

# Factors controlling decomposition rates of fine root litter in temperate forests and grasslands

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## Abstract

**Background and aims** Fine root decomposition contributes significantly to element cycling in terrestrial ecosystems. However, studies on root decomposition rates and on the factors that potentially influence them are fewer than those on leaf litter decomposition. To study the effects of region and land use intensity on fine root decomposition, we established a large scale study in three German regions with different climate regimes and soil properties.

**Methods** In 150 forest and 150 grassland sites we deployed litterbags (100 µm mesh size) with standardized litter consisting of fine roots from European beech in forests and from a lowland mesophilous hay meadow in grasslands. In the central study region, we compared decomposition rates of this standardized litter with root litter collected on-site to separate the effect of litter quality from environmental factors.

**Results** Standardized herbaceous roots in grassland soils decomposed on average significantly faster ( $24 \pm 6$  % mass loss after 12 months, mean  $\pm$  SD) than beech roots in forest soils ( $12 \pm 4$  %;  $p < 0.001$ ). Fine root decomposition varied among the three study regions. Land use intensity, in particular N addition, decreased fine root decomposition in grasslands. The initial lignin:N ratio explained 15 % of the variance in grasslands and 11 % in forests. Soil moisture, soil temperature, and C:N ratios of soils together explained 34 % of the variance of the fine root mass loss in grasslands, and 24 % in forests.

**Conclusions** Grasslands, which have higher fine root biomass and root turnover compared to forests, also have higher rates of root decomposition. Our results further show that at the regional scale fine root decomposition is influenced by environmental variables such as soil moisture, soil temperature and soil nutrient content. Additional variation is explained by root litter quality.

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## Introduction

Decomposition of plant litter is one of the main processes driving nutrient and carbon (C) cycling in terrestrial ecosystems and constitutes a major source of atmospheric CO<sub>2</sub> (Hobbie 1992). While most of the previous studies on decomposition have focused on aboveground plant litter, recent isotopic analyses and assessments of root and shoot biomarkers indicate that root-derived C is retained more efficiently in soils and microorganisms than are C inputs from above ground plant litter (Kramer et al. 2010; Mendez-Millan et al. 2010). Due to the close proximity, root decomposition products are generally better incorporated into soil aggregates and more easily adsorbed to mineral surfaces than the ones of aboveground litter (Rasse et al. 2005; Sanaullah et al. 2011). Thus, plant litter, consisting of dead roots, is a major source of soil organic matter, the largest terrestrial pool of C (Schmidt et al. 2011).

Fine root decomposition rates do not mirror those of aboveground plant litter in temperate ecosystems (Bird and Torn 2006; Hobbie et al. 2010; Silver and Miya 2001), due to differences in the chemical composition of both litter types (Kögel-Knabner 2002), and because aboveground litter experiences different environmental conditions compared to belowground litter (Hobbie et al., 2010). Yet, only few experimental studies have been designed to identify predictors of fine root decomposition (Chen et al. 2002; Cusack et al. 2009; Fahey et al. 1988; Ostertag and Hobbie 1999), and even fewer extend beyond a single ecosystem (e.g. Long-term Intersite Decomposition Experiment (LIDET), Harmon et al. 2009). Previous studies have shown that root litter quality (mainly lignin, nitrogen (N), C and calcium (Ca) content) and climate factors such as temperature and precipitation are the primary controls of root decomposition. (Chen et al. 2002; Hobbie 2005; Silver and Miya 2001). However, little is known about how the direct and indirect effects of land use and management may influence fine root decomposition.

In grasslands, management practices such as fertilization, mowing and grazing can alter plant community structure, litter inputs and soil properties (Dickinson and Polwart 1982; Hobbie 2005; Semmartin et al. 2008). Some studies found that fertilization increases the decomposition of plant litter, due to higher N availability (Carreiro et al. 2000; Hobbie and Vitousek 2000). In contrast, other studies observed either no significant change or a suppression of decomposition rates as a

response to higher fertilization regimes (Bryant et al. 1998). The contrasting responses to N additions across studies may be partly explained by differences in site-specific atmospheric N deposition and by litter quality (Knorr et al. 2005). However, it is unclear how N fertilization alters root decomposition. Long-term grazing and mowing activities may affect root decomposition directly by altering the plant species composition and soil decomposer communities (Bardgett et al. 1998), or indirectly through changes in soil properties such as bulk density and soil moisture (e.g. through trampling) (Sankaran and Augustine 2004; Taboada and Lavado 1988).

In forests, thinning and harvesting can lead to soil degradation via soil compaction (Berthrong et al. 2009; Liao et al. 2010), which in turn might alter fine root decomposition rates due to changes in the aeration of the soil and the volumetric soil moisture content. Further, these practices as well as tree species selection may lead to variations in soil C and nutrient contents which may result in changes of the decomposer community. Higher temperature and moisture in more heavily thinned forests can further result in faster root decomposition although responses may vary among forests of different dominant tree species (Cortina and Vallejo 1994; Prescott et al. 2000). The goal of this study was to quantify root litter decomposition rates in differently managed temperate grasslands and forests, and to disentangle the effects of litter quality from environmental site effects.

In terrestrial ecosystems, decomposition of plant litter is commonly assessed using the litterbag method, which consists of enclosing plant tissue of known mass and chemical composition in a screened container (Bocock and Gilbert 1957) that is placed in or on top of the soil. When adopting the litterbag approach some artefacts need to be considered, such as microclimate alterations and exclusion of specific fauna size classes due to the selection of small mesh sizes. Nevertheless, the litterbag method is helpful to study decomposition especially if comparing decomposition rates occurring in different ecosystems and regions in a consistent way (Kurz-Besson et al. 2005). Meta-analyses have been used in previous studies to synthesize root decomposition (for example Silver and Miya 2001). The main caveat of the meta-analysis approach is that it compares decomposition rates derived from smaller scale root decomposition studies which adopted different methodologies (i.e. diverse litterbag mesh sizes, enclosure of root litter with diverse quality, and deployment at different soil depth). To study fine root decomposition at a

large spatial scale in a comparative way, we used a standardized methodology for a range of ecosystems with different soil types, land uses and climate.

Our main objective was to quantify root litter decomposition after 12 months in differently managed temperate grasslands and forests. We were further interested in understanding whether environmental site conditions (including the microbial decomposer abundance) have a stronger impact on decomposition than differences in root litter quality between grasslands and forests. We deployed litterbags containing standardized herbaceous fine roots in 150 grassland plots and European beech roots in 150 forest plots in three German study regions managed with different land use intensities. Moreover, we compared the mass loss rates of standardized root litter with the rates for root litter collected on-site for one of the three regions (50 forest and 50 grassland plots). We hypothesized that root decomposition rates of standardized root litter would vary between the three study regions, reflecting differences in climate and soil properties. We also hypothesized that decomposition of on-site collected root litter would be affected by management practices, specifically fertilization, grazing and mowing activities in grasslands, and by different disturbance intensity and tree species selection in forests.

