https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

Biogeosciences Discussions
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Evaluation of bacterial glycerol dialkyl glycerol tetraether and ^{2}H-^{18}O biomarker proxies along a Central European topsoil transect

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https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

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Keywords

Leaf wax *n*-alkanes, hemicellulose sugars, pH, temperature, CBT, MBT', precipitation $\delta^2H/\delta^{18}O$, relative humidity

Abstract

Molecular fossils, like bacterial branched glycerol dialkyl glycerol tetraethers (brGDGTs), and the stable isotopic composition of biomarkers, such as δ^2H of leaf wax-derived *n*-alkanes ($\delta^2H_{n\text{-alkane}}$) or $\delta^{18}O$ of hemicellulose-derived sugars ($\delta^{18}O_{sugar}$) are increasingly used for the reconstruction of past climate and environmental conditions. Plant-derived $\delta^2H_{n\text{-alkane}}$ and $\delta^{18}O_{sugar}$ values record the isotopic composition of plant source water ($\delta^2H/\delta^{18}O_{source\text{-}water}$), which usually reflects mean annual precipitation ($\delta^2H/\delta^{18}O_{precipiation}$), modulated by evapotranspirative leaf water enrichment and biosynthetic fractionation. Accuracy and precision of respective proxies should be ideally evaluated at a regional scale. For this study, we analysed topsoils below coniferous and deciduous forests, as well as grassland soils along a Central European transect in order to investigate the variability and robustness of various proxies, and to identify effects related to vegetation. Soil pH-values derived from brGDGTs correlate reasonably well with measured soil pH-values, but systematically overestimate them ($\Delta pH = 0.6 \pm 0.6$). The branched vs. isoprenoid tetraether index (BIT) can give some indication whether the pH reconstruction is reliable. Temperatures derived from brGDGTs overestimate mean annual air temperatures slightly ($\Delta T_{MA} = 0.5°C \pm 2.4$). Apparent isotopic fractionation ($\varepsilon_{n\text{-alkane/precipitation}}$ and $\varepsilon_{sugar/precipitation}$) is lower for grassland sites than for forest sites due to "signal damping", i.e. grass biomarkers do not record the full evapotranspirative leaf water enrichment. Coupling $\delta^2H_{n\text{-alkane}}$ with $\delta^{18}O_{sugar}$ allows to reconstruct the stable isotopic composition of the source water more accurately than without the coupled approach ($\Delta\delta^2H$ = ~-21‰ ±22 and $\Delta\delta^{18}O$ = ~-2.9‰ ±2.8). Similarly, relative humidity during daytime and vegetation period (RH_{MDV}) can be reconstructed using the coupled isotope approach (ΔRH_{MDV} = ~-17 ±12). Especially for coniferous sites, reconstructed RH_{MDV} values as well as source water isotope composition underestimate the measured values. This can be likely explained by understory grass vegetation at the coniferous sites contributing significantly to the *n*-alkane pool but only marginally to the sugar pool in the topsoil. The large uncertainty likely reflect the fact that biosynthetic fractionation is not constant, as well as microclimate variability. Overall, GDGTs and the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach have great potential for more quantitative paleoclimate reconstructions.

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

1 Introduction

Information about the variability and consequences of past climate changes is a prerequisite for precise predictions regarding the present climate change. Molecular fossils, so called biomarkers, climate proxies have great potential to enhance our understanding about variations of past climate and environmental changes. Lipid biomarkers in particular, are increasingly used for paleoclimate and environmental reconstructions (e.g. Brincat et al., 2000; Eglinton and Eglinton, 2008; Rach et al., 2014; Romero-Viana et al., 2012; Schreuder et al., 2016). However strengths and limitations of respective proxies need known (Dang et al., 2016). For this, calibrations using modern reference samples are essential.

Terrestrial branched glycerol dialkyl glycerol tetraethers (brGDGTs) that are synthesized in the cell membranes of anaerobe heterotrophic soil bacteria (Oppermann et al., 2010; Weijers et al., 2010) have great potential for the reconstruction of past environmental conditions (e.g. Coffinet et al., 2017; Schreuder et al., 2016; Zech et al., 2012), although some uncertainties exist. Calibration studies suggest that the relative abundance of the individual brGDGTs varies with mean annual air temperature (T_{MA}) and soil pH (Peterse et al., 2012; Weijers et al., 2007), at least across large, global climate gradients or along pronounced altitudinal gradients (Wang et al., 2017). However, in arid regions the production of brGDGT is limited, while isoprenoidal GDGTs (iGDGTs) produced by archaea provide the dominant part of the overall soil GDGT pool (Anderson et al., 2014; Dang et al., 2016; Dirghangi et al., 2013; Wang et al., 2013; Xie et al., 2012). The ratio of brGDGTs vs. isoprenoid GDGTs (BIT) can be used as indication whether a reconstruction of T_{MA} and pH will be reliable. Moreover, Mueller-Niggemann et al. (2016) revealed an influence of the vegetation cover on the brGDGT producing soil microbes. From field experiments, it is known, that vegetation type and mulching practice strongly effect soil temperature and moisture (Awe et al., 2015; Liu et al., 2014). Thus, multiple factors can be expected to influence soil microbial communities and GDGT production. So far, little is known about the variability of GDGT proxies on a regional scale, and a calibration study with small climate gradient but with different vegetation types might be useful.

Compound specific stable hydrogen isotopes of leaf wax biomarkers, such as long chain *n*-alkanes ($\delta^2H_{n\text{-alkanes}}$) record the isotopic signal of precipitation and therefore past climate and environmental conditions (Sachse et al., 2004, 2006). However, various influencing factors are known all along the way from the moisture source to leaf waxes (Pedentchouk and Zhou, 2018 and Sachse et al., 2012 for review). One is the evapotranspiration of leaf water (Feakins and Sessions, 2010; Kahmen et al., 2013; Zech et al., 2015), which is strongly driven by relative air humidity (RH; e.g. Cernusak et al., 2016 for review). In addition, a strong precipitation signal is known to be incorporated into long chain leaf waxes (Hou et al., 2008; Rao et al., 2009; Sachse et al., 2004). In paleoclimate studies, it is often not feasible to disentangle between the evapotranspirative enrichment from the precipitation signal. Zech et al. (2013) proposed to couple $\delta^2H_{n\text{-alkane}}$ results with oxygen stable isotopes of hemicellulose-derived sugars ($\delta^{18}O_{sugar}$). Assuming constant biosynthetic fractionation factors (ε_{bio}) for the different compound classes (*n*-alkanes and hemicellulose sugars), the coupling enables the reconstruction of the isotopic composition of leaf water, RH and $\delta^2H/\delta^{18}O$ of plant source water ($\approx \delta^2H/\delta^{18}O$ of precipitation; Tuthorn et al., 2015). So far, a detailed evaluation of this approach on the European scale, as well as concerning possible effects related to vegetation changes is missing.

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

We analysed topsoil samples under coniferous, deciduous and grassland vegetation along a Central European transect in order to estimate the variability of the biomarker proxies. More specifically, we aim to test whether:

(i) the vegetation type has an influence on the brGDGT proxies, the $\delta^2H_{n\text{-alkane}}$ and the $\delta^{18}O_{sugar}$ stable isotopic composition, as well as on reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ and RH.

(ii) the published brGDGT proxies used for reconstructing mean annual temperature and soil pH are sensitive enough to reflect the medium changes in temperature and soil pH along our transect.

(iii) the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach faithfully reflects $\delta^2H/\delta^{18}O$ of precipitation and RH along the transect.

2 Material and methods

2.1 Geographical setting and sampling

In November 2012, we collected topsoil samples (0-5 cm depth) at 16 locations along a transect from Southern Germany to Southern Sweden (Fig. 1A) and distinguished between sites with coniferous forest (con, n = 9), deciduous forest (dec, n = 14) and grassland (grass, n = 6) vegetation cover (for more details see Schäfer et al. (2016) and Tab. S1).

