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Research Paper

Short-term effects of site preparation practices for afforestation on soil properties

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Abstract: A field study was conducted at Harwood Forest, NE England to investigate the effects of forest management practices (drainage, mounding and fertilization) on soil properties from 2006 to 2008. The experiment was laid out in a factorial split plot design on grassland in peaty gley soil. Mounding increased soil bulk density, while drainage reduced carbon concentration in 0–10 cm layer. Soil organic carbon concentration in the 0–10 cm layer was increased by fertilisation. Mounding did not have any effect in soil organic carbon. The concentration of nitrogen in the in 0–10 cm was significantly reduced by drainage and was not affected by mounding or fertilisation. Soil microbial biomass carbon was not affected by drainage, mounding or fertilisation. Ammonium (NH_4^+) was significantly increased by mounding and fertilisation, while none of the treatment affected nitrate (NO_3^-) availability.

Keywords: Drainage, Fertilisation, mounding, peaty gley soil organic carbon, nitrogen

Introduction

Soils are home to a substantial, if not the largest, proportion of biodiversity in terrestrial ecosystems, and form the basis for all forest growth (Bauhus et al., 2002; Hopmas et al., 2005), play a crucial role in the carbon, nutrient (Neill et al., 1997; Bauhus et al., 2002; Hopmas et al., 2005) and hydrological cycles (Bauhus et al., 2002; Hopmas et al., 2005). World's soils constitute a significant reservoir of carbon (C) in both the organic and mineral forms; thereby play a crucial role in mitigating or contributing C to the atmosphere (Zerva and Mencuccini 2005a). Globally, soils are the largest C pool in the terrestrial environment (Wang and Amundson 1999) and contain more than two thirds of the total C stored in living plants (Schlesinger, 1990; Harmon et al., 1990; Schimel, 1995) and almost twice the amount in the atmosphere (Schlesinger, 1990; Schimel 1995).

Successful establishment of forest plantations in peatland sites characterised by poor drainage, low temperatures and poor soil fertility (e.g. Bubier et al., 1998; Strom and Christensen 2007) necessitated provisions for subsurface drainage. Land use changes related to forest management practices can affect the C cycle and C storage in soils (Parfitt et al., 2003) as well as the cycling of nitrogen (N) in soils. The interaction between C and N cycles may influence C storage (Conant et al., 2001) and regulate N availability (Parfitt et al., 2003). These possible soil organic carbon (SOC) losses have detrimental effects on soil structure and soil fertility, contributing to global warming by further increasing the atmospheric concentration of carbon dioxide (CO_2) (von Lützow and Kögel-Knabner 2009).

Afforestation or the conversion of historically treeless areas into forests is a rapidly spreading land-use change which has potential to sequester carbon (Berthrong, 2009). In the UK forest plantations are often established on former grasslands on peaty gley soils that require drainage and mounding to lower the water table and prepare planting spots (Mojeremane 2009). These practices may change the soil's physical, chemical and biological properties (Jurgensen et al., 1997; Merino et al., 1998) and influence the amount, quality and distribution of soil organic matter (SOM) (Paul et al., 2002). Machinery used in forest management operations such as site preparation affect C and N by compacting the soil, thereby altering properties such as bulk density (McNabb et al., 2001; Xu et al., 2002).

Effects of site preparation for afforestation or replanting on soil C and N are not only important because C and N are often the major factors determining soil quality but also because soil act as a C source or sink on a global scale (Johnson and Curtis, 2001). Site preparation and timber harvesting activities may turn soils into sources of C to the atmosphere (Detwiler and Hall 1988). The main site preparation for afforestation in the UK used to be mechanically lowering the soil water table depth by open drainage ditches. Currently, the construction of new drainage channels is discouraged, but the practice of excavating old drainage channels upon replanting is still widespread practice (Mojeremane et al., 2012). While ploughing is discouraged and current best practices favours the use of mounding or surface scarification, there is little qualitative information on the impact of these practices on the soil properties and C and N concentrations. Based on the UK's prevailing environmental conditions, we hypothesised that site preparation would increase soil bulk density and decrease C and N. The objective of this this research was to assess the above mentioned hypothesis.

