**Biogeosciences
Discussions**

On the choice of the driving temperature for eddy-covariance carbon dioxide flux partitioning

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Influences of temperature observations on CO$_2$ flux partitioning

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Abstract

Networks that merge and harmonise eddy-covariance measurements from many different parts of the world have become an important observational resource for ecosystem science. Empirical algorithms have been developed which combine direct observations

5   of the net ecosystem exchange of carbon dioxide with simple empirical models to disentangle photosynthetic (GPP) and respiratory fluxes (R_{eco}). The increasing use of these estimates for the analysis of climate sensitivities, model evaluation, and calibration demands a thorough understanding of assumptions in the analysis process and the resulting uncertainties of the partitioned fluxes. The semi-empirical models used in

10 flux partitioning algorithms require temperature observations as input, but as respiration takes place in many parts of an ecosystem, it is unclear which temperature input – air, surface, bole, or soil at a specific depth – should be used. This choice is a source of uncertainty and potential biases.

In this study we analysed the correlation between different temperature observations

15 and nighttime NEE (which equals nighttime respiration) across FLUXNET sites to understand the potential of the different temperature observations as input for the flux partitioning model. We found that the differences in the correlation between different temperature data streams and nighttime NEE are small and depend on the selection of sites. We investigated the effects of the choice of the temperature data by running

20 two flux partitioning algorithms with air and soil temperature. We found the time lag (phase shift) between air and soil temperatures explains the differences in the GPP and R_{eco} estimates when using either air or soil temperatures for flux partitioning. The impact of the source of temperature data on other derived ecosystem parameters was estimated, and the strongest impact was found for the temperature sensitivity. Over-

25 all, this study suggests that the choice between soil or air temperature must be made on site-by-site basis by analysing the correlation between temperature and nighttime NEE. We recommend using an ensemble of estimates based on different temperature

observations to account for the uncertainty due to the choice of temperature and to assure the robustness of the temporal patterns of the derived variables.

1 Introduction

Eddy-covariance measurements have contributed strongly to our understanding of
5 ecosystem responses to climate with respect to water, carbon, and energy fluxes (Law et al., 2002; Falge et al., 2002; Teuling et al., 2010; Mahecha et al., 2010, and many more).

To analyse changes in the observed net ecosystem exchange (NEE) with respect to the underlying processes photosynthesis and respiration, NEE is often partitioned
10 into gross primary production (GPP) and ecosystem respiration (R_{eco}). This procedure is usually based on semi-empirical models of respiration, which use temperature as a driver. Respiratory fluxes are generated by many components of an ecosystem, e.g. leaves, branches, stems, but also below-ground, by soil organisms and roots. The temperature driving the respiration processes varies between the different locations where
15 they take place. Moreover eddy-covariance systems observe the flux above the canopy and therefore not at the time when the fluxes form but rather delayed by the transport time from the location of respiration to the sensor (Phillips et al., 2011). This heterogeneity within the ecosystem influences the relationship between observed fluxes and temperature: it can appear as noise or hysteresis patterns may occur. Trying to account
20 for this heterogeneity to improve model performance strongly increases the complexity and, therefore, the number of parameters in the flux partitioning model. However, keeping the model's structure simple is an important goal in order to remain as close as possible to the data, to avoid additional assumptions, and to avoid additional sources of uncertainty. Currently, frequently used flux-partitioning algorithms choose only one
25 temperature stream, and it remains unclear which temperature (air, T_{air} or soil, T_{soil}, in a specific soil depth, or the surface temperature, T_{surf}) is the most appropriate (Subke and Bahn, 2010). Moreover hysteresis can lead to systematic under- or overestimation

in flux-partitioning algorithms that selectively fit only daytime or nighttime data and then extrapolate the model in time.