2.2 Database of instrumental climate variables and isotope composition of precipitation

Climate data was derived from close-by weather observation stations operating by the regional institutions (Deutscher Wetterdienst (DWD) for Germany, Danmarks Meteorologiske Institut (DMI) for Denmark and the Sveriges Meteorologiska och Hydrologiska Institute (SMHI) for Sweden). The DWD provides hourly data for each station (DWD Climate Data Center, 2018b), enabling not only the calculation of T_{MA}, but also of the mean annual relative air humidity (RH_{MA}), mean temperature and relative air humidity during the vegetation period (T/RH_{MV}), and of daytime temperature and relative humidity averages over the vegetation period (T/RH_{MDV}). In addition, annual precipitation observations were used to derive the mean annual precipitation amount (P_{MA}; DWD Climate Data Center, 2018b). From the DMI, the respective climate variables were derived from published technical reports (Cappelen, 2002; Frich et al., 1997; Laursen et al., 1999). The SMHI provides open data from which we derived the climate variables for the Swedish sites (Swedish Meteorological and Hydrological Institute, 2018). For more details about the climate database used for calculations and comparisons, the reader is referred to Tab. S2.

For comprising German precipitation $\delta^2H/\delta^{18}O$ along the transect, we realized a regionalisation (called $\delta^2H/\delta^{18}O_{GIPR}$) using online available data from 34 German GNIP stations, 4 Austrian ANIP stations and the Groningen GNIP station (van Geldern et al., 2014; IAEA/WMO, 2018; Stumpp et al., 2014; Umweltbundesamt GmbH, 2018), following the approach of Schlotter (2007). However, instead of the multivariate regression procedure applied by Schlotter (2007), we used a random forest approach (Hothorn et al., 2006; Strobl et al., 2007, 2008) to describe the relationship of squared latitude, latitude, longitude and altitude vs. long term weighted means of precipitation $\delta^2H/\delta^{18}O$, and realized the prediction for the study sites. For the Danish

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

and Swedish sites, such a procedure was not possible. Hence, the annual precipitation $\delta^2H/\delta^{18}O$ values were derived from the Online Isotopes in Precipitation Calculator (OIPC, version 3.1), therefore called $\delta^2H/\delta^{18}O_{OIPC}$ (Bowen, 2018; Bowen and Revenaugh, 2003; IAEA/WMO, 2015). The finally used $\delta^2H/\delta^{18}O_{GIPR/OIPC}$ data are given in Tab. S1.

The T_{MA} along the transect ranges from 5.3 to 10.6°C, and P_{MA} ranges from 554 to 1769 mm (Fig. 1B). Precipitation $\delta^2H/\delta^{18}O$ shows moderate changes along the transect, $\delta^2H_{GIPR/OIPC}$ varies between -52 and -79‰, and $\delta^{18}O_{GIPR/OIPC}$ ranges from -7.4 to -10.9‰ (Fig. 1C).

Correlations between $\delta^{18}O_{GIPR/OIPC}$ and P_{MA}, altitude of the locations, T_{MA} are given in the supplementary material (Fig. S1 to S3), along with a $\delta^2H_{GIPR/OIPC}$ vs. $\delta^{18}O_{GIPR/OIPC}$ scatter plot (Fig. S4).

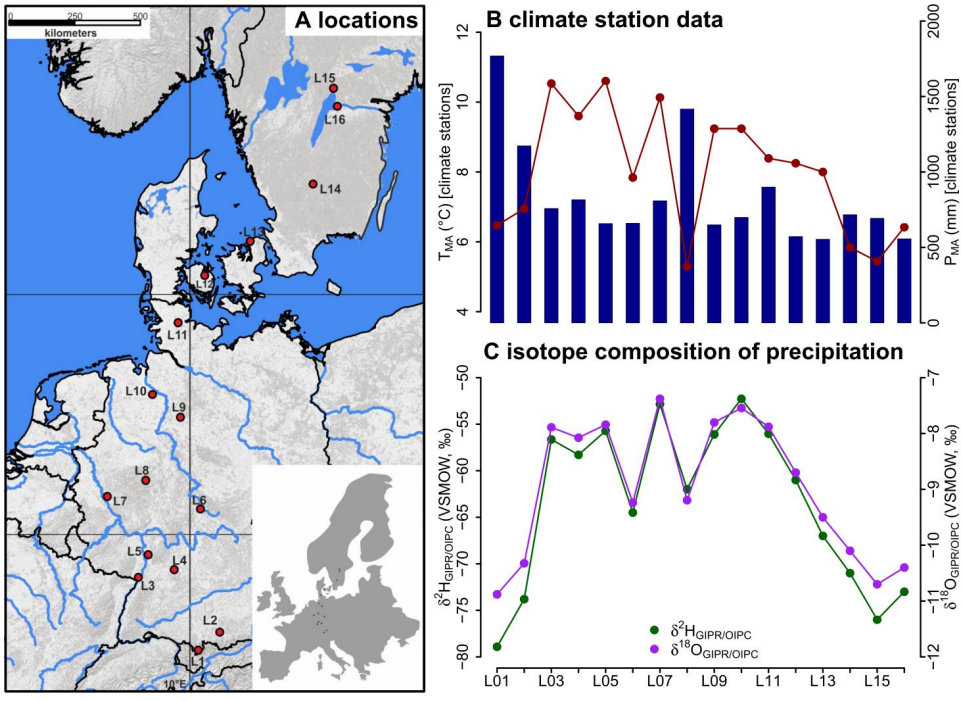


Fig. 1. (A) Sample locations (red dots, map source: US National Park Service), (B) variations of mean annual air temperature (T_{MA}) and mean annual precipitation (P_{MA}) derived from close-by climate station data, and (C) hydrogen and oxygen stable isotope composition of precipitation ($\delta^2H_{GIPR/OIPC}$ and $\delta^{18}O_{GIPR/OIPC}$, respectively) as derived for the sampled transect locations (see section 2.2 GIPR $\delta^2H/\delta^{18}O$ generation procedure). The reader is referred to section 2.2 (and Tab. S1 and S2) for database and reference information of data plotted in (B) and (C).

2.3 Soil extractions and analysis

2.3.1 GDGTs and pH

A detailed description of sample preparation for lipid analysis can be found in Schäfer et al. (2016). Briefly, 1–6 g freeze-dried and grounded soil sample was microwave extracted with 15

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

ml dichloromethane (DCM)/methanol (MeOH) 9:1 *(v:v)* at 100°C for 1 h. Extracts were separated over aminopropyl silica gel (Supelco, 45 µm) pipette columns. The nonpolar fraction (including *n*-alkanes) was eluted with hexane and further purified over $AgNO_3$ coated silica pipette columns (Supelco, 60-200 mesh) and zeolite (Geokleen Ltd.). The GDGT-containing fraction was eluted with DCM:MeOH 1:1 *(v:v)*, re-dissolved in hexane/isopropanol (IPA) 99:1 *(v:v)* and transferred over 0.45 µm PTFE filters into 300 µl inserts. For quantification, a known amount of a C_{46} diol standard was added after transfer. The samples were analysed at ETH Zurich using an Agilent 1260 Infinity series HPLC–atmospheric chemical pressure ionization mass spectrometer (HPLC–APCI-MS) equipped with a Grace Prevail Cyano column (150 mm × 2.1 mm; 3 µm). The GDGTs were eluted isocratically with 90% A and 10% B for 5 min and then with a linear gradient to 18% B for 34 min at 0.2 ml min^{-1}, where A=hexane and B=hexane/isopropanol (9:1, *v:v*). Injection volume was 10 µl and single ion monitoring of $[M+H]^+$ was used to detect GDGTs.

