

Rhizosphere priming of barley with and without root hairs



Johanna Pausch^{a,*}, Sebastian Loepmann^a, Anna Kühnel^b, Kelsey Forbush^c,
Yakov Kuzyakov^{a,d}, Weixin Cheng^c

^a Department of Soil Science of Temperate Ecosystems, Georg-August-University Göttingen, Germany

^b Department of Soil Science, Technical University of Munich, Freising, Germany

^c Department of Environmental Studies, University of California, Santa Cruz, CA, USA

^d Department of Agricultural Soil Science, Georg-August-University Göttingen, Germany

ARTICLE INFO

Article history:

Received 8 February 2016

Received in revised form

11 May 2016

Accepted 14 May 2016

Keywords:

Soil CO₂ efflux

Root morphology

Root hairless barley mutant

Isotope labeling

Enzyme activities

ABSTRACT

The influence of plant roots and the associated rhizosphere activities on decomposition of soil organic matter (SOM), the rhizosphere priming effect, has emerged as a crucial mechanism regulating global carbon (C) and nitrogen (N) cycles. However, the role of root morphology in controlling the rhizosphere priming effect remains largely unknown. To investigate the link between root hairs, a critical part of the entire root morphology, and the rhizosphere priming effect, we grew a barley wild type and a barley mutant without root hairs in a greenhouse and continuously labeled them with ¹³C-depleted CO₂. Soil CO₂ efflux was measured during tillering and head-emergence stages of plant growth. Based on its δ¹³C signature, total CO₂ was partitioned for root-derived and SOM-derived CO₂, and the SOM decomposition primed in the rhizosphere was calculated. Soil microbial biomass C and N, and the activities of six extracellular enzymes (β-cellobiohydrolase, β-glucosidase, acid phosphatase, β-xylosidase, leucin-aminopeptidase, and N-acetyl-β-glucosaminidase) were measured to test the effects of root hairs.

During the early stage of development (tillering), when plants were sufficiently supplied with nutrients, the barley mutant without root hairs produced more shoot biomass. In contrast, high C costs for root-hair formation likely reduced the growth of the barley wild type. At this stage, the wild type with regular root hairs produced a positive rhizosphere priming effect (69% increase), but the mutant without root hairs produced a negative priming effect on SOM decomposition (28% decline). At the head-emergence stage, when nutrients were scarce, plant biomass production of the mutant was reduced, probably due to inefficient nutrient uptake in the absence of root hairs. At this stage, both barley types produced positive rhizosphere priming effects (72% and 209% increase for the wild type and the mutant, respectively) and the microbial biomass was higher for both planted soils compared to the tillering stage. Extracellular enzymes responsible for the decomposition of stable SOM had higher activities in cases of positive priming effects. Overall, root hairs played an important role in regulating rhizosphere priming.

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1. Introduction

Soil CO₂ is one of the largest fluxes in the global C cycle, approximately ten-fold greater than CO₂ emissions from fossil fuel combustion (Schlesinger and Andrews, 2000; Amundson, 2001). The majority of this flux results from the decomposition of soil organic matter (SOM) and litter by microbes (Kuzyakov, 2006). In recent years, there is an emerging view that, in addition to

temperature and moisture, carbon substrate availability is a key factor controlling SOM turnover (Fontaine et al., 2007; Paterson and Sim, 2013). These changes in the rate of SOM turnover following the input of easily decomposable substrates for microorganisms are termed 'priming effects'.

While decaying leaf and root litter provides some labile substrate for soil microbes, the majority of the labile substrate in soils comes from roots. For example, some studies have reported that SOM decomposition may be 380% greater in soils with roots compared to unplanted soils (positive rhizosphere priming effects; RPE) (Cheng et al., 2014). Accordingly, the magnitude of RPE may control C fluxes at the ecosystem level and influence ecosystem

* Corresponding author. Department of Soil Science of Temperate Ecosystems, University of Göttingen, Büsgen-Institute, Büsgenweg 2, 37077 Göttingen, Germany.
E-mail address: jpausch@gwdg.de (J. Pausch).

feedbacks to climate (Cheng et al., 2014; Finzi et al., 2015).

The explanation for most of the reported positive RPE is microbial activation, i.e. the stimulation of growth and activity induced by root-derived substrates. Microbes utilize this energy subsidy to produce extracellular enzymes (exoenzymes) that enhance the release of nutrients from SOM (Blagodatskaya and Kuzyakov, 2008). While microbes benefit from the nutrients released through enhanced decomposition, plants may benefit too – suggesting that RPE may be an evolutionary stable strategy (Cheng et al., 2014).

Living roots release numerous available low molecular weight substrates such as sugars, carboxylic acids and amino acids throughout the soil profile and over the course of the growing season (Nguyen, 2003; Jones et al., 2009). These substrates are not homogeneously distributed along the root segments but are rather released in distinct areas, mainly at the root tips (McDougall and Rovira, 1970; Nguyen, 2003; Dennis et al., 2010; Pausch and Kuzyakov, 2011). For this reason, root morphology (e.g. lateral root formation, number of root tips, root hair formation) may largely impact exudation (Nguyen, 2003), and may, hence, be decisive for rhizosphere priming effects. The root morphology, in turn, is mainly controlled by the nutrient availability in the soil since changes in root architecture can alter the capacity of plants to take up nutrients (López-Bucio et al., 2003). Several strategies have been developed by plants to increase the uptake of limited nutrients from the soil. An efficient strategy to acquire limited nutrients is the production of root hairs, which could differ in numbers, density and length between plant species depending on the kind of nutrients and nutrient availability in the soil (Jungk, 2001). The substantial contribution of root hairs to plant nutrition and accompanied therewith nutrient shortages in the rhizosphere and high energy supply to microbes through exudation, as well as direct and indirect enhancement of enzyme activities (Spohn and Kuzyakov, 2014) may be crucial for rhizosphere priming effects. A barley mutant lacking root hairs completely was discovered by Gahoonia et al. (2001). This mutant enabled us to study the role of root hairs for rhizosphere priming effects.

In the present experiment a barley wild type with root hairs and the root hairless mutant were grown under controlled conditions. Rhizosphere priming effects, i.e. changes in the rate of SOM decomposition, were indicated by an increase or decrease of SOM-derived CO₂ production in planted compared to an unplanted soil. By continuously labeling plants with ¹³C-depleted CO₂, we were able to differentiate root-derived CO₂ from SOM-derived CO₂ and calculate RPE as the difference in SOM-derived CO₂ between planted and unplanted soils. To investigate the influence of plant age, the soil CO₂ efflux was trapped at two plant growth stages of (tillering and head-emergence). Microbial parameters (microbial biomass C and nitrogen (N), enzyme activities) were analyzed to assess changes of microbial activities.

