge data from the two scales

177 Given that *Calluna* and sedges were common dominant vegetation types in both the plot and
178 drain scale data it is possible to examine the impact of scale on [DOC] and variability. The
179 pore water [DOC] in plots dominated by *Calluna* was less than that of water sampled from
180 drains dominated by *Calluna*, as indicated by the mean, median, and minimum (Table 1),
181 however, the differences were not statistically significant. In contrast the pore water [DOC]
182 in plots dominated by sedges was weakly ($p < 0.10$) higher than that of water sampled from
183 drains dominated by sedges, as indicated by the mean, medium and minimum (Table 1). The
184 CV and IQR of [DOC] were greater in the plot data for *Calluna* and lower in the sedge data
185 in comparison with the national survey data and the F-test indicated the variance was
186 statistically significant greater for the plot data sedge data ($p < 0.01$) but the variances were
187 similar for the *Calluna* data (Table 1).

188

189 **4. Discussion**

190 Our data, collected at two scales and from sites throughout England and Scotland, indicate
191 that there were significant differences in DOC associated with different vegetation types
192 (Table 1). *Calluna* was consistently associated with the highest [DOC], sedges yielded
193 intermediate [DOC] and *Sphagnum* low [DOC]. To our knowledge only Vestgarden *et al.*
194 (2010) have undertaken a field study directly examining the role of vegetation on [DOC],
195 finding that the relative [DOC] of soil water under *Sphagnum*, *Molinia* and *Calluna* patches
196 varied with depth and season. Given differences in methodologies these results cannot be
197 directly compared to those of our study, although it is interesting to note that *Calluna* is
198 associated with the higher [DOC] in both, except at depth. Although not examining the effect
199 of vegetation type on [DOC] directly, Fenner and others (2007b) attributed an increase in
200 [DOC] during an elevated carbon dioxide experiment to a change from predominantly

201 *Sphagnum* to *Juncus effusus* vegetation cover; this is in agreement with the findings of our
202 study.

203

204 The differences in [DOC] of *Calluna* and sedges sampled at the two scales contrast: *Calluna*
205 was associated with higher values at the drain scale and sedges with higher [DOC] at the plot
206 scale. The variability in [DOC] was greatest for *Calluna* at the plot scale and for sedges at the
207 drain scale. Explanations for these differences cannot be resolved in this study but may relate
208 to the sampling windows (June to July 2008 for the plot data and February to December 2006
209 for the drain data), processing of the DOC between the pore water and drainage channels
210 related to differences in the DOC character, the exact vegetation community in the plots and
211 drain catchments, and the connectivity of the pore water to drainage channels.

212

213 Although to our knowledge only Vestgarden *et al.* (2010) have reported a relationship
214 between [DOC] and vegetation type in the field the importance of vegetation type in
215 controlling [DOC] has been shown in soil cores (Neff and Hooper 2002; Vestgarden and
216 Austnes 2009) and indirectly through the use gradients, such as ground wetness (Wickland
217 and others 2007), by changing the amount of litter (Lajtha and others 2005) or by considering
218 different peatland types (Chanton and others 2008). Both Neff and Hooper (2002) and
219 Vestgarden and Austnes (2009) concluded that vegetation cover and composition was at least
220 as important as climatic conditions, which provides evidence that vegetation management
221 may provide a means of ameliorating against the increasing trend in DOC, depending on the
222 success of manipulation and sufficient response times. Furthermore, given that land
223 management activities have a substantial impact on peatland vegetation coverage, both the
224 extent and the species composition, the changes in [DOC] observed in association with land
225 management (Armstrong and others 2010; Clay and others 2009; Evans and others 2006b;

226 Worrall and others 2007a; Worrall and others 2007b; Yallop and Clutterbuck 2009) may be
227 attributable to the effect of vegetation.