## Material and methods

### Study regions, management data, vegetation survey

Our study was carried out in the Biodiversity Exploratories (Fischer et al. 2010) which comprise a variety of

forests and grasslands under different land use intensities in three regions within Germany. The Schwäbische Alb (ALB) is situated in south-western Germany, the Hainich-Dün (HAI) in central Germany, and the Schorfheide-Chorin (SCH) in north-eastern Germany. Each study region includes 50 forest plots and 50 grassland plots. In SCH 27 grassland plots have been established on organic soils (Histosols and Gleysols) and 23 on mineral soils. The three study regions differ in climate, altitude, and soil characteristics (Table 1).

In grasslands a land use intensity index (LUI) was calculated using landowner surveys. The LUI summarizes the individual land uses by summing up values for fertilization (kg N per hectare per year), mowing (number of cuts per hectare per year), and grazing intensities (livestock units per hectare per year) which have been normalized by the mean of the region (Blüthgen et al. 2012). To evaluate land use and disturbance intensity in the forests we used the LUDI index (Luyssaert et al. 2011). This index combines values of stand density and diameter at breast height for a relatively unmanaged forest and different management schemes, in conjunction with self-thinning values.

In 2009, we recorded vegetation data in 20 m × 20 m forest plots in spring and summer (for details see Boch et al. 2013), and in 4 m × 4 m grassland plots only once in summer (for details see Socher et al. 2013). We identified all vascular plant species and estimated their percentage cover. To assess the diversity of the vascular plant species in forest plots, we combined the spring and summer records to consider early and late emerging plants. For both, grassland and forest sites, we calculated the “Shannon Index” as a measure of plant species

**Table 1** Main geographical and environmental characteristics of the three study regions: Schwäbische Alb, Hainich-Dün, Schorfheide-Chorin

	Schwäbische Alb	Hainich-Dün	Schorfheide-Chorin
Location	SW Germany	Central Germany	NE Germany
Coordinates	N 48° 26' E 9° 23'	N 51° 9' E 10° 28'	N 53° 0' E 13° 46'
Area [km <sup>2</sup> ]	~422	~1,300	~1,300
Soil type forest	Cambisol (eutric)- Leptosol	Luvisol	Cambisol (dystric)
Soil type grassland	Leptosol-Cambisol	Cambisol-Stagnosol- Vertisol	Histosol-Gleysol-Cambisol- Luvisol-Albeluvisol
Altitude a.s.l [m].	460–860	285–550	3–140
Mean annual temperature (MAT) [°C]	6.0–7.0	6.5–8.0	8.0–8.5
Mean annual precipitation [mm]	700–1,000	500–800	500–600

diversity (Shannon 1948) (Eq. 1):

$$H = -\sum_{i=1}^n pi \ln pi \quad (1)$$

where  $H$  corresponds to entropy in this case equivalent to the Shannon Index,  $pi$  is the percentage cover of individuals in the sampling area represented by species  $i$  and is assessed by the quotient of number of individuals of species  $i$  ( $N_i$ ) and the total number of individuals ( $N$ ). Thus, the maximum diversity possible for  $N$  individuals occurs when all species have the same percentage cover when each individual belongs to a different species. We further distinguished annual and perennial species and calculated their number per plot.

### Soil temperature and moisture

Soil temperature and soil moisture were measured continuously in each plot every 10 min with ground surface temperature sensors (Meier NT Type 2021, Zwönitz, Germany) and soil humidity probes (DeltaT ML2X, Cambridge, UK) at a soil depth of 10 cm. In this study, we used the 6 and 12 months averages of soil temperature and volumetric soil moisture. We also calculated the number of days during which the soil was frozen ( $-0\text{ }^{\circ}\text{C}$ ) for each plot. Due to gap periods during the measurements, complete data on soil temperature was only available for 187 and soil moisture for 274 of the 300 plots.

### Root collection, abiotic soil properties and microbial biomass

In each of the 300 plots we collected 14 mineral soil cores with a split tube sampler (diameter of 5 cm) along two 20 m transects in grasslands and 40 m transects in forests in May 2011. Organic layers in forests and aboveground plant parts in grasslands were removed before coring. We then prepared a composite sample from the 14 mineral soil cores by mixing the upper 10 cm of the mineral soil. Roots were removed from the composite sample, cooled to  $4\text{ }^{\circ}\text{C}$  for soil analysis (root biomass, texture, pH and C and N concentrations,) and frozen to  $-20\text{ }^{\circ}\text{C}$  for microbial biomass measurements, and transported to the laboratory. Soil samples were all sieved to  $<2\text{ mm}$ .

Microbial biomass C was estimated by chloroform-fumigation-extraction (Vance et al. 1987). In brief, 10 g soil (fresh weight) of a homogeneous subsample of each plot was fumigated under vacuum with ethanol-free

chloroform in a desiccator for 24 h. After removing the chloroform, samples were extracted by adding 40 ml of a 0.5 M  $\text{K}_2\text{SO}_4$  solution (1:4 w/v soil/extractant ratio), shaken for 30 min at  $250\text{ rev min}^{-1}$  on a horizontal shaker and centrifuged for 30 min at 4,422 g. A second subsample of 10 g was treated similarly but without fumigation for the estimation of 0.5 M  $\text{K}_2\text{SO}_4$  solution extractable organic C. Organic C in the supernatants was measured with a DOC/TN-analyser (Multi N/C 2100S, Analytik Jena, Jena, Germany). Extractable organic C content of the non-fumigated samples was subtracted from C content of the fumigated samples and resulted in extractable microbial biomass. For estimation of total microbial biomass a  $k_{\text{ec}}$ -factor of 0.45 was used (Joergensen 1996).

Soil samples were air dried. The dry biomass of the washed fine roots was weighed after oven-drying the root samples at  $40\text{ }^{\circ}\text{C}$ . To evaluate the soil texture we determined the percentage of sand (2–0.063 mm), silt (0.063–0.002 mm) and clay ( $<0.002\text{ mm}$ ) in the soil samples. We separated soil particles and size classes by sieving and sedimentation procedures (DIN-ISO 11277). We determined the pH values of our composite soil samples in duplicate using a 0.01 M  $\text{CaCl}_2$  solution. The soil solution ratio was 1:2.5.

Soil subsamples were ground in a ball mill (RETSCH MM200, Retsch, Haan, Germany). Total C and N concentrations were determined by dry combustion in an elemental analyser (VarioMax, Hanau, Germany). To evaluate the concentration of organic C in each soil sample we determined the amount of inorganic C by removing all organic carbon at a temperature of  $450\text{ }^{\circ}\text{C}$  for 16 h, and subtracted this value from the total C concentration.