The collection and harmonisation of eddy-covariance observations from stations all over the world (www.fluxdata.org) and methodological developments now allow for global gridded datasets of carbon fluxes based on local eddy-covariance flux measurements to be derived (Beer et al., 2010). In addition to the value of having a global estimate based on observations, these global gridded data sets are highly promising for land surface model validation and improvement (Bonan et al., 2011). For such studies precise uncertainty estimates strongly increase the usefulness of the data. The uncertainty of the GPP estimate has been partly considered in some studies by including estimates based on daytime and nighttime data, but both algorithms rely on air temperature (and not soil temperature) as a driver for respiration (Beer et al., 2010; Lasslop et al., 2010).

The goal of our study is to improve our understanding of potential biases and uncertainties in flux partitioning algorithms as affected by the choice of the driving temperature and with this our understanding on how to best use these estimates. We first illustrate how differences between air and soil temperature can influence flux partitioning algorithms. We investigate the potential of air and soil temperatures as drivers for the Lloyd and Taylor (Lloyd and Taylor, 1994) respiration model across FLUXNET sites. We quantify the uncertainty and potential biases arising from the choice of the driving temperature for respiration. We further try to attribute the differences between the annual flux components derived with the air and soil temperatures to statistical measures of the relationship between air and soil temperatures and to vegetation structure. Moreover, we quantify the difference in ecosystem parameters derived during the flux partitioning using either air or soil temperature.

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2 Methods

2.1 Data

We used data from the FLUXNET "La Thuile" data base (www.fluxdata.org, version of December 2007) containing 976 site-years of half hourly eddy-covariance data. The data are CO_2 storage change corrected, spike filtered, u_*-filtered (u_* is the friction velocity), and subsequently gap-filled (Papale et al., 2006). A flux partitioning algorithm is applied that partitions the net exchange into GPP and R_{eco} (Reichstein et al., 2005). In spite of this high level of harmonisation the derivation of half hourly fluxes from the high frequency raw data still varies from site to site (Mauder et al., 2008). For the analysis only sites providing the necessary data with a high quality, i.e. the data necessary to apply the flux partitioning and retrieve reliable annual sums were used. We could include 270 yr from 90 sites.

We used air and soil temperature observations; air temperature is usually measured above the canopy close to the eddy-covariance system. The depth of soil temperature observations varies between sites (see Appendix A). When multiple soil temperatures were available we used the observation closest to the surface, assuming that the top layer is usually the largest source of respiration in soils. In addition to these two temperature data streams we computed the surface temperature, T_{surf}, based on longwave radiation for sites measuring the radiation components (see Table 1). The surface temperature can be derived as:

$$T_{surf} = \left(\frac{L_{out} - (1 - \epsilon) L_{in}}{\sigma_B \epsilon} \right)^{1/4} \tag{1}$$

which is based on the Stefan-Boltzmann law. Here, σ_B denotes the Stefan-Boltzmann constant, L_{out} is the longwave outgoing radiation, L_{in} is the longwave incoming radiation, and ϵ is the emissivity of the surface, here set to 0.98, which is a common value for green vegetation (Humes et al., 1994). The correlation analysis here is based on

the growing season only, thus minimising a potential effect of the seasonality of the emissivity.

For the definition of the growing season (or carbon uptake period) several methods for the extraction of the growing season from GPP time-series are available. These methods include curve fitting with logistic functions of the time series and the use of derivatives of the smoothed time-series. The use of simple thresholds has proved to be well-suited and straightforward for this kind of application, particularly when the growing season is extracted from a GPP time series (e.g. Richardson et al., 2010). Here, we scaled GPP, such that the 0.05 quantile was zero and the 0.95 quantile was equal to one. Then we extracted all the values between the first and the last days in which GPP was higher than 0.2 (i.e. 20 % of the 0.95 quantile–0.05 quantile range). The threshold of 20 % was considered as a well-suited threshold, and it is similar to values reported in literature for similar application (e.g. Richardson et al., 2010 uses 25 % of all GPP data).