Materials and Methods

Site description

The field experiment was initiated in May 2006 at Harwood forest, NE England (55°10'N, 2°3'W, 200–400 m asl) on seasonally waterlogged peaty gley with a superficial organic layer varying between 15–40 cm. The forest covers 4000 ha dominated by even aged Sitka spruce (*Picea sitchensis* (Bong.) Carr.) stands. The average annual precipitation and temperature at Harwood are 950 mm and 7.6°C, respectively (Zerva and Mencuccini, 2005a). The establishment of the forest started in the 1930s with planting on moorland and upland rough pasture. Mounding at 2 × 2 m spacing is now used for new planting and restocking

(Mojeremane et al., 2012). The present experiment was established on unplanted unimproved grassland situated between two second rotation Sitka spruce stands. The grassland is dominated by *Festuca ovina* and *Deschampsia flexuosa* with *Calluna vulgaris* and occasionally *Eriophorum vaginatum* and had been used to graze domestic stock a year prior to the study started (Mojeremane, 2009). There was no evidence of drainage present at the site prior to the experiment (Mojeremane, 2009).

Experimental design and layout

The field layout comprised of a full factorial split-plot design with six plots measuring 30 × 8 m each established in May 2006 (Mojeremane, 2009). Three plots were selected at random and drained by an excavator following local standard practices. The open drainage ditches were placed 1.5 m from plot edges and excavated to a depth of 65–70 cm (Mojeremane et al., 2012). About 10-m-wide buffer strips isolated drained and undrained plots. Four subplots (8 × 6 m) isolated by 2-m-wide buffer strips were established in each main plot (Mojeremane et al., 2012). Two randomly selected subplots were mounded, while the remainder were left unmounded with the mounds being ~40-cm wide and 30-cm deep (Mojeremane et al., 2012). Fertilisation was carried out in one mounded and one unmounded subplot randomly chosen within each subplot in a crossed design with mounding (Mojeremane et al., 2012) by applying a compound fertilizer with 81 kg N ha⁻¹, 72 kg P ha⁻¹ and 35 kg ha⁻¹ (Taylor, 1991). Hence, the main plots allowed testing for drainage effects, whereas the subplots allowed testing for mounding, in isolation or combined (Mojeremane et al., 2010, 2012). Each treatment was replicated three times (Mojeremane et al., 2010).

Soil sampling

Soil sampling was done during November 2006, February and August 2007, February and June 2008. Soil samples were collected from four randomly selected locations within subplots. A manually driven square soil corer (5×5cm) was used to obtain samples from 0–20 cm depth (November 2006, and February 2008). August 2007 and June 2008 samples were collected from 0–10, 10–20, and 20–30 cm depth. Soil samples were bulked by subplot to make composite samples and taken to the laboratory in black polythene bags for storage in a freezer (–4°C) awaiting analyses.

Soil pH and bulk density determination

Soil samples were analysed for pH in a 1:2.5 soil/water ratio by a combination glass electrode (Wall and Hytönen, 2005; Xue et al., 2006). The bulk density was determined by the core method (Grossman and Reinsch, 2002) using 5.4 cm diameter and 6 cm long cores. Bulk density was calculated according to Eliot et al. (1999).

$$P_b = M/V$$

Where, P_b is the bulk density (g cm⁻³), M is the dry mass of a given soil sample (g) and V its fresh volume (cm³).

Soil Chemical analyses

For soil organic carbon (SOC) and total N analyses were performed on August 2007 and June 2008 soil samples. Soil samples were passed through a 4 mm sieve and oven dried to constant weight at 60°C. Dried samples were passed through a 2mm sieve using hand applied pressure before grinding in a Ball Mill to pass through a 0.5mm sieve. Both SOC and N concentrations were determined by the dry combustion method (Nelson and Summers, 1982) using a C/N analyser (Carlo-Erba, N 2500). The mass of C and N in soil samples was calculated using the following equation:

$$M_c = M_d \times C/N (\%) / 100$$

Where, M_c is total mass of C or N in the sample; M_d is dry matter of sample; and C or N is the percentage obtained from C/N analyser. C and N concentrations were expressed in g kg⁻¹.