### Chemical composition of fine roots

To identify the initial quality, subsamples of the root litter were ground in a ball mill (RETSCH MM200, Retsch, Haan, Germany). Total C and N concentrations were determined using an elemental analyzer (Vario EL, Elementar, Hanau, Germany). Concentrations of Ca, magnesium (Mg), aluminium (Al) and phosphorus (P) were measured using inductively coupled plasma—optical emission spectrometry (ICP-OES, Optima 3300 DV, Perkin Elmer, Norwalk, USA), after 50 mg of root material were diluted in 3 ml of  $\text{HNO}_3$  65 %, (Merck, Darmstadt, Germany) and microwave digested at high pressure (Multiwave, Anton Paar, Graz, Austria) (Raessler et al. 2005). Lignin and holocellulose content (ideally composed by cellulose and hemicellulose) were

estimated from thermogravimetric analysis in Argon atmosphere (TGA/SDTA851<sup>o</sup> Mettler Toledo, GmbH, Giessen, Germany) (Yang et al. 2005). Temperature programming consisted of an initial isothermal phase at a temperature of 100 °C for 5 min, a dynamic phase with a heating rate of 40 °C min<sup>-1</sup> from 100 °C to 1,000 °C, and a second isothermal phase for 10 min at a temperature of 1,000 °C. To prevent heat and mass transfer limitations, small samples (5 to 10 mg) were used. Weight loss and heating rate were continuously recorded. We calculated the amounts of lignin and holocellulose by dividing the biomass pyrolysis in the following ranges: < 220 °C, moisture evolution; 220–400 °C, holocellulose decomposition; > 400 °C, lignin decomposition as indicated by Yang et al. (2005). As reference materials we used pure lignin (alkali, low sulfonate content) and cellulose powder.

#### Root litter decomposition experiments

We conducted two litterbag decomposition experiments using two different litter types: 1) we studied decomposition of standardized root litter at all 300 plots in forests and grasslands of the three regions, and 2) we used on-site collected material only in plots of the HAI region. As standard material for the forest sites we used fine roots collected from 2 year old European beech saplings (*Fagus sylvatica* L.) grown in sand. As standard material for the grassland sites, we used fine roots which we collected from a 16 m<sup>2</sup> area in a lowland mesophilous hay meadow belonging to the alliance Arrhenatherion elatoris W. Koch (Isserstedt, Thuringia, Germany, 50° 57' 30.3" N 11° 31' 20.5" E).

We removed the mineral-soil particles attached to the roots, by carefully washing the roots with distilled water in a 63 µm sieve. In addition, we suspended the root material in a tray containing distilled water to remove the more adherent mineral particles. We separated the fine roots (<2 mm diameter) from the coarse roots and dried all samples at 40 °C to constant weight in a force-air oven. We put 0.5 g of dry fine root litter (mass selected according to average local root biomass distributions in top-soils) into a 10 cm × 10 cm litterbag made of a 100 µm polyester mesh screening to allow micro-faunal decomposition (Schwegmann Filtrations-Technik GmbH, Grafschaft-Gelsdorf, Germany). The variation of the initial weight was (0.508±0.006 g). We individually labelled each litterbag with a stainless

steel label which was placed inside the enclosure and measured the total mass of each litterbag.

For each of the two experimental set-ups we tied together three litterbags, representing replicates containing the same material type (standard litter and on-site collected litter) for every collection time. In October 2011, we placed them approximately 3 m apart from meteorological stations. The litterbags were distributed vertically into a 10 cm deep slit in the mineral soil. After 6 months (in April 2012) and after 12 months (October 2012) respectively we collected three litterbags at each site containing standardized material. In addition, we sampled three litterbags containing on-site collected material in all of the 100 HAI plots in October 2012. The collected litterbags were transported to the laboratory where we gently removed the ingrown material and cleaned the fine root-litter from adherent soil particles. After drying at 40 °C we calculated the fine root decomposition rates (mass loss in %) for each plot as the mean mass loss of the three litterbags collected at each collection date (standard deviations for the three replicates ranged between 0.1 and 3.5 % after 6 months of decomposition and between 0.1 and 8.2 % after 12 months of decomposition). We further used the negative exponential single-pool decomposition model (Olson 1963) to estimate the root litter decomposition rates (*k*), though we recognize that this represents only initial decomposition rates (12 months) and that these slow over time in most decomposition experiments (e.g. Sun et al. 2012).

#### Statistics

We conducted statistical analyses with R, version 3.0.2 (R Development Core Team 2013). Throughout the manuscript we present data as means±standard error. Two way analysis of variance (ANOVA) accompanied by Holm's test was used to examine statistical differences between the mass loss of standardized fine root litter (after 6 and 12 months), soil moisture, soil temperature, soil properties, land use and root biomass, in grassland and forest plots among the regions (in the SCH study region we additionally distinguished between organic soils (Histosols and Gleysols) and mineral soils). Significant differences between fine root litter chemistry

of grasslands and forests, and between standardized and on-site collected root material were tested with Student's *t*-test.

To detect the environmental predictors of fine root decomposition in grasslands and forests, we applied general linear models (GLM). In grasslands we conducted three separate GLMs, one grouping all three study regions, one for the mineral soils in the three study regions and one for SCH to evaluate the difference between organic and mineral soils. To check whether complex interactions between explanatory variables were present and to select which variables should be included in the GLM and in which order, before conducting each GLM we fitted a tree model (Crawley 2007). In the tree model analysis we included the following variables: soil properties (pH, organic C content, C:N ratio and texture, microbial biomass), climate (soil temperature and moisture), land use (LUI and LUDI), plant diversity (Shannon index) and number of perennial species. We calculated the variance inflation factor (VIF) between the variables selected from the fitting of the tree models to prevent multicollinearity. We manually simplified all GLMs to the minimum adequate model by using stepwise selection of variables until all terms contained in the model were significant (Crawley 2007). We then checked the model assumptions using the diagnostic plot function in R (Crawley 2007).

To assess whether environmental variables have a stronger impact on decomposition than root litter chemistry differences within grassland and forest sites, we used multiple regression analysis. As for the GLMs, we fitted a tree model (one for grasslands and one for forests) before conducting multiple regressions. Root litter chemistry (content of C, N, Ca, Mg, Al, P, holocellulose, C:N and lignin:N ratio) was included in the tree models in addition to the environmental variables. Quadratic terms of each variable were included in the model to check for non-linear responses.

We used partial linear regression to determine the amounts of variations explained by land use and the other environmental variables on decomposition of standardized fine root litter in grasslands (Legendre 2008). We used this method also to distinguish between the variation explained by environmental and root-litter chemistry variables on fine root decomposition in grasslands and

forests of the HAI region. The variables we adopted in the partial linear regression were the ones which were selected from the minimum adequate GLM and the multiple regressions.

## Results

### Soil characteristics and root biomass

The soils of the ALB study region showed higher volumetric soil moisture content than the ones in HAI and SCH for both grasslands and forests during 6 and 12 months of decomposition. Soil temperatures were higher in SCH than in the ALB and HAI regions (Table 2). The soils in the HAI study region were on average frozen for more days during the 12 months period (7 days) than ALB (5 days) and SCH (4 days).

In grassland soils, we found on average higher clay contents, pH values, C:N ratios and microbial biomass than in forest soils (Table 3). Soil properties also differed between study regions (Table 3). Organic C and microbial C were positively related to clay contents ( $r=0.68$ ,  $r=0.70$ ) in mineral soils. Microbial biomass was also positively correlated with the soil organic C content ( $r=0.65$ ). The organic soils of the SCH contained higher soil organic C and microbial biomass compared to the mineral soils of the study region. In grasslands, fine root biomass was highest in the mineral soils of the SCH followed by HAI, the organic soils in SCH, and then ALB, whereas in the forests root biomass was greater in the HAI study region than in SCH and ALB.