Maximum leaf area index (LAI) and soil organic carbon (SOC) data were used in the analysis of differences in air or soil temperatures for flux partitioning, when these data were available on www.fluxdata.org.

A detailed set of soil temperature observations at Hyytiälä in Finland, including 5 measurement depths at 5 locations around the eddy-covariance tower, was used to investigate the influence of measurement depth and the number of soil temperature sensors.

2.2 Flux partitioning algorithms

Flux partitioning algorithms divide the net carbon exchange into photosynthetic uptake and respiratory release based on the fact that photosynthesis is zero during the night, i.e. without light. Most algorithms rely on simple empirical models, but neural network approaches or more complex land surface models have also been applied (Desai et al., 2008). Here two widely used approaches are included: one that fits nighttime NEE data with a respiration model and another that avoids nighttime data and uses

a combination of a light response curve and a respiration model to fit daytime NEE. Both flux partitioning algorithms described here are also available as online tools at http://www.bgc-jena.mpg.de/~MDIwork/eddyproc/.

2.2.1 Nighttime data based estimate

The first flux partitioning algorithm follows the description in Reichstein et al. (2005). The algorithm fits a respiration model to the measured nighttime NEE data and extrapolates the optimised model to daytime periods using the respective temperature observations of the day. An Arrhenius-type model after Lloyd and Taylor (1994) is used to derive and extrapolate the temperature dependence of R_{eco}:

$$R_{eco} = r_b \exp \left(E_0 \left(\frac{1}{T_{ref} - T_0} - \frac{1}{T_{obs} - T_0} \right) \right) \tag{2}$$

where r_b [µmol C m^{-2} s^{-1}] is the base respiration at the reference temperature T_{ref} [°C], set to 15°C), E_0 [°C] the temperature sensitivity, T_{obs} [°C] is the observed temperature and parameter T_0 [°C] is set to -46.02°C as in Lloyd and Taylor (1994). For E_0 a constant value for each year is derived while r_b was estimated every 5 days using a 15 days window (as in Reichstein et al., 2005). The difference between modelled R_{eco} and measured NEE yields the estimate for GPP. For this algorithm, the absolute flux magnitude and therefore the parameter r_b is mainly determined by the mean nighttime flux.

2.2.2 Daytime data based estimate

The daytime data based estimate uses an approach described in detail in Lasslop et al. (2010). It is based on a light response curve extended to account for the temperature sensitivity of respiration by including the Lloyd and Taylor model and a drought limitation

of GPP.

$$NEE = -\frac{\alpha \beta R_g}{\alpha R_g + \beta} + r_b \exp\left(E_0 \left(\frac{1}{T_{ref} - T_0} - \frac{1}{T_{obs} - T_0}\right)\right) \qquad (3)$$

where α [µmol C J^{-1}] is the canopy light utilisation efficiency and represents the initial slope of the light response curve, β [µmol C m^{-2} s^{-1}] is the maximum CO$_2$ uptake rate of the canopy at light saturation (see Eq. 4), and Rg is the global radiation [W m^{-2}]. The limitation of GPP due to high water vapour pressure deficit (VPD) is included by reducing the maximum carbon uptake with an exponential function for VPD values higher than 10 hPa (VPD$_0$). The strength of this decrease is determined by the parameter k.

$$\beta = \begin{cases} \beta_0 \cdot \exp\left(-k\left(VPD - VPD_0\right)\right) & \text{if } VPD > VPD_0 \\ \beta_0 & \text{otherwise} \end{cases} \qquad (4)$$

T_{ref} and T_0 were fixed as in the nighttime data based approach. The parameter E_0 was estimated using nighttime data of 12 days ($R_g < 4\,\text{W m}^{-2}$), then E_0 was fixed and r_b, α, β and k were derived from daytime data using a 4 days window. The estimation was repeated every second day and yields time series of the empirical ecosystem parameters. The r_b parameter is mainly determined by the morning and evening observations as they have the strongest influence in the model range where radiation, and therefore the contribution of GPP, approaches zero.