The pH of the samples was measured in the laboratory of the Soil Biogeochemistry group, Institute of Agronomy and Nutritional Sciences, Martin-Luther-University Halle-Wittenberg, in a 1:3 soil:water *(w/v)* mixture.

2.3.2 $\delta^2H_{n\text{-alkane}}$

The hydrogen isotopic composition of the highest concentrated *n*-alkanes (*n*-C_{25}, *n*-C_{27}, *n*-C_{29}, *n*-C_{31}, and *n*-C_{33}) was determined using a TRACE GC Ultra Gas Chromatography connected to a Delta V Plus Isotope Ratio Mass Spectrometer via a ^2H pyrolysis reactor (GC-^{2}H-Py-IRMS; Thermo Scientific, Bremen, Germany) at the ETH Zurich. The compound-specific $^2H/^1H$ ratios were calibrated against an external standard with C_{15} – C_{35} homologues. External standard mixtures (A4 mix from A. Schimmelmann, University of Indiana) were run between the samples for multipoint linear normalization. The H^+_3 factor was determined on each measurement day and was constant throughout the periods of the sample batches. Samples were analysed in duplicates, and results typically agreed within 4% (average difference = 1.4%). All δ^2H values are expressed relative to the Vienna Standard Mean Ocean Water (V-SMOW).

2.3.3 $\delta^{18}O_{sugar}$

Hemicellulose sugars were extracted and purified using a slightly modified standard procedure (Amelung et al., 1996; Guggenberger et al., 1994; Zech and Glaser, 2009). Briefly, myoinositol was added to the samples prior to extraction as first internal standard. The sugars were released hydrolytically using 4M trifluoroacetic acid for 4 h at 105°C, cleaned over glass fiber filters and further purified using XAD and Dowex columns. Before derivatization with methylboronic acid (Knapp, 1979), the samples were frozen, freeze-dried, and 3-O-methylglucose in dry pyridine was added as second internal standard. Compound-specific hemicellulose sugar ^{18}O measurements were performed in the laboratory of the Soil Biogeochemistry group, Institute of Agronomy and Nutritional Sciences, Martin-Luther-University Halle-Wittenberg, using GC-^{18}O-Py-IRMS (all devices from Thermo Fisher Scientific, Bremen, Germany). Standard deviations of the triplicate measurements were 1.4‰ (over 29 investigated samples) for arabinose and xylose, respectively. We focus on these two hemicellulose-derived neutral sugars

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

arabinose and xylose as they strongly predominate over fucose in terrestrial plants, soils and sediments (Hepp et al., 2016 and references therein). Rhamnose concentrations were too low to obtain reliable $\delta^{18}O$ results. All $\delta^{18}O$ values are expressed relative to the Vienna Standard Mean Ocean Water (V-SMOW).

2.4 Theory and Calculations

2.4.1 Calculations used for the GDGT-based reconstructions

The branched and isoprenoid tetraether (BIT) index is calculated according to Hopmans et al. (2004), for structures see Fig. S5:

$$BIT = \frac{Ia+IIa+IIIa}{Ia+IIa+IIIa+crenarchaeol}. \quad (1)$$

The cyclopentane moiety number of brGDGTs correlates negatively with soil pH (Weijers et al., 2007), which led to the development of the cyclization of branched tetraethers (CBT) ratio. CBT and the CBT based pH (pH_{CBT}) were calculated according to Peterse et al. (2012):

$$CBT = -\log\frac{Ib+IIb}{Ia+IIa}, \quad (2)$$

$$pH_{CBT} = 7.9 - 1.97 \times CBT. \quad (3)$$

The number of methyl groups in brGDGTs correlates negatively with T_{MA} and soil pH (Peterse et al., 2012; Weijers et al., 2007). Thus, the ratio of the methylation of branched tetraethers (MBT) ratio and the CBT ratio can be used to reconstruct T_{MA}. We use the equation given by Peterse et al. (2012):

$$MBT' = \frac{Ia+Ib+Ic}{Ia+Ib+Ic+IIa+IIb+IIc+IIIa}, \quad (4)$$

$$T_{MA} = 0.81 - 5.67 \times CBT + 31.0 \times MBT'. \quad (5)$$

2.4.2 Calculations and concepts used for the coupled δ^2H-$\delta^{18}O$ approach

The apparent fractionation is calculated according to Cernusak et al. (2016):

$$\varepsilon_{n\text{-alkane/precipitation}} = \left(\frac{\delta^2H_{n\text{-alkane}} - \delta^2H_{GIPR/OIPC}}{1+\delta^2H_{GIPR/OIPC}/1000}\right), \quad (6)$$

$$\varepsilon_{sugar/precipitation} = \left(\frac{\delta^{18}O_{sugar} - \delta^{18}O_{GIPR/OIPC}}{1+\delta^{18}O_{GIPR/OIPC}/1000}\right). \quad (7)$$

The isotopic composition of leaf water ($\delta^2H/\delta^{18}O_{leaf\ water}$) can be calculated using ε_{bio} for $\delta^2H_{n\text{-alkane}}$ (-160‰, Sachse et al., 2012; Sessions et al., 1999) and $\delta^{18}O_{sugar}$ (+27‰, Cernusak et al., 2003; Schmidt et al., 2001):

$$\delta^2H_{leaf\ water} = \left(\frac{1000+\delta^2H_{n\text{-alkane}}}{1000+\varepsilon_{bio}\ (n\text{-alkane})}\right) \times 10^3 - 1000, \quad (8)$$

$$\delta^{18}O_{leaf\ water} = \left(\frac{1000+\delta^{18}O_{sugar}}{1000+\varepsilon_{bio}\ (sugar)}\right) \times 10^3 - 1000. \quad (9)$$

Zech et al. (2013) introduced the conceptual model for the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach in detail. Briefly, the coupled approach is based on the following assumptions (illustrated in Fig. 8): (i) The isotopic composition of precipitation, which is set to be equal to the plant source water, typically plots along the global meteoric water line (GMWL; $\delta^2H = 8 \times \delta^{18}O + 10$) in a

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

$\delta^{18}O$ vs. $\delta^{2}H$ space (Craig, 1961); (ii) Source water uptake by plants does not lead to any fractionation (e.g. Dawson et al., 2002), and significant evaporation of soil water can be excluded; (iii) Evapotranspiration leads to enrichment of the remaining leaf water along the local evaporation line (LEL; Allison et al., 1985; Bariac et al., 1994; Walker and Brunel, 1990), compared to the source water taken up by the plant; (iv) The biosynthetic fractionation is assumed to be constant. In addition, isotopic equilibrium between plant source water (~ weighted mean annual precipitation) and the local atmospheric water vapour is assumed. Further assumption concerns the isotope steady-state in the evaporating leaf water reservoir. The coupled approach allows for reconstructing the isotopic composition of plant source water ($\delta^{2}H/\delta^{18}O_{source\text{-}water}$) from the reconstructed leaf water, by calculating the intercepts of the LELs with the GMWL (Zech et al., 2013). The slope of the LEL (S_{LEL}) can be assessed by the following equation (Gat, 1971):

$$S_{LEL} = \frac{\varepsilon_2^* + C_k^2}{\varepsilon_{18}^* + C_k^{18}}, \quad (10)$$

where ε^* are equilibrium isotope fractionation factors and C_k are kinetic fractionation factors. The latter equals to 25.1‰ and 28.5‰, for C_k^2 and C_k^{18}, respectively (Merlivat, 1978). The equilibrium fractionation factors can be derived from empirical equations (Horita and Wesolowski, 1994) by using T_{MDV} values. For two Danish sites T_{MDV} are not available, instead T_{MV} is used here (section 2.2 and Tab. S2).