### Fine root initial chemical composition

C concentrations, estimated lignin content and lignin:N ratios were lower, and P and holocellulose content were higher in root samples collected in grasslands than those collected in the forest sites (Table 4). Variability in the chemical quality of roots was greater for root material collected across 50 plots (both for forest and grassland) compared to standard material derived from a single site (grassland) or species (forest). The standardized grassland root litter differed significantly from fine root litter collected in the HAI grassland plots, with

**Table 2** Soil moisture and temperature over 6 and 12 months, in all study regions (mean  $\pm$  SD). Soil temperature is expressed in  $^{\circ}\text{C}$  and soil moisture is expressed in percentage of volumetric water content (%VWC). Significant differences between study regionsare indicated by lowercase letters, and between land use types by capital letters according to Holm's test ( $p < 0.05$ ) ( $n = 274$  for soil moisture,  $n = 187$  for soil temperature)

Land use	Study region	Soil moisture		Soil temperature	
		6 months %	12 months %	6 months $^{\circ}\text{C}$	12 months $^{\circ}\text{C}$
Forest	All plots	26 $\pm$ 10 <sup>A</sup>	26 $\pm$ 10 <sup>A</sup>	7 $\pm$ 5 <sup>A</sup>	9 $\pm$ 4 <sup>A</sup>
	Schwäbische Alb	34 $\pm$ 7 <sup>a</sup>	34 $\pm$ 7 <sup>a</sup>	5 $\pm$ 3 <sup>a</sup>	7 $\pm$ 2 <sup>a</sup>
	Hainich-Dün	29 $\pm$ 6 <sup>b</sup>	29 $\pm$ 6 <sup>b</sup>	5 $\pm$ 1 <sup>a</sup>	9 $\pm$ 2 <sup>ab</sup>
	Schorfheide-Chorin	15 $\pm$ 3 <sup>c</sup>	15 $\pm$ 4 <sup>c</sup>	10 $\pm$ 5 <sup>b</sup>	10 $\pm$ 4 <sup>b</sup>
Grassland	All plots	37 $\pm$ 9 <sup>B</sup>	34 $\pm$ 9 <sup>B</sup>	7 $\pm$ 4 <sup>A</sup>	11 $\pm$ 4 <sup>B</sup>
	Schwäbische Alb	41 $\pm$ 4 <sup>d</sup>	40 $\pm$ 4 <sup>d</sup>	5 $\pm$ 2 <sup>a</sup>	8 $\pm$ 2 <sup>a</sup>
	Hainich-Dün	33 $\pm$ 6 <sup>a</sup>	29 $\pm$ 4 <sup>b</sup>	6 $\pm$ 2 <sup>a</sup>	11 $\pm$ 3 <sup>b</sup>
	Schorfheide-Chorin	34 $\pm$ 11 <sup>a</sup>	29 $\pm$ 10 <sup>b</sup>	12 $\pm$ 6 <sup>c</sup>	14 $\pm$ 4 <sup>c</sup>

lower N ( $p < 0.05$ ) and holocellulose content ( $p < 0.001$ ). The higher N content and variability for the roots collected on-site compared to the standardized material in grasslands probably results from the wide range of fertilization regimes. The Ca and Al concentrations were lower in the standardized compared to on-site collected root material (Table 4). The standardized root material used in forests differed from the average of the fine root litter collected on the HAI forest sites, with higher C ( $p < 0.001$ ) and lignin ( $p < 0.05$ ) content, possibly due to variable contributions of understory roots in

on-site collected samples. In addition, standardized beech roots had lower Mg, Ca and Al content than the root material collected on site in the HAI forest plots ( $p < 0.05$ ).

Fine root mass loss in the standard litter across three regions

Herbaceous standardized fine root litter deployed in grasslands lost more mass ( $p < 0.01$ ) after 12 months (24 $\pm$ 6 % mass loss) compared to the mass loss of tree standardized fine root litter in forests (12 $\pm$

**Table 3** Soil properties and dry root biomass among different land uses and study regions (mean  $\pm$  SD). Significant differences between study regions are indicated by lowercase letters and between forests and grasslands by capital letters according to Holm's test ( $p < 0.05$ )

Land use	Study region	n	Sand g kg <sup>-1</sup>	Clay g kg <sup>-1</sup>	pH	Total N g kg <sup>-1</sup>	Organic C g kg <sup>-1</sup>	C:N	Microbial biomass $\mu\text{g g}^{-1}$	Root biomass g m <sup>-2</sup>
Forest	All plots	150	329 $\pm$ 387 <sup>A</sup>	281 $\pm$ 203 <sup>A</sup>	4 $\pm$ 1 <sup>A</sup>	3 $\pm$ 2 <sup>A</sup>	40 $\pm$ 20 <sup>A</sup>	15 $\pm$ 3 <sup>A</sup>	436 $\pm$ 308 <sup>A</sup>	98 $\pm$ 90 <sup>A</sup>
	Schwäbische Alb	50	60 $\pm$ 46 <sup>a</sup>	496 $\pm$ 105 <sup>a</sup>	5 $\pm$ 1 <sup>a</sup>	5 $\pm$ 1 <sup>a</sup>	53 $\pm$ 14 <sup>a</sup>	13 $\pm$ 1 <sup>a</sup>	737 $\pm$ 220 <sup>a</sup>	67 $\pm$ 56 <sup>a</sup>
	Hainich-Dün	50	58 $\pm$ 18 <sup>a</sup>	301 $\pm$ 99 <sup>b</sup>	5 $\pm$ 1 <sup>b</sup>	3 $\pm$ 1 <sup>b</sup>	36 $\pm$ 10 <sup>ab</sup>	13 $\pm$ 1 <sup>a</sup>	449 $\pm$ 214 <sup>b</sup>	155 $\pm$ 120 <sup>b</sup>
	Schorfheide-Chorin	50	871 $\pm$ 61 <sup>b</sup>	45 $\pm$ 19 <sup>c</sup>	3 $\pm$ 0.1 <sup>c</sup>	1 $\pm$ 0.2 <sup>b</sup>	21 $\pm$ 5 <sup>b</sup>	19 $\pm$ 3 <sup>b</sup>	123 $\pm$ 38 <sup>c</sup>	73 $\pm$ 45 <sup>a</sup>
Grassland	All plots	150	190 $\pm$ 228 <sup>B</sup>	378 $\pm$ 192 <sup>B</sup>	7 $\pm$ 1 <sup>B</sup>	7 $\pm$ 5 <sup>B</sup>	72 $\pm$ 64 <sup>B</sup>	10 $\pm$ 1 <sup>B</sup>	656 $\pm$ 291 <sup>B</sup>	175 $\pm$ 149 <sup>B</sup>
	Schwäbische Alb	50	57 $\pm$ 45 <sup>a</sup>	536 $\pm$ 134 <sup>d</sup>	6 $\pm$ 1 <sup>d</sup>	6 $\pm$ 1 <sup>a</sup>	65 $\pm$ 14 <sup>a</sup>	10 $\pm$ 1 <sup>c</sup>	748 $\pm$ 210 <sup>a</sup>	35 $\pm$ 30 <sup>d</sup>
	Hainich-Dün	50	58 $\pm$ 23 <sup>a</sup>	423 $\pm$ 130 <sup>e</sup>	7 $\pm$ 1 <sup>e</sup>	4 $\pm$ 1 <sup>ab</sup>	46 $\pm$ 12 <sup>a</sup>	10 $\pm$ 1 <sup>c</sup>	594 $\pm$ 169 <sup>d</sup>	215 $\pm$ 87 <sup>e</sup>
	Schorfheide-Chorin mineral soils	23	643 $\pm$ 101 <sup>c</sup>	137 $\pm$ 93 <sup>f</sup>	6 $\pm$ 1 <sup>d</sup>	2 $\pm$ 1 <sup>b</sup>	21 $\pm$ 4 <sup>b</sup>	10 $\pm$ 1 <sup>c</sup>	236 $\pm$ 62 <sup>e</sup>	246 $\pm$ 122 <sup>f</sup>
	organic soils	27	324 $\pm$ 179 <sup>d</sup>	216 $\pm$ 119 <sup>e</sup>	7 $\pm$ 1 <sup>e</sup>	15 $\pm$ 2 <sup>d</sup>	170 $\pm$ 102 <sup>c</sup>	11 $\pm$ 2 <sup>c</sup>	906 $\pm$ 338 <sup>f</sup>	180 $\pm$ 127 <sup>e</sup>