2.3 Statistical analysis

The correlation between the different temperatures and nighttime NEE was performed for 12 days moving windows. This is the window size that was used in the Lasslop et al. (2010) approach for the estimation of the temperature sensitivity and is similar to the 15 days used in Reichstein et al. (2005), a scale in which the spectra of fluxes show a minimum (Stoy et al., 2009). For each time window we computed the correlation between nighttime NEE and the temperature. We used T_{air}, T_{soil}, the average of the two,

T_{mean}, T_{surf} and an optimized temperature (T_{opt}). T_{opt} was computed as the weighted average of T_{soil} and T_{air}, with a weight optimised to maximize the correlation with night-time NEE.

The use of T_{opt} introduced an additional parameter compared to the use of T_{air} or T_{soil}. Therefore, for the selection of the temperature driver that best explains the variability of night-time NEE (i.e. the best linear model that relates NEE with T_{air}, T_{soil} or T_{opt}) we used the Akaike's Information Criterion (AIC) (Akaike, 1973). AIC is a well-known metric for model selection (Anderson et al., 2000). AIC is a measure of the trade-off between the goodness-of-fit (model explanatory power) and model complexity (number of parameters). Hence, using AIC we can evaluate if the increase in model performance introduced by adding an additional parameter is overcome by an introduction of additional uncertainty.

$$AIC = n \cdot \log(RSS/n) + 2 \cdot (p + 1) \tag{5}$$

where RSS is the residual sum of squares, p is the number of parameters and n is the sample size. Given a dataset, several competing models (e.g. in our study different models including different temperature drivers, T_{air}, T_{soil} and T_{opt}) can be ranked according to their AIC. The model having the lowest AIC is considered the best.

For the flux partitioning the software package PV-Wave (Visual Numerics, 2005) was used. For the statistical analysis the software packages R (R Development Core Team, 2010), and Matlab were used. The lag between the different temperatures was computed by using a cross-correlation function. The time lag was identified when the correlation was at a maximum within a window of 24 h. The degree of dampening was computed as 1 minus the slope of the regression between T_{air} and T_{soil}.

3 Results and discussion

3.1 Effects of phase shift and dampening in temperature data on the model fit

The general understanding of how air temperature relates to soil temperature is that the amplitudes of soil temperature are dampened and its phase is shifted against air tem-
perature due to heat transport processes (Hillel, 2004; Monteith and Unsworth, 1996). Figure 1 shows the effects of dampening and phase shifts on the model fit based on synthetic data. The synthetic respiration rate is the output of the Lloyd and Taylor model driven by the original synthetic temperature time series. If the temperature is only dampened, the two temperatures could still be used as proxies for each other and
would only result in different estimates of the temperature sensitivity and base respira-
tion, while the modelled respiration would be almost identical. If the two temperatures are lagged not only are the parameters affected; modelled respiration is lagged (Subke and Bahn, 2010) and, the amplitude is decreased. For most of the FLUXNET sites used here we found a time lag of less than 12 h (Fig. 2a). The dampening is considerable for
most of the sites (Fig. 2b). Due to the frequent occurrence of lags across FLUXNET sites, differences between flux partitioning estimates using one or the other tempera-
ture can be expected. As these lags can propagate into the GPP and R_{eco} estimates, they should be used carefully when exploring lags between R_{eco} and GPP or between the estimates and environmental drivers. In this analysis the maximim acceptable lag
was set to 24 h. Without this limitation only a few sites/years showed a lag longer than 24 h. These were sites/years where temperature was poorly correlated with R_{eco}. This phase shift directly propagates into the flux partitioning estimates. Therefore there is high uncertainty at the hourly timescale but more confidence in estimates using aggre-
gated data (i.e. daily or monthly) when analysing lags or correlations with other fluxes
or environmental drivers.