In a $\delta^{18}O$-$\delta^{2}H$ diagram, the distance of the leaf water from the GMWL define the deuterium-excess of leaf water ($d_{leaf\text{-}water} = \delta^{2}H_{leaf\text{-}water} - 8 \times \delta^{18}O_{leaf\text{-}water}$, according Dansgaard, (1964); Fig. 8). To convert $d_{leaf\text{-}water}$ into mean RH during daytime and vegetation period (RH_{MDV}), a simplified Craig-Gordon model can be applied (Zech et al., 2013):

$$RH = 1 - \frac{\Delta d}{\varepsilon_2^* - 8 \times \varepsilon_{18}^* + C_k^2 - 8 \times C_k^{18}}, \quad (11)$$

where Δd is the difference in $d_{leaf\text{-}water}$ and the deuterium-excess of source water ($d_{source\text{-}water}$).

2.5 Statistics

In the statistical analysis we checked sample distributions for normality (Shapiro and Wilk, 1965) and for equal variance (Levene, 1960). If normality and equal variances are given, we perform an Analysis of Variance (ANOVA). If that is not the case, we conduct the non-parametric Kruskal-Wallis Test. ANOVA or Kruskal-Wallis are used to find significant differences (a=0.05) between the vegetation types (deciduous, conifer and grass).

In order to describe the relation along a 1:1 line, the coefficient of correlation (R^2) was calculated as $R^2 = 1 - \sum(\text{modeled} - \text{measured})^2 / \sum(\text{measured} - \text{measured mean})^2$. The small r^2 is taken as coefficient of correlation of a linear regression between a dependent (y) and explanatory variable(s). The root mean square error (RMSE) of the relationships was calculated as $RMSE = \sqrt{\left(\frac{1}{n} \cdot \sum(\text{modeled} - \text{measured})^2\right)}$. All data plotting and statistical analysis was realized in R (version 3.2.2; R Core Team, 2015).

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

3 Results and Discussion

3.1 GDGT concentrations

GDGT Ia has the highest concentration under all vegetation types, followed by GDGT IIa and GDGT IIIa (Fig. 2). GDGT Ib, IIb and Ic occur in minor, GDGT IIc and IIIb only in trace amounts. GDGT IIIc was below the detection limit in most of the samples (Tab. S3). Although other studies document an influence of the vegetation cover on soil temperature and soil water content, which control the microbial community composition in soils (Awe et al., 2015; Liu et al., 2014; Mueller-Niggemann et al., 2016), we find no statistically different pattern of the individual brGDGTs.

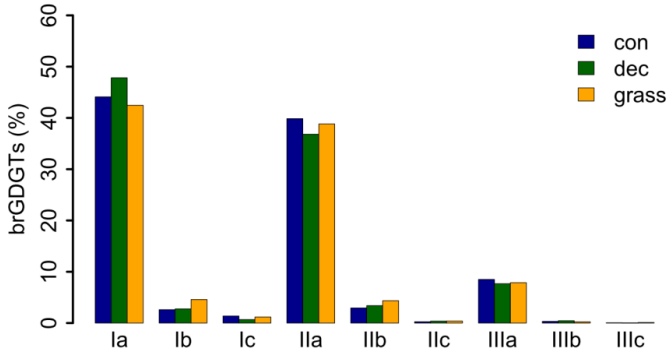


Fig. 2. Mean concentrations of individual brGDGTs as percentage of all brGDGTs for the three investigated types. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=14); grass = grassland sites (n=6).

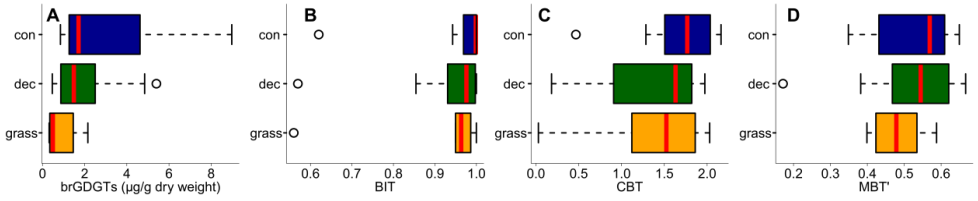
Total concentrations of brGDGTs range from 0.32 to 9.17 µg/g dry weight and tend to be highest for the coniferous samples and lowest for the grasses (Fig. 3A, Tab. S3). Bulk brGDGT concentrations lie within ranges of other studies examining soils of mid latitude regions (Huguet et al., 2010b, 2010a; Weijers et al., 2011). Similar concentrations in coniferous and deciduous samples imply that brGDGT production does not strongly vary in soils below different forest types. The grass samples show lower brGDGT concentrations compared to the forest samples, but this is probably mainly due to ploughing of the grass sites and hence admixing of mineral subsoil material. Anyhow, the differences in brGDGT concentrations are not significant (p-value = 0.06).

3.2 BIT index

Most of the samples have a BIT index higher than 0.9 (Fig 3B and Tab. S3). The BIT-values are typical for soils in humid and temperate climate regions (Weijers et al., 2006). However, outliers exist. The most likely source of iGDGTs in soils are Thaumarchaeota, i.e. aerobe ammonia oxidizing archaea producing Crenarchaeol and its regioisomer (Schouten et al., 2013 and references therein), precipitation amounts drop below 700-800 mm (Dang et al., 2016; Dirghangi et al., 2013). The P_{MA} data of our sampling sites mostly show precipitation > 550 mm (Fig. 1B), but one has to be aware that this data is based on the climate station nearest to the respective sampling locations and microclimate effects, such as sunlight exposure, canopy

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

306 cover or exposition might have a pronounced influence on the brGDGT vs. iGDGT distribution.
307 Mueller-Niggemann et al. (2016) found higher BIT indices in upland soils compared to paddy
308 soils and stated that the management type also influences BIT values in soils. Along our
309 transect, grass sites tend to have slightly lower BIT-values than forest sites, probably due to the
310 absence of a litter layer and hence, no isolation mechanism preventing evaporation of soil water.
311 Anyhow, differences between vegetation types are not significant (p-value = 0.32).



312

313 **Fig. 3.** (A) Total concentrations of brGDGTs in µg g^{-1} dry weight, as well as (B) BIT, (C) CBT
314 and (D) MBT'. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites
315 (n=14); grass = grassland sites (n=6). Box plots show median (red line), interquartile range
316 (IQR) with upper (75%) and lower (25%) quartiles, lowest whisker still within 1.5IQR of lower
317 quartile, and highest whisker still within 1.5IQR of upper quartile, dots mark outliers.

318

319 **3.3 CBT-derived pH**

320 The CBT ratio shows a pronounced variation independent of vegetation type with values
321 between 0.03 and 2.16 (Fig 3C). The coniferous samples tend to be highest, but the differences
322 between vegetation types are not significant (p-value = 0.48). The CBT index can be related to
323 pH in acidic and/or humid soils (e.g. Dirghangi et al., 2013; Mueller-Niggemann et al., 2016;
324 Peterse et al., 2012; Weijers et al., 2007) but might be an indicator of soil water content and
325 hence, precipitation in more arid and alkaline soils (e.g. Dang et al., 2016). There is a
326 pronounced correlation between CBT and soil pH (Fig. 4), which is in good agreement with
327 other studies from mid latitude regions where precipitation is relatively high (Anderson et al.,
328 2014 and references therein). Moreover, the CBT to pH relationship in terms of slope and
329 intersect in our dataset (CBT = -0.47 × pH + 3.5, r^2 = 0.7, p-value < 0.0001, n = 29) is well
330 comparable to the correlation described for the global calibration dataset of Peterse et al. (2012)
331 (CBT = -0.36 × pH + 3.1, r^2 = 0.7, p-value < 0.0001, n = 176).