228

229 Effect of vegetation type on physical environment conditions

230 Vegetation controls the physical environmental conditions by effecting the water table depth
231 and temperature. Lower water tables are associated with higher DOC production given the
232 increased aerobic zone (Clark and others 2009; Clymo 1987). It is commonly accepted that
233 water table influences the vegetation type (Heathwaite and others 1993) and that blanket
234 peatland topography may also impact the local water tables (Holden 2005a), but different
235 physiological characteristics of different plant functional types can cause differences in water
236 table depths in the same hydrological setting by effecting hydraulic conductivities (Holden
237 2005b; Holden 2009; Holden and others 2001) and evapotranspiration. Vegetation primarily
238 impacts the water table depth by affecting the evapotranspiration rate as different plant
239 functional types have different transpiration rates, intercept varying amounts of water which
240 then evaporates, and afford different surface cover and therefore influence evaporation direct
241 from the peatland surface (Gilman 1994; Lafleur and others 2005; Schouwenaars 1993).
242 While difficult to isolate (Kim and Verma 1996; Schouwenaars 1993), the differences in
243 interception losses between vegetation types are likely to dominate over differences in
244 evaporation and transpiration given ground cover is often high and transpiration losses are
245 limited over the short-growing season in upland areas (Wright and Harding 1993). This
246 reasoning suggests that lower water table depths will be associated with shrubs and sedges (as
247 found by McNamara (2008)), and therefore high [DOC] would be expected in soil water
248 under shrubs and low [DOC] from soil water sampled from under *Sphagnum*, as was found in
249 our data (Figure 2).

250

251 The other physical control of vegetation on DOC is soil temperature with significantly
252 different temperatures measured under different vegetation types (McNamara and others
253 2008). Higher soil temperatures, up to a threshold, are associated with higher decomposition
254 rates (Cole and others 2002; Davidsons and Janssens 2006; Dioumaeva and others 2003),
255 increased exudation of DOC from roots (Uselman and others 2004), and evapotranspiration.
256 However our data show the highest [DOC] were associated with *Calluna* (Figure 2), whereas
257 McNamara (2008) recorded the lowest temperatures under *Calluna*, thus we hypothesise that
258 temperature is not a causal mechanism of the association between vegetation and [DOC].
259 This is in agreement with larger scale studies examining the role of temperature on [DOC]
260 (Freeman and others 2001a; Pastor and others 2003).

261

262 Effect of vegetation on biotic conditions

263 There are a range of mechanisms by which the biotic conditions influence [DOC]. Different
264 rates and quality of root exudates (Yan and others 2008) contribute to [DOC] directly but also
265 stimulate microbial activity (Freeman and others 2004), are a potential biological control on
266 [DOC]. However, given the limited data on the amount and quality of root exudates and the
267 consumption of DOC during microbial respiration it is not possible to assess the relationship
268 between root exudation and pore water and channel [DOC].

269

270 Differences in [DOC] and composition between different litters have been determined in the
271 laboratory (Cleveland and others 2004; Moore and Dalva 2001) and thus is an established
272 mechanism by which vegetation type may influence [DOC]. The differences in [DOC]
273 leached from different litters can be attributed to the chemical controls of vegetation over
274 [DOC] due to litter and root exudation, including those associated with pH (Kuhry et al 1993;

275 Clymo, 1987), compounds which inhibit decomposition (Turetsky 2003; Freeman and others
276 2001b), and nutrient concentrations (Aerts and others 1999; Coulson and Butterfield 1978).

277

278 Finally, microbial assemblages and soil fauna populations have been found to vary with
279 vegetation type (Artz and others 2007; Artz and others 2008; Coulson and Butterfield 1978;
280 Standen and Latter 1977). As both microbe and soil fauna decompose organic matter their
281 numbers and types will influence [DOC]. Quantification of DOC produced by organisms is
282 difficult to obtain (Møller and others 1999) given the dependencies, however, higher
283 numbers of soil fauna have been associated with *Calluna* (Coulson and Butterfield 1978;
284 Standen and Latter 1977) which was characterised by high [DOC] in our study.

285 Consequently, we suggest that the numbers and type of microbes and fauna are an important
286 driver in the [DOC]-vegetation type relationship.

287

288 **5. Conclusion**

289 We conclude that there is a correlative relationship, which was apparent in soil water sampled
290 from plots and in water sampled from drains within a national survey, between vegetation
291 type and [DOC]. Higher [DOC] were associated with *Calluna* and lower concentrations with
292 sedges and *Sphagnum*. Potential mechanisms of causality, based on the different physical and
293 biotic conditions associated with different plant functional types, were identified in the
294 literature, including water table depth, temperature, microbial assemblages, root exudates and
295 litter quantity and quality. We recognise that there are topographic and hydrological controls
296 that determine where different plant types grow within a peatland, but these plants in turn
297 impact local water tables, peat growth and local topography. Further research effort is
298 required to elucidate the dominant causal drivers between plant functional traits and DOC
299 production and transport. Furthermore, these results from the two scales raise the possibility