3.2 Correlation between nighttime NEE and temperature

During the nighttime, NEE equals ecosystem respiration as photosynthesis is zero in the absence of photosynthetically-active radiation (PAR). The linear correlation between nighttime NEE and temperature is a first indication of the explanatory power of temperature as the main driver in a respiration model. Individual temperature data streams cannot perfectly represent the variability of respiration within an ecosystem. In the correlation analysis we tried to identify the temperature that is the best proxy for mapping the variability of the ecosystem respiration. We compared the median of the correlations within a 12-day time window of nighttime NEE with temperature observations, i.e. T_{soil}, T_{air}, the average of the two, T_{mean}, and T_{opt} allowing improvement of the model performance with a weight parameter (Fig. 3) during the growing season. The correlation between nighttime NEE and temperature shows little difference between the different temperature observations. The distribution is bimodal with mostly positive correlations, but negative correlations also exist. All sites with negative correlations either have a subtropical Mediterranean climate or a climate characterised by hot warm summers. This suggests that in these sites the NEE – temperature relationship is strongly influenced by drought effects. For these sites the dynamics of the estimated respiration are not reliable within the 12-day time window as temperature sensitivity is forced to be positive in both algorithms considered here. It is likely that moisture affects the relation between NEE and temperature either directly by limiting respiration or by changing the transport processes of heat and CO$_2$. Leaf fall has been found to change the sign of the correlation between NEE and temperature by strongly increasing substrate availability (Pilegaard et al., 2011). Although the difference in the mean correlation of soil and air temperature is small, we observed a significantly stronger correlation with T_{air} in the FLUXNET sites with positive correlations. On the other hand, the variation of the correlation coefficients between sites is smaller for T_{soil}. The largest part of ecosystem respiration is soil respiration, which usually contributes more than 60 % (Bolstad et al., 2004; Janssens et al., 2001; Zha et al., 2006), it is surprising that the correlation with

air temperature is not significantly lower than the correlation with soil temperature. A reason why this is not captured in our correlation analysis might be the variability of temperature in the soil and the lags due to transport. During winter the correlation with T_{soil} is expected to be higher as the vegetation is inactive, and a larger part of the respi-
ration takes place in the soil (Kolari et al., 2009). The highest median of the correlations across sites with nighttime NEE was observed for the optimised temperature T_{opt}. The optimised temperature includes an additional parameter which also introduces additional uncertainty. In spite of the penalty in the calculation of the AIC score for the additional parameter, the optimised temperature still performs slightly better compared
to only T_{soil} or only T_{air} (Fig. 4), as the lower the Akaike criterion, the better the performance. For a subset of sites, observations of the longwave radiation were available and T_{surf} could be considered. For this subset, we again observed the optimised temperature had the greatest correlation with nighttime NEE (median $= 0.367$); the differences in the correlation between the other temperature streams were not significant.
For T_{mean} the median correlation was 0.346, followed by T_{surf} derived from longwave radiation (0.339) (no figure). T_{soil}, for this subset, showed a stronger correlation (0.332) than the air temperature (0.325). The correlation coefficients were very similar, and the mean of the correlations were not significantly different for this subset of sites.

The correlations do not clearly distinguish between air and soil temperature. To gain
additional insight we analysed the weights of the optimised temperature. To remove the effect of the different variances of soil and air temperature we normalised the temperature time series before computing the weights. The weights for the soil temperature are then clearly lower than 0.5 suggesting that the air temperature time series contributed more strongly to the optimised temperature (Fig. 5). For more specific studies,
for instance using only specific vegetation types, using soil temperature could be advantageous as the correlation with soil temperature is higher for deciduous broadleaf forests, for example, while for grasslands or wetlands the correlation is greater with air temperature (see Fig. 6). Strong negative correlations occur less often with T_{soil}. Overall, the correlation analysis suggests that neither air nor soil temperature is clearly

better suited, but analysing the correlation of nighttime NEE with temperature can yield results indicating one is better than the other for specific datasets or sites. The higher or lower correlations cannot be interpreted as there being a greater or lesser contribution of the one or other ecosystem component to the total flux. However, in flux partitioning or gap filling procedures the temperature which explains more of the variability in an eddy-covariance time series should be considered the most useful.