We hypothesize that the rhizosphere priming effect depends on root morphology. More specifically, a better nutrient acquisition of the wild type with root hairs through a higher total root surface area resulted in a larger plant biomass production, thus, leading to higher exudation and higher positive RPE. We also expect that the higher the positive priming is the more active are exoenzymes responsible for the decomposition of more stable substrates. Plant age is known to play an important role for the intensity of priming (Fu and Cheng, 2002; Pausch et al., 2013). Due to different growth pattern and nutrient demands, we hypothesize that plant age influences rhizosphere priming on SOM decomposition differently for the hairless barley mutant and the barley wild type with root hairs.

2. Material and methods

2.1. Experimental setup

Two barley (*Hordeum vulgare* L.) types, a wild type (cv. optic; WT) and a root-hairless mutant called bald root barley (*brb*, Gahoonia et al., 2001, Fig. 1), were grown in a greenhouse and were continuously labeled with ¹³C-depleted CO₂ (Cheng and Dijkstra, 2007). The plants were exposed to the tracer from the emergence of the first leaf till the end of the experiment. Briefly, a constant CO₂ concentration of 400 ± 5 ppm and a constant δ¹³C value of about –18‰ was maintained inside the greenhouse over the course of the experiment by regulating the flow of ¹³C-depleted CO₂ (99.9% CO₂, δ¹³C of –38‰) from a tank and setting CO₂-free air flow rate proportional to the leakage rate (300 L/min) of the greenhouse (Zhu and Cheng, 2012; Pausch et al., 2013). The CO₂-free air was produced from compressed air passed through six soda lime columns (20 cm diameter, 200 cm length) filled with approximately 40 kg soda lime (pellets made of NaOH and Ca(OH)₂ mixture) each. The CO₂-free air flow was set at 120 L/min. The CO₂ concentration inside the greenhouse was continuously monitored by an infra-red gas analyzer (Model LI-820, Li-COR, Lincoln, NE, USA) and stabilized at 400 ± 5 ppm by computer-controlled CO₂ injection from the tank. A fan was used to ensure a uniform distribution of the CO₂ inside the greenhouse. For the duration of the experiment, the δ¹³C value of the greenhouse air was measured every three days during the light period by pumping air through a glass airstone immersed in 50 ml of 0.5 M NaOH solution. The CO₂ trapping efficiency was nearly 100% as checked by an infra-red gas analyzer (Model LI-6262, Li-COR, Lincoln, NE, USA). An aliquot of the sample was precipitated with SrCl₂ as SrCO₃ using the method described by Harris et al. (1997) and analyzed for δ¹³C (relative to PDB standard) using a PDZ Europa ANCA-GSL elemental analyzer interfaced to a PDZ Europa 20–20 isotope ratio mass spectrometer (Sercon Ltd., Cheshire, UK). The mean δ¹³C value of the CO₂ in the greenhouse air was –18.2 ± 0.3‰.

The two barley types were grown in PVC pots (15 cm diameter, 40 cm height, equipped with an inlet tube at the bottom for aeration and soil CO₂ trapping). A nylon bag filled with ~1500 g sand was placed at the bottom of each PVC pot to improve air circulation. Each pot was filled with about 7 kg sieved (<2 mm) soil. The soil was taken from the plough horizon (top 20 cm) of a sandy loam (Mollisol) from a farm on the campus reserves of the University of California, Santa Cruz. The soil contained 1.18 ± 0.01% organic C and 0.13 ± 0.001% N, had δ¹³C and δ¹⁵N values of –26.45 ± 0.07‰ and 7.12 ± 0.02‰, respectively, and a pH value of 5.8. All filled pots were wetted to 20% gravimetric soil moisture content (equivalent of 80% of the water holding capacity) with deionized water.

For each barley type, 10 pots were set up. In addition, 8 unplanted pots (unplanted soil; US) were prepared (in total 28 pots). The seeds were presoaked overnight and 6–8 barley seeds were planted per pot. The inlet tube at the bottom of each pot was connected to an aquarium pump to aerate the pots. This was done 2 times during the dark period to avoid contamination of the growth chamber δ¹³C signal with that of soil-derived CO₂ during the assimilation period.

The soil moisture content was measured gravimetrically and adjusted daily to 80% of the water holding capacity. To maintain homogeneous soil moisture and good soil structure, water was added through perforated tubes buried at the center of the pot (inner diameter 0.32 cm, total length 20 cm, buried length 10 cm). The location of the pots in the greenhouse was changed weekly by mixing them randomly to guarantee similar growing conditions for the plants. The day time air temperature inside the greenhouse was maintained at 23 °C by two air conditioning units. The night time

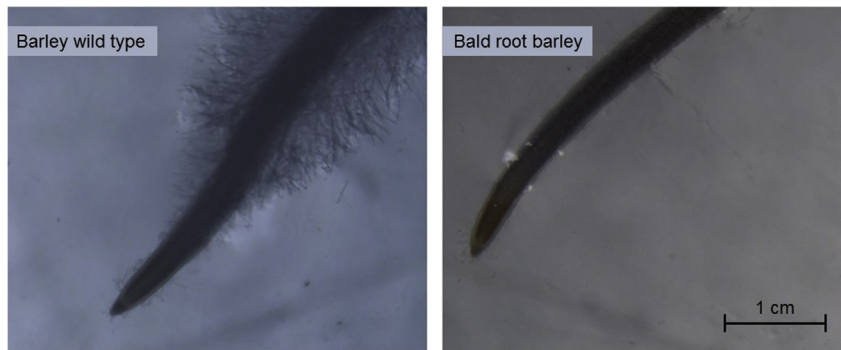


Fig. 1. Light microscopy images of the barley wild type and the root-hairless barley mutant germinated in deionized water.

temperature was kept above 17 °C. Artificial lighting (1100 W lights, P.L. Light Systems, Beamsville, ON) was used to ensure an adequate light intensity throughout the experiment. The light intensity was kept above 900 W m⁻². The photoperiod was set from 4:30a.m. to 4:30p.m. The relative air humidity was kept at 45% by a dehumidifier (Kenmore Elite 70 pint, Sears, Chicago, IL, USA).

