

Dear Editor and Reviewer,

We appreciate your support and the valuable comments which made the revision easier and improved the manuscript. Please find our answers below, point by point highlighted in red. Please find attached the manuscript after the answers in which the changes made according to the suggestions are also highlighted in red.

This paper summarizes findings of a study on total soil respiration and its components in a grassland over various moist and dry periods during one growing season. The authors used near-continuous automatic measurements of CO₂ fluxes and their isotopic composition to distinguish between sources of CO₂ in the soil. Those sources included respiration by heterotrophic microorganisms, and respiration by roots + rhizosphere organisms and by mycorrhizae that together compose autotrophic respiration. The results showed a decrease in the autotrophic contribution to R_s during drought, mainly because of a greatly reduced flux from roots + rhizosphere. The main issues raised in earlier reviews were addressed by the authors as effectively as possible (see points 1 and 2 below), yet some additional revisions still need to be performed.

1. Spatial replication and capture of spatial variation: Authors made an effort to rotate measurement chambers among different locations. Thus, they obtained a minimal set of 2-3 replications during each of the 5 selected periods. While this approach seems to make optimal use of their measurement capacities, there still remains a spatial mismatch between the larger undisturbed area dedicated to measurements of total R_s and the strip used for exclusion treatments.

The separated positioning of the tubes (at the edge of the undisturbed area) was chosen to avoid the soil disturbances in the undisturbed area. The experimental area was ~10m² and it was included in an about 10 x 10 m area where the vegetation and surface exposure were the same. Therefore, the tubes were not placed at the edge of a vegetation patch but at the artificial edge of the measured part. Supposedly, soil properties and functioning were similar within the experimental area.

In addition, the aim of the larger area no. 9 is unclear. These issues need to be clarified. One of the root exclusion tubes was larger (area 9 in Fig. S1, 60 cm in diameter) including one of the root- and mycorrhiza exclusion tubes. The aim of this larger root-excluded area was to study the effect of the distance from the mesh on the mycorrhizal filament length (not the aim of this study), but there were no significant differences in CO₂ efflux and mycorrhizal filament length between the measured position of this larger area and other tubes.

In general, the above-mentioned mismatch may not be fatal, considering the short distances between the control and the treated areas. Maybe more importantly, root and root + mycorrhizae exclusion involved major disturbance of the soil, which might be of greater consequences than the spatial issues (this points should be discussed, see below).

Setting up the experimental area in 2010 caused a large disturbance in soil structure, that is why we did not use directly the measured CO₂ fluxes of the tubes (R_{re}, R_{rme}), but used the isotopic signals to estimate their share in the soil CO₂ efflux of the undisturbed area (R_{soil}).

2. Calculated isotopic signatures and error propagation: The isotopic values and the error terms seem reasonable now, considering the large variation inherent in such measurements. However, the estimation of the isotopic signatures of autotrophic respiration and its components is not uniform. As to Fig. S4, $\delta^{13}\text{C}$ of R_{mycrhiz} and R_{rhizo} are -28.9 and -28.6‰, respectively. In Fig. 2 and p12 l19, the opposite was presented, with $\delta^{13}\text{C}$ of R_{mycrhiz} being higher than that of R_{rhizo}. The second version makes more sense, if $\delta^{13}\text{C}$ of

Rmyc is -27.2‰, because only in that way can the isotopic signature of Rmycrhiz be combined by those of Rhizo and Rmyc.

Yes, you are right, the caption of Fig. S4 was wrong, it is corrected now.

3. $\delta^{13}\text{C}$ of Reco is lower by about 1‰ than $\delta^{13}\text{C}$ of Rs, which indicates that $\delta^{13}\text{C}$ of plant shoot respiration is lower than $\delta^{13}\text{C}$ of Reco (as obvious also from autotrophic belowground respiration, calculated as Rmycrhiz). The isotopic signature of shoot respiration might be constrained by results from isotopic analysis of shoot biomass. Moreover, it seems that shoot respiration could be estimated in the same way as the autotrophic belowground components, which would add an important aboveground component to the paper.

Yes, we agree, $\delta^{13}\text{C}_{\text{Reco}}$ was the lowest among the measured isotopic signals, so it can be assumed that the isotopic signal of the above-ground respiration could be the most depleted $\delta^{13}\text{C}$. $\delta^{13}\text{C}$ value of the above-ground component could be estimated by plotting $\delta^{13}\text{C}_{\text{Reco}}$ values against the $\text{R}_{\text{soil}}/\text{R}_{\text{eco}}$ ratio. Unfortunately, R_{eco} measurements have poor data coverage during the study period, only 21% of the night time measurements provided usable values. This was due to the frequent still wind conditions during night time. While still wind conditions do not support R_{eco} measurements, the Keeling-plot approach works well at the same time due to the high concentration gradient. Therefore measurements by these two methods do not overlap well during this study.

We tried to use gapfilled R_{eco} for the estimation of the isotopic signature of the above-ground parts but the regression was not significant, therefore we could not use the obtained y-intercept value for further calculations. Without the isotopic signals of the above-ground respiration it was not possible to estimate its share in ecosystem respiration.

Drought induced decrease in GPP (or photosynthesis) was mentioned in the text, but no data were presented and or relevant paper was cited.

In our manuscript Fig. 1b contains daily minimum NEE values showing the changes in the photosynthesis of the vegetation. Periods of vegetation activity (active, drying, etc) were distinguished considering the changes in NEE (P12 L12).

Our former findings showed strong decline in GPP in response to drought in this grassland (Nagy et al., 2007), we referred to it in the introduction together with other papers and put it also into the discussion section according to your suggestions.

4. Mean respiration rates could be summarized in a table, for each period and for the entire season. This table could not only contain the measured, but also calculated absolute rates and their error term (using the relative data).

We put Table 2 into the manuscript containing these average values and their error terms and added a few lines to the results.

5. The Discussion unnecessarily repeats many results, and a large part of this section was dedicated to numerical comparisons with results from other publications. Only the most important results should be repeated at the beginning of the Discussion, and comparisons with findings of other studies may be noted in the margin. The bulk of this section should be dedicated to explaining the obtained results, suggesting mechanisms underlying the findings, outlining potential consequences, etc. While it shortly mentions such consequences, the main part of the conclusion again repeats the results.

Parts of the discussion section have been rewritten according to these suggestions and a few lines have been added to the section emphasizing the conclusions of the study.

6. Potential drawbacks of the methods and their consequences should also be mentioned. E.g.,

the great disturbance of the soil to exclude roots and mycorrhizae might be expected to affect decomposition rates of SOM and, thus, Rh. In addition, microbial activity was greatly reduced by root + mycorrhizae exclusion as compared with the undisturbed treatment. How might this fact affect Rh?

Our approach combined the root- and root- and mycorrhiza exclusion treatments with isotopic measurements. The aim of this combination was to assess the contributions by the heterotrophic and autotrophic components in soil CO_2 efflux of the undisturbed soil.

Indeed, the insertion of the tubes caused some mixing of soil layers (4 layers were separated during the excavation of the tubes and packed back layer by layer) and disintegration of soil aggregates, also affecting soil water content and temperature as compared to the undisturbed soil. Moreover, R_{het} could be greatly reduced due to the lack of roots and mycorrhizal filaments.

That is why the measured respiration rates of the root- and mycorrhiza excluded tubes were not used directly for calculations of the component contributions, but could be used to calculate the isotopic signal of the autotrophic and heterotrophic components.

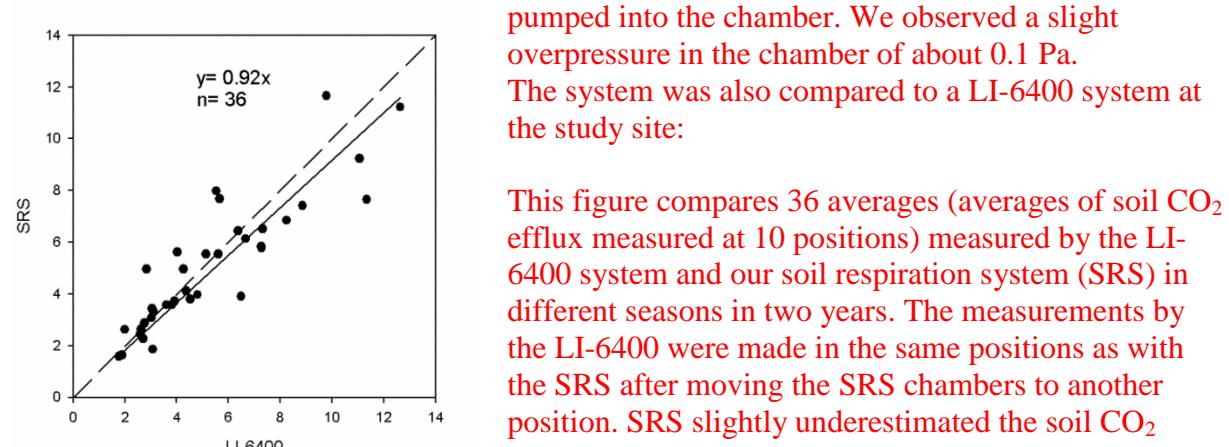
All of the estimated contributions to soil CO_2 efflux by rhizospheric, mycorrhizal fungi and heterotrophic components were applied for the undisturbed soil.