Determination of soil Microbial biomass C

Soil microbial biomass carbon (MBC) was determined from the November 2006, February and August 2007 samples using the previously published chloroform fumigation extraction method (Brookes et al., 1985; Vance et al., 1987). About 20 g of soil were fumigated with ethanol-free chloroform for 24 hrs in a vacuum oven containing a vial with soda lime. Both the fumigated samples and their non-fumigated counterparts (control) were extracted with 80 mL of 0.5M K₂SO₄ on a reciprocal shaker set at 100 rev. min⁻¹ for an hour. Solutions were then transferred into 50 ml tubes and centrifuged at 4000 rev. min⁻¹ for 10 minutes. The supernatant was transferred to 20 ml plastic vials and filtered through 0.45µm Millipore filters. Inorganic C was removed from the supernatant by acidifying to a pH of 2 using a concentrated phosphoric acid and purging with N₂ to degas samples. Organic C in the extract was analysed in an automated total OC analyser (DC-80, Sartec Ltd., Kent, England) with UV-persulphate oxidation and IR detector (Wu et al., 1990). Microbial biomass was calculated using previously published method (Wu et al., 1990; Jorgensen and Mueller, 1996; Jorgensen, 1996). The MBC was calculated as follows:

$$\text{Microbial biomass C} = E_c / K_{EC}$$

Where E_c = (OC extracted from fumigated soil) – (OC extracted from non-fumigated soils) and K_{EC} = 0.45.

Determination of Inorganic N

For analyses of inorganic N (NH₄⁺ and NO₃⁻) August 2007 and February 2008 soil samples were used. Soil was sieved through a 2 mm aperture sieve followed by weighing of 5 g samples into glass bottles. About 100 mL of 1M KCl was added to each sample, sealed and the solution was thoroughly mixed on an orbital shaker set at 150 rev. min⁻¹ for an hour. The solution was then filtered through a filter paper, Ashless Paper 2, and the extracts analysed for NH₄⁺ and NO₃⁻ in a continuous flow Series 3 Auto analyser system.

Determination of above-ground plant biomass evaluation

Above-ground plant biomass was measured in June 2007. Two 1×1 m quadrants were randomly established in each subplot and all plants within quadrants were clipped at ground level using pruning shears and bagged in black polythene bags for

transportation to the laboratory where they were transferred into paper bags and oven dried to constant weight at 80°C. The plant dry biomass (DM) was expressed in tons per hectare of dry mass ($t\ ha^{-1}\ DM$).

Statistical analysis

All data were checked for normality and transformed when required. Data were analysed using the general linear model (GLM) for analysis of variance. The GLM included the effects of drainage, mounding and fertilisation entered as fixed factors and plot entered as a random factor. The initial GLM included all possible second and third order interactions. If interactions were found not significant, they were excluded and model run again without them to confirm the significance of the main factors. In case of significant interactions, the data set was split and separate analysis run for each combination. All analyses were run in Minitab 15 using GLM procedure.

Results

Soil bulk density and pH

The soil bulk density significantly increased with soil depth across treatments ($P=0.0001$). The soil bulk density ranged from 0.12–0.15, 0.18–0.23 and 1.02–1.13 $g\ cm^{-3}$ in the 0–10, 10–20 and 20–30 soil depths across treatments, respectively. Soil bulk density from both sampling occasions was not affected by drainage or fertilisation. Soil bulk density was significantly increased by mounding in the 0–10 cm depth in August 2007 ($P=0.05$) and June 2008 ($P=0.001$). Mounding also increased the soil bulk density in the 10–20 cm depth in both sampling dates (all $P=0.01$). Soil pH varied significantly with soil depth ($P=0.0001$) and was not affected by drainage, mounding or fertilisation. Soil pH varied from 3.9–4.0, 3.8–3.9 and 4.1–4.2 in the 0–10, 10–20 and 20–30cm depth, respectively.

Soil organic carbon

SOC concentrations are shown in Table 1. SOC concentration varied with depth on both sampling dates ($P = 0.0001$). SOC in 0–10 cm depth was significantly decreased by drainage in August 2007 ($P = 0.03$) and June 2008 ($P = 0.04$) but not the 10–20 and 20–30 cm depths. There was a significant decrease in SOC in the fertilised treatment in 0–10 cm layer ($P = 0.02$) in August 2007 only. In none of the sampling occasions was SOC affected by mounding.