Soil is a very heterogeneous part of the ecosystem, and the representativeness of soil measurements can be a limiting factor. We performed an additional correlation analysis for one site, Hyytiälä, where observations of soil temperature at five observation depths at five locations around the tower were provided. The analysis showed that the correlation between temperature and nighttime NEE decreases on average with increasing observation depth (see Fig. 7). The differences between the correlations at the first two depths are already statistically significant ($p = 0.03$). This supports our decision to always use the upper soil temperature observation and emphasises the importance of measurements in the very top layer (2 cm). For many sites the uppermost soil temperature measured at 5 cm depth (see Appendix A). As the measurement depth is not completely harmonised within FLUXNET, differences in the soil temperature measurement depth can explain part of the spread of the correlations between nighttime NEE and temperature across sites. The correlations of temperature measurements with nighttime NEE from different locations (at a similar depth) are very similar in their magnitude and variability (see Fig. 7b). Representativeness of the soil temperature measurements is therefore not a critical issue at this site.

3.3 Difference in flux components derived using air and soil temperature

Both flux partitioning algorithms were run for the FLUXNET data using either air or soil temperature. When comparing the median of the differences in the annual estimates of the flux components derived with air or soil temperature the two algorithms applied differ: while for the Lasslop et al. (2010) approach only the median of the differences in GPP estimates across FLUXNET sites differed significantly from zero, for the

Reichstein et al. (2005) approach both flux components showed a small difference of ≈ 20 g C m^{-2} yr^{-1} (Fig. 8). The estimates using T_{air} were higher. The nighttime and daytime data based estimates were compared in a previous study (Lasslop et al., 2010). The range of these deviations due to the choice of the algorithm were about twice as high as the differences found here due to the choice of the temperature observations, however, the mean difference between the algorithms was not significant across the FLUXNET sites.

These GPP datasets were the basis for global estimates derived by upscaling eddy covariance data to global gridded estimates. Assuming a constant bias of 25 g C m^{-2} yr^{-1} for all sites a rough estimate for the effect of the choice of the temperature stream can be estimated as 3.2 Pg C yr^{-1} for the global GPP of 123 Pg C yr^{-1} (Beer et al., 2010). This is a conservative estimate as one would expect that the true value is between the two estimates.

The different outcomes of the two algorithms were further analysed by means of the averaged diurnal cycles. For the Lasslop et al. (2010) approach for R_{eco} the positive differences during nighttime and the negative differences during daytime cancelled out. The GPP differences of the same approach were zero at night; during day the differences were negative on average and summed up (Fig. 9). For the Reichstein et al. (2005) approach the residuals for both estimates were zero on average during night and the mostly negative differences during daytime summed up and caused the difference in the annual estimates. The methodological difference between the algorithms that can explain this effect is the estimation of the base respiration parameter, that determines the flux magnitude at the reference temperature. While the nighttime based approach fits this parameter with all the nighttime data, this parameter is most strongly determined by morning and evening observations for the daytime based approach. During morning and evening hours the temperature is closer to the mean temperature, while the nighttime values are at the lower range of observed temperature values. Thus, the temperature sensitivity parameter of the nighttime data based approach is used to extrapolate only to higher temperatures while the flux magnitude is fixed such

that the mean nighttime flux equals the respiration flux during night. For the daytime data based approach the respiration flux magnitude is mainly determined by the morning and evening values and the algorithm relies on the temperature sensitivity estimate to compute fluxes with lower temperature during night, but also for higher temperatures during daytime. Thus, in this approach when a bias in the temperature sensitivity parameter occurs, the errors during the cooler night can be cancelled out by estimates for the warmer day. On the other hand observations during morning and evening have a high relative uncertainty (absolute uncertainty/flux magnitude), and can be an explanation for the higher MAD of the differences in annual R_{eco} (Fig. 8) and the higher spread of the differenced diurnal cycle. The half hourly values of the averaged diurnal cycle show differences up to 1.5 µmol C m^{-2} s^{-1} between the soil and air temperature estimate which is consistent with other studies (Kolari et al., 2009; Ibrom et al., 2006).