2.2. Measurements

2.2.1. Soil CO₂ efflux

Soil CO₂ efflux from each pot was measured at two growth stages of barley, 29–30 days after planting (DAP) at tillering (T1) and 64–65 DAP at head-emergence (T2), by using a closed-circulation CO₂ trapping system (Cheng et al., 2003; Pausch et al., 2013). Prior to each CO₂ trapping the pots were sealed with non-toxic silicone rubber (GI-1000, Silicones Inc., NC, USA) added directly to the soil surface.

Soil CO₂ trapping was performed on 4 replicates each of the unplanted soil, and the barley with and without root hairs at T1. At T2, CO₂ was trapped from 6 replicates of planted pots and 4 replicates of unplanted pots. Shortly before CO₂ trapping, the CO₂ inside the pots was removed by circulating the isolated air through a soda lime column (3 cm diameter, 50 cm length) for 40 min. Then CO₂ produced in the sealed pots was trapped for 24 h in 400 ml of 0.5 M NaOH solution. Four blanks were included to correct the total inorganic C content for possible contamination from carbonate in the NaOH stock solution and from sample handling (Cheng et al., 2003; Pausch et al., 2013). An aliquot of each NaOH solution was analyzed for total inorganic carbon using a Shimadzu TOC-5050A Total Organic Carbon Analyzer. Another aliquot was precipitated as SrCO₃ (Harris et al., 1997) and analyzed for δ¹³C by means of a continuous flow isotope ratio mass spectrometer as described above.

2.2.2. Shoot, root and soil analyses

After each CO₂ sampling the pots were destructively harvested. The shoots were cut at the base. The soil of each pot was pulled out and the roots were separated by hand-picking. Subsamples of about 1 kg soil were stored in a freezer (−18 °C) until further analysis. Shoots, root, and soil samples were dried at 60 °C for 3 days, weighed, ground in a ball mill and measured for δ¹³C and δ¹⁵N using a Carlo Elba 1108 elemental analyzer interfaced to a Thermo-Finishing Delta Plus XP isotope ratio mass spectrometer at the Isotope Facility of University of California-Santa Cruz.

Dissolved N (DN) which is extractable with K₂SO₄ was determined as described below (2.2.3). Extractable phosphorus (bioavailable inorganic *ortho*-phosphate) was determined on dried and ground soil samples by the Bray-Method at the Analytical Laboratory, University of California, Davis (<http://anlab.ucdavis.edu/>).

2.2.3. Microbial biomass C and N

Soil microbial biomass C (MBC) and soil microbial biomass N (MBN) were determined on all soil samples by the chloroform-fumigation-extraction method described by Vance et al. (1987) with the modification that fumigated and non-fumigated soil samples (7.5 g) were extracted for 1 h with 30 mL of 0.05 M K₂SO₄ solution. The samples were filtered (Quantitative Paper FT-3-101-09, Sartorius) and the extracts were analyzed for total organic C and N by means of a multi N/C analyzer (multi N/C analyzer 2100S, Analytik Jena). Total N content of the non-fumigated extracts were used as a measure of available N. The difference between the extracts of fumigated and non-fumigated samples gave the amount of chloroform-labile C and N (hereafter referred to as MBC and MBN). We noted that these values did not correspond to total amount of MBC and MBN as the extraction efficiency was not taken into account. Reported conversion factors *k_{ec}* (or *k_{en}* for N) ranged from less than 0.2 to 0.45 among different soils (Wu et al., 1990; Dictor et al., 1998; Bailey et al., 2002). Thus, for the purpose of comparing treatment effects and avoiding biases of conversion factors, the data presented in this study were not corrected by conversion factor.

2.2.4. Enzyme assays

To determine the activities of the enzymes β-cellobiohydrolase (exo-1,4-β-glucanase, EC 3.2.1.91), β-glucosidase (EC 3.2.1.21), acid phosphatase (EC 3.1.3.2), β-xylosidase (EC 3.2.2.27), leucine-aminopeptidase (LAP) (EC 3.4.11.1), and N-acetyl-β-glucosaminidase (chitinase, EC 3.2.1.52), we used 4-methylumbelliferyl-β-D-cellobioside, 4-methylumbelliferyl-β-D-glucopyranoside, 4-methylumbelliferyl-phosphate, 4-methylumbelliferyl-7-β-D-xylopyroniside, L-leucine-7-amino-4-methylcoumarin hydrochloride and 4-methylumbelliferyl N-acetyl-β-D-glucosaminide, respectively. The soil suspension was dispersed by an ultrasonic disaggregator (50 J s⁻¹ for 120 s) after addition of half a gram of soil to 50 ml sterile water in autoclaved jars (De Cesare et al., 2000). While stirring the soil suspension 50 μl aliquots were withdrawn and dispensed in 96-well microplates (Brand pureGrade, black). Buffer (80 ml) was added (0.1 M MES buffer, pH 6.1 for carbohydrases and phosphatase, 0.05 M TRIZMA buffer, pH 7.8 for leucine-aminopeptidase) (Marx et al., 2001, 2005; Loeppmann et al., 2016).

We added 100 μl of series concentrations of substrate solutions (20, 40, 60, 80, 100, 200, 400 μmol substrate g⁻¹ soil) to the wells and kept the temperature at 21 °C. The micro-plates were agitated and measured fluorometrically (excitation 360 nm; emission 450 nm) after 1 h, 2 h, and 3 h incubation with an automated fluorometric plate-reader (Wallac 1420, Perkin Elmer, Turku, Finland). Fluorescence was converted into an amount of MUB (4-methylumbelliferone) or AMC (7-amino-4-methylcoumarin), according to specific standards, which had been prepared in sub-

samples from the various soil suspensions.

2.3. Calculations

The contribution of CO₂ derived from SOM decomposition ($C_{SOM-DEIVED}$, mg C kg⁻¹ soil day⁻¹) to total soil respiration was calculated using a linear two-source isotopic mixing model:

$$C_{SOM-DEIVED} = C_{TOTAL} \frac{\delta^{13}C_{TOTAL} - \delta^{13}C_{ROOT-DEIVED}}{\delta^{13}C_{SOM-DEIVED} - \delta^{13}C_{ROOT-DEIVED}} \quad (1)$$

$$C_{ROOT-DEIVED} = C_{TOTAL} - C_{SOM-DEIVED} \quad (2)$$

where C_{TOTAL} is the total CO₂ efflux of the planted soil (mg C kg⁻¹ soil day⁻¹) and $\delta^{13}C_{TOTAL}$ the corresponding $\delta^{13}C$ value (‰). $\delta^{13}C_{SOM-DEIVED}$ is the $\delta^{13}C$ value of CO₂ from SOM decomposition measured in the unplanted soils (‰) (Table 1). Note, as priming may promote the decomposition of specific compounds of SOM with varying $\delta^{13}C$, the $\delta^{13}C_{SOM-DEIVED}$ could slightly be biased.