300 of managing vegetation to control [DOC] to reduce carbon losses from peatlands and to
301 reduce water treatment costs for potable supplies. Such management might include actively
302 spreading *Sphagnum* to encourage its propagation and raising water tables to encourage
303 *Sphagnum* regeneration at the expense of more shrubby vegetation such as *Calluna*. Once
304 *Sphagnum* establishes, species like *Calluna* may be reduced by overgrowth with *Sphagnum*
305 and by the wetter surface conditions within the peatland. These management strategies would
306 be in line with those undertaken in many peatland restoration projects and may be cost-
307 effective in the long-term for organisations such as water companies to buy into as both
308 capital and operational treatment costs for DOC and water discolouration at water treatment
309 works are high. The data we present also has important implications for paired catchment
310 studies which examine the impact of treatments, such as grazing, burning, or restoration
311 activity, and highlights the necessity to assess vegetation cover in addition to the standard
312 morphological and hydrological variables.

313

314 **8. Acknowledgements**

315 We would like to thank Yorkshire Water for funding the research. The views expressed
316 herein are not necessarily those of Yorkshire Water. Brenda Cookson, Paddy Keenan and
317 David Ashley are thanked for their assistance in the field and laboratory. Nick Ostle and
318 Susan Waldron provided useful comments on the manuscript. The support of a Philip
319 Leverhulme Prize awarded to Joseph Holden is also gratefully acknowledged as this provided
320 the time for his contributions to the paper.

321 Figure 1. Map of national survey sampling sites.

322

323 Figure 2. [DOC] of soil water samples taken from different vegetation plots.

324

325 Figure 3. [DOC] of water sampled from surface drains dominated by *Calluna*, mixed
326 vegetation and sedges within a national survey.

327

328 9. References

- 329 Aerts, R.; Verhoeven, J.T.A.; Whigham, D.F. Plant-mediated controls on nutrient cycling in
330 temperate fens and bogs. *Ecology*. 80:2170-2181; 1999
- 331 Armstrong, A.; Holden, J.; Kay, P.; Francis, B.; Foulger, M.; Gledhill, S.; McDonald, A.T.;
332 Walker, A. The impact of peatland drain-blocking on dissolved organic carbon loss
333 and discolouration of water; results from a national survey. . *Journal of Hydrology*;
334 2010
- 335 Artz, R.R.E.; Anderson, I.C.; Chapman, S.J.; Hagn, A.; Schloter, M.; Potts, J.M.; Campbell,
336 C.D. Changes in fungal community composition in response to vegetational
337 succession during the natural regeneration of cutover peatlands. *Microbial Ecology*.
338 **54**:508-522; 2007
- 339 Artz, R.R.E.; Chapman, S.J.; Siegenthaler, A.; Mitchell, E.A.D.; Buttler, A.; Bortoluzzi, E.;
340 Gilbert, D.; Yli-Petays, M.; Vasander, H.; Francez, A.J. Functional microbial
341 diversity in regenerating cutover peatlands responds to vegetation succession. *Journal*
342 *of Applied Ecology*. 45:1799-1809; 2008
- 343 Chanton, J.P.; Glaser, P.H.; Chasar, L.S.; Burdige, D.J.; Hines, M.E.; Siefel, D.I.; Tremblay,
344 L.B.; Cooper, W.T. Radiocarbon evidence for the importance of surface vegetation on
345 fermentation and methanogenesis in contrasting types of boreal peatlands. *Global*
346 *Biogeochemical Cycles*. 22:GB4022; 2008
- 347 Clark, J.; Evans, C.; Bottrell, S.; Monteith, D.; Ratcliffe, M.; Chapman, P. Have increased
348 dissolved organic carbon (DOC) losses from UK peat and organo-mineral soils been
349 driven by the decline in acid rain? *Proceedings of the 13th International Peat*
350 *Congress*. Tullamore, Ireland; 2008a
- 351 Clark, J.M.; Ashley, D.; Wagner, M.; Chapman, P.J.; Lane, S.N.; Evans, C.D.; Heathwaite,
352 A.L. Increased temperature sensitivity of net DOC production from ombrotrophic peat
353 due to water table draw-down. *Global Change Biology*. 15:794-807; 2009
- 354 Clark, J.M.; Lane, S.N.; Chapman, P.J.; Adamson, J.K. Link between DOC in near surface
355 peat and stream water in an upland catchment. *Science of The Total Environment*.
356 404:308-315; 2008b
- 357 Clay, G.D.; Worrall, F.; Fraser, E.D.G. Effects of managed burning upon dissolved organic
358 carbon (DOC) in soil water and runoff water following a managed burn of a UK
359 blanket bog. *Journal of Hydrology*. 367:41-51; 2009
- 360 Cleveland, C.C.; Neff, J.C.; Townsend, A.R.; Hood, E. Composition, dynamics and fate of
361 leached dissolved organic matter in terrestrial ecosystems: results from a
362 decomposition experiment. *Ecosystems*. 7:275-285; 2004