Table 1. Effects of drainage, mounding and fertilisation on SOC and N concentration

Sampling date	Parameter	Soil depth	Treatment					
			Drained	Undrained	Mounded	Unmounded	Fertilised	Unfertilised
August 2007	Total C ($g\ kg^{-1}$)	0–10	419.2±13.4a	458.2±5.6b	436.8±13.7a	440.7±9.6a	451.2±7.5a	426.3±14.0b
		10–20	448.3±23.9a	452.0±15.5a	447.6±26.3a	452.7±11.0a	459.9±13.7	440.4±24.6
		20–30	37.5±4.4a	37.2±4.6a	39.7±4.9a	35.1±3.8a	36.8±3.6	38.0±5.2
	Total N ($g\ kg^{-1}$)	0–10	18.3±0.8a	20.8±0.3b	19.5±0.7a	19.6±0.7a	19.4±0.5a	19.7±0.9a
		10–20	15.3±0.7a	15.4±0.7a	15.7±0.9a	15.0±0.5a	15.9±0.6a	14.8±0.8a
		20–30	1.26±0.2a	1.23±0.1a	1.4±0.2a	1.1±0.1a	1.2±0.1a	1.3±0.2a
	C/N ratio	0–10	23.1±0.6a	22.1±0.5a	22.5±0.6a	22.7±0.6a	23.5±0.5a	21.8±0.5b
		10–20	29.2±0.8a	29.7±0.6a	28.5±0.7a	30.3±0.5a	29.2±0.6a	29.7±0.7a
		20–30	30.1±1.2a	30.0±1.1a	29.0±1.2	31.0±1.1a	30.7±1.3a	29.4±0.9a
June 2008	Total C ($g\ kg^{-1}$)	0–10	429.4±8.1a	460.9±5.02b	446.2±8.1a	444.2±8.3a	449.0±7.9a	441.3±8.4a
		10–20	444.2±16.5a	452.0±12.2a	452.2±17.7a	443.9±10.4a	447.2±15.0a	449.0±14.1a
		20–30	35.8±3.8a	41.9±7.1a	39.9±4.4a	37.7±6.9a	35.9±3.3a	41.8±7.4a
	Total N ($g\ kg^{-1}$)	0–10	17.1±0.3a	18.4±0.2b	17.8±0.3b	17.7±0.3a	17.9±0.3a	17.6±0.3a
		10–20	17.8±0.7a	18.0±0.5a	18.1±0.7a	17.7±0.4a	17.9±0.6a	17.9±0.6a
		20–30	1.0±0.2a	1.2±0.3a	1.2±0.2a	1.1±0.3a	1.0±0.1a	1.2±0.3a
	C/N ratio	0–10	25.1±0.01a	25.0±0.01a	25.1±0.0a	25.1±0.0a	25.1±0.0a	25.1±0.0a
		10–20	25.1±0.03a	25.1±0.02a	25.1±0.0a	25.1±0.0a	25.0±0.0a	25.1±0.0a
		20–30	40.1±2.4a	38.6±2.2a	38.7±2.6a	40.1±1.9a	38.9±1.8a	39.9±2.8a

Values are mean± standard error. Different letters in bold following values within lines denotes a statistically significant difference between drained and undrained, mounded and unmounded, fertilised and unfertilised treatment ($P<0.05$).

Soil nitrogen

Total soil N concentration is shown in Table 1. Total soil N concentration varied significantly with soil depth ($P = 0.0001$) and was higher in the 0–10 and 10–20 cm soil depth compared to the 20–30 cm depth. Drainage significantly increased soil N

concentration only in the 0–10 cm soil depth in August 2007 ($P = 0.04$). In June 2008, the total soil N was significantly decreased by drainage in the 0–10 cm depth ($P = 0.04$) but not in the 0–20 and 20–30 cm soil depths. In none of the sampling dates was total N affected by fertilisation or mounding. Soil C:N ratio varied significantly with soil depth ($P = 0.0001$). The C:N ratio was significantly increased by fertilisation only in the 0–10 cm depth in August 2007 ($P=0.01$). In none of the sampling dates was the C:N ratio affected by drainage or mounding.

Soil microbial biomass C

The mean values for soil MBC are shown in Table 2. The undrained had slightly higher MBC than drained treatment in November 2006. In February 2007 soil MBC was marginally increased by drainage ($P = 0.6$). Similarly, soil MBC was slightly higher (but not significant) in the drained than undrained treatment in August 2007. The sampling dates differed significantly ($P = 0.0001$) in soil MBC. Soil MBC was slightly higher (but not significant) in the fertilised than unfertilised treatment during all sampling occasions. The mounded treatment had slightly higher (but not significant) MBC than the unmounded treatment during all the sampling occasions.