$C_{ROOT-DEIVED}$ is the root-derived CO₂ in the planted soils (mg C kg⁻¹ soil day⁻¹) with $\delta^{13}C_{ROOT-DEIVED}$ as the corresponding $\delta^{13}C$ value (‰) (Table 1). To consider isotopic fractionation, we accounted for ¹³C differences between the isotopic composition of roots and that of root-derived CO₂ (Pausch et al., 2013). The fractionation factor (f) was taken from Zhu and Cheng (2011) and was -0.87‰. This fractionation factor was measured for wheat (*Triticum aestivum*) and was chosen since barley and wheat are both belonging to the *Poaceae* family and are both monocotyledons with similar properties. Root-derived CO₂ was calculated based on the assumption of an equal isotopic fractionation for the barley wild type and the mutant.

$\delta^{13}C_{ROOT-DEIVED}$ was calculated by correcting the $\delta^{13}C$ value of the root ($\delta^{13}C_{ROOT}$) by a fractionation factor (f):

$$\delta^{13}C_{ROOT-DEIVED} = \delta^{13}C_{ROOT} + f \quad (3)$$

The RPE on SOM decomposition was calculated by subtracting the CO₂ flux of the unplanted soil ($C_{SOM-DEIVED}(US)$) from the SOM-derived CO₂ flux of the planted soil ($C_{SOM-DEIVED}(WT, brb)$).

$$RPE = C_{SOM-DEIVED}(WT, brb) - C_{SOM-DEIVED}(US) \quad (4)$$

The RPE was related to total root biomass (gDW) as well as expressed as percentage of basal respiration of the unplanted soil.

2.4. Statistics

The values presented in the figures and tables are given as means ± standard errors. A one-way analysis of variance (ANOVA) was conducted to test for significant differences in all measured data between the unplanted soil, and barley with and without root hairs by calculating the ANOVA separately for each sampling date. The significance of differences between individual means was

obtained by a *post hoc* unequal N HSD test. Moreover, rhizosphere priming values were tested for significant deviation from zero by a *t*-test. All statistical analyses were performed with the statistical package STATISTICA for Windows (version 7.0; StatSoft Inc., OK, USA).

For enzyme analyses, we used a non-linear regression (Michaelis-Menten kinetics) to estimate the kinetic parameter V_{max} (Marx et al., 2001). Each soil sample was measured as an analytical triplicate. The kinetic parameters were fitted by minimizing the least-square sum using GraphPad Version 6 software (Prism, USA). Parameter optimization was restricted to the applied model equation as indicated by maximum values of statistic criteria: r^2 . Outliers were identified by the ROUT method, based on the False Discovery Rate (FDR) (Motulsky and Brown, 2006). A multiple *t*-test was applied to test for differences in enzyme activities between the unplanted soil, and the barley with and without root hairs and between T1 and T2. Statistical significance was determined using the Holm-Sidak method ($P \leq 0.05$).

3. Results

3.1. Plant biomass and $\delta^{13}C$ and $\delta^{15}N$ values

The main difference between the two barley types was the production of plant biomass; shoots and roots. The mutant (*brb*), completely lacking root hairs, produced higher shoot biomass during the tillering stage (T1) but root biomass was not significantly different compared to the wild type (WT). However, at head-emergence stage (T2) root biomass of the mutant was reduced compared to the wild type (WT) (Table 2).

At T1, C and N contents of shoots were on average $34.3 \pm 0.4\%$ and $6.1 \pm 0.1\%$, respectively. At T2 a higher C content of $38.2 \pm 0.4\%$ was measured while the N content decreased to $1.7 \pm 0.1\%$ compared to T1. This led to a much higher C/N ratio at T2 compared to T1 (Table 2). Similarly, the C content of roots was higher at T2 ($33.6 \pm 1.1\%$) compared to T1 ($22.1 \pm 2.0\%$). The N content of roots, however, was similar between the two sampling times. The C/N ratio of roots increased from 13.9 ± 0.6 at T1 to 25.3 ± 1.4 at T2.

The plants were successfully labeled with ¹³C-depleted CO₂ as shown by the $\delta^{13}C$ values of shoots and roots (Table 2). The $\delta^{13}C$ value of shoots was $-41.4 \pm 0.1\%$ at T1 which was lower than at T2 ($-39.3 \pm 0.3\%$). The $\delta^{13}C$ value of roots did not differ between T1 and T2. Interestingly, the $\delta^{15}N$ value of roots increased between T1 and T2 and was about 2.6‰ higher at T2 (Table 2).

3.2. Soil N and P contents

Available P did not differ between planted and unplanted soils likely because of the high P availability of the soil used in this study (Table 3). However, nutrient uptake by plants led to lower dissolved N (DN). The DN content was reduced by ~46% in planted soil compared to the unplanted soil at tillering. At head-emergence

Table 1

End member values (±SEM) used in two-source isotopic mixing models, in order to calculate the contribution of SOM-derived CO₂ to total soil CO₂ of the planted treatments, at tillering (T1) and head-emergence stage (T2) for unplanted soil (US), the barley wild type (WT) and the root-hairless barley mutant (*brb*). The sample size is given in parentheses.

Sampling time	Treatment	Root-derived CO ₂ [‰]	SOM-derived CO ₂ of the unplanted soil [‰]
T1	US		-24.20 ± 0.51 (4)
	WT	-38.65 ± 0.28 (3)	
	<i>brb</i>	-39.65 ± 0.36 (4)	
T2	US		-24.90 ± 0.22 (4)
	WT	-38.57 ± 0.22 (6)	
	<i>brb</i>	-39.29 ± 0.21 (4)	

Table 2
Plant (shoot and root) biomass, C and N contents, C/N ratios, and $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values (\pm SEM) 30 (T1) and 65 (T2) days after planting. Significant differences ($P \leq 0.05$) between the treatments are indicated by different lowercase letters. The asterisk indicates significant differences between T1 and T2.