**Table 4** Chemical quality of the standardized fine root litter and the fine root litter collected on site in the Hainich-Dün and used in this study (mean  $\pm$  SD). For the standardized material 20 replicates of fine root material were analysed. Significant differencesbetween fine roots collected on-site and standardized material for grasslands and forests are indicated by lowercase letters according to Holm's test ( $p < 0.05$ )

Fine root material	Land use	n	C	N	P	Ca	Mg	Al	Holocellulose	Lignin	C:N	Lignin:N
			mg/g	mg/g	mg/g	mg/g	mg/g	mg/g	mg/g	mg/g	mg/g	mg/g
Standardized	Forests	20	470 $\pm$ 13 <sup>a</sup>	11 $\pm$ 2 <sup>a</sup>	1 <sup>a</sup>	5 <sup>a</sup>	1 <sup>a</sup>	6 $\pm$ 0 <sup>a</sup>	418 $\pm$ 6 <sup>a</sup>	433 $\pm$ 7 <sup>a</sup>	49 $\pm$ 6 <sup>a</sup>	40 $\pm$ 1 <sup>a</sup>
	Grasslands	20	433 $\pm$ 17 <sup>b</sup>	8 $\pm$ 1 <sup>b</sup>	2 <sup>b</sup>	6 $\pm$ 1 <sup>a</sup>	3 <sup>b</sup>	5 <sup>a</sup>	585 $\pm$ 7 <sup>b</sup>	251 $\pm$ 5 <sup>b</sup>	53 $\pm$ 3 <sup>b</sup>	31 $\pm$ 1 <sup>b</sup>
Collected on-site	Forest	50	448 $\pm$ 32 <sup>c</sup>	11 $\pm$ 2 <sup>a</sup>	1 <sup>a</sup>	10 $\pm$ 5 <sup>b</sup>	2 $\pm$ 2 <sup>b</sup>	7 $\pm$ 3 <sup>b</sup>	424 $\pm$ 40 <sup>a</sup>	361 $\pm$ 35 <sup>c</sup>	41 $\pm$ 7 <sup>c</sup>	34 $\pm$ 6 <sup>c</sup>
	Grasslands	50	400 $\pm$ 24 <sup>d</sup>	11 $\pm$ 3 <sup>a</sup>	2 $\pm$ 1 <sup>b</sup>	9 $\pm$ 4 <sup>b</sup>	2 $\pm$ 3 <sup>b</sup>	8 $\pm$ 3 <sup>b</sup>	510 $\pm$ 32 <sup>b</sup>	259 $\pm$ 13 <sup>b</sup>	38 $\pm$ 9 <sup>d</sup>	24 $\pm$ 5 <sup>b</sup>

4 %). The mass loss of fine root litter in both land uses nearly doubled from the first collection (6 months, over winter) to the second (a full year, including summer; Table 5). The mass loss of fine root litter was significantly faster in the SCH study region followed by ALB and HAI for both collection times (Table 5). Mass loss of standardized fine root material after 6 and 12 months of decomposition was correlated in grasslands ( $r=0.51$ ) and forests ( $r=0.64$ ) (Fig. 1). These not so high correlations may be related to the comparison of different root litterbags collected at different times in the same plots. We based our subsequent analyses on the mass loss of fine roots after 12 months decomposition.

#### Predictors of fine root mass loss (standardized litter)

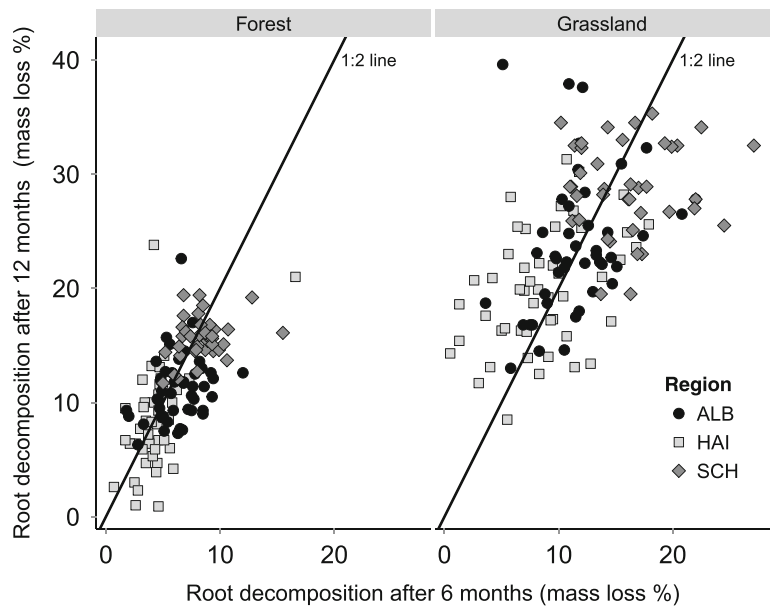
Differences between study regions influenced fine root decomposition in grasslands and forests (Table 6b, 6d). Within all grasslands, the mass loss of the standardized fine root material was further influenced by the total organic C content present in the soil, the soil moisture and the LUI (adjusted  $r^2=0.39$ ) (Table 6a). Partial linear regression analysis showed that the LUI explained 4 % of the variation while organic C, study region, and soil moisture together explained 24 % of the variation. The effect of organic C and soil moisture was not observed in the model including only the mineral soils of the three study regions (Table 6b). In fact,

**Table 5** Mass loss of standard fine root litter and of the fine root litter collected on-site after 6 and 12 months of decomposition (mean  $\pm$  SD). Significant differences between study regions areindicated by lowercase letters, and between forests and grasslands by capital letters according to Holm's test ( $p < 0.05$ )