363 Clymo, R.S. Interactions of Sphagnum with water and air. in: Hutchinson T.C., Meema K.M.,
364 eds. Effects of Atmospheric Pollutants on Forests, Wetlands and Agricultural
365 Ecosystems. Heidelberg: Springer; 1987

366 Cole, L.; Bardgett, R.D.; Ineson, P.; Adamson, J.K. Relationships between enchytraeid
367 worms (Oligochaeta), climate change, and the release of dissolved organic carbon
368 from blanket peat in northern England. *Soil Biology and Biochemistry*. 34:599-607;
369 2002

370 Coulson, J.C.; Butterfield, J. An investigation of the biotic factors determining the rates of
371 plant decomposition on blanket bog. *Journal of Ecology*. 66:631-650; 1978

372 Davidsons, E.A.; Janssens, I.A. Temperature sensitivity of soil carbon decomposition and
373 feedbacks to climate change. *Nature*. 440:165-173; 2006

374 De Deyn, G.B.; Cornelissen, J.H.C.; Bardgett, R.D. Plant functional traits and soil carbon
375 sequestration in contrasting biomes. *Ecology Letters*. 11:516-531; 2008

376 Dinsmore, K.J.; Billett, M.F.; Skiba, U.M.; Rees, R.M.; Drewer, J.; Helfter, C. Role of the
377 aquatic pathway in the carbon and greenhouse gas budgets of a peatland catchment.
378 *Global Change Biology*; 2010

379 Dioumaeva, I.; Trumbore, S.; Schuur, E.A.G.; Goulden, M.L.; Litvak, M.; Hirsch, A.I.
380 Decomposition of peat from upland boreal forest: Temperature dependence and
381 sources of respired carbon. *Journal of Geophysical Research*. 108:8222; 2003

382 Dorrepaal, E. Are plant growth-form-based classifications useful in predicting northern
383 ecosystem carbon cycling feedbacks to climate change? *Journal of Ecology*. 95:1167-
384 1180; 2007

385 Evans, C.; Monteith, D.; Reynolds, B.; Clark, J. Buffering of recovery from acidification by
386 organic acids. *Science of the Total Environment*. 404:316-325; 2008

387 Evans, C.D.; Chapman, P.J.; Clark, J.M.; Monteith, D.T.; Cresser, M.S. Alternative
388 explanations for rising dissolved organic carbon export from organic soils. *Global
389 Change Biology*. 12:2044-2053; 2006a

390 Evans, C.D.; Monteith, D.T.; Cooper, D.M. Long-term increases in surface water dissolved
391 organic carbon: Observations, possible causes and environmental impacts.
392 *Environmental Pollution*. 137:55-71; 2005

393 Evans, M.; Warburton, J.; Yang, J. Eroding blanket peat catchments: Global and local
394 implications of upland organic sediment budgets. *Geomorphology*. 79:45-57; 2006b

395 Fenner, N.; Freeman, C.; Lock, M.A.; Harmens, H.; Reynolds, B.; Sparks, T. Interactions
396 between elevated CO₂ and warming could amplify DOC exports from peatland
397 catchments. *Environmental Science & Technology*. 41:3146-3152; 2007a

398 Fenner, N.; Ostle, N.; McNamara, N.; Sparks, T.; Harmens, H.; Reynolds, B.; Freeman, C.
399 Elevated CO₂ Effects on Peatland Plant Community Carbon Dynamics and DOC
400 Production. *Ecosystems*. 10:635-647; 2007b

401 Freeman, C.; Evans, C.D.; Monteith, D.T.; Reynolds, B.; Fenner, N. Export of organic carbon
402 from peat soils. *Nature*. 412 785; 2001a