Sampling time		T1		T2	
Treatment		WT	<i>brb</i>	WT	<i>brb</i>
Shoot	Biomass [$\text{g}_{\text{DW}} \text{pot}^{-1}$]	3.03 \pm 0.13b	3.70 \pm 0.07a	38.87 \pm 2.38a*	30.70 \pm 2.54a*
	C content [%]	34.08 \pm 0.74a	34.51 \pm 0.31a	38.67 \pm 0.21a*	37.69 \pm 0.69a*
	N content [%]	6.15 \pm 0.06a	6.08 \pm 0.09a	1.47 \pm 0.10a*	1.87 \pm 0.16a*
	C/N	5.54 \pm 0.08a	5.67 \pm 0.05a	26.72 \pm 2.09a*	20.64 \pm 1.95a*
	$\delta^{13}\text{C}$ [‰]	-41.45 \pm 0.05a	-41.40 \pm 0.10a	-38.75 \pm 0.10a*	-39.75 \pm 0.42a*
	$\delta^{15}\text{N}$ [‰]	5.23 \pm 0.28a	5.07 \pm 0.21a	6.13 \pm 0.40a	5.67 \pm 0.50a
Root	Biomass [$\text{g}_{\text{DW}} \text{pot}^{-1}$]	1.65 \pm 0.04a	1.88 \pm 0.43a	12.30 \pm 1.21a*	7.37 \pm 0.90 b*
	C content [%]	21.13 \pm 2.69a	23.04 \pm 3.18a	32.14 \pm 0.96a*	34.99 \pm 1.83a*
	N content [%]	1.61 \pm 0.35a	1.68 \pm 0.26a	1.27 \pm 0.11a	1.47 \pm 0.15a
	C/N	13.89 \pm 1.29a	13.88 \pm 0.46a	25.94 \pm 1.75a*	24.57 \pm 2.47a*
	$\delta^{13}\text{C}$ [‰]	-37.84 \pm 0.21a	-38.78 \pm 0.36a	-37.70 \pm 0.27a	-38.46 \pm 0.33a
	$\delta^{15}\text{N}$ [‰]	3.61 \pm 0.63a	2.51 \pm 0.27a	5.87 \pm 0.25a*	5.53 \pm 0.33a*
Shoot/Root		1.91 \pm 0.11a	2.21 \pm 0.35a	3.37 \pm 0.63a	4.25 \pm 0.31a*

Table 3
Dissolved nitrogen (DN), available phosphorus (Bray-P), and chloroform-labile microbial biomass C and N, at tillering (T1) and head-emergence stage (T2) for unplanted soil (US), the barley wild type (WT) and the root-hairless barley mutant (*brb*). Significant differences ($P \leq 0.05$) between the treatments are indicated by different lowercase letters. The asterisk indicates significant differences between T1 and T2.

Sampling time	T1			T2		
	US	WT	<i>brb</i>	US	WT	<i>brb</i>
K ₂ SO ₄ -extractable DN [mg N kg^{-1} soil]	104.62 \pm 4.68a	57.13 \pm 1.34 b	57.74 \pm 3.18b	nd	15.00 \pm 1.71a*	15.61 \pm 1.59a*
Bray-P [mg P kg^{-1} soil]	66.71 \pm 1.87a	62.74 \pm 0.69a	66.79 \pm 1.20a	68.35 \pm 1.58a	60.82 \pm 2.52a	59.92 \pm 1.70a*
Chloroform-labile MBC [mg C kg^{-1} soil]	67.60 \pm 5.41a	62.03 \pm 2.72a	66.61 \pm 6.26a	81.25 \pm 7.55a	101.46 \pm 14.42a*	92.21 \pm 7.16a*
Chloroform-labile MBN [mg N kg^{-1} soil]	8.92 \pm 1.77a	6.72 \pm 1.81a	8.66 \pm 0.39a	10.21 \pm 1.93a	13.19 \pm 1.33a*	14.49 \pm 1.21a*
MBC/MBN	8.70 \pm 2.18a	11.41 \pm 2.71a	7.84 \pm 1.13a	9.47 \pm 3.05a	7.67 \pm 0.56a	6.48 \pm 0.63a

stage, only 14% of the initial DN content (unplanted soil at T1) remained in the planted soils (Table 3).

3.3. Microbial biomass C and N and enzyme activities

MBC and MBN (chloroform-labile C and N) was similar between the unplanted soil, and the barley with and without root hairs (Table 3). However, at the head-emergence stage of plant growth (T2) MBC and MBN increased for both barley types compared to the tillering stage (T1).

At tillering stage, the wild type had lower activities of β -glucosidase, β -cellobiohydrolase (not significant), and acid phosphatase, while the activity of β -xylosidase and chitinase were higher compared to the unplanted soil (Fig. 2). In contrast, the activities of β -glucosidase, β -cellobiohydrolase, and LAP did not differ between *brb* and the unplanted soil, but a higher activity of β -xylosidase and acid phosphatase was measured. At head-emergence stage, both barley types induced lower activity rates of β -glucosidase and LAP (yet not significant for *brb*), while the activities of β -xylosidase and chitinase increased through planting. β -cellobiohydrolase and acid phosphatase activities of the planted soils were similar to that of the unplanted soil at T2.

3.4. CO₂ efflux partitioning

Total soil CO₂ efflux was influenced by planting and by the barley genotype as well as by sampling time. At T2 all planted soils showed higher total soil CO₂ efflux (sum of SOM- and root-derived CO₂) compared to the unplanted soils (Fig. 3).

SOM-derived CO₂ was higher for WT (29.2 \pm 0.6 mg C kg⁻¹ soil day⁻¹) compared to *brb* (12.5 \pm 1.8 mg C kg⁻¹ soil day⁻¹) at T1 (Fig. 3, top). Moreover, at T2 both barley types had higher SOM-derived CO₂ (24.3 \pm 3.0 mg C kg⁻¹ soil day⁻¹ for WT and

43.8 \pm 9.7 mg C kg⁻¹ soil day⁻¹ for *brb*) compared to the unplanted soil (14.2 \pm 0.3 mg C kg⁻¹ soil day⁻¹). However, this was only statistically significant ($P > 0.05$) for the root-hairless mutant. While SOM-derived CO₂ remained relatively constant for the unplanted soil and the WT between T1 and T2, the *brb* showed higher SOM-derived CO₂ at T2.

Root-derived CO₂ consists of CO₂ released from root respiration per se and of CO₂ released through microbial decomposition of rhizodeposits. Root-derived CO₂ positively correlates with root biomass ($R^2 = 0.99$, data not shown) (Pausch et al., 2013). At tillering, root-derived CO₂ did not differ between the barley wild type and the mutant. However, root-derived CO₂ increased at head-emergence for both barley types with increasing root biomass (Fig. 3 (bottom), Table 2). Moreover, the lower root biomass of the *brb* at T2 compared to WT is reflected in a slight, but not significant, lower root-derived CO₂.