Table 2. Effects of drainage, mounding and fertilisation on soil microbial biomass C (mg g^{-1})

Sampling date	Treatment					
	Drained	Undrained	Mounded	Unmounded	Fertilised	Unfertilised
November 2006	1.54±0.28a	2.14±0.13a	1.92±0.25a	1.77±0.22a	1.74±0.22a	1.95±0.25a
February 2007	2.69±0.30a	1.98±0.21a	2.57±0.30a	2.11±0.24a	2.30±0.34a	2.38±0.2a
August 2007	3.50±0.07a	2.85±0.23a	3.27±0.20a	3.08±0.19a	3.06±0.17a	3.29±0.21a

Values are mean±standard error. Different letters following values within lines denotes a statistically significant difference between drained and undrained, mounded and unmounded, fertilised and unfertilised.

Inorganic N

The inorganic N (NH_4^+ and NO_3^-) concentrations are shown in Table 3. In none of the sampling dates was ammonium (NH_4^+) affected by drainage. Ammonium concentration was significantly increased by both fertilisation and mounding in August 2007 (all $P=0.01$). In none of the sampling occasions was nitrate (NO_3^-) affected by drainage mounding or fertilisation.

Table 3. Effects of drainage, mounding and fertilisation on inorganic N (mg g^{-1})

Sampling date	Variable	Treatment					
		Drained	Undrained	Mounded	Unmounded	Fertilised	Unfertilised
August 2007	NH_4^+	0.09±0.02a	0.06±0.02a	0.10±0.02a	0.06±0.02b	0.21±0.02a	0.05±0.01b
	NO_3^-	0.02±0.01a	0.04±0.01a	0.03±0.01a	0.03±0.01a	0.02±0.01a	0.04±0.01a
February 2008	NH_4^+	0.02±0.00a	0.03±0.00a	0.03±0.01a	0.02±0.00a	0.02±0.00a	0.03±0.01a
	NO_3^-	0.02±0.00a	0.02±0.00a	0.02±0.00a	0.02±0.00a	0.02±0.00a	0.02±0.00a

Values are mean±standard error. Different letters following values within lines denotes a statistically significant difference between drained and undrained, mounded and unmounded and fertilised and unfertilised.

Above plant biomass

Standing above plant biomass assessed as dry mass is shown in Table 4. Above-ground plant biomass was significantly increased by drainage ($P = 0.04$) and fertilisation ($P = 0.002$). Above-ground plant biomass was not affected by mounding.

Table 4. Effects of drainage, mounding and fertilisation on standing above plant biomass (t ha^{-1} DM)

		Treatment				
Drained	Undrained	Mounded	Unmounded	Fertilised	Unfertilised	
7.760.35a	6.03±0.30b	6.84±0.36a	6.95±0.47a	7.53±0.37a	6.26±0.37b	

Values are mean±standard error. Different letters following values within lines denotes a statistically significant difference between drained and undrained, mounded and unmounded and fertilised and unfertilised.

Discussion

Soil bulk density and pH

Bulk density increased with soil depth, which is consistent with findings of others (Tamminen and Starr, 1994; Zerva, 2004). Tamminen and Starr (2004) investigated the relationship between bulk density and organic matter content, soil structural properties and depth and found that density increased with soil depth and remained uniform at soil depth greater than 20 cm. Bulk density was not affected by drainage or fertilisation but was significantly increased by mounding in the 0–10 cm depth. The increase is consistent with results of other studies which reported that the bulk density in compacted soils increased after site preparation and timber harvesting (Cullen et al., 1991; Johnson et al., 1991; Merino et al., 1998; McNabb et al., 2001). The increase in bulk density in the mounded treatment was probably enhanced by the combined influence of compaction caused by machinery (Banco-Canqui et al., 2004; Mojeremane, 2009) and decrease in SOC concentration (Li et al., 2007).