Fine root litter	Land use	Study region	n	Mass loss after 6 months (%)	Mass loss after 12 months (%)	k-values (year <sup>-1</sup> )
Standardized	Forest	All plots	150	6 $\pm$ 3 <sup>A</sup>	12 $\pm$ 4 <sup>A</sup>	0.12 $\pm$ 0.5 <sup>A</sup>
		Schwäbische Alb	50	6 $\pm$ 2 <sup>a</sup>	11 $\pm$ 3 <sup>a</sup>	0.12 $\pm$ 0.03 <sup>a</sup>
		Hainich-Dün	50	4 $\pm$ 2 <sup>b</sup>	8 $\pm$ 5 <sup>b</sup>	0.09 $\pm$ 0.05 <sup>b</sup>
		Schorfheide-Chorin	50	8 $\pm$ 2 <sup>c</sup>	16 $\pm$ 2 <sup>c</sup>	0.17 $\pm$ 0.02 <sup>c</sup>
	Grassland	All plots	150	12 $\pm$ 5 <sup>B</sup>	24 $\pm$ 6 <sup>B</sup>	0.27 $\pm$ 0.8 <sup>B</sup>
		Schwäbische Alb	50	12 $\pm$ 4 <sup>d</sup>	23 $\pm$ 7 <sup>d</sup>	0.27 $\pm$ 0.07 <sup>d</sup>
		Hainich-Dün	50	9 $\pm$ 4 <sup>c</sup>	20 $\pm$ 5 <sup>c</sup>	0.22 $\pm$ 0.06 <sup>c</sup>
		Schorfheide-Chorin mineral soils organic soils	23 27	15 $\pm$ 3 <sup>c</sup> 17 $\pm$ 4 <sup>e</sup>	26 $\pm$ 3 <sup>f</sup> 31 $\pm$ 3 <sup>g</sup>	0.32 $\pm$ 0.04 <sup>f</sup> 0.37 $\pm$ 0.04 <sup>g</sup>
Collected on site	Forest	Hainich-Dün	50	-	8 $\pm$ 4 <sup>Cb</sup>	0.09 $\pm$ 0.05 <sup>Cb</sup>
	Grassland	Hainich-Dün	50	-	19 $\pm$ 7 <sup>De</sup>	0.21 $\pm$ 0.06 <sup>De</sup>



**Fig. 1** Comparison between the mass loss of standardized fine root litter after 6 and 12 months of decomposition across all study regions, for both grasslands and forests



when excluding the organic soils present in the SCH, fine root mass loss was explained by differences in study regions and LUI. The separation between organic and mineral soils in the SCH explained most of the variance of fine root mass loss in this study region followed by the LUI index (Table 6c). We checked the effect of fertilization, grazing and mowing intensities on fine root mass loss within the grasslands of all three study regions. The standardized fine roots decomposed faster in unfertilized plots than fertilized plots both after six months (unfertilized grasslands:  $12 \pm 4$  %; fertilized grasslands:  $10 \pm 4$  %,  $p < 0.01$ ) and after 12 months (unfertilized grasslands:  $25 \pm 6$  %; fertilized grasslands:  $22 \pm 6$  %,  $p < 0.01$ ). Fine root mass loss did not differ significantly between grazed and not grazed plots and mowed and not mowed plots. In forests we did not observe an effect of land use on fine root decomposition.

Fine root mass loss for on-site collected litter in the HAI region

Despite differences in the initial fine root litter quality, the average mass loss of fine roots during the first 12 months of decomposition was similar between on-site collected root litter and the

standardized root material in the HAI study region both for grasslands ( $\sim 20$  % mass loss) and forests ( $\sim 8$  % mass loss) (Table 5). Overall, the correlation between mass lost by on-site collected root litter versus standardized root litter was poor ( $r = 0.44$  in grasslands and  $r = 0.10$  in forests) (Fig. 2).

Predictors of fine root mass loss (on-site collected litter)

According to our analysis fine root decomposition in grasslands was primarily positively related to the lignin:N ratio of roots and to soil C:N ratios (Table 7a). We further observed a relation between fine root decomposition and soil moisture and soil temperature. The best linear model for predicting fine root mass loss in grasslands was highly significant ( $p < 0.001$ ) with an adjusted  $r^2$  value of 0.55. We observed that in the grasslands the lignin:N ratio of fine roots was negatively correlated to the LUI index ( $r = -0.64$ ) (Fig. 3). In grasslands the lignin:N ratio explained 15 % of the variation of fine root decomposition; while the sum of the environmental variables explained 34 % of the variation (Fig. 4a).

In forests, fine root decomposition was negatively related to the lignin:N ratio of roots, followed by soil moisture and temperature (Table 7b). The best linear

**Table 6** Result of the best predictive model on decomposition rates (mass loss %) of standardized fine root litter a) in all grasslands (mineral and organic soils, 150 plots) b) in the mineral soil of the grasslands c) in the grasslands of the Schorfheide-Chorin (mineral and organic soils, 50 plots) d) forests (150 plots), according to ANCOVA analysis. \*\*\* $p < 0.001$ , \*\* $p < 0.01$ , \* $p < 0.05$

Coefficients	Df	Mean Sq	F value
a) Mass loss (standardized material, grasslands)			
Organic C	1	119.6	4.9*
Study region	2	742.7	30.4***
Soil Moisture	1	140.3	5.7*
LUI	1	191.1	7.8**
Residuals	109	24.5	
b) Mineral soils Schwäbische Alb, Hainich-Dün, Schorfheide-Chorin. Mass loss (standardized material, grasslands)			
Study region	2	330.42	12.30**
LUI	1	176.9	0.01*
Residuals	92	30.2	
c) Schorfheide-Chorin. Mass loss (standardized material, grasslands)			
Mineral vs Organic soil	1	236.7	26.27***
LUI	1	50.2	5.57**
Residuals	38	9.1	
d) Mass loss (standardized material, forests)			
Study region	2	563.5	53.35***
Residuals	132	10.6	

**Table 7** Result of the best predictive model on decomposition rates (mass loss %) of fine roots collected on-site in the a) Hainich-Dün grasslands and b) Hainich-Dün forests, according to multiple regression analysis. \*\*\* $p < 0.001$ , \*\* $p < 0.01$ , \* $p < 0.05$