332 However, there are some outliers in the CBT-pH correlation, which need a further examination
333 (see locations grass L04, dec L10 and dec L12 as marked in Figs. 4 and 5). The outliers show
334 lower BIT indices (< 0.85, Tab. S3). Even though the data from the nearest climate station
335 suggest no abnormal P_{MA}. Local effects such as differences in the amount of sunlight exposure,
336 nutrient availability for brGDGT producing organisms or, most likely soil water content might
337 influence the brGDGT production at these locations (Anderson et al., 2014; Dang et al., 2016).
338 A lower BIT index as well as a lower CBT occur when soil water content decreases (Dang et
339 al., 2016; Sun et al., 2016) or when aeration is high and less anoxic microhabitats for GDGT
340 producing microbes exist (e.g. Dirghangi et al., 2013).

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Preprint. Discussion started: 29 May 2019

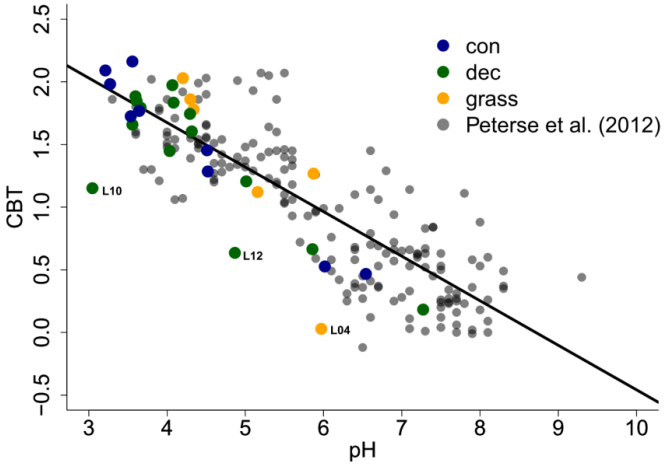

Fig. 4. CBT to pH relationship in our dataset in comparison to the global calibration dataset from Peterse et al. (2012) (CBT = -0.36 × pH + 3.1, r^2 = 0.7, p-value < 0.0001, n = 176, black line). Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=14); grass = grassland sites (n=6).

As the CBT and pH are similarly correlated in our dataset and the global dataset of Peterse et al. (2012), the CBT-derived pH correlated well with the actual pH (Fig. 5A; R^2 = 0.3). Expressed as ΔpH (CBT-derived pH - measured pH), there is a tendency that the GDGTs result in an overestimation of the real pH for the forest sites (Fig. B). Yet a Kruskal-Wallis test shows no statistically significant difference between the vegetation types, with a p-value of 0.13. The overall ΔpH of 0.6 ±0.6 shows that the reconstruction of soil pH using brGDGTs works well along this transect.

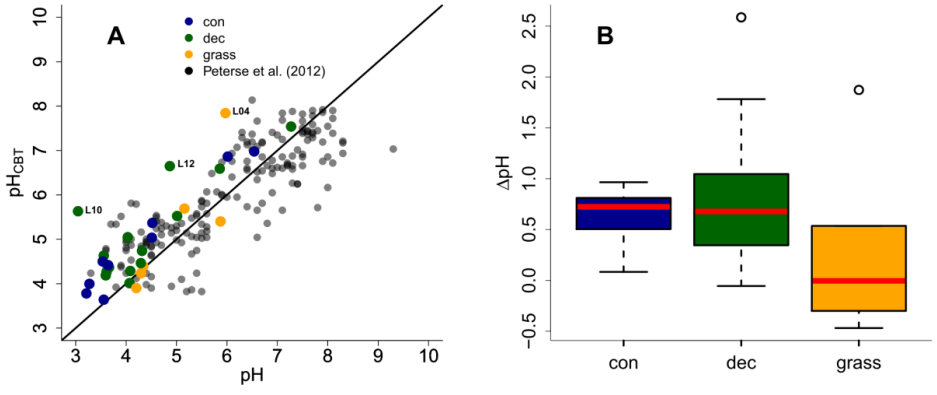

Fig. 5. (A) Correlation between measured pH and reconstructed soil pH (pH_{CBT}) from our transect data in comparison to the global calibration dataset from Peterse et al. (2012) (R^2 = 0.7, RMSE = 0.75, n = 176). Black line indicates the 1:1 relationship. (B) Boxplots of ΔpH (refers to pH_{CBT}-pH). Box plots show median (red line), interquartile range (IQR) with upper (75%)

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

and lower (25%) quartiles, lowest whisker still within 1.5IQR of lower quartile, and highest whisker still within 1.5IQR of upper quartile, dots mark outliers. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=14); grass = grassland sites (n=6).

3.4 MBT'-CBT-derived T_{MA} reconstructions

The MBT' shows high variability with values ranging from 0.17 to 0.67 no statistical differences between vegetation types (p-value = 0.54; Fig. 3D, Tab. S3). When comparing reconstructed (MBT'-CBT-derived) T_{MA} with climate station T_{MA}, the data plot close to the 1:1 line, and fit well into the global dataset of Peterse et al. (2012) (Fig. 7A). The ΔT_{MA} reveal an overall offset of 0.5°C ±2.4 and there is no statistically difference between vegetation types (Fig. 7B). The standard deviation in ΔT_{MA} of ±2.4 is well in line with the RMSE of 5.0 for the global calibration dataset (Peterse et al., 2012).

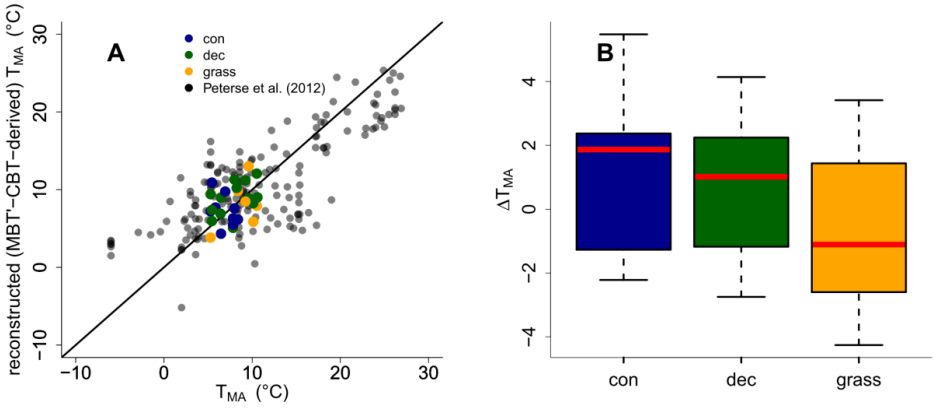

Fig. 6. (A) Correlation between climate station T_{MA} and reconstructed (MBT'-CBT-derived) T_{MA}. For comparison, the global calibration dataset from Peterse et al. (2012) is shown. The black line indicates the 1:1 relationship. (B) Boxplots of ΔT_{MA} (refers to reconstructed T_{MA}-T_{MA} from climate stations) in the different vegetation types from our transect study. Box plots show median (red line), interquartile range (IQR) with upper (75%) and lower (25%) quartiles, lowest whisker still within 1.5IQR of lower quartile, and highest whisker still within 1.5IQR of upper quartile, dots mark outliers. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=14); grass = grassland sites (n=6).

3.5 Apparent fractionation of δ^2H and $\delta^{18}O$ in the different vegetation types

The δ^2H values could be obtained for *n*-alkanes C_{27}, C_{29} and C_{31} in all samples and additionally at two locations for *n*-C_{25} and *n*-C_{33} at six other locations. The $\delta^2H_{n\text{-alkane}}$ values, calculated as mean of *n*-C_{25} to *n*-C_{31} δ^2H, ranges from -156 to -216‰. Pooled standard deviations show an overall average of 3.6‰. The $\delta^{18}O_{sugar}$ values, calculated as the area weighted means for arabinose and xylose, ranges from 27.7 to 39.4‰. The average weighted mean standard deviation is 1.4‰. The compound-specific isotope data is summarized along with the calculations in Tab. S4.