On a root dry weight basis, root-derived CO₂ was similar in the two barley types and between sampling dates (Fig. 3, inlet). However, there was a trend of less root-derived CO₂ at head-emergence (Fig. 3, inlet).

3.5. Rhizosphere priming effect

During the early stage of plant growth (T1), rhizosphere priming was largely influenced by the barley genotype. While SOM decomposition was increased for the wild type by 69% compared to the unplanted soil (difference to zero $P = 0.002$), it decreased for the root-hairless barley by 28% (difference to zero $P = 0.051$; Fig. 4, right y-axis). At the head-emergence stage (T2), both barley types showed positive priming effects with even higher intensity under the mutant. Rhizosphere priming was enhanced for the wild type compared to the unplanted soil by 72% (difference to zero $P = 0.020$). The highest positive priming effect was measured for

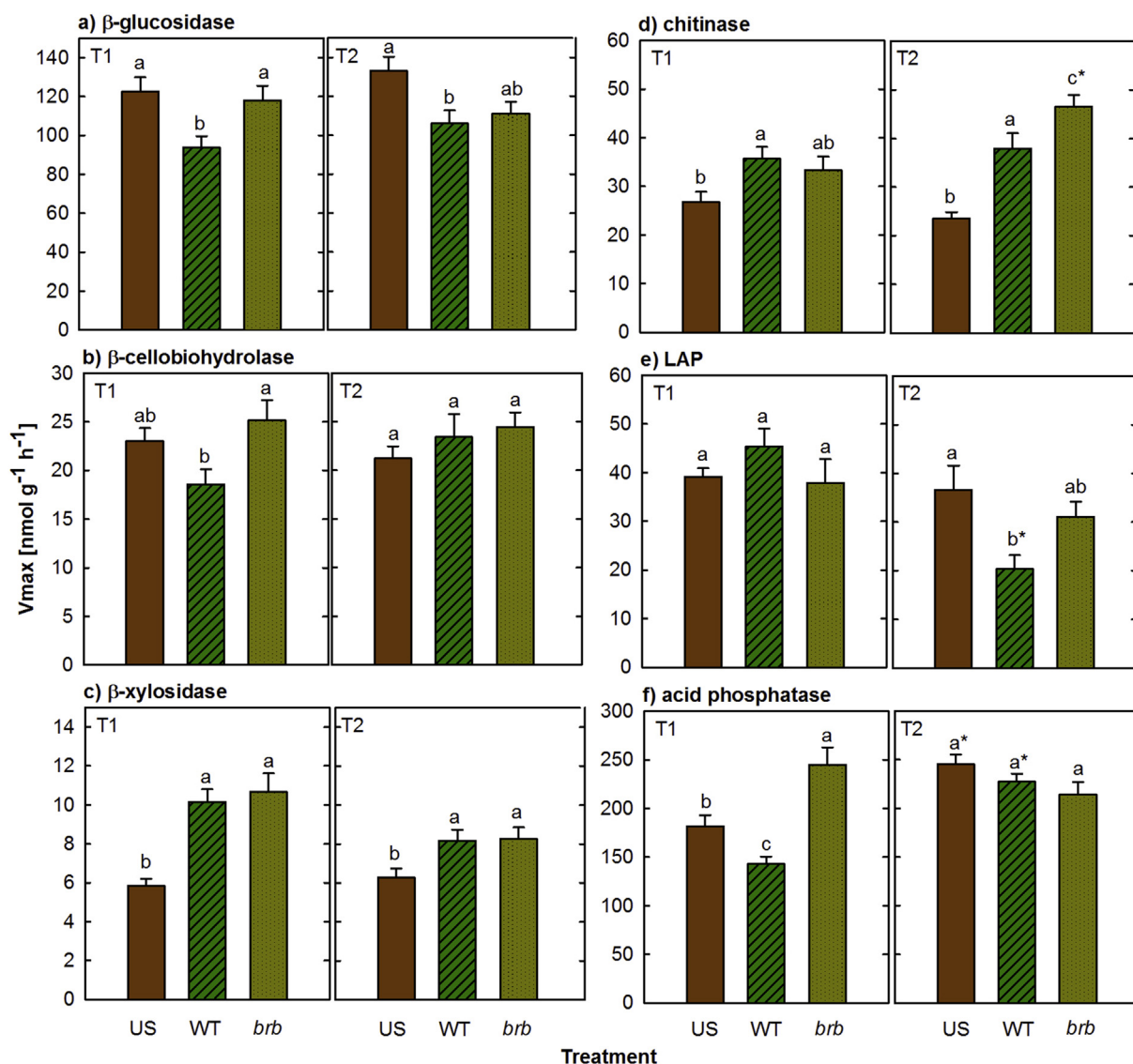


Fig. 2. Potential enzyme activities for a) β-glucosidase, b) β-cellobiohydrolase, c) β-xylosidase, d) chitinase, e) leucine-aminopeptidase (LAP), and f) acid phosphatase (±SEM) at tillering (T1) and head-emergence stage (T2) for unplanted soil (US), the barley wild type (WT) and the root-hairless barley mutant (*brb*). Bars labeled by different lowercase letters indicate significant differences ($P \leq 0.05$) between the treatments at one sampling date. Significant differences between T1 and T2 are indicated by an asterisk.

the barley mutant lacking root hairs, amounting to 209% of the unplanted soil (difference to zero $P = 0.055$).

To account for root properties effects, a specific RPE was calculated by relating total primed C to root biomass (Fig. 4, left y-axis). The specific RPE was highest for the wild type at the early stage of plant growth with 47.9 ± 1.1 mg C g⁻¹ root day⁻¹ (difference to zero $P = 0.096$). A negative priming effect for the *brb* was measured at T1 with -22.2 ± 8.5 mg C g⁻¹ root day⁻¹ (difference to zero $P = 0.055$). At the head-emergence stage (T2) the WT showed with 5.7 ± 2.0 mg C g⁻¹ root day⁻¹ (difference to zero $P = 0.035$), a lower specific RPE than during tillering (T1). In contrast, the barley mutant shifted from negative priming at T1 to a positive RPE at T2 (28.2 ± 9.7 mg C g⁻¹ root day⁻¹; difference to zero $P = 0.062$).