3.4 Inter site patterns in the difference of the estimates

Here we analysed whether parameters that are related to the decoupling of soil and atmosphere can explain the variability of the effect that the choice of temperature has on the result of the flux partitioning algorithm. As the algorithms are exactly the same and only the temperature input is changing, the relationship between air and soil temperature could be an indicator for the magnitude of the difference. To characterise the relationship between air and soil temperature we used the lag between the two. The relation between the differences and the temperature lag was stronger for the nighttime based method (Fig. 10). The correlation was less significant for the daytime based method. The relationship for the daytime data based R_{eco} was of opposite direction. The flux components of the daytime based method are less strongly related as GPP is not computed as the residual but is based on the light response. Including this additional model seems to make the method less prone to biases. Other indicators for the strength of the decoupling between soil and air temperatures are the maximum LAI (LAI$_{max}$) of the site and the soil organic carbon content (SOC). A high LAI decouples air and soil processes and dampens the differences in soil temperature (Zheng et al.,

1993), while a very low LAI and direct insulation can heat up the soil faster than the surrounding air. Soil organic carbon has a low heat conductivity and high heat capacity, which can increase the lag between air and soil temperature. Therefore we would expect a correlation between values of LAI$_{max}$ or SOC and the differences in GPP and

5   R_{eco} due to the choice of the temperature stream. The correlation between LAI$_{max}$ was of similar magnitude as for the temperature lag (Fig. 11). For the SOC the correlation was not significant (not shown). Unfortunately the high heterogeneity of the soil and the not fully harmonised measurement protocols decrease the comparability across sites and might obscure existing patterns.

10   When separating the differences into vegetation types we found a general tendency for the difference to increase with increasing vegetation height (see Fig. 12). The pattern was less clear for the Lasslop et al. (2010) approach.

3.5 Influence on ecosystem parameters

As eddy-covariance data are increasingly used to derive ecosystem parameters or to

15 constrain and evaluate process based land surface models for the water and carbon cycle (Bonan et al., 2011; Knorr and Kattge, 2005; Mahecha et al., 2010; Reichstein et al., 2003b), we analysed the effects of the choice of temperature for three ecosystem parameters: temperature sensitivity of ecosystem respiration, base respiration and maximum carbon uptake rate. The largest differences were found for the temperature

20 sensitivity (Fig. 13). Because the variance of the air temperature is higher than of soil temperature it yields lower estimates for the temperature sensitivity when fitted to the same NEE data. While in the short term a lower temperature sensitivity estimated for air temperature would be compensated by the higher variability of the air temperature, as in the case of the flux partitioning, a long term trend will increase both temperatures,

25 air and soil, in a similar way. Thus extrapolating this parameter to other time scales, for instance by including this parameter in a process based model for the use in climate projections, requires caution (Mahecha et al., 2011; Graf et al., 2011). The base respiration is significantly higher for soil temperature, which might be due to a parameter

correlation of the base respiration with the temperature sensitivity. The sign of the difference depends on the relation between the mean of the temperature observations used and the reference temperature. It can be minimised by setting the reference temperature to the mean observed temperature.

5   The maximum carbon uptake rate was lower value with the soil temperature estimate, as the GPP estimate computed using soil temperature was lower. This should be considered when optimising or evaluating models with estimates derived with the algorithms presented here.

4   Conclusions

10 In this study we analysed the uncertainty of GPP and R_{eco} estimates caused by the choice between air or soil temperature observations using two