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

Apparent fractionation ($\varepsilon_{n\text{-alkane/precipitation}}$) is on the order of -120 to -150‰, i.e. a bit less than the biosynthetic fraction of -160‰. This implies that evapotranspirative enrichment is ~ 10 to 40‰ (Fig. 7A). $\varepsilon_{n\text{-alkane/precipitation}}$ is lower for grass sites compared to the forest sites. Differences are significant between deciduous and grass sites (p-value = 0.005). This finding supports the results of other studies (Kahmen et al., 2013; Liu and Yang, 2008; McInerney et al., 2011), and can be named "signal damping". Grasses do not only incorporate the evaporatively-enriched leaf water only but also unenriched leaf water in the growth and differentiation zone of grasses (Gamarra et al., 2016; Liu et al., 2017).

The grass-derived hemicellulose sugar biomarkers do not fully record the evapotranspirative enrichment of the leaf water, either, as indicated by lower apparent fractionation ($\varepsilon_{sugar/precipitation}$) in Fig. 7B. The differences are significant between forest and grass sites (p-value < 0.005). This is in agreement with a study on cellulose extracted from grass blades (Helliker and Ehleringer, 2002), and again, the "signal damping" can be explained with incorporation of enriched leaf water and non-enriched stem water.

Based on the comparison of evapotranspirative enrichment between forest and grass sites, the "signal damping" can be quantified to be ~ 31% for the hemicellulose sugars, and ~ 49% for the *n*-alkanes. This is in agreement with other studies that reported a loss of 22% of the leaf water enrichment for hemicellulose sugars (Helliker and Ehleringer, 2002) and 39 to 62% loss of the leaf water enrichment for *n*-alkanes (Gamarra et al., 2016).

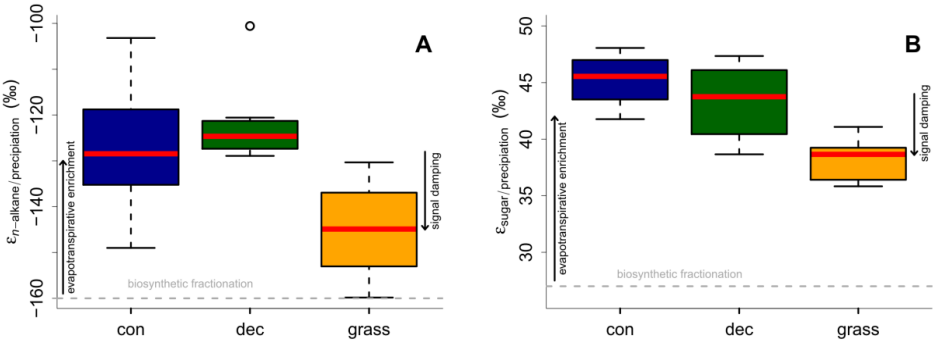

Fig. 7. Apparent fractionation (A) $\varepsilon_{n\text{-alkane/precipitation}}$ and (B) $\varepsilon_{sugar/precipitation}$. Biosynthetic fractionation factors according to section 2.4.2. Box plots show median (red line), interquartile range (IQR) with upper (75%) and lower (25%) quartiles, lowest whisker still within 1.5IQR of lower quartile, and highest whisker still within 1.5IQR of upper quartile, dots mark outliers. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=11 and 14 for *n*-alkanes and sugars, respectively); grass = grassland sites (n=4 and 6 for *n*-alkanes and sugars, respectively). The figure conceptually illustrates the effect of biosynthetic fractionation and evapotranspirative enrichment as well as "signal damping".

3.6 $\delta^2H/\delta^{18}O_{source\text{-}water}$ reconstructions

The δ^2H versus $\delta^{18}O$ diagram shown in Fig. 8 graphically illustrates the reconstruction of $\delta^2H/\delta^{18}O_{leaf\text{-}water}$ (colored dots) from $\delta^2H_{n\text{-alkane}}/\delta^{18}O_{sugar}$ (crosses), as well as the reconstruction

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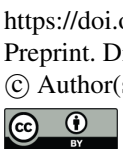
Preprint. Discussion started: 29 May 2019



of $\delta^2H/\delta^{18}O_{source\text{-}water}$ (black dots). For reconstructing $\delta^2H/\delta^{18}O_{source\text{-}water}$, LELs with an average slope of 2.8 ±0.1 (Eq. 10) can be generated through every leaf water point and the intercepts of these LELs with the GMWL.

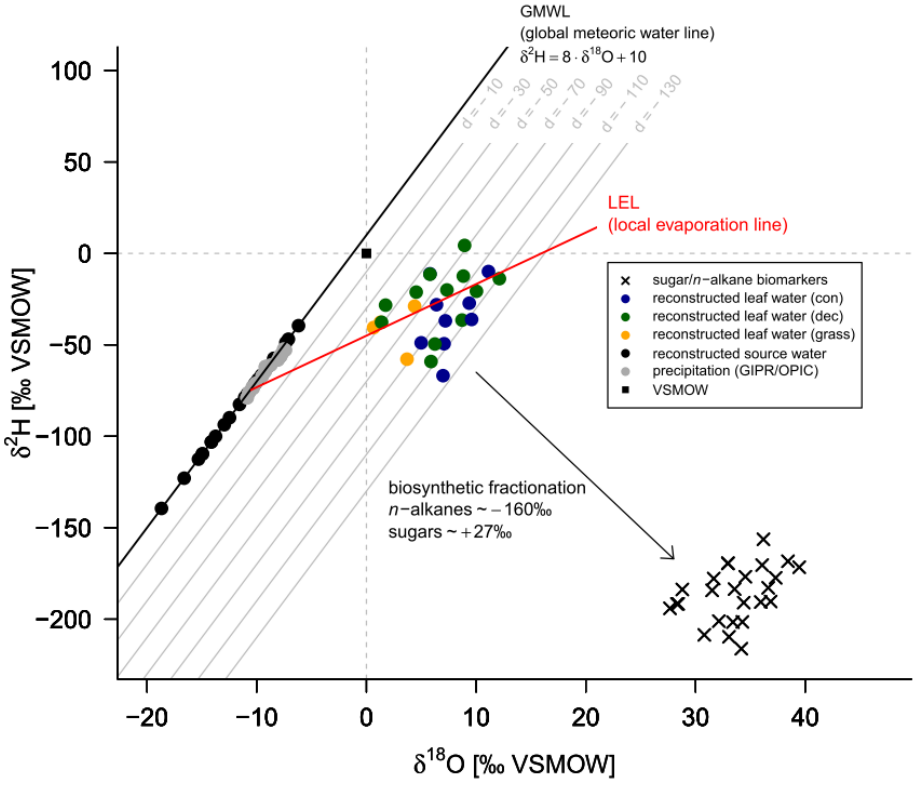


Fig. 8. δ^2H vs. $\delta^{18}O$ diagram illustrating $\delta^2H_{n\text{-alkane}}$ and $\delta^{18}O_{sugar}$, reconstructed $\delta^2H/\delta^{18}O_{leaf\text{-}water}$ (according Eqs. 8 and 9) and reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ in comparison to GIPR/OIPC-based $\delta^2H/\delta^{18}O_{precipitation}$. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=11); grass = grassland sites (n=4).

The reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ results can be compared with the $\delta^2H/\delta^{18}O_{GIPR/OIPC}$ data (Fig. 9). This comparison reveals that the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach yields more accurate $\delta^2H/\delta^{18}O_{source\text{-}water}$ results than hitherto applied $\delta^2H_{n\text{-alkane}}$ single isotope approaches. However, the range of the reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ values is clearly larger than in $\delta^2H/\delta^{18}O_{GIPR/OIPC}$ values. δ^2H is systematically underestimated by ~ 21‰ ±22 (Fig. 9B) and $\delta^{18}O$ by ~ 2.9‰ ±2.8 (Fig. 9D). The type of vegetation seems to be not particularly relevant (p-value = 0.18 for $\Delta\delta^2H$ and p-value = 0.34 for $\Delta\delta^{18}O$). Nevertheless, the systematic offsets tend to be lowest for the decidous sites ($\Delta\delta^2H/\Delta\delta^{18}O$ is closer to zero with ~-5‰ ±15 and ~-1.1‰ ±2.1), followed by grass sites (~-14‰ ±20 and ~-2.1‰ ±2.6). In comparison, the coniferous sites show the largest offsets (~-23‰ ±26 for $\Delta\delta^2H$ ~-3.0‰ ±3.3 for $\Delta\delta^{18}O$). Differences are, however, not statistically significant. The systematic offset and the large variability might have

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

more specific reasons, and we suggest that this is related to the type of vegetation. Deciduous trees produce lots of leaf waxes and sugars (e.g. Prietzel et al., 2013; Zech et al., 2012a), and all biomarkers reflect and record the evapotranspirative enrichment of the leaf water (e.g. Cernusak et al., 2016; Tuthorn et al., 2014). The coupled approach and the leaf water reconstruction based on the *n*-alkane and sugar biomarkers thus works well. However, coniferous trees produce quite low amounts of *n*-alkanes (Diefendorf and Freimuth, 2016; Zech et al., 2012a), while sugar concentrations are as high as in other vascular plants (e.g. Hepp et al., 2016; Prietzel et al., 2013). For the coniferous soil samples this means that the *n*-alkanes stem most likely from the understory whereas the sugars originate from grasses and coniferous needles. When the understory is dominated by grass species then the *n*-alkane biomarkers do not record the full leaf water enrichment signal, whereas the sugars from the needles do. The reconstructed leaf water for the coniferous sites is therefore too negative concerning δ^2H, and reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ values thus also become too negative (Fig. 8). Concerning the grass sites the following explanation can be found. Correcting for "signal damping" makes the reconstructed leaf water points more positive and shifts them in Fig. 8 up and right. As the "signal damping" is stronger for δ^2H than for $\delta^{18}O$ the corrected leaf water points are now above the uncorrected ones. The corrected leaf water points leads to more positive reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ values for the grass sites.

Vegetation type specific rooting depths could partly cause the overall high variability in reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$. Deep rooting species most likely use the water from deeper soil horizons and/or shallow ground water, which is equal to the (weighted) mean annual precipitation (e.g. Herrmann et al., 1987). Shallow rooting plants take up water from upper soil horizons, which is influenced by seasonal variations in $\delta^2H/\delta^{18}O_{precipiation}$ and by soil water enrichment (Dubbert et al., 2013). Thus, the overall assumption that the source water of the plants reflects the local (weighted) mean precipitation might be not fully valid for all sites. Moreover, a partly contribution of root-derived rather than leaf-derived sugar biomarkers in our topsoil samples is very likely. This does, by contrast, not apply for *n*-alkanes, which are hardly produced in roots (Zech et al., 2012b and the discussion).

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Preprint. Discussion started: 29 May 2019

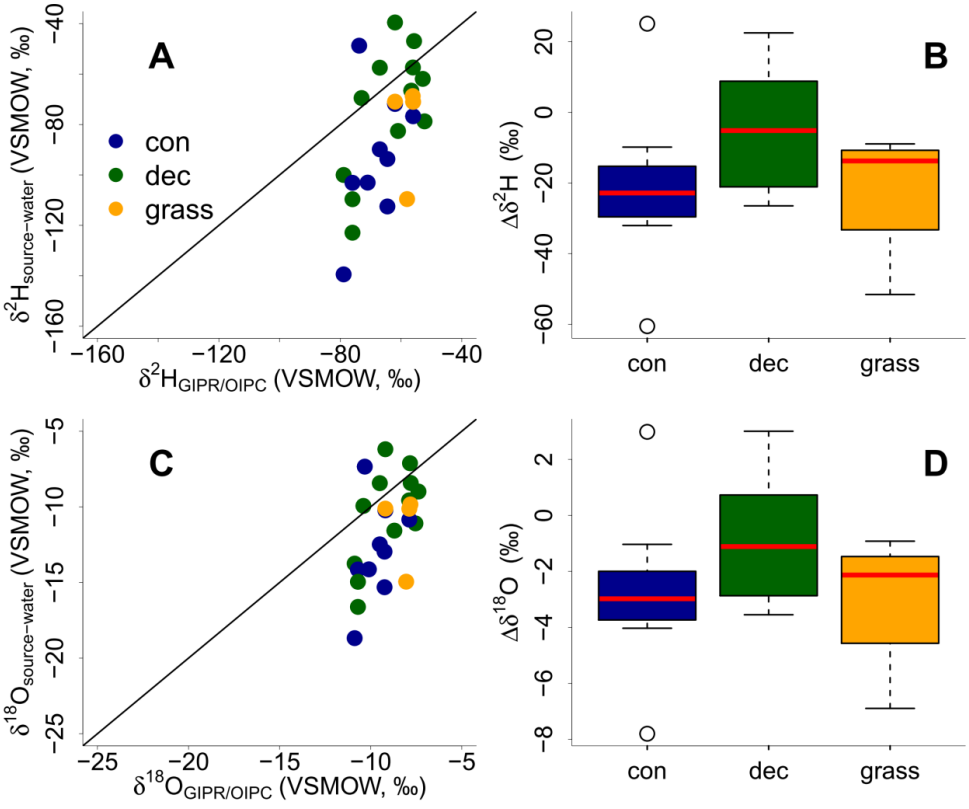

Fig. 9. Correlation of reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ vs. precipitation $\delta^2H/\delta^{18}O_{GIPR/OIPC}$ (A and C). Black lines indicate 1:1 relationship. Differences between reconstructed source water and precipitation ($\Delta\delta^2H/\delta^{18}O$ = $\delta^2H/\delta^{18}O_{source\text{-}water}$ - $\delta^2H/\delta^{18}O_{GIPR/OIPC}$) for the three different vegetation types (B and D). Box plots show median (red line), interquartile range (IQR) with upper (75%) and lower (25%) quartiles, lowest whisker still within 1.5IQR of lower quartile, and highest whisker still within 1.5IQR of upper quartile. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=11); grass = grassland sites (n=4).

Moreover, the high variability within the vegetation types could be caused by variability in ε_{bio} of ^2H in *n*-alkanes, as well as ^{18}O in sugars. There is an ongoing discussion about the correct ε_{bio} for ^{18}O in hemicellulose sugars (Sternberg, 2014 vs. Zech et al., 2014), and ε_{bio} is probably not constant over all vegetation types. This translates into errors concerning leaf water reconstruction and thus for reconstructing $\delta^2H/\delta^{18}O_{source\text{-}water}$ values (Eq. 9 and Fig. 8). Likewise, the ε_{bio} values reported in the literature for ^2H of *n*-alkanes can be off from -160‰ by tens of permille (Feakins and Sessions, 2010; Tipple et al., 2015; Feakins et al., 2016; Freimuth et al., 2017). The degree to which hydrogen originates from NADPH rather than leaf water is important, because NADPH is more negative (Schmidt et al., 2003). The wide range in biosynthetic ^2H fractionation factors is therefore also related to the carbon and energy metabolism state of plants (Cormier et al., 2018).

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

3.7 RH reconstruction

Reconstructed RH_{MDV} ranges from 34 to 74%, while RH_{MDV} from climate station data range from 61 to 78% (Fig. 10A). Biomarker-based values thus systematically underestimate the station data (ΔRH_{MDV} = -17% ±12; Fig. 10B). Yet the offsets are much less for deciduous tree and grass sites (ΔRH_{MDV} = -10% ±12 and -7% ±9, respectively). The offsets for the coniferous sites are -30% ±11, and significantly larger than for the deciduous and grass sites (p-values < 0.05).