7. Automatic soil respiration system: More information is needed on the function of the little chambers, as request by other reviewers. Are the chambers automatically closed during measurements? What was the measurement frequency? How could it be guaranteed that soil enclosed by the chambers received an appropriate amount of rain through the holes? What were the temperature differences between the chambers and the environment? The flow rate of the air through the chamber was relatively high. The potential generation of a vacuum should be discussed, which might have forced air out of the ground. More importantly, because the chambers were very shallowly introduced into the ground (according to Nagy et al. 2011), such a vacuum could possibly suck air from the environment.

The soil respiration system was an open dynamic system, the chambers had no closing-opening mechanism. The main advantages of the system are the small chambers avoiding the regular cutting of the above-ground parts of the vegetation (chambers can be placed between the plants). The small size (radius of the chambers is 2.5 cm) also supports the balancing of any developing gradient of the driving variables (e.g. soil water content) between chamber inside and outside. Chamber walls exceeded the chamber top by 3 mm directing precipitation to the vent holes.

The system was calibrated against known fluxes on a calibration tank in the CzechGlobe Institute, Brno in 2010 ($F_{\text{SRS}} = 0.98 \times F_{\text{calibration_tank}}$, $R^2 = 0.92$, $n = 86$). During the measurements we used flow rates of about 300 ml min^{-1} and the reference air was taken from close to the soil (10 cm), therefore the air above the surface was pumped into the chamber. We observed a slight overpressure in the chamber of about 0.1 Pa.

The system was also compared to a LI-6400 system at the study site:



efflux compared to the LI-6400. 1:1 line is shown (dashed line).

8. The English language needs improvement. Most urgently, the title should be edited. The term “ratio” was used inappropriately at several locations in the text. It should be replaced by “fraction” or “percentage”, e.g. in the title of subchapter 3.3, at the beginning of the Discussion and in the y-axis title of Fig. 3.

Thanks for your remark, we corrected the terms.

The manuscript went through a professional English language editing we hope its English is satisfactory now.

9. Additional comments

P3, l19: This sentence is expected to appear towards the end of the Introduction, together with the study’s objectives. Additionally, reference to GPP as part of an objective raises expectations of data presentation, but no data on GPP were shown here.

P3, l24: The statement on the percentage of plant species associated with AMF should be referenced.

P5, l24: Rather Fig. S1.

P6, l18: Acronyms must to be explained at first appearance, here and elsewhere.

P7, l17: Please turn these short and repeating sentences into on coherent sentence.

P7, l23: It is unclear where reference air was sampled.

p14, l21: Neither GPP nor leaf photosynthesis rates were presented in this work. Therefore, a study relevant for this site needs to be cited.

Fig. 1: The meaning of the horizontal lines needs to be explained.

Fig. 2: All parts of the box plots should be explained.

Fig. 4: Please indicate the statistical analysis used for this figure.

Thanks for your remarks, these problems have been corrected in the text.

Autotrophic component of soil respiration is repressed by drought more than the heterotrophic one in dry grasslands

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Abstract

Summer droughts projected to increase in Central Europe due to climate changes strongly influence the carbon cycle of ecosystems. Persistent respiration activities during drought periods are responsible for a significant carbon loss, which may turn the ecosystem from a sink into a source of carbon. There are still gaps in our knowledge regarding the characteristic changes taking place in the respiration of the different components of the ecosystem in response to drought events.

In the present study, we combined a physical separation of soil respiration components with continuous measurements of soil CO₂ efflux and its isotopic (¹³C) signals at a dry grassland site in Hungary. The physical separation of soil respiration components was performed by means of inox meshes and tubes inserted into the soil. The root-excluded and root- and mycorrhiza excluded treatments served to measure the isotopic signals of the rhizospheric, mycorrhizal fungi and heterotrophic components, respectively.

In the dry grassland investigated in the study the three components of the soil CO₂ efflux decreased at different rates under drought conditions. During drought the contribution made by the heterotrophic components was the highest (54±8%; mean±SE). Rhizospheric component was the most sensitive to soil drying with its relative contribution to the total soil respiration dropping from 66±7% (non-stressed) to 35±17% (mean±SE) under drought conditions. According to our results the heterotrophic component of soil respiration is the major contributor to the respiration activities during drought events in the dry grassland ecosystem studied.

1 Introduction

Grassland ecosystems respond forcefully to drought events via substantial reduction of their primary production (GPP, Hoover et al., 2014; Parton et al., 2012; Reichstein et al., 2013). In contrast, below-ground respiration is not so strongly affected (van der Molen et al., 2011; Yang and Zhou, 2013) but tends to be reduced as well under drought (Balogh et al., 2011; Suseela and Dukes, 2013). Soil respiration is the second largest component of carbon cycling in grasslands and returns as much as 50–90% of annual GPP back to the atmosphere (Bahn et al., 2008). Thus, the magnitude of soil respiration can turn the carbon budget from a net sink into a net source in dry years (Nagy et al., 2007). **Here we address the question whether under drought this is primarily a function of autotrophic respiration declining along with the soil drying while heterotrophic respiration remains less affected.**

According to climate change scenarios the frequency of droughts is expected to increase in Central Europe (Prudhomme et al., 2014) where dry grassland ecosystems represent one of the major land use types. It is well known that there is a need for better mechanistic models to address the effects of climatic extremes on carbon fluxes (e.g. Blagodatsky and Smith, 2012). However, progress has so far been limited due to the high complexity of responses given by the different ecosystem respiration components to the climatic drivers.

Soil organic matter (SOM) and litter derived respiration is considered to belong to the heterotrophic soil respiration component (Moyano et al., 2009). Their decomposition is attributed mainly to soil bacteria and fungi and has about 50% share in the total soil respiration in dry grasslands (Bao et al., 2010; Chen et al., 2009; Gomez-Casanovas et al., 2012). On the other hand, some of the soil fungi relying on recent photosynthetic assimilates also contribute to the autotrophic respiration component. Arbuscular mycorrhizal fungi (AMF) are obligatory symbiont soil fungi, forming intimate mutualistic associations in 70–90% of the plant species in grasslands (**Hiiesalu et al., 2014**). About 10-20% of the assimilated C may be attributed to AMF in exchange for acquiring water and essential nutrients for plant productivity (van der Heijden et al., 2015). Therefore soil respiration includes components of an autotrophic-heterotrophic continuum from roots through the root-associated fungi (rhizospheric and mycorrhizal) to non-root-associated (heterotrophic) microbial components.

Belowground CO₂ production by the autotrophic and heterotrophic components show large diel and seasonal variability (Fassbinder et al., 2011; Moyes et al., 2010). The drivers behind

all this are not fully revealed and the role of soil microbes in the process is still not clear mainly due to the diversity of soil biota (Bardgett et al., 2008). Moreover, drivers of CO₂ production frequently interact with each other (Balogh et al., 2015; Vargas et al., 2010), hampering the partitioning of the total CO₂ efflux into components. Studies found a stronger effect of photosynthesis than that of temperature on root respiration (Gomez-Casanovas et al., 2012; Heinemeyer et al., 2012; Hopkins et al., 2013). Both autotrophic and heterotrophic components were shown to be sensitive to water shortages (Carbone et al., 2011; Moyano et al., 2013). The autotrophic component was found to be dominant over the heterotrophic one during drought periods in a Mediterranean woodland ecosystem (Casals et al., 2011) but we have limited information about grasslands of shallow rooted herb species regarding the dominant source of carbon during drought periods.