403 Freeman, C.; Ostle, N.; Kang, H. An enzymic 'latch' on a global carbon store. *Nature*.
404 409:149; 2001b

405 Freeman, C.; Fenner, N.; Ostle, N.J.; Kang, H.; Dowrick, D.J.; Reynolds, B.; Lock, M.A.;
406 Sleep, D.; Hughes, S.; Hudson, J. Export of dissolved organic carbon from peatlands
407 under elevated carbon dioxide levels. *Nature*. 430:195-198; 2004

408 Gilman, K. *Hydrology and Wetland Conservation*. Chichester: John Wiley; 1994

409 Gorham, E. Northern Peatlands: Role in the Carbon Cycle and Probable Responses to
410 Climatic Warming. *Ecological Applications*. 1:182-195; 1991

411 Heathwaite, A.L.; Göttlich, K.H.; Burmeister, E.G.; Kaule, G.; Grospietsch, T.H. Mires:
412 definitions and form. in: Heathwaite A.L., Göttlich K.H., eds. Mires: Processes,
413 Exploitation and Conservation. Chichester: Wiley; 1993

414 Holden, J. Peatland hydrology and carbon cycling: why small-scale process matters.
415 Philosophical Transactions of the Royal Society A. 363:2891-2913; 2005a

416 Holden, J. Piping and woody plants in peatlands: cause or effect? . Water Resources
417 Research. 41:W06009; 2005b

418 Holden, J. Flow through macropores of different size classes in blanket peat. . Journal of
419 Hydrology. 364:342-348; 2009

420 Holden, J.; Burt, T.P. Runoff production in blanket peat covered catchments. Water
421 Resources Research. 39:1191; 2003

422 Holden, J.; Burt, T.P.; Cox, N.J. Macroporosity and infiltration in blanket peat: the
423 implications of tension disc infiltrometer measurements. Hydrological Processes.
424 15:289-303; 2001

425 Hruska, J.; Kram, P.; McDowell, W.H.; Oulehle, F. Increased Dissolved Organic Carbon
426 (DOC) in Central European Streams is Driven by Reductions in Ionic Strength Rather
427 than Climate Change or Decreasing Acidity. Environmental Science & Technology.
428 43; 2009

429 Kim, J.; Verma, S.B. Surface exchange of water vapour between an open sphagnum fen and
430 the atmosphere. Boundary-Layer Meteorology. 79:243-264; 1996

431 Kuhry, P.; Nicholson, B.J.; Gignac, L.D.; Vitt, D.H.; Bayley, S.E. Development of Sphagnum
432 -dominated peatlands in boreal continental Canada. Canadian Journal of Botany.
433 71:10-22 1993

434 Lafleur, P.M.; Hember, R.A.; Admiral, S.W.; Roulet, N.T. Annual and seasonal variability in
435 evapotranspiration and water table at a shrub-covered bog in southern Ontario,
436 Canada. Hydrological Processes. 19:3533-3550; 2005

437 Lajtha, K.; Crow, S.E.; Yano, Y.; Kaushal, S.S.; Sulzman, E.; Sollins, P.; Spears, J.D.H.
438 Detrital controls on soil solution N and dissolved organic matter in soils: a field
439 experiment. Biogeochemistry. 76:261-281; 2005

440 McNamara, N.P.; Plant, T.; Oakley, S.; Ward, S.; Wood, C.; Ostle, N. Gully hotspot
441 contribution to landscape methane (CH₄) and carbon dioxide (CO₂) fluxes in a
442 northern peatland. Science of The Total Environment. 404:354-360; 2008

443 Mitchell, G.; McDonald, A.T. Discoloration of Water by Peat Following Induced Drought
444 and Rainfall Simulation. Water Research. 26:321-326; 1992

445 Møller, J.; Miller, M.; Kjølner, A. Fungal-bacterial interaction on beech leaves: influence on
446 decomposition and dissolved organic carbon quality. Soil Biology and Biochemistry.
447 31:367-374; 1999

448 Monteith, D.T.; Stoddard, J.L.; Evans, C.D.; de Wit, H.A.; Forsius, M.; Hogasen, T.;
449 Wilander, A.; Skjelkvale, B.L.; Jeffries, D.S.; Vuorenmaa, J.; Keller, B.; Kopacek, J.;
450 Vesely, J. Dissolved organic carbon trends resulting from changes in atmospheric
451 deposition chemistry. Nature. 450:537-540; 2007