4. Discussion

During the tillering stage of plant growth, SOM decomposition was enhanced (positive priming) in soils with the barley wild type by 69% compared to the basal respiration of the unplanted soil. The

data was within the range of priming results published for wheat, another monocotyledon plant from the *Poaceae* family. Wheat showed positive priming effects ranging from 42% of the unplanted soil (28 days old wheat) (Cheng and Johnson, 1998; Cheng et al., 2014) to 75% for 30 days old wheat (Pausch et al., 2013). Positive priming effects could be explained by the 'microbial activation hypothesis' (Kuzyakov, 2002; Cheng and Kuzyakov, 2005), which assumes that the activity and growth of microorganisms is enhanced through metabolizing labile substrates (e.g. root exudates), further leading to an accelerated SOM turnover. In the presence of labile plant C microbes start decomposing SOM to acquire N ('Microbial nitrogen mining' Crain et al., 2007).

Interestingly, the barley mutant lacking root hairs showed a complete opposed effect on SOM turnover. SOM decomposition was reduced by 28% compared to the unplanted soil; hence, the *brb* induced negative priming at the tillering stage. As both barley types produced same amounts of roots at the tillering stage, our result point to root morphology (here the presence or absence of root hairs) as a main determinant for RPE. Negative priming effects were

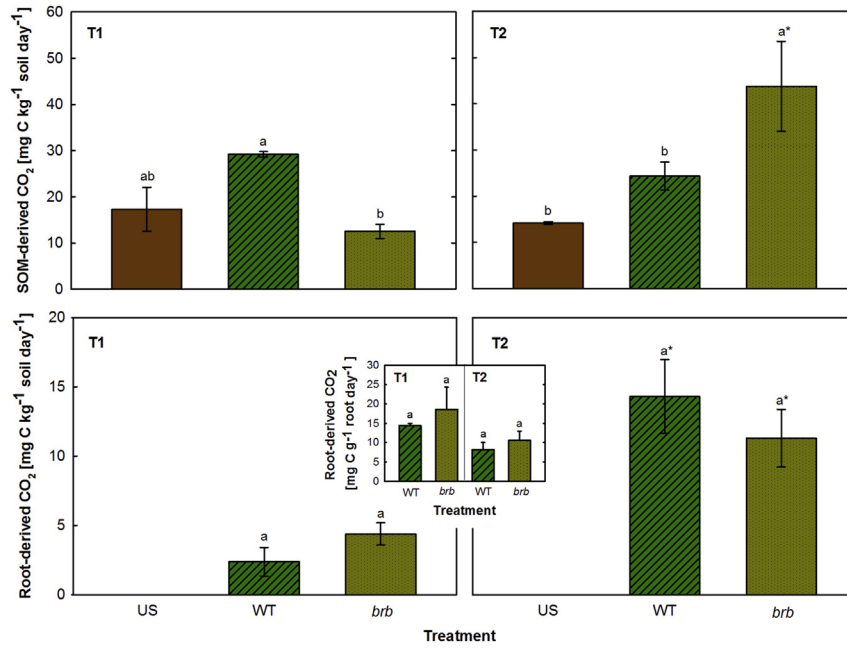


Fig. 3. SOM-derived CO₂ (top) and root-derived CO₂ (bottom) at T1 (left) and T2 (right). Bars labeled by different lowercase letters indicate significant differences ($P \leq 0.05$) between the treatments at one sampling date. Significant differences between T1 and T2 are indicated by an asterisk. The inset shows the root-derived CO₂ per g root for both barley types and sampling stages.

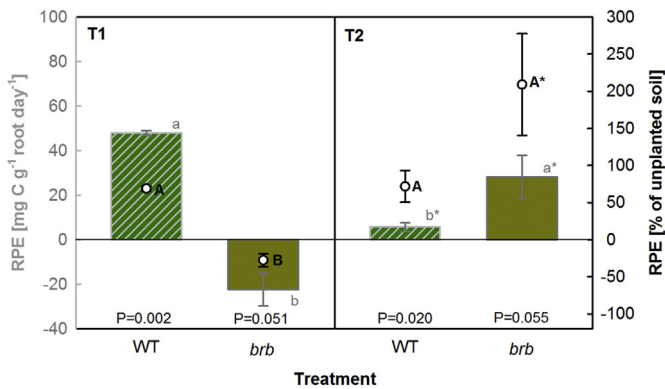


Fig. 4. Rhizosphere priming effects per root dry weight and time (\pm SEM) (left axis, bars) and as % of the unplanted soil (\pm SEM) (right axis, dots) at T1 and T2 in treatments with a barley wild type (WT) and a root hairless barley mutant (*brb*). Bars or dots labeled by different letters indicate significant differences ($P \leq 0.05$) between the treatments. Significant differences between T1 and T2 are indicated by asterisk. The P values from a t-test to test for significant deviation of RPE (% of unplanted soil) from zero are shown.

observed in short-term experiments (Cheng et al., 2014) and were explained either by 1) 'Preferential substrate utilization' (PSU), i.e., microorganisms, not limited in N, can switch from the decomposition of SOM to the decomposition of easily available rhizodeposits or by 2) 'Microbial competition hypotheses'. The latter suggests that microbes and plants compete for nutrients and thus, microbial growth decreases, thereby, depressing SOM decomposition (Kuzakov, 2002; Cheng and Kuzakov, 2005). At tillering, when mineral nutrients were still abundant, the activity of extracellular enzymes measured in soils with the root-hairless mutant did not differ or even increased (β -xylosidase, acid phosphatase) compared to the unplanted soil. This may point to PSU.

The present study suggests that at the tillering stage, root morphology plays a major role for rhizosphere priming effects. It is

likely, that the extension of the rhizosphere by root hairs accelerated SOM decomposition.

At the head-emergence stage, both barley types induced positive rhizosphere priming effects. When referred to the unplanted soil, the wild type primed 72%, while the *brb* increased SOM-decomposition by 209% compared to the unplanted soil. In a recent study Mwfulirwa et al. (2016) investigated barley genotypes and reported negative priming at day 19 after planting while all genotypes induced positive priming after 27 days of growth when nutrients are becoming scarce. Overall, a higher root biomass per pot (7 times higher for WT between T1 and T2, and 3 times higher for *brb*) at increasing N limitation (about 7 times as lower at T2 as the initial value of US at T1) triggers the positive priming effects at T2 in the present study. Since the dissolved N is highly reduced at T2, it is probable that microorganisms start mining for N to meet their N demand and thus, decompose SOM more intensively. This effect is clearly indicated by the $\delta^{15}\text{N}$ values of the plants. $\delta^{15}\text{N}$ values increase with SOM stabilization (Kramer et al., 2003), hence, more stabilized SOM pools are likely enriched in ^{15}N . For both barley types the $\delta^{15}\text{N}$ values of roots were higher at the head-emergence stage indicating a higher N gain from more stabilized SOM sources and hence positive rhizosphere priming effects.