The widely used separation techniques (trenching and girdling) are not considered suitable for grasslands (Epron, 2009), thus the physical separation of the soil CO₂ efflux components via root exclusion is hardly feasible without seriously disturbing the soil structure and the root system. A viable option, however, is the use of stable isotopic signatures ($\delta^{13}\text{C}$) of soil respiration to estimate the relative contributions of the main components (Carbone et al., 2011; Hopkins et al., 2013). Although diel patterns in $\delta^{13}\text{C}$ may also be subject to biases in the measuring methods (Fassbinder et al., 2011; Midwood and Millard, 2011), seasonal changes are expected to reflect the changes in the contributions of source components rather than the changes in the isotopic signals of the component itself (Knohl et al., 2005). However, SOM $\delta^{13}\text{C}$ can also change during the year with fresh plant material being more depleted in ¹³C than the older SOM components (Bowling et al., 2002), therefore fresh litter may contribute to the decreasing $\delta^{13}\text{C}$ of the heterotrophic component. Drying of the surface layers can also modify $\delta^{13}\text{CO}_2$ since heterotrophic respiration could be restricted to the deeper layers of the soil (Moyes et al., 2010). Drying of the soil can also change the amount of CO₂ produced in the topsoil layer (Balogh et al., 2015) by allowing greater atmospheric invasion and thereby enriching soil air in ¹³C (Phillips and Nickerson, 2010). The disequilibrium between the measured isotopic composition and the isotopic composition of the respiratory source could be significant especially in tracer experiments (Gamnitzer et al., 2011) but it is assumed to be less pronounced in open chamber measurements due to the steady-state diffusion (Nickerson et al., 2013).

Uncertainties in estimating the contributions of soil respiration components could be reduced by a combination of different methodologies (Risk et al., 2012). The question we are asking

is: which of the investigated soil respiration components (autotrophic - including rhizospheric and mycorrhizal fungi - and heterotrophic components) of the dry grasslands dominates during drought? Our hypothesis was that autotrophic respiration would be reduced linearly with photosynthesis, whereas heterotrophic respiration might not be affected as strongly, resulting in a net loss of C from the soil carbon reservoir. In order to achieve our goals we used an experimental approach based on the physical separation of soil respiration components combined with measurements of soil CO₂ efflux and its isotopic (¹³C) signal.

2 Methods

2.1 Site description

The vegetation at the Bugac site (46.69° N, 19.6° E, 114 m above sea level) is a dry sandy grassland dominated by *Festuca pseudovina*, *Carex stenophylla* and *Cynodon dactylon* and it was under extensive grazing for 20 years prior to our study. Ten-year mean annual precipitation (2004-2013) was 575 mm and the mean annual temperature reached 10.4 °C. The soil is a chernozem type sandy soil with high organic carbon content (Balogh et al., 2015).

2.2 Spatial separation of soil CO₂ efflux components

In 2010 ten soil cores (160 mm in diameter and 800 mm in depth, one of them 600 mm in diameter) were excavated. The roots were removed and the root-free soil was packed back - layer by layer - into PVC tubes with the same dimensions. Four tubes were used to exclude both roots and mycorrhiza. Walls of another 6 tubes were partially removed and replaced by inox mesh (40 µm mesh size) to exclude roots while ensuring that the mycorrhiza filaments can grow into the tubes (Moyano et al., 2007). These root-free and root- and mycorrhiza-free soil cores were settled at a distance of 6 m from the eddy covariance tower to the south direction (Fig. S1.). The distance between the soil cores/tubes was 50 cm.

Soil CO₂ efflux and its isotopic signal were measured in plots:

- with undisturbed soil (various positions, 36 positions in total within a ~4 m² plot): total soil respiration, R_{soil}, δ¹³C_{Rsoil},
- without roots and arbuscular mycorrhizal fungi (4 spatial replications) = heterotrophic component only, R_{rme}, δ¹³C_{Rrme},

- with root-excluded soil (6 spatial replications) = without roots, but with arbuscular mycorrhizal fungi, R_{re} , $\delta^{13}C_{Re}$.

2.3 Gas exchange measuring systems

Three different gas exchange systems were used in our study: eddy-covariance system (EC), automated soil respiration measuring system (SRS) connected to an isotopic CO_2 analyser (cavity ring-down spectroscopy system, CRDS). The experimental area was in the EC footprint (Supplementary material Fig. S1) but the size of the EC flux footprint area was larger by several orders of magnitude than the area covered by the SRS. Care was taken during the establishment of the experiment to select a plot with the same average soil characteristics and vegetation cover as found in the EC footprint area. Hence, the net ecosystem exchange (NEE) and evapotranspiration (ET) estimates obtained in this way can be considered representative also for the small-scale SRS and isotope measurements.

Data from 15th May 2013 to 12th November 2013 (182 days) were analysed in the present study.

2.3.1 Eddy covariance setup

The EC system at the Bugac site measured the CO_2 and H_2O fluxes continuously from 2002. In dry years the grassland can turn into a net carbon source (Nagy et al., 2007) but the longer-term annual sums of NEE showed it to be a net sink, ranging from -171 to $+106\text{ g C m}^{-2}\text{ yr}^{-1}$ (Pintér et al., 2010) with a $-100\text{ g C m}^{-2}\text{ yr}^{-1}$ average.

The EC system consists of a CSAT3 sonic anemometer (Campbell Scientific, USA) and a Li-7500 (Licor Inc, USA) open-path infra-red gas analyser (IRGA), both connected to a CR5000 data logger (Campbell Scientific, USA) via an SDM (synchronous device for measurement) interface. Additional measurements used in this study were: air temperature and relative humidity (HMP35AC, Vaisala, Finland), precipitation (ARG 100 rain gauge, Campbell, UK), global radiation (dual pyranometer, Schenk, Austria), incoming and reflected photosynthetically active radiation (SKP215, Campbell, UK), volumetric soil moisture content (CS616, Campbell, UK) and soil temperature (105T, Campbell, UK). These measurements were performed as described by Nagy et al. (2007) and Pintér et al. (2010). Fluxes of sensible and latent heat and CO_2 were processed using an IDL program after Barcza et al. (2003) adopting the CarboEurope IP methodology. For a detailed description of data processing and gap-filling see Nagy et al. (2007) and Farkas et al. (2011).

2.3.2 Soil respiration system

The 10 chamber automated soil respiration system was set up in July 2011. The system is an open dynamic one, consisting of an SBA-4 infrared gas analyser (PPSystems, UK), pumps, flow meters (D6F-01A1-110, Omron Co., Japan), electro-magnetic valves, and PVC/metal soil chambers (Fig. S3). The chambers were 10.4 cm high with a diameter of 5 cm, covering a soil surface area of 19.6 cm². The flow rate through the chambers was 300 ml min⁻¹, replacing the air in the chamber in 40 seconds. The PVC chambers were enclosed in a white metal cylinder with 2 mm airspace in between to stabilize the chamber and to prevent warming by direct radiation. Four vent holes with a total area of 0.95 cm² were drilled on the top of the chambers. Vent holes also served to allow precipitation to drip into the chambers. Chamber walls exceeded the chamber top by 3 mm directing precipitation to the vent holes. The system caused minor disturbances in the soil structure and the spatial structure of the vegetation. It was applied without cutting the leaves/shoots of the plants, so it did not disturb transport processes taking place inside the plant stems and roots. It was suitable for continuous, long-term unattended measurements of soil CO₂ efflux and was also used in previous experiments (Balogh et al., 2015; Nagy et al., 2011). The soil respiration chambers contained no standing aboveground plant material. Before the study the system was tested on a calibration tank (CzechGlobe, Brno, Czech Republic) against known fluxes ($F_{SRS} = 0.98 \times F_{calibration_tank}$, $R^2 = 0.92$, $n = 86$) and it was also compared to a LI-6400 system at the study site ($F_{SRS} = 0.92 \times F_{LI6400}$, $R^2 = 0.92$, $n = 36$).

Other studies (Nickerson et al., 2013; Risk et al., 2011) also used this chamber size, arguing that these chambers could be placed between the plants in grasslands, while larger chambers might create a non-representative surface due to the cutting necessary for placing the chambers on the ground (Risk et al., 2011).

R_{soil} was measured by 6 SRS chambers, while R_{rme} and R_{re} were measured by 2-2 SRS chambers, respectively.

2.3.3 Isotopic (¹³CO₂) measurements

A Picarro G1101-i gas analyser (CRDS, Picarro Inc., CA, USA) was attached to the soil respiration system from May to November in 2013. This CRDS system measured the isotopic composition inside the chambers and in the reference air. Reference air was sampled 10 cm above the surface in the foliage of plants. The SRS sequentially measured each of the 10 chambers for 3 minutes. Every second chamber was additionally probed for isotopic signature

measurements by the CRDS (3 minutes), followed by reference air measurements for another 3 minutes. Thus, the isotopic measurements of 5 chambers took 30 minutes in a single cycle. The CRDS integration time was set at 10 seconds, thus the CRDS provided 18 measurement points per chamber per cycle. Although the system response of the CRDS was clearly slower than the response of the SRS, the 3-minute duration was long enough to obtain robust results. Since CRDS followed the 3-minute intervals of SRS measurements no additional grace time had to be considered for the isotopic measurements.