452 Moore, T.R.; Dalva, M. Some controls on the release of dissolved organic carbon by plant
453 tissues and soils. Soil Science. 166:38-47; 2001

454 Neff, J.C.; Hooper, D.U. Vegetation and climate controls on potential CO₂, DOC and DON
455 production in northern latitude soils. Global Change Biology. 8:872-884; 2002

456 Nilsson, M.; Sagerfors, J.; Buffam, I.; Laudon, H.; Eriksson, T.; Grelle, A.; Klemetsson, L.;
457 Weslien, P.; Lindroth, A. Contemporary carbon accumulation in a boreal oligotrophic
458 minerogenic mire; a significant sink after accounting for all C-fluxes. Global Change
459 Biology. 14:2317-2332; 2008

460 Pastor, J.; Solin, J.; Bridgham, S.D.; Updegraff, K.; Harth, C.; Weishampel, P.; Dewey, B.
461 Global warming and the export of dissolved organic carbon from boreal peatlands.
462 OIKOS. 100:380-386; 2003

463 Ringqvist, L.; Öborn, I. Copper and zinc adsorption onto poorly humified Sphagnum and
464 Carex peat. Water Research. 36:2233-2242; 2002

465 Roulet, N.T.; Lafleur, P.M.; Richard, P.J.H.; Moore, T.R.; Humphreys, E.R.; Bubier, J.
466 Contemporary carbon balance and late Holocene carbon accumulation in a northern
467 peatland. Global Change Biology. 13:397-411; 2007

468 Schouwenaars, J.M. Hydrological differences between bogs and bog-relicts and
469 consequences for bog restoration. Hydrobiologia. 265:217-224; 1993

470 Skjelkvåle, B.L.; Stoddard, J.L.; Jeffries, D.S.; Tørseth, K.; Høgåsen, T.; Bowman, J.;
471 Mannio, J.; Monteith, D.T.; Mosello, R.; Rogora, M.; Rzychon, D.; Vesely, J.;
472 Wieting, J.; Wilander, A.; Worsztynowicz, A. Regional scale evidence for
473 improvements in surface water chemistry 1990-2001. Environmental Pollution.
474 137:165-176; 2005

475 Standen, V.; Latter, P.M. Distribution of a population of *Cognettia sphagnetorum*
476 (*Enchytraeidae*) in relation to microhabitats in a blanket bog. Journal of Animal
477 Ecology. 46:213-229; 1977

478 StataCorp. Stata Statistical Software: Release 10. Texas: Stata Press; 2007

479 Stoddard, J.L.; Karl, J.S.; Deviney, F.A.; DeWalle, D.R.; Driscoll, C.T.; Herlihy, A.T.;
480 Kellogg, J.H.; Murdoch, P.S.; Webb, J.R.; Webster, K.E. Response of surface water
481 chemistry to the Clean Air Act Amendments of 1990. North Carolina, United States:
482 Environmental Protection Agency; 2003

483 Tipping, E. Sources and ages of dissolved organic matter in peatland streams: evidence from
484 chemistry mixture modelling and radiocarbon data. Biogeochemistry; 2010

485 Tipping, E.; Hurley, M.A. A model of solid-solution interactions in acid organic soils, based
486 on the complexation properties of humic substances. European Journal of Soil
487 Science. 39:505-519; 1988

488 Tranvik, L.J.; Jansson, M. Climate change (Communication arising): Terrestrial export of
489 organic carbon. Nature. 415:861-862; 2002

490 Turetsky, M.R. The Role of Bryophytes in Carbon and Nitrogen Cycling. The Bryologist.
491 106:395-409; 2003

492 Uselman, S.M.; Qualls, R.G.; Thomas, T.B. Effects of increased atmospheric CO₂,
493 temperature, and soil N availability on root exudation of dissolved organic carbon by
494 a N-fixing tree (*Robinia pseudoacacia* L.). Plant and Soil. 222:191-202; 2004

495 Vestgarden, L.S.; Austnes, K. Effects of freeze-thaw on C and N release from soils below
496 different vegetation in a montane system: a laboratory experiment. Global Change
497 Biology. 15:876-887; 2009