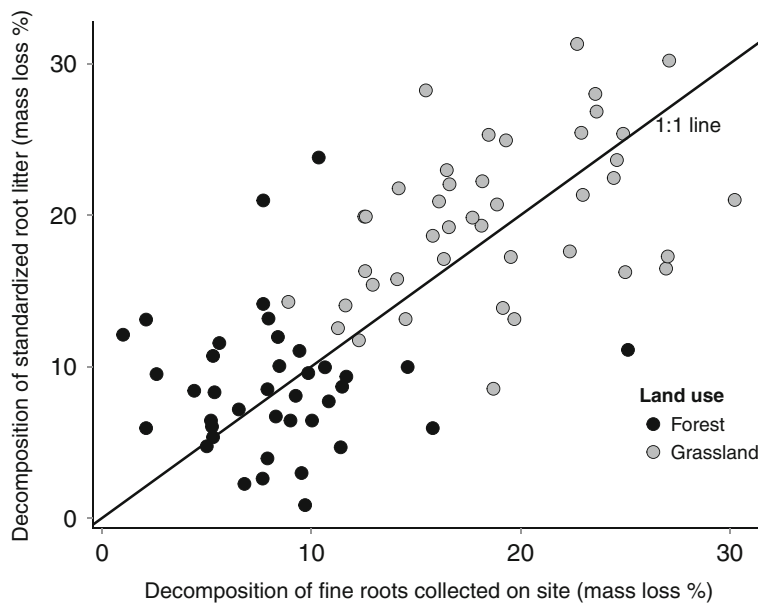
Coefficients	Estimate	Std. Error	t value
a) Mass loss (on-site collected material, grasslands)			
Intercept	-233.80	95.83	-2.44*
Lignin/N ratio	49.40	15.90	3.11**
C/N ratio soil	25.26	9.83	2.57*
Soil Moisture	8.60	3.08	2.79**
(Soil Temperature) <sup>2</sup>	0.18	0.7	2.58*
C:N ratio soil : Soil Moisture	-0.97	0.32	-3.05**
Soil moisture : Soil temperature	0.14	0.05	2.68*
b) Mass loss (on-site collected material, forests)			
Intercept	84.88	17.87	4.75***
Lignin/N ratio	-0.18	0.06	-2.89**
Soil Moisture	-1.44	0.37	-3.94***
Soil Temperature	-10.04	2.82	-3.57**
(Soil Temperature) <sup>2</sup>	0.27	0.09	2.72*
Soil moisture : Soil temperature	0.16	0.04	3.9***

Residual standard error: 3.262 on 25° of freedom, Residual standard error: 2.431 on 35° of freedom

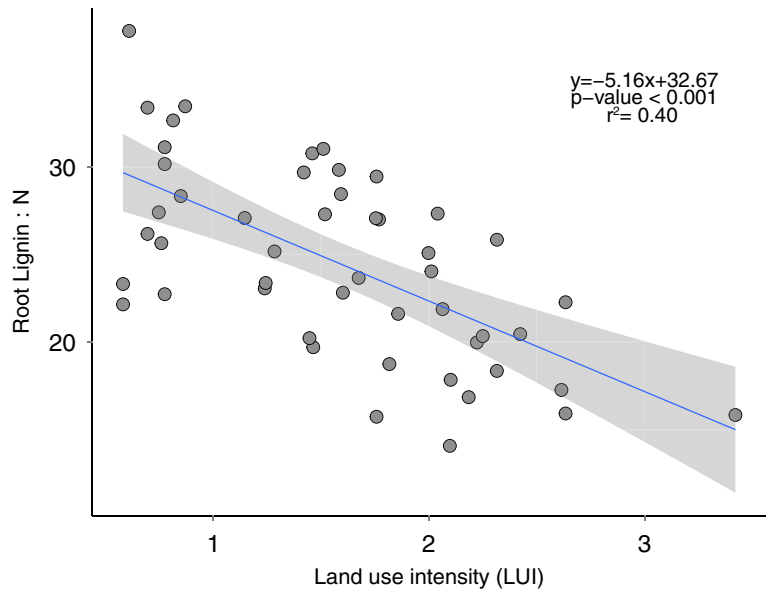
model for predicting fine root decomposition in grasslands was highly significant ( $p < 0.001$ ) with an adjusted

$r^2$  value of 0.42. We further observed a decline in the lignin:N ratio with the number of tree species present on

**Fig. 2** Relationship between the mass loss of fine roots collected on-site and fine roots used as standardized material in the Hainich-Dün study region after 12 months of decomposition



**Fig. 3** Regression between the lignin:N ratio and the land use intensity index (LUI) in the Hainich-Dün study region

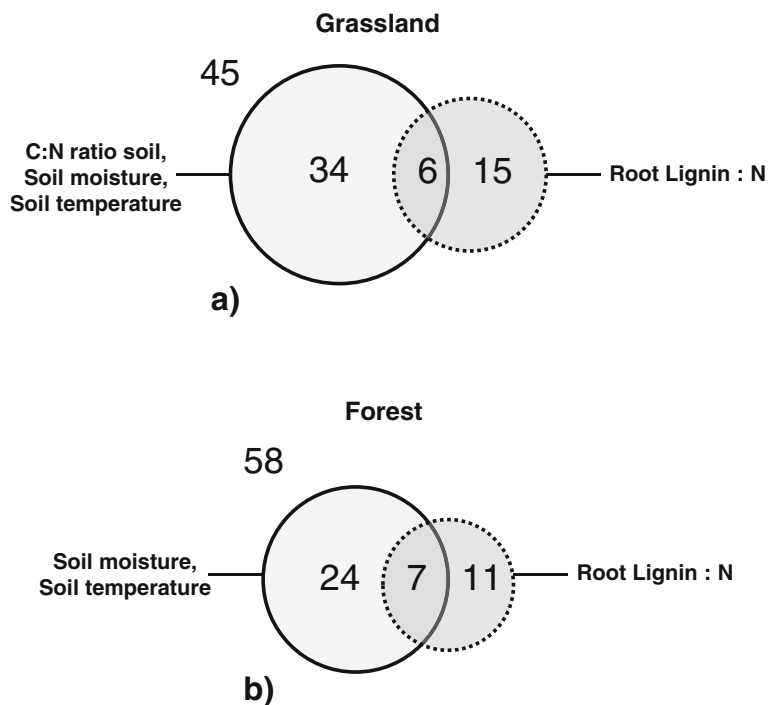


site ( $r=-0.31$ ,  $p<0.05$ ). Root lignin:N ratios explained 11 % of the decomposition of fine roots, while the sum of the environmental variables explained 24 % of the variance (Fig. 4b).

**Discussion**

Our root decomposition study shows, in accordance with other experiments using the litterbag method, that

**Fig. 4** Venn diagrams representing the partition of the variation of decomposition in response to environmental variables (continuous line) and litter chemistry (dashed line) for grassland (a) and forest ecosystems (b) in the Hainich-Dün region. The total variation for each diagram is 100. The intersection represents the amounts of variation explained by two different linear models we used for the analysis (the first model only including root litter quality and the second only including the environmental variables). The unexplained variation (residual variation) is represented by the number outside the circles. Partition of variation is expressed in %. Adapted from Legendre (2008)



substantial decomposition can occur in litterbags in grasslands and forests, even in the absence of mechanical breakdown by macrofauna. Overall we observed relatively low rates of mass loss during the first year of decomposition. One factor to explain this is the relatively cold and dry conditions in temperate ecosystems. Parton et al. (2007) observed that leaf and root decomposition were slowest in cold dry regions such as tundra and boreal forests and fastest in tropical regions. We cannot exclude that absolute values might be affected by the experimental set-up (i.e. mesh size), but as all litterbags were the same, our discussion will focus on relative differences between study sites and root litter quality. In any case, with this work in three large regions in Germany we mostly provide an indication of the initial decomposition rates of fine roots and the factors which control this process in central European temperate grasslands and forests.

In comparison to previous studies on aboveground litter decomposition in temperate ecosystems (Butenschon et al. 2011; Heim and Frey 2004), fine roots seem to lose mass at lower rates than leaves. Other studies also found that roots decompose more slowly than leaf litter of different species or ecosystem types (Vivanco and Austin 2006). Lower decomposition rates of root litter in comparison to aboveground litter may be a result of different nutritional requirements of decomposers above versus belowground. Environmental differences above and belowground could also overcome litter chemistry effects on decomposition (Hobbie et al. 2010).