Although this sampling scheme provides very good temporal coverage (replication in time), it is not perfectly addressing spatial variability and hence the position of each of the chambers was moved 11 times to randomly selected locations during the study period (i.e., every 2–3 weeks) to obtain sequential spatial replications for each plot type (undisturbed, root-excluded, root- and mycorrhizal fungi excluded; see Supplementary material Figs S1 and S2). More precisely, $\delta^{13}\text{C}_{\text{Rsoil}}$ was measured by 3 chambers at 36 (3 chambers x 12 positions) randomly selected positions within the experimental area (undisturbed soil, Supplementary material Fig. S1). $\delta^{13}\text{C}_{\text{Rre}}$ was measured by 1 chamber which was moved to positions 1, 3, 5, 6, 8, 9 during the study period (Supplementary material Fig. S2). $\delta^{13}\text{C}_{\text{Rrme}}$ was measured by 1 chamber which was moved to positions 2, 4, 7, 10 during the study period (Supplementary material Fig. S2).

Since contributions by the different soil CO₂ efflux components were estimated for five different periods within the study period distinguished by NEE, SWC values and isotopic signals (see Results), data for each estimation originated from 2–3 spatial replications.

2.4 Data processing and modelling

Data processing and statistical analysis were done in R (R Core Team 2014). Before calculating daily averages of $\delta^{13}\text{C}$ values a filtering method was applied to each dataset. Out of each 180-second-long measurement on a certain chamber, the first 70 s (to measure a steady state signal) and the last 20 s were cut and the remaining values were used for further calculations. As reference and chamber air were measured sequentially reference values during chamber measurements were estimated by linear interpolation between the neighbouring reference sequences.

After the interpolation, $\delta^{13}\text{C}$ values of the soil CO₂ efflux were calculated using the isotopic mass balance approach in each plot:

$$\delta^{13}\text{C}_R = \frac{\delta^{13}\text{C}_{\text{out}} \times c_{\text{out}} - \delta^{13}\text{C}_{\text{in}} \times c_{\text{in}}}{c_{\text{out}} - c_{\text{in}}} \quad (1)$$

where $\delta^{13}\text{C}_{\text{out}}$ and $\delta^{13}\text{C}_{\text{in}}$ are the isotopic signature of the outgoing and incoming air of the chamber and c_{out} and c_{in} are the CO_2 concentrations of the outgoing and incoming air of the chamber, respectively.

$$\delta^{13}\text{C} = \frac{R_{\text{sample}}}{R_{\text{standard}}} - 1 \quad (2)$$

and R stands for the $^{13}\text{C}:^{12}\text{C}$ isotope ratio of the sample and the international VPDB standard (0.011182), respectively.

Individual measurements were filtered out by using a moving-window procedure if the investigated value (at the window center) was outside the range of the mean \pm median absolute deviation of the values in a 10 day moving window. This filtering procedure left an overall data availability of 68-70%. Daily averages were calculated using the remaining data. To determine the isotopic signature of the ecosystem respiration (R_{eco}), Keeling plots were constructed by plotting the night-time $\delta^{13}\text{C}$ values measured 10 cm over the surface against the inverse of the CO_2 concentration. The extrapolated y-intercept of the linear regression was used as $\delta^{13}\text{C}_{\text{Reco}}$ values.

Total soil CO_2 efflux was separated isotopically into its components. We defined the components following the terminology presented by Moyano et al. (2009):

Heterotrophic respiration= microbial respiration from litter and SOM decomposition.

Autotrophic respiration= mycorrhizospheric respiration including rhizospheric and mycorrhizal fungi components.

Rhizospheric respiration= respiration of roots and root-associated microorganisms in the rhizosphere, not including mycorrhizal fungi.

Two-source mixing models were used to estimate the fraction (a) of the rhizospheric and (b) mycorrhizospheric components based on the measured isotopic signals:

$$\delta^{13}\text{C}_{\text{Rsoil}} = a \times \delta^{13}\text{C}_{\text{Rrhizo}} + (1 - a) \times \delta^{13}\text{C}_{\text{Rre}} \quad (3)$$

$$\delta^{13}\text{C}_{\text{Rsoil}} = b \times \delta^{13}\text{C}_{\text{Rmycrhiz}} + (1 - b) \times \delta^{13}\text{C}_{\text{Rrme}} \quad (4)$$

where $\delta^{13}\text{C}_{\text{Rsoil}}$ is the $\delta^{13}\text{C}$ of the total soil CO_2 efflux, $\delta^{13}\text{C}_{\text{Rre}}$ is the $\delta^{13}\text{C}$ of the root-excluded soil, $\delta^{13}\text{C}_{\text{Rrme}}$ is the $\delta^{13}\text{C}$ of the root- and mycorrhiza excluded soil (heterotrophic respiration), a is the fraction of the rhizospheric component (R_{rhizo}) and b is the fraction of the mycorrhizospheric component (R_{mycrhiz}) to the total soil efflux. According to these equations $1-b$ represents the ratio of heterotrophic respiration component to the total soil efflux and $b-a$ represents the ratio of mycorrhizal fungi component.

$\delta^{13}\text{C}_{\text{Rrhizo}}$ value was estimated by plotting $\delta^{13}\text{C}_{\text{Rsoil}}$ values against the $R_{\text{re}}/R_{\text{soil}}$ ratio (Supplementary material Fig. S3b). Since $R_{\text{re}}/R_{\text{soil}}$ is hypothetically zero when only

rhizospheric respiration is present, y-intercept of the linear regression was assumed as $\delta^{13}\text{C}_{\text{Rrhizo}}$. $\delta^{13}\text{C}_{\text{Rmycrhiz}}$ was estimated using the same approach (Supplementary material Fig. S3a), $\delta^{13}\text{C}_{\text{Rsoil}}$ values were plotted against the $\text{R}_{\text{rme}}/\text{R}_{\text{soil}}$ ratio and y-intercept of the linear regression was assumed as $\delta^{13}\text{C}_{\text{Rmycrhiz}}$. Similarly, $\delta^{13}\text{C}_{\text{Rre}}$ values were plotted against the $\text{R}_{\text{rme}}/\text{R}_{\text{re}}$ ratio and y-intercept of the linear regression was assumed as $\delta^{13}\text{C}_{\text{Rmyc}}$ (Supplementary material Fig. S4c) but this value was not used in further calculations.

Contributions of rhizospheric, mycorrhizal fungi and heterotrophic respirations to total soil respiration were calculated by the mixing models applied on subsets (periods) of the dataset of the total study period. Estimated values of rhizospheric (R_{rhizo}), mycorrhizal fungi (R_{myc}) and heterotrophic respiration (R_{het}) were calculated by multiplying the measured R_{soil} rates (total soil respiration) with the estimated fractional contributions (F) of each component as follows:

$$\text{R}_{\text{het}} = \text{R}_{\text{soil}} \times F_{\text{het}} \quad (5)$$

$$\text{R}_{\text{rhizo}} = \text{R}_{\text{soil}} \times F_{\text{rhizo}} \quad (6)$$

$$\text{R}_{\text{myc}} = \text{R}_{\text{soil}} \times F_{\text{myc}} \quad (7)$$

where, F_{het} , F_{rhizo} and F_{myc} are the fraction of the heterotrophic, rhizospheric and mycorrhizal respiration in total soil respiration, respectively.

2.5 Microbial investigations

Soil samples for the microbial investigations were taken after the gas exchange measurements in May 2014 to avoid the disturbance of the measurements by sampling the soil. Sampling date was chosen considering the maximum of the carbon sequestration capacity of the investigated grassland (Nagy et al., 2007). 5-5 samples were taken from 5 soil layers (0-10 cm, 10-20 cm, 20-30 cm, 30-40 cm and 40-50 cm) in each plot.

Determination of AM fungal hyphal length in the soil was based on the methods of Bååth and Söderström (1979) using separation by wet-sieving and centrifugation. The separated fungal hyphae were stained using agar solution (0.75%) containing trypan blue (0.05%) then dried for 24 h at 70°C. The hyphal length was measured in the dried agar film by the intersection method (Tennant, 1975) under a binocular microscope.

The fluorescein diacetate (FDA) hydrolysis assay was used to estimate the total microbial activity in soil samples and expressed as mg fluorescein released kg⁻¹ dry soil (Adam and Duncan, 2001).

2.6 Uncertainty assessment

Isotopic signals of soil respired CO_2 were studied extensively but several uncertainties related to the different methods were also revealed. Steady-state methods were found to provide more robust estimates than static chambers but still charged with biases (e.g. diffusive fractionation, Nickerson and Risk 2009). Open systems have the advantage of unattended automatic measurement collecting large amount of data but are less sensitive to small isotopic differences (Midwood and Millard, 2011).