Plant phenology plays a major role for the magnitude of rhizosphere priming effects. The specific rhizosphere priming (RPE per root dry weight) was reduced for the barley wild type when comparing between the tillering and head-emergence growth stages. A reduction of RPE at later growth stages has also been reported for wheat after flowering (Cheng et al., 2003). Young plants translocate higher proportions of assimilated C belowground than older plants (reviewed by Nguyen, 2003). Root-derived CO₂ per unit of root mass was higher at tillering compared to head-emergence (yet not significant).

Moreover, the C/N ratio of roots was about twice as high at the head-emergence stage compared to the tillering and C/N ratios of shoots were even three to four times higher when comparing the

two sampling dates. Thus, the large N demand of the plant is likely to induce the positive priming effects measured at T2. At the head-emergence stage, when nutrients are becoming scarce, the barley mutant without root hairs probably suffered from the inefficiency in nutrient uptake as indicated by a lower root biomass compared to the wild type. However, in contrast to the specific RPE of the wild type, which was reduced between the sampling dates most likely because of reduced allocation of assimilates belowground (Nguyen, 2003), the root hairless mutant showed the opposed effect. Specific RPE increased between the tillering and head-emergence stages to positive values, and even exceeds the RPE of the wild type. The inefficiency of the hairless mutant in nutrient uptake may have increased rhizodeposition due to a faster decay of roots, induced by insufficient supply of nutrients.

Root hairs may contribute 70–90% to total root surface area (Bucher, 2007), and therefore root hairs are crucial for nutrient uptake of the plant. Especially the uptake of phosphorus, which is highly immobile in soils, is promoted by root hairs. Phosphorus is quantitatively the second most limiting nutrient for plant growth after N (Lambers et al., 2006). Gahoonia and Nielsen (2003) showed a much stronger P-depletion zone around the root hairs of a barley wild type than for the root hairless mutant. Under P limitation, higher phosphatase activity in the soil was shown to increase the transformation of organic phosphates into available forms (Gahoonia et al., 2001; Paterson, 2003; Olander and Vitousek, 2000). In our study, the activity of acid phosphatase at tillering is lower for the wild type but higher for the mutant without root hairs compared to the unplanted soil. This pattern diminished at the head-emergence stage where phosphatase activity did not differ between unplanted soil and barley with and without root hairs. These contradictory results are likely explained by the high P availability of our soil.

The activity of the C-cycling associated enzymes, β -glucosidase and β -cellobiohydrolase, responsible for the decomposition of relatively labile C molecules (simple sugars, starch, cellulose), was lower for the wild type compared to the unplanted soil and the root hairless mutant at T1. At T2, both barley types showed lower β -glucosidase activities compared to the unplanted soil (yet not significant for *brb*) indicating lower microbial investments in C-cycling enzymes (Allison et al., 2011). Planting increased the activity of β -xylosidase and chitinase at both growth stages. The decomposition of the hemicellulose xylan, a major constituent of litter, is catalyzed by β -xylosidase into more available carbohydrates (Sinsabaugh and Moorhead, 1994; Loepmann et al., 2016). The chitinase (N-acetyl-glucosaminidase) degrades chitin (unbranched polymer of N-acetyl-D-glucosamine), which is found in bacterial and fungal cells (Beier and Bertilsson, 2013). Chitin is an important source of organic N in soil, as it is one of the most abundant polymers on earth and contains about 6% N in relatively recalcitrant form (Ekenler and Tabatabai, 2002; Duo-Chuan, 2006; Kelly et al., 2011). In N-poor microsites, i.e. at low concentrations of mineral N, the production of N-acquiring exoenzymes such as amino-peptidases (e.g. LAP) and chitinases (e.g. N-acetyl-glucosaminidase) are stimulated to obtain more N from organic forms (Olander and Vitousek, 2000; Weintraub and Schimel, 2005; Kelly et al., 2011). When mineral N is becoming scarce, microbes decompose more labile forms of N-containing organic matter first because less energy is required (Kelly et al., 2011). Thereafter, according to microbial life strategy, several microbial guilds may shift their enzyme production (Schimel and Schaeffer, 2012) from enzymes responsible for degradation of relatively labile substrate (e.g. LAP, cleaving of peptide bonds in proteins) to enzymes decomposing relatively recalcitrant substrates (e.g. chitinase, hydrolysis of chitooligosaccharides into N-acetylglucosamine) to meet the metabolic N demand (Kelly et al., 2011). In this experiment, the LAP

activity was reduced for both barley types between T1 and T2 (statistically significant only for WT). In contrast, chitin is produced by microorganisms and chitinase activity was shown to be enhanced by the presence of the rhizosphere (Geisseler et al., 2010). Microbial biomass was higher at T2 than at T1 for the planted soils. Thus, a higher microbial growth and death led to the release of chitin into the soil, which induced the high chitinase activities of both barley types at T2. The shift from enzymes degrading labile substrates to enzymes that decompose more recalcitrant forms of N is a strong indication for priming effects on SOM decomposition.

5. Conclusions

The present study suggests that rhizosphere priming effects are intimately linked to root morphology, e.g. root hairs. While the barley wild type with root hairs induced positive priming during tillering (69% above unplanted soil), the mutant without root hairs suppressed SOM decomposition by 28%. At head-emergence, microbial biomass increased for both planted soils compared to the tillering stage; and both barley types showed positive priming effects (accelerated SOM decomposition). The SOM decomposition rate under the hairless mutant barley even exceeds that of the wild type despite lower root biomass (72% priming for the wild type, 209% priming for the mutant). In cases of positive priming, chitinase and β -xylosidase activities increased suggesting accelerated decomposition of stable SOM. Future research emphasis should be placed on potential mechanisms linking root morphology and microbial activity with rhizosphere priming effects.

Acknowledgements

This study was supported by grants from the German Research Foundation (PA 2377 1/1; FOR 918) and German Academic Exchange Service (DAAD) within a PPP program (57051794), and a grant from the US National Science Foundation (Grant No. 1354659). The seeds for the experiment were kindly supplied by Dr. Timothy George (Ecological Sciences, The James Hutton Institute, Dundee, Scotland UK). We would like to thank Jenna Merrilees, Susann Enzmann, and Karin Schmidt for their help during the setup and analyses of the experiment.

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