Soil organic carbon

Soils are the major reservoir of C in terrestrial ecosystems (Henderson, 1995) and research has shown that soil C is negatively affected by land use changes and soil management practices (Batjees, 1996; Ross et al., 1999; Post and Kwon, 2000; Prentice et al., 2000). Drainage reduced soil organic C in the 0–10 cm depth in this study. This result is in agreement with previous studies conducted in peaty gley soil which demonstrated that drainage and ploughing used in afforestation and replanting decrease soil C in peaty gley soils (Zerva and Mencuccini 2005a; Zerva et al., 2005). It has been demonstrated that drainage increase organic matter decomposition and enhances C losses to the atmosphere as CO_2 fluxes (e.g. Martikainen et al., 1995; Nykänen et al., 1995). The water table depth in this study was lowered by drainage, which increased soil temperature and improved aeration

(Mojeremane, 2009). Changes in these environmental variables have been shown to create aerobic conditions that stimulate the soil microbial activity and enhance organic matter decomposition (Smith *et al.*, 1994; Trettin *et al.*, 1995; Olson *et al.*, 1996; Merino *et al.*, 1998; Zerva *et al.*, 2005; Tate *et al.*, 2006) which caused C losses as CO₂ in the present study (Mojeremane *et al.*, 2012). SOC losses from drained organic soils have been attributed to oxidation and enhanced soil respiration linked to improved aeration and increased temperature (Raich and Schlesinger, 1992; Rey *et al.*, 1992; Euskirchen *et al.*, 2003; Saiz *et al.*, 2006). It is also possible that drainage increased the production of highly decomposable fine roots (Thomas *et al.*, 1996) in the present study, thereby simulating soil microbial activity and enhancing organic matter decomposition rates (Lohila *et al.*, 2003; Kuzykov and Cheng, 2004). The decrease in SOC observed in the drained treatment was lower when compared to studies conducted elsewhere, probably because of differences in climate, soil type, the intensity of drainage and the time since the site was drained (Mojeremane, 2009).

Fertilisation increased SOC in August 2007, similar to effects observed in fertilised forest soils (Berg and Matzner, 1997; Franklin *et al.*, 2003; Foereid *et al.*, 2004; Olsson *et al.*, 2005; Jandl *et al.*, 2007). Increased SOC in fertilised soils have been attributed to the suppression of ligninolytic enzymes of soil microbes and by chemical stabilisation (Arnebrant *et al.*, 1996; Jandl *et al.*, 2007). The effect of fertilisation was not significant in June 2008, probably because N uptake by plants (Vitousek and Matson, 1985; Emmett *et al.*, 1991), loss of N as N₂ and N₂O (Robertson *et al.*, 1987; Sitaula *et al.*, 1995) and NO₃⁻ leaching (Vitousek and Matson, 1985; Smith *et al.*, 1994) exhausted the N pool in the soil.

Soil nitrogen

Soil N in 0–10 cm depth was significantly affected by drainage but not fertilisation or mounding. Drainage decreased N concentration, which is consistent with others who reported that timber harvesting and mechanical site preparation decrease N in forest soils (Smith *et al.*, 1994; Merino *et al.*, 1998). Decreased N in the drained treatment could probably be attributed to increased soil temperature and aeration which created aerobic conditions that favoured the mineralisation of N in the SOM and increased N uptake by plants or losses as N₂ and N₂O emissions or dissolved nitrates (Mojeremane, 2009). Prior studies demonstrated that site preparation practices increased leaching in upland and boreal forest soils (Nieminen, 1998; Mannerkoski *et al.*, 2005; Piirainen *et al.*, 2007) which probably occurred in the present study.

Soil microbial biomass carbon

Soil microbes play a critical role in mediating feedbacks between terrestrial ecosystems and global climate change (Dooley and Treseder, 2012). They regulate the transfer of C from terrestrial ecosystems to the atmosphere through organic matter decomposition in soil (Swift *et al.*, 1979). They also regulate soil nutrients via organic matter mineralisation and solubilisation of soil minerals (Mazzarino *et al.*, 1993; Franzluebbers *et al.*, 1994; Fritze *et al.*, 1994; Blazier *et al.*, 2005), especially in infertile natural and agricultural systems (Yao *et al.*, 2000). The soil microorganism populations and activity may be reduced in infertile waterlogged soils such as peaty gley soil in the present study site. Lowering the water table of saturated soils for forestry and agriculture uses through drainage increases soil temperature, oxygen and nutrient availability (Lieffers and Rothwell, 1987; Lieffers, 1988) and may favour populations and activity of soil microorganisms. However, the activity of soil microorganisms measured as MBC was not affected by drainage, mounding or fertilization in the present study.