In our study $\delta^{13}\text{C}_{\text{Reco}}$ estimates were independent of chamber related biases, using night-time $\delta^{13}\text{CO}_2$ and CO_2 concentration data of the free air over the surface for the calculation (Keeling-plot approach). This approach gave similar results to the chamber-based measurements, providing also partial verification of the latter ones. Moreover, isotopic measurements were independent on soil CO_2 efflux measurements, since IRGA and CRDS systems took different air samples from the same soil chambers. Isotopic data together with CO_2 efflux rates were collected during 1980 measurement cycles on 182 days in order to have robust estimates of isotopic signals.

A C4 grass (*Cynodon dactylon*) was also present in the study site potentially modifying the $\delta^{13}\text{C}$ of the respired CO_2 . Its cover was about 10% in the pasture (Koncz et al., 2014) but it was less frequent (i.e. less than 5%) in the experimental area. Calculated uncertainties of the relative contributions of each components (rhizospheric, mycorrhizal fungi and heterotrophic) contain the uncertainty due to a possible 5% contribution by the C4 grass. The isotopic signal of CO_2 efflux by the C4 plant was supposed to be -14‰.

In order to estimate the uncertainty of the measurements and estimated contributions by the different components to the total soil respiration random errors of each factor (CO_2 concentrations, isotopic compositions, model fit errors and possible C4 contribution) were propagated by Gaussian error propagation (Lo, 2005).

3 Results

3.1 Meteorological conditions, NEE, ET, soil CO_2 efflux, $\delta^{13}\text{C}$ of CO_2 efflux

The end of May and the beginning of June was the most productive period in the year due to ample water availability with the lowest NEE (strongest carbon sink activity) and highest evapotranspiration (ET) values being measured in this period (Fig. 1a). It rained only a few times from the end of June to 19th August (total precip: 10 mm) and the accompanying high temperature resulted in drought. Daily minimum NEE was around zero at the end of July and

in August. Rain events after the drought period had significant effects on soil CO₂ effluxes (Fig. 1c). There was a second active period following autumn rains but CO₂ uptake and ET were smaller than in May or June.

R_{soil} was the highest among the soil CO₂ effluxes, while R_{rme} was the lowest, the average CO₂ effluxes in the whole study period were 5.0±2.1, 3.8±1.6 and 2.6±1.2 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ (mean±SD) in R_{soil}, R_{re} and R_{rme}, respectively (Table 2). R_{re} was sometimes higher than R_{soil}, especially shortly after rain events. The lowest daily average total soil CO₂ efflux was measured in 15th August (2.22 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), while the lowest daily average R_{re} and R_{rme} values were observed in 2nd October (1.25 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$) and 2nd November (1.04 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), respectively. The highest values of soil CO₂ effluxes were measured in May in all treatments (R_{soil}, R_{re} and R_{rme}). Sudden increases in R_{re} and R_{rme} were observed shortly after rain events but R_{soil} showed slower (but more persistent) response to precipitation (Fig. 1c).

Isotopic signature of R_{eco} was the lowest in May and June, increased in July and August and decreased again in October and November (Fig. 1e). $\delta^{13}\text{C}_{\text{Reco}}$ showed clear responses to precipitation pulses with sudden declines being observed during the rain events. Chamber-based $\delta^{13}\text{C}_{\text{Rsoil}}$ showed similar changes during the study period. $\delta^{13}\text{C}_{\text{Rrme}}$ and $\delta^{13}\text{C}_{\text{Rre}}$ showed large scatter during the whole study period with no clear and detectable trends (Fig. 1d). Differences between $\delta^{13}\text{C}_{\text{Rsoil}}$ and $\delta^{13}\text{C}_{\text{Rrme}}$ were the largest in the active period and the smallest under drought conditions.

According to the NEE, SWC values and isotopic signals we distinguished 5 periods within the study period: an active period from 15th May to 20th June, a drying (stress development) period from 21st June to 22nd July, a drought period from 23rd July to 19th August, a wetting (stress release) period from 20th August to 16th September and a re-greening (recovery) period from 17th September to the end of the study period (11th November) (Fig. 1).

3.2 $\delta^{13}\text{C}$ of the respiration components

Fig 2. shows the measured and estimated $\delta^{13}\text{C}$ values of the different soil CO₂ efflux components. $\delta^{13}\text{C}_{\text{Rrme}}$ was the highest, while $\delta^{13}\text{C}_{\text{Rsoil}}$ was the lowest, suggesting that it was the rhizospheric respiration that was the most substantially depleted, while heterotrophic respiration was the least depleted in ¹³C. Mean values of $\delta^{13}\text{C}_{\text{Reco}}$, $\delta^{13}\text{C}_{\text{Rsoil}}$, $\delta^{13}\text{C}_{\text{Rre}}$ and $\delta^{13}\text{C}_{\text{Rrme}}$ were -27.9±0.5‰, -26.8±1.3‰, -26.4±1.8‰ and -25.7±2‰ (mean±SE), respectively. The estimated isotopic signals of the respiration of mycorrhizospheric

($\delta^{13}\text{C}_{\text{Rmycrhiz}}$), rhizospheric ($\delta^{13}\text{C}_{\text{Rrhizo}}$) and mycorrhizal fungi components ($\delta^{13}\text{C}_{\text{Rmyc}}$) were $-28.6 \pm 1.6\text{‰}$, $-28.9 \pm 1.7\text{‰}$ and $-27.2 \pm 2.3\text{‰}$ (estimate \pm SE), respectively (Fig. 2).

36% of the variation in $\delta^{13}\text{C}_{\text{Rsoil}}$ was explained by SWC ($\delta^{13}\text{C}_{\text{Rsoil}} = -0.1267 \times \text{SWC} - 25.537$, $R^2 = 0.36$, $P < 0.0001$), while only 3% of the variation of $\delta^{13}\text{C}_{\text{Rrme}}$ was explained by SWC and there was no correlation between $\delta^{13}\text{C}_{\text{Rre}}$ and SWC. Similar results were obtained between T_s and the isotopic signals but the correlation was weaker ($\delta^{13}\text{C}_{\text{Rsoil}} = 0.1056 \times T_s - 28.588$, $R^2 = 0.11$, $P < 0.0001$). Daily minimum NEE (NEEmin, Fig. 1b) explained 29% of the variation in $\delta^{13}\text{C}_{\text{Rsoil}}$ ($\delta^{13}\text{C}_{\text{Rsoil}} = 0.0941 \times \text{NEEmin} - 26.245$, $R^2 = 0.29$, $P < 0.0001$) but no correlation was found between NEEmin and $\delta^{13}\text{C}_{\text{Rrme}}$ and between NEEmin and $\delta^{13}\text{C}_{\text{Rre}}$.

3.3 Fraction of the different components in total soil respiration during the vegetation period

Two end-member mixing models (eq. 3 and 4) were used to estimate the relative contributions of rhizospheric, mycorrhizal fungi and heterotrophic components to total soil respiration during the study period. The estimated contributions by the different components were $50 \pm 6\%$, $13 \pm 8\%$ and $37 \pm 6\%$ (mean \pm SE) for the rhizospheric, mycorrhizal fungi and heterotrophic components, respectively. The autotrophic component (mycorrhizospheric component) of soil respiration showed significant decrease during the drying and drought periods. Rhizospheric component was the most sensitive to drying and drought. Average contributions by the rhizospheric component to total soil CO_2 efflux decreased from $66 \pm 7\%$ (mean \pm SE) in the active period to $35 \pm 13\%$ during the drought period (Fig. 3). After drought rhizospheric contributions increased again and become dominant during the re-greening period in autumn $63 \pm 7\%$ (mean \pm SE). During the transient (drying and wetting) periods the rhizospheric contributions to the total soil CO_2 efflux were $38 \pm 11\%$ and $46 \pm 8\%$, respectively. Relative mycorrhizal contributions were between 8-21% during the whole study period, with the highest contribution ($21 \pm 11\%$; mean \pm SE) during the wetting period. Heterotrophic contributions to soil respiration were the lowest in the active period ($21 \pm 7\%$) and the highest under drought ($54 \pm 13\%$) (Fig. 3).

Changes in soil CO_2 effluxes showed similar responses to drying and drought conditions as isotopic signals. Average R_{soil} decreased by 60% (referenced to the average during the active period) as a response to drought, while R_{re} and R_{rme} showed declines of 56 and 52% respectively, suggesting that declines in root respiration were substantially larger than those in R_{soil} (60%).

The estimated rates of rhizospheric, mycorrhizal fungi and heterotrophic components (eq. 5-7) are shown in Table 2. Pearson correlation coefficients pairing the estimated respiration rates and their possible driving variables (NEE, Ts, SWC) showed significant negative correlation between R_{rhiz} and NEE ($R = -0.94$, $p < 0.05$) and a significant positive correlation between R_{rhiz} and SWC ($R = 0.82$, $p < 0.05$). R_{het} changed with Ts but the correlation was not significant.