498 Vestgarden, L.S.; Austnes, K.; Strand, L.T. Vegetation control on DOC, DON, and DIN
499 concentrations in soil water from a montane system, southern Norway. Boreal
500 Environment Research. 15; 2010

501 Vuorenmaa, J.; Forsius, M.; Mannio, J. Increasing trends of total organic carbon
502 concentrations in small forest lakes in Finland from 1987 to 2003. Science of The
503 Total Environment. 365:47-65; 2006

504 Ward, S.E.; Bardgett, R.D.; McNamara, N.P.; Ostle, N.J. Plant functional group identity
505 influences short-term peatland ecosystem carbon flux: evidence from a plant removal
506 experiment. Functional Ecology. 23:454-462; 2009

507 Wickland, K.P.; Neff, J.C.; Aiken, G.R. Dissolved organic carbon in Alaskan boreal forest:
508 sources, chemical characteristics, and biodegradability. Ecosystems. 10:1323-1340;
509 2007

510 Worrall, F.; Armstrong, A.; Adamson, J.K. The effects of burning and sheep-grazing on
511 water table depth and soil water quality in a upland peat. *Journal of Hydrology*. 339:1-
512 14; 2007a

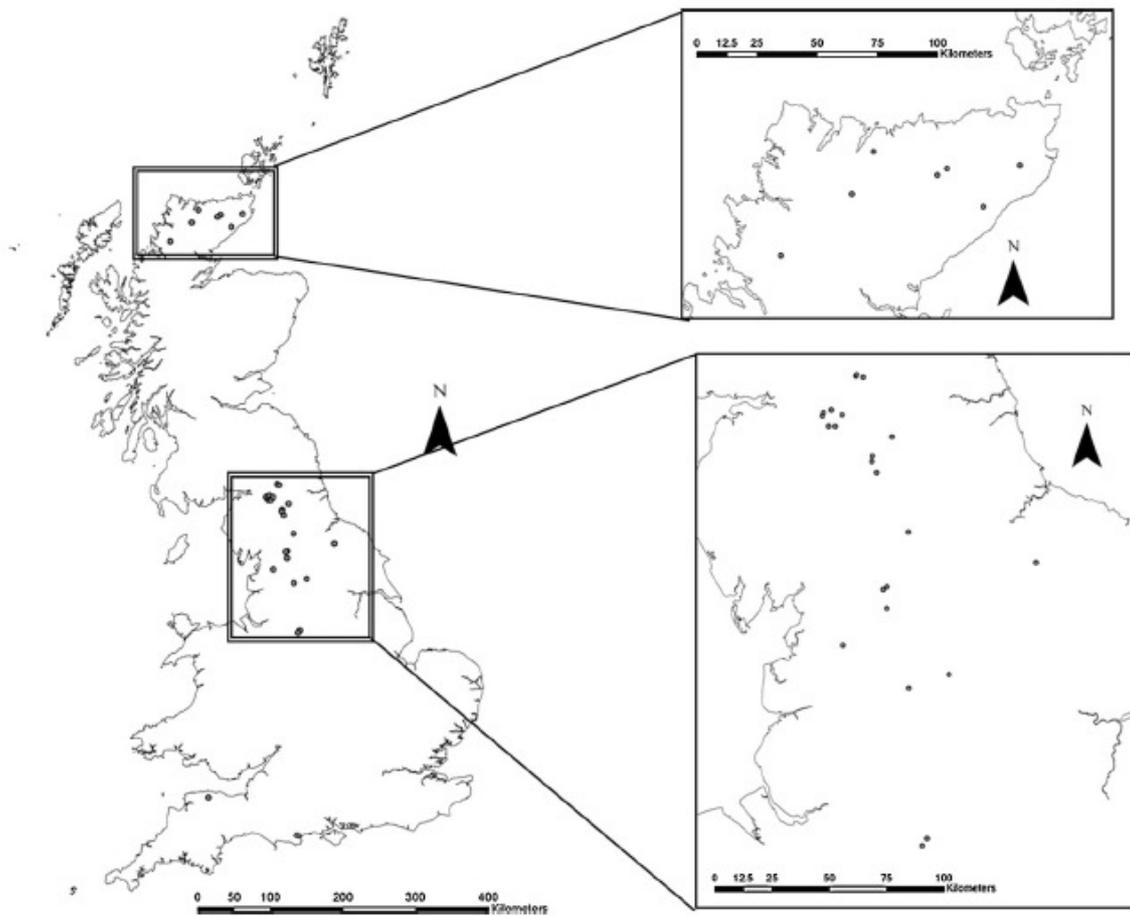
513 Worrall, F.; Armstrong, A.; Holden, J. Short-term impact of peat drain-blocking on water
514 colour, dissolved organic carbon concentration and water table depth. *Journal of*
515 *Hydrology*. 337:315-325; 2007b