#### Fine root decomposition rates in grasslands and forests

Overall, herbaceous roots (deployed in grasslands) decomposed twice as fast than roots from forests (deployed in forests) in all study regions (Table 5). This was true for both standardized and on-site collected litter. No common material was incubated in both grasslands and forests so this result can be related to differences in environmental conditions or in litter quality. We noted, for instance, higher soil moisture in grasslands than in forests (Table 3). Direct influences of moisture contents on fine root respiration, have been previously observed by Chen et al. (2000) for unsaturated soils. In grasslands, we further observed higher microbial biomass contents and higher pH values.

We also observed differences in fine root litter quality between herbaceous plants and trees (Table 4). For

example greater lignin content in tree roots can control mass loss rates during the first stages of decomposition by its high resistance to enzymatic attack, as well as through physical interference with the decomposition of other cell wall fractions (Alexander 1977). Results of the global meta-analysis on fine root decomposition performed by Silver and Miya (2001) have also shown that decomposition rates of roots belonging to tree species were lower than those of graminoid roots due to lower Ca content and higher lignin:N ratios in tree roots.

Overall, we found up to 2 times higher stocks of fine roots and roughly 2 times faster decomposition rates in grasslands than in forests. These observations together with the results of recent studies in the same regions which have shown younger C in fine roots (Solly et al. 2013), and in the unprotected fraction of the soil organic matter (Herold et al. 2014) in grasslands, indicate faster fine root derived C cycling in grasslands than in forests. Interestingly these differences do not extend to the mineral associated fraction of the soil organic matter.

#### Site effects on fine root decomposition rates

As hypothesized, standardized root decomposition varied among the three study regions, with the highest rates (in grassland and forest) in the SCH study region, followed by ALB and HAI (Table 5). The decomposition patterns were the same for herbaceous and beech standardized root material deployments, indicating that the trends were not related to litter quality, but to differences in the soil decomposition environment between the study regions. HAI is intermediate in average climate and most soil properties (Table 2, Table 3), but has the lowest overall decomposition rates. The only factor we found that might explain this is that the soils in HAI were frozen for a longer number of days compared to ALB and SCH. It has been shown that in soils with temperatures below 0 °C the microbial activity is slowed down. This is due to the rapid decline of unfrozen water content which can decrease the diffusion of substrates, nutrients, and waste products (Dioumaeva et al. 2002; Mikan et al. 2002; Ostroumov and Siegert 1996). Thus, also the degradation of fine roots may be slowed down, especially during the winter season in temperate ecosystems. In grasslands, we further observed considerably higher fine root decomposition in organic soils (Histosols and Gleysols present in the SCH study region) than in mineral soils in the SCH study region. This was probably related to the different soil properties, such

as soil moisture and nutrient contents, which may in turn influence the soil biota which directly degrades the root litter.

After the removal of variance specific of the study regions and differences in organic and mineral soils, we observed that the decomposition of standardized fine root litter in grasslands decreased with increasing land use intensity. However, in comparison to other environmental properties which explained together 24 % of fine root decomposition, land use intensity explained only a small proportion of variation (4 %). The negative relation observed between root decomposition and land use intensity is in our study mainly affected by the addition of N by fertilization. The lower need to mineralize organic N might slow down decomposition rates (Fog 1988). Knorr et al. (2005) reported that litter decomposition is inhibited by N additions when fertilization rates exceed by 2 to 20 times the atmospheric N deposition level. Since our fertilized plots received a maximum of  $140 \text{ kg N ha}^{-1} \text{ yr}^{-1}$  and the average bulk N deposition levels for our study regions are on average  $10 \text{ kg N ha}^{-1} \text{ yr}^{-1}$  (Schwarz et al. 2014), it is possible that the amount of added fertilizer inhibited the rates of root decomposition in fertilized ecosystems. In forests we did not observe land use effects on fine root decomposition.

#### Relevance of environmental site conditions and root litter chemistry on fine root decomposition

The lignin:N ratio explained the largest amount of variability (15 % in grasslands and 11 % in forests) of the mass loss of on-site collected fine root-litter in the grasslands and forests of the HAI region (Fig. 4). The remaining variability was explained by the sum of other environmental factors such as soil moisture and soil temperature, but also by the soil C:N ratios in grasslands (all environmental variables explained 34 % of the variance in grasslands and 24 % in forests) (Fig. 4). Although the lignin:N ratio itself merely describes the proportions of lignin to N without providing information about how lignin and N are distributed in plant organs, previous studies have shown that it is a valuable predictor for root decomposition (Bardgett et al. 1998; Hobbie 2005; Scheffer and Aerts 2000). Berg (1984) also concluded that the dominant factors for initial mass loss rates of root litter in a Scots pine forest are the relative amount of nutrients together with the initial lignin content. In particular, plant litter with higher N

concentrations decomposes faster than its lower nitrogen counterpart, while plant litter with high levels of lignin decomposes slower (Berg 1984; Janssens et al. 2010). While in forest ecosystems we observed a decline of the decomposition rates for increasing lignin:N ratio of the litter, in grasslands we found the opposite relationship. The detected patterns may reflect the variation in the initial root chemistry among the fine roots collected on-site in grasslands and forests (Hobbie et al. 2010; Silver and Miya 2001). For instance we found high variability of N concentration in fine roots of grasslands (table 4). The lignin concentrations were instead relatively constant. On the other hand in forest sites the lignin concentrations in tree roots encompassed a relatively large variability.

In the grasslands of the HAI region we observed a negative relation between the LUI index and the lignin:N ratio of fine roots collected on-site (Fig. 3). This indicates that land use can affect fine root decomposition through its influence on litter quality. In forests, we observed a negative correlation between the lignin:N ratio and the number of tree species per plot. However, this may be a function of our study design, as most of our plots are covered by pure European beech or mixed forests with more than one species and only a small number are pure Norway spruce (*Picea abies* (L.) H. Karst.) or Scots pine (*Pinus sylvestris* L.) forests. Therefore, higher lignin:N ratio in plots with a smaller number of tree species may be due to the higher amounts of lignin contained in European beech fine roots in comparison to, for example, coniferous tree species (Hobbie et al. 2010).

## Conclusions

We observed that fine root decomposition in temperate grasslands is two times faster than in temperate forests within the range of measured values. This finding together with other observations of older C in fine roots and in the unprotected fractions of soil organic matter in forests overall indicate slower C turnover in forests compared to grasslands. In both grasslands and forests the decomposition patterns of standardized fine root litter were different for the three study regions, indicating that the trends are influenced by differences in environmental properties such as the soil biota and the soil microclimate. Land use intensity in grasslands, in particular N additions, also had an influence on fine root

decomposition, though this amount of variation was small compared to the other abiotic factors. Within one study region, environmental variables explained the decomposition of both standardized and on-site collected litter. Additional variation for the on-site collected litter was explained by root lignin:N ratio, which in our study was influenced by land use; for example N addition in grasslands and tree species composition in forests.

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