3.4 Microbial biomass and activity

Hyphal length (on dry soil weight basis) was significantly lower in the upper layers of root- and mycorrhiza excluded soil than in undisturbed soil, while it was significantly higher in root-excluded plots at 10-20 cm depth. Hyphal length in the root-excluded soil was similar to undisturbed soil in the other soil layers. Fluorescein values were significantly lower in all soil layers in the root- and mycorrhiza excluded plots than in the undisturbed soil. Fluorescein values in the root-excluded plots were also lower than in undisturbed soil but this difference was not significant (Fig. 4).

4 Discussion

Our approach combined the root- and root- and mycorrhiza exclusion treatments with isotopic measurements. The aim of this combination was to assess the contributions made by the heterotrophic and autotrophic components in soil CO_2 efflux of the undisturbed soil. Although the root- and mycorrhiza exclusion caused large disturbances in soil structure by inserting the tubes into the soil, we used these treatments only for identifying the isotopic signals of the investigated components. All of the estimated contributions to soil CO_2 efflux by rhizospheric, mycorrhizal fungi and heterotrophic components were applied for the undisturbed soil.

4.1 Estimated contributions made by the different components to the total soil CO_2 efflux and effect of drought on CO_2 effluxes and $\delta^{13}\text{C}$ values

While the percentages of the autotrophic component in the total soil CO_2 efflux were $63 \pm 6\%$ on average (rhizospheric and mycorrhizal fungi components, $50 \pm 6\%$ and $13 \pm 8\%$, respectively) being much higher than the average percentage of the heterotrophic ($37 \pm 6\%$) component, the contributions by the different components showed significant changes during the growing season. In other studies conducted in grassland ecosystems the estimated yearly

average ratio of the autotrophic component was found to be lower accounting for 38-52% of the total soil respiration (Bao et al., 2010; Heinemeyer et al., 2012), while reaching 74% during the growing season in a prairie grassland (Gomez-Casanovas et al., 2012) and 60-74% in an arid perennial grassland (Carbone et al., 2008). Our study was conducted from May to the beginning of November, therefore we can assume considering the lower vegetation activity in the dormant season (Nagy et al., 2007) that the contribution of the autotrophic component could be lower while that of the heterotrophic component higher for the whole year than the estimations for the growing season.

Soil CO₂ effluxes decreased in all treatments (R_{soil} , R_{re} , R_{rme}) under dry conditions, the largest decline being observed in total soil respiration (R_{soil}), therefore a strong response of the autotrophic component to drought could be assumed. The measured isotopic signals also showed decreasing autotrophic contributions to CO₂ efflux during soil drying. $\delta^{13}\text{C}_{\text{Rsoil}}$ showed negative responses to SWC and was more enriched when SWC was low, while $\delta^{13}\text{C}$ of the root- and mycorrhiza-excluded respiration (R_{rme}) showed no response. Since $\delta^{13}\text{C}_{\text{Reco}}$ was the lowest of the measured isotopic signals it can be assumed that the isotopic signals of the above-ground respiration could be the most depleted $\delta^{13}\text{C}$. Therefore, the observed increase in $\delta^{13}\text{C}_{\text{Reco}}$ and $\delta^{13}\text{C}_{\text{Rsoil}}$ values during the drying period and during the drought also showed the decline of both the above- and below-ground autotrophic components. The same phenomenon was shown by the modelling results with the smallest contribution made by the rhizospheric component estimated for the drought period (35±13%; mean±SE), while the highest for the active period (66±7%; mean±SE). Fractions of the heterotrophic respiration were the highest during drought (54±13% mean±SE) and the mycorrhizal fungi respiration showed only a small decrease during drought compared to the active period (from 13±10%; to 11±18%;), suggesting that the non root-associated microbes and mycorrhizal filaments were less sensitive to water shortages than the rhizosphere. Soil aggregates are expected to provide micro-habitats for soil organisms that should be moist enough for those organisms to thrive even under drought (Davidson et al., 2012). Since there was an absence in plant photosynthetic supply during drought period, mycorrhizal fungi component is expected to use stored carbon for respiration (van der Heijden et al., 2008).

Low $\delta^{13}\text{C}_{\text{Rsoil}}$ and $\delta^{13}\text{C}_{\text{Reco}}$ values were measured in the wetting and re-greening periods due to the drought-induced fall of the fresh litter to the surface as fresh plant material could be more depleted than the old litter (Bowling et al., 2002). The declines in $\delta^{13}\text{C}_{\text{Rsoil}}$ and $\delta^{13}\text{C}_{\text{Reco}}$ immediately after the rain events during drying and drought periods could also be explained

by the wetting of the litter layer, exposing relatively fresh substrate to degradation for short periods. This phenomenon could also cause an overestimation in contributions made by the depleted components (rhizospheric) during rain events. Since the rhizospheric contribution estimated for the re-greening period was high it is assumed that this result was obtained partly due to the increased amount of fresh litter. Similar results were obtained in a tallgrass prairie by Gomez-Casanovas et al. (2012), where the autotrophic components were more sensitive to soil drying than the heterotrophic ones. In contrast, Carbone et al. (2008) found more sensitive response by the heterotrophic component in an arid (<150 mm annual precipitation) perennial grassland. Fractions of autotrophic components were reported to increase in response to drought in a woodland ecosystem, supposing that the signature of the recent photosynthetic supply became enriched during drought and that could also explain the increase in the soil respired CO₂ (Casals et al., 2011). A drought induced increase in $\delta^{13}\text{C}$ of root respiration of trees was also assumed in a recent study (Risk et al., 2012), suggesting that the isotopic signals of the assimilates, thereby the signals of the autotrophic component might also increase. In our study, $R_{\text{rme}}/R_{\text{soil}}$ showed significant positive correlation with $\delta^{13}\text{C}_{\text{Rsoil}}$ (the regression was used to estimate $\delta^{13}\text{C}_{\text{Rmycrhiz}}$, Supplementary material Fig S4), so $\delta^{13}\text{C}_{\text{Rsoil}}$ was high if the fraction of heterotrophic CO₂ efflux to the total soil CO₂ efflux was found to be high. Moreover, NEEmin values were close to zero during drought (average daily minimum NEE was $-0.91 \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$) showing the lack of the photosynthetic supply in this period. Photosynthetic CO₂ uptake of this vegetation was found to be sensitive to drought conditions (Nagy et al., 2007) and it can act as a driver of the soil CO₂ production and efflux (Balogh et al., 2015). The observed strong correlation between the estimated rhizospheric respiration and NEE can also be explained by the interacting effects of drought and photosynthetic supply of respiration. These findings support that in the grasslands under study the autotrophic respiration component was more sensitive to soil drying and its activity determined the isotopic signals of the total soil respiration during the study period.

According to these studies and to our results we can assume that the different vegetation types may respond differently to drought: woodlands may increase their autotrophic contribution while grasslands may decrease it (Casals et al., 2011; Gomez-Casanovas et al., 2012; Risk et al., 2012). Plants with different rooting habits have different water availability during dry periods (van der Molen et al., 2011), which could explain the differences between the different ecosystems in their response to drought.

4.2 Measured and estimated isotopic signals of the soil respiration components

Measured and calculated $\delta^{13}\text{C}$ values of the different respiration components showed differences similar to the ones reviewed by Bowling et al. (2008). $\delta^{13}\text{C}_{\text{Reco}}$ (containing also the signal from above ground green biomass) was the most depleted, while $\delta^{13}\text{C}_{\text{Rrme}}$ (heterotrophic components only) was the least depleted. $\delta^{13}\text{C}$ of the root- and mycorrhiza excluded respiration was similar to SOM $\delta^{13}\text{C}$ measured in a previous study (Denef et al., 2013): -25‰ and -26‰ in the topsoil layers (without the litter layer). CO_2 effluxes from mycorrhizal fungi were expected to be more enriched in ^{13}C relative to the total soil respiration (about +3‰, Bowling et al. 2008). Estimated $\delta^{13}\text{C}$ of mycorrhizal fungi component was $-27.2 \pm 2.3\text{‰}$ (estimate \pm SE), which is 1.7‰ higher than the rhizospheric component ($-28.9 \pm 1.7\text{‰}$; estimate \pm SE).

In our study neither $\delta^{13}\text{C}_{\text{Rrme}}$ values (heterotrophic respiration), nor $\delta^{13}\text{C}_{\text{Rre}}$ values (heterotrophic+mycorrhizal fungi respiration) showed correlation with SWC but $\delta^{13}\text{C}_{\text{Rsoil}}$ (total soil respiration) showed significant negative correlation with SWC. We can assume that $\delta^{13}\text{C}$ of heterotrophic respiration was not influenced by SWC changes during the growing season as it was found also by other studies (Phillips and Nickerson, 2010; Risk et al., 2012). Furthermore, the lack of correlation with the present study also suggests that soil moisture induced changes in diffusivity (disequilibrium effect due to changing soil moisture) were not large enough to affect the measured $\delta^{13}\text{C}$ values.

4.3 Microbial investigations

High hyphal density was maintained in R_{re} plots and low but still significant microbial activities (SOM decomposition) were detected in R_{rme} plots, therefore the measured $\delta^{13}\text{C}$ values characterized the sources of the root-free ($\delta^{13}\text{C}_{\text{Rre}}$) and root- and mycorrhiza-free ($\delta^{13}\text{C}_{\text{Rrme}}$) soils. The fact that very high amounts of hyphae were found in the root-excluded soil in the 10-20 cm layer proved that mycorrhizal fungi filaments were able to penetrate through the inox mesh and supported significant microbial activity. Grasses have extensive fibrous root systems with moderate to high levels of mycorrhizal colonization (van der Heijden et al., 2015). The range of AM hyphal lengths found in this study (1.9–8.8 m g⁻¹ soil) were similar to those reported in the literature (e.g. Mumme and Rillig 2008). The higher hyphal densities found in root-free soil might have been related to the higher availability of SOM-derived nutrients and to more space without the roots (i.e. lack of competition).

According to our results significant amount of CO₂ was respired from mycorrhizal filaments in the undisturbed soil, having a 12-31% share in the respiration carried out by the autotrophic component.

Values of fluorescein in root-excluded plots were similar to those measured in the undisturbed soil probably because hyphae of AM fungi provide an increased area for interaction with other microorganisms (hyphosphere, Andrade et al. 1997), but were much lower in root- and mycorrhiza excluded soil. These results support the component estimations showing the significant activities of root-associated microorganisms.

5 Conclusions

In the dry grasslands investigated in our study all three components of the soil CO₂ effluxes decreased, following different dynamics under drought conditions. Both the measured CO₂ effluxes and the isotopic signals showed similar results regarding the component responses. The strongest decrease in response to drought was seen in rhizospheric respiration (relative contribution to the total respiration decreased from 66±7% to 35±13%; mean±SE), while the relative contribution to the total soil respiration by the heterotrophic components increased during soil drying. During drought the contribution of the heterotrophic component was found to be the highest (54±8%; mean±SE). Mycorrhizal fungi respiration had its highest share in soil respiration (21±11%; mean±SE) in the wetting period after drought. According to these results the autotrophic component of the soil respiration is more sensitive to drought than the heterotrophic one in the dry grassland ecosystem studied. Thus, carbon source activities during drought periods identified by NEE measurements originated from carbon sources already stored, thereby decreasing the carbon content of the soil.

Drought events are expected to be more frequent in Central Europe in the future, and it is expected that the productivity of grassland ecosystems may strongly respond to projected dryness, influencing the carbon cycle of the ecosystems. Since potential productivity is generally linked to soil carbon content a pronounced decrease in soil organic matter due to the enhanced activity of the heterotrophic component under drought may directly affect the long term productivity of grasslands.

Author contributions

J. Balogh, M. Papp, K. Pintér and Z. Nagy conceived and designed the experiment, M. Papp, K. Pintér and K. Posta performed the experiment, J. Balogh, Sz, Fóti, W. Eugster and Z. Nagy analyzed the data and wrote the paper, but all co-authors contributed to writing.

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Table 1 Measured and estimated CO₂ effluxes and isotopic signals in this study

	CO ₂ efflux	Isotopic signals
measured	R _{eco} , R _{soil} , R _{re} , R _{rme}	δ ¹³ C _{Reco} , δ ¹³ C _{Rsoil} , δ ¹³ C _{Rre} , δ ¹³ C _{Rrme}
estimated	R _{rhizo} , R _{myc} , R _{het}	δ ¹³ C _{Rmychorhiz} , δ ¹³ C _{Rrhizo} , δ ¹³ C _{Rmyc}

Table 2 Mean measured (undisturbed soil and tubes) and estimated respiration rates for the different periods ($\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$) with propagated uncertainties.

period	measured (mean \pm SD)			estimated (mean \pm SE)		
	R_{soil}	R_{re}	R_{rme}	R_{het}	R_{myc}	R_{rhizo}
active	7.7 \pm 1.6	5.1 \pm 1.5	3.9 \pm 1.1	1.7 \pm 1.1	1.0 \pm 1.2	5.1 \pm 1.1
drying	5.7 \pm 2.0	3.8 \pm 1.5	2.9 \pm 0.6	2.9 \pm 0.7	0.6 \pm 0.9	2.1 \pm 0.8
drought	3.2 \pm 1.1	2.3 \pm 0.4	1.9 \pm 0.4	1.7 \pm 0.5	0.3 \pm 0.6	1.1 \pm 0.5
wetting	4.8 \pm 1.7	4.3 \pm 1.5	2.6 \pm 1.2	1.5 \pm 0.6	1.0 \pm 0.7	2.3 \pm 0.7
re-greening	3.8 \pm 1.0	2.4 \pm 0.8	2.0 \pm 1.1	1.1 \pm 0.4	0.3 \pm 0.5	2.4 \pm 0.4
total study period	5.0 \pm 2.1	3.8 \pm 1.6	2.6 \pm 1.2	1.8 \pm 0.6	0.7 \pm 0.8	2.6 \pm 0.7