516 Worrall, F.; Harriman, R.; Evans, C.D.; Watts, C.D.; Adamson, J.; Neal, C.; Tipping, E.;
517 Burt, T.; Grieve, I.; Monteith, D.; Naden, P.S.; Nisbet, T.; Reynolds, B.; Stevens, P.
518 Trends in Dissolved Organic Carbon in UK Rivers and Lakes. *Biogeochemistry*.
519 70:369-402; 2004

520 Wright, I.R.; Harding, R.J. Evaporation from natural mountain grassland. *Journal of*
521 *Hydrology*. 145:267-283; 1993

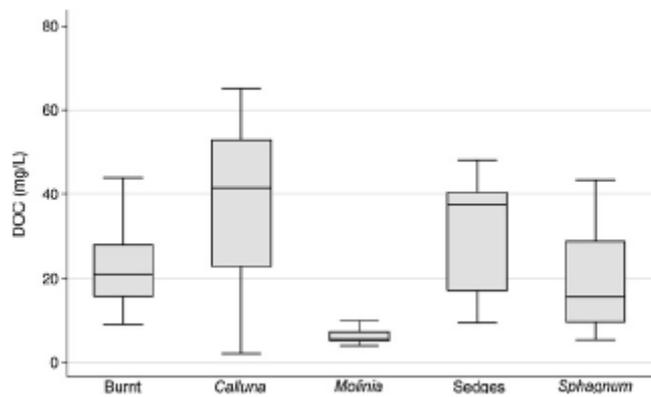
522 Yallop, A.R.; Clutterbuck, B. Land management as a factor controlling dissolved organic
523 carbon release from upland peat soils 1: spatial variation in DOC productivity.
524 *Science of The Total Environment*. 407:3803-3813; 2009

525 Yan, W.; Artz, R.R.E.; Johnson, D. Species-specific effects of plants colonising cutover
526 peatlands on patterns of carbon source utilisation by soil microorganisms. *Soil*
527 *Biology and Biochemistry*. 40:544-549; 2008
528
529

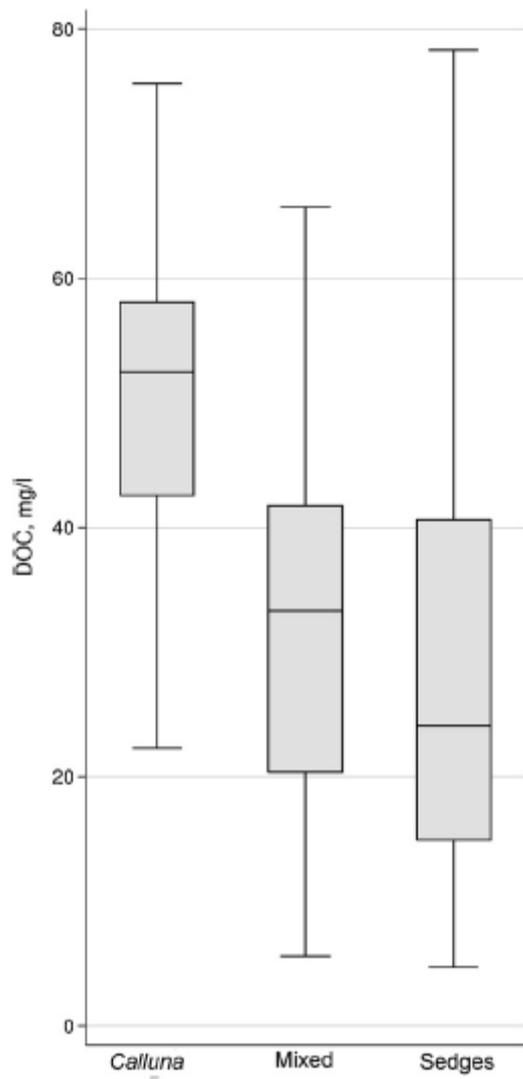


530
531
532
533

A. Arnórsson et al. / Limnology



534
535



536