The MBC was slightly increased by drainage in February and August 2007. Low soil pH and fertility observed in the present study site may have affected the soil microbial population and activity. The lowest MBC was observed in November and February (winter months), suggesting that the ecosystem activity was low during the winter months and nutrients requirement by soil microorganisms and plants was probably met by background nutrient levels of soils (Insam *et al.*, 1989). In the late summer (August 2007) the soil microbial biomass in all treatment increased with soil temperature (Mojeremane, 2009). It seems, therefore that effects of treatments may have been masked by seasonal effects, which is consistent with others who observed that fluctuations in soil temperatures affected MBC (Lynch and Panting, 1982; Sarathchandra *et al.*, 1988, 1989).

MBC in the fertilised treatment was slightly lower (but not significant) than in the control which is in agreement with others who reported lower MBC in fertilised grasslands than control treatment (Yates *et al.*, 1997; Bardgett and Cook, 1998). This result is comparable to several other studies which failed to detect significant effects of fertilisation on MBC (Castro *et al.*, 1994; Vose *et al.*, 1995; Sarathchandra *et al.*, 2001). In contrast, others reported increased MBC in fertilised forest (Hobbie, 2000; Vestgarden, 2001) and agricultural soils (Lynch and Panting 1982; Hesebe *et al.*, 1985). Soil pH in the present study was low (3.8–4.2) (Mojeremane, 2009) and could have negatively affected MBC (Shah *et al.*, 1990; Nodar *et al.*, 1992). Acidic soils have been shown to favour fungal populations relative to their bacterial counterparts (Nodar *et al.*, 1992).

Inorganic N

Lowering the water table depth increases the mineralisation of N and subsequent nutrient availability to plants in peaty soils (Grootjans *et al.*, 1985; Updegraff *et al.*, 1995; Bridgham *et al.*, 1998). Our results show that inorganic NH₄⁺ and NO₃⁻ were not affected by drainage probably because N mineralisation was compensated by increased losses. Ammonium availability was increased by mounding in August 2007 probably due to increase in net mineralisation of the organic matter on top of mounds. This was not surprising because changes in the microtopography related to mounds and hollows after mounding modifies a number of important environmental variables (Liechty *et al.*, 1997), such as soil temperature and moisture (Nohrstedt, 2000) as well as the thickness and distribution of organic and mineral soil layers (Beatty and Stone, 1986; Schaeztl *et al.*, 1990). Mounding buried the soil organic layers beneath the mineral soil of mounds and increased soil temperature in 0–5 cm depth (Mojeremane, 2009) and aeration. These changes may have favoured microbial activity on the top of mounds and accelerated N mineralisation. Nitrate was not affected by mounding in this study.

Fertilisation increased NH₄⁺ in August 2007 and the increase was not evident at the end of study (June 2008), probably due to increased uptake by plants or losses as oxides of nitrogen in year one of study (Mojeremane, 2009). Nitrate availability was not affected by fertilisation and there is a possibility that NO₃⁻ was leached to deeper layers or lost through drainage water (Baker and Johnson, 1981; Bergstrom and Brink, 1986; Mojeremane, 2009).

Standing above-ground plant biomass

Standing above-ground biomass production measured a year after establishment of the experiment was increased by drainage. The increase probably resulted from increased soil temperature and improvement in aeration of the root zone which favours root growth and nutrient availability. Water-saturated soils like those in the present study site have been shown to negatively affect plant growth by limiting root and shoot growth (Huang et al., 1994; McDonald et al., 2001; Malik et al., 2001, 2002). The increase in plant biomass in the present study could be attributed to increased substrate temperature and oxygen availability after drainage, which increased nutrient availability since organic matter decomposition is enhanced under aerobic conditions (Clymo, 1984). Plant biomass was also increased by fertilisation. The uptake of nitrogen, potassium and phosphorus by plants from the applied fertiliser may have enhanced their growth. Our results are in agreement with other studies conducted in Calluna dominated vegetation on heathland which reported increased plant growth application of N (Aerts et al., 1991; Caporn et al., 1995; Uren et al., 1997; Carroll et al., 1999). Increased above-plant biomass in the fertilised treatment in the present study may suggest that the site is poor in nutrient.

Conclusion

Soil organic carbon and nitrogen in the present study site were decreased by drainage in 0–10 cm soil layer. Lowering the water table by drainage increases oxygenation and soil temperature and thus increases the mineralisation of SOM and N in the organic matter which resulted in the loss of C and N to the atmosphere. Fertilisation increased SOC accumulation in the 0–10 cm layer. Soil microbial biomass carbon and inorganic N were not affected by the imposed treatments due to acidic soil conditions and poor soil fertility.

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