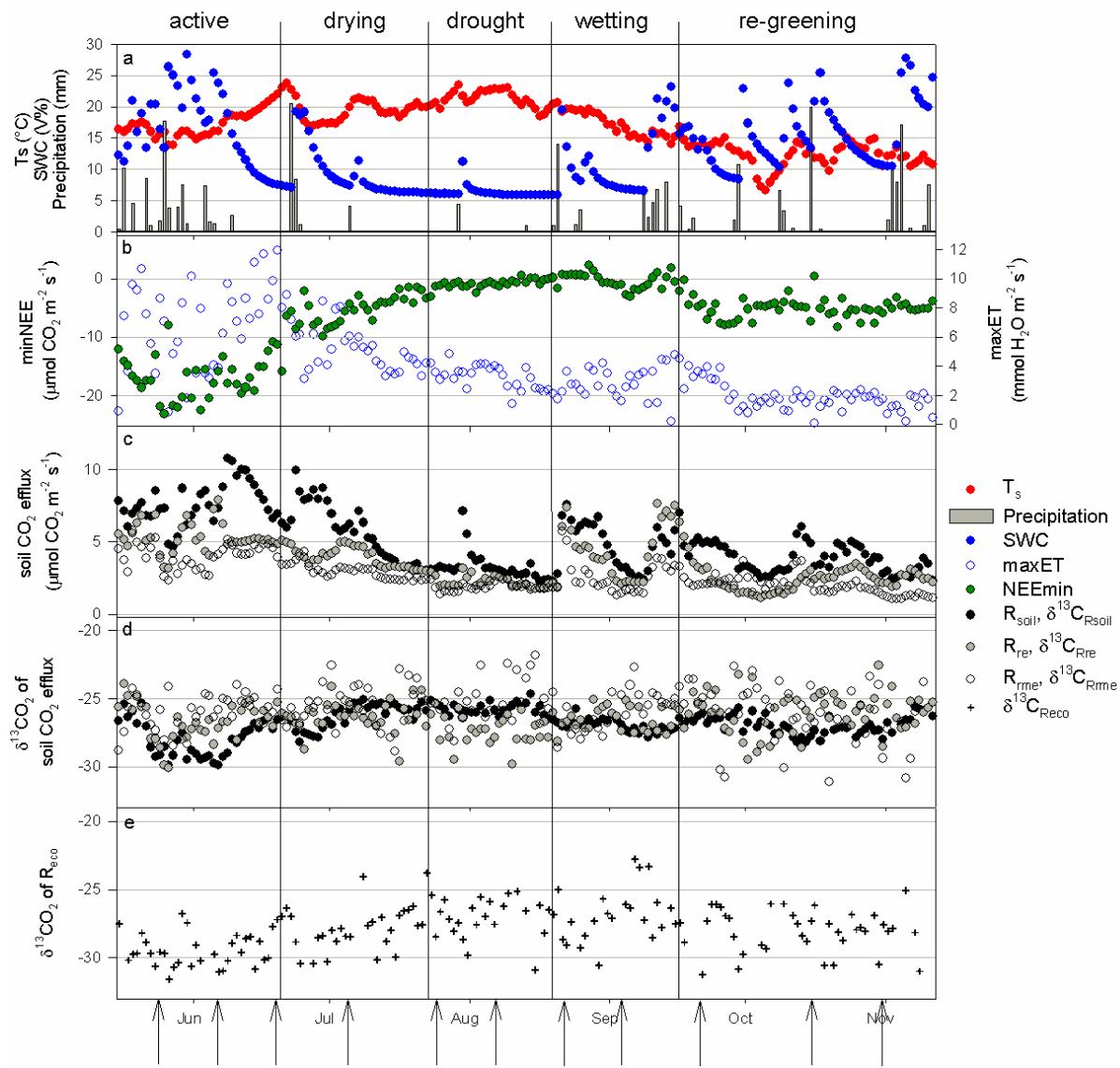


Fig. 1 (a) Daily averages of soil temperature (T_s), soil water content (SWC) at 5 cm depth and daily sum of precipitation, (b) daily minimum half-hourly NEE and maximum half-hourly ET, (c) daily averages of CO_2 efflux in undisturbed soil (R_{soil}), root-excluded soil (R_{re}) and root- and mycorrhizal fungi excluded soil (R_{rme}), (d) daily averages of $\delta^{13}\text{C}$ of soil CO_2 efflux in undisturbed soil ($\delta^{13}\text{C}_{\text{Rsoil}}$), root-excluded soil ($\delta^{13}\text{C}_{\text{Rre}}$) and root- and mycorrhizal fungi excluded soil ($\delta^{13}\text{C}_{\text{Rrme}}$) and (e) daily averages of $\delta^{13}\text{C}$ of ecosystem respiration ($\delta^{13}\text{C}_{\text{Reco}}$) during the study period in 2013, at Bugac site. Arrows indicate the positions changes of the soil chambers. **Gray horizontal lines** show Y major values.

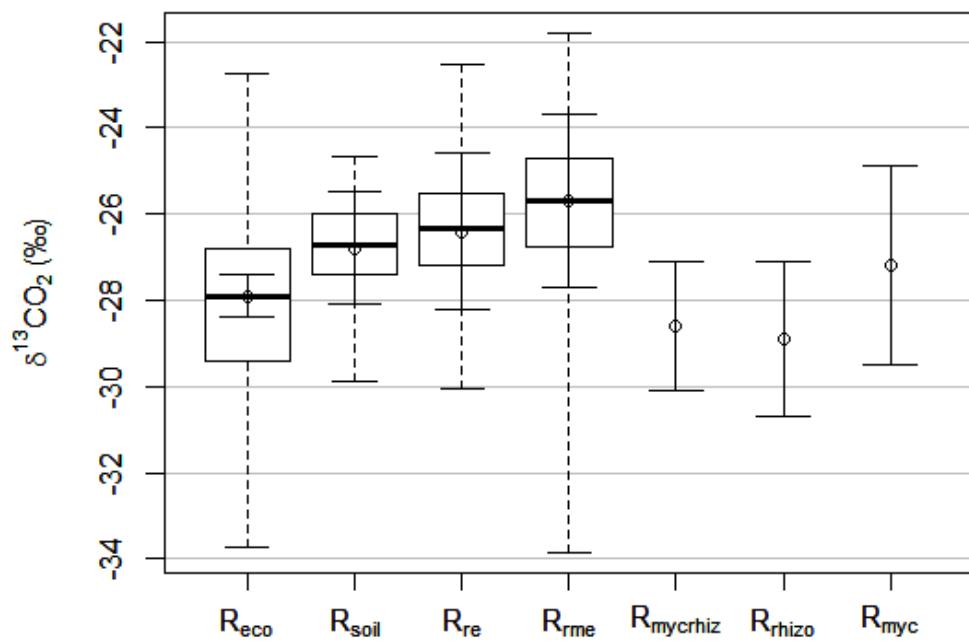


Fig. 2 Measured (R_{eco} , R_{soil} , R_{re} , R_{rme}) and estimated (R_{mycrhiz} , R_{rhizo} , R_{myc}) $\delta^{13}\text{C}$ values of the respiration components. Horizontal black lines in boxes show medians and dashed whiskers show data extremes. Open circles and solid whiskers show means \pm propagated standard errors. Gray horizontal lines show Y major values.

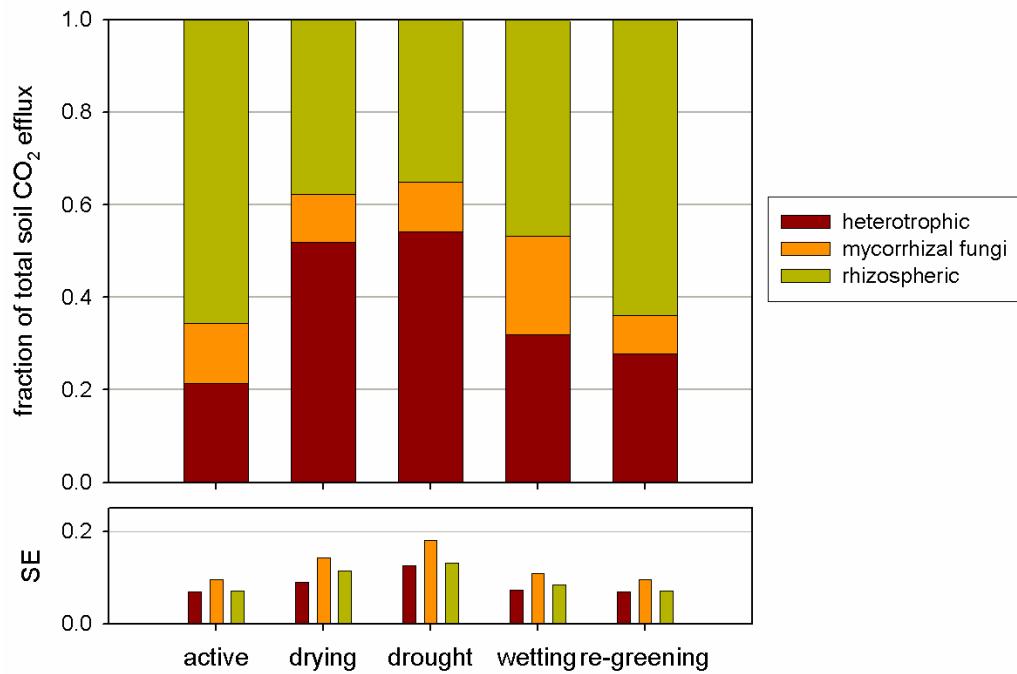


Fig. 3 Relative contributions made by rhizospheric, mycorrhizal fungi and heterotrophic components to the total soil respiration in the different parts of the vegetation period (15/05/2013-12/11/2013) at Bugac site. Propagated uncertainties of each estimate are shown in the lower panel. **Gray horizontal lines show Y major values.**

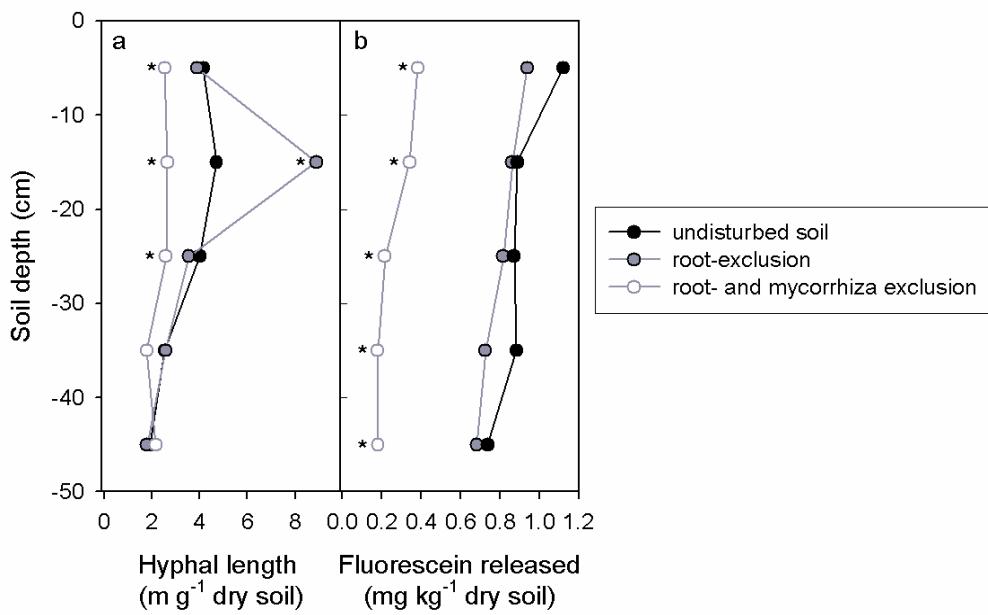


Fig. 4 (a) Mean hyphal length (m g^{-1} dry soil) and (b) mean microbial activity expressed as fluorescein released (mg kg^{-1} dry soil) in the undisturbed soil, root-exclusion and root- and mycorrhiza exclusion in different soil depths. Asterisks denote significant differences from undisturbed soil **determined by the Tukey honest significant difference test**.