Too low reconstructed RH_{MDV} values for the coniferous sites make sense in view of the previously discussed option that soils contain *n*-alkanes from the understory (which is dominated by grass species), while sugars stem from needles and grasses. As explained earlier already, the "signal damping" leads to too negative reconstructed $\delta^2H_{leaf\text{-}water}$ (whereas $\delta^{18}O$ is affected less by the "signal damping"), and too negative $\delta^2H_{leaf\text{-}water}$ translates into overestimated d-excess and underestimated RH values. In Fig. 8, a correction for this require moving the coniferous leaf water data points upwards towards more positive δ^2H values, thus the distance between the leaf water and the source water is reduced.

The underestimation of RH for the deciduous and grass sites could be partly associated with the use of the GMWL as baseline for the coupled $\delta^2H_{n\text{-}alkane}$-$\delta^{18}O_{sugar}$ approach. The deuterium-excess of the LMWLs is generally lower than the +10‰ of the GMWL, while the slopes of the LMWLs are well comparable to the GMWL (Stumpp et al., 2014). In addition, if soil water evaporation occurred before water uptake by the plants, this would lead to an underestimation of biomarker-based RH_{MDV} values. It can be furthermore assumed that plant metabolism is highest during times with direct sunshine and high irradiation, i.e. during noon at sunny days. The relevant RH could therefore be lower than the climate station-derived RH_{MDV}. Indeed, already climate station RH_{MDV} is considerable lower than RH_{MA} and RH_{MV} (Tab. S1).

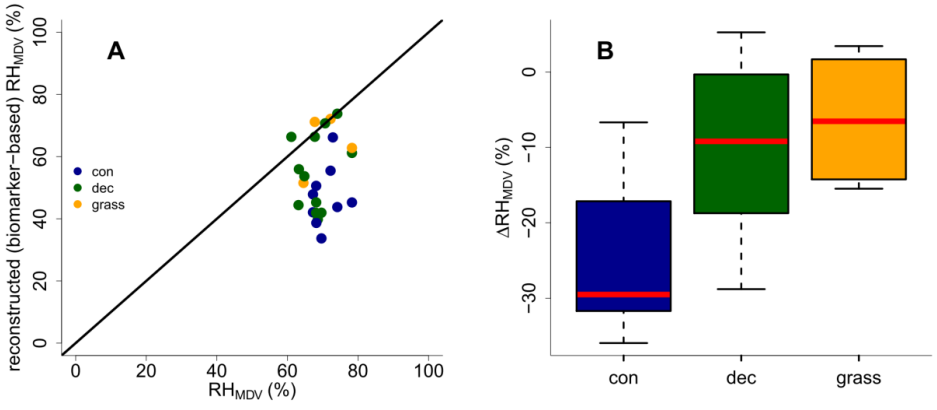

Fig. 10. (A) Comparison of reconstructed (biomarker-based) RH_{MDV} values and climate station RH_{MDV} data. The black line indicates the 1:1 relationship. (B) Differences between reconstructed and climate station RH_{MDV} values (ΔRH_{MDV} = reconstructed – climate station RH_{MDV}) for the three different vegetation types along the transect. Abbreviations: con = coniferous forest sites (n=9); dec = deciduous forest sites (n=11); grass = grassland sites (n=4).

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

The uncertainty of reconstructed RH_{MDV} values are large for all three investigated vegetation types, and again these uncertainties are probably also related to ε_{bio}, which is most likely not constant as assumed for our calculations. Moreover, microclimate variability is underestimated in our approach. As mentioned in sections 2.4.2 and 3.6, in the coupled approach not only the source water of the plants is equated with (weighted) mean annual precipitation, but also an isotopic equilibrium between the source water and the (local) atmospheric water vapour is assumed. However, in areas with distinct seasonality this might be not fully valid. To account for this lack of equilibrium between precipitation and local atmospheric water vapour, apparent ε values can be calculated with data from Jacob and Sonntag, (1991). As shown by Hepp et al. (2018) those values can be used to achieve alternative RH reconstructions based on the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach. Such calculated RH_{MDV} values are on average 1.5% more negative than the original values. However, this difference in RH is far below the analytical uncertainties of the compound-specific biomarker isotope analysis.

Finally, the integration time of the investigated topsoils has to be discussed. Unfortunately, no ^{14}C dates are available for the soil samples. However, most likely the organic matter has been built up over a longer timescale than the available climate data, which is used for comparison. In combination with vegetation changes/management changes throughout that period, this could surely lead to a less tight relationship of the reconstructions compared to the climate station data. Root input of arabinose and xylose seems to be of minor relevance in our topsoil samples. Otherwise, the reconstructed $\delta^{18}O_{sugar}$ values would be too negative resulting in RH_{MDV} overestimations, which is not observed.

4 Conclusions

We were able to show that

(i) the vegetation type does not significantly influence the brGDGT concentrations and proxies, yet the coniferous sites tend to have higher brGDGT concentrations, BIT indices and CBT/MBT' ratios, while grass sites tend to be lowest.

(ii) CBT faithfully records soil pH with a median ΔpH of 0.6 ±0.6, The CBT overestimates the real pH particularly at the forest sites.

(iii) CBT-MBT'-derived T_{MA} reflect the climate station-derived T_{MA} values with a median ΔT_{MA} of 0.5°C ±2.4, but again slightly too high reconstruction for the forest sites were observed.

(iv) differences in the apparent fractionation between the investigated vegetation types are caused by "signal damping", i.e. the grasses do not see and record the full evaporative enrichment of leaf water.

(v) the reconstructed $\delta^2H/\delta^{18}O_{source\text{-}water}$ reflects the $\delta^2H/\delta^{18}O_{GIPR/OIPC}$ with a systematic offset for δ^2H of ~-21‰ ±22 and for $\delta^{18}O$ of ~-2.9‰ ±2.8 (based on overall medians of $\Delta\delta^2H/\delta^{18}O$). This is caused by too negative reconstructions for coniferous and grass sites. For coniferous sites, this can be explained with *n*-alkanes originating from understory grasses, and for the grass sites the "signal damping" more effect δ^2H than $\delta^{18}O$. This leads to too negative reconstructed $\delta^2H_{leaf\text{-}water}$ values and thus to too negative $\delta^2H/\delta^{18}O_{source\text{-}water}$ reconstruction.

https://doi.org/10.5194/bg-2019-197
Preprint. Discussion started: 29 May 2019

(vi) reconstructed (biomarker-based) RH_{MDV} values tend to underestimate climate station-derived RH_{MDV} values (ΔRH_{MDV} = ~ -17% ±12). For coniferous sites the underestimations are strongest, which can be explained with understory grasses being the main source of *n*-alkanes for the investigated soils under coniferous forests.

Overall, our study highlights the great potential of GDGTs and the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach for more quantitative paleoclimate reconstructions. Taking into account effects of different vegetation types improves correlations and reconstructions. This holds particularly true for the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach, which is affected by "signal damping" of the grass vegetation. Assuming constant biosynthetic fractionation is likely a considerable source of uncertainty. Climate chamber experiments would be very useful to further evaluate and refine the coupled $\delta^2H_{n\text{-alkane}}$-$\delta^{18}O_{sugar}$ approach, because uncertainties related to microclimate variability can be reduced. Field experiments like ours suffer from the fact that biomarker pools in the sampled topsoils may have been affected by past vegetation and climate changes.

Acknowledgements

We thank L. Wüthrich, H. Veit, T. Sprafke, A. Groos (all University of Bern), A. Kühnel (Technical University of Munich) for constructive discussions and statistical advices, and M. Schaarschmidt (University of Bayreuth), C. Heinrich and M. Benesch (Martin-Luther-University Halle-Wittenberg) for laboratory assistance during $\delta^{18}O_{sugar}$ analysis and pH measurements, respectively. The Swiss National Science Foundation (PP00P2 150590) funded this research. J. Hepp greatly acknowledges the support by the German Federal Environmental Foundation (DBU) in form of his PhD-fellowship.

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Preprint. Discussion started: 29 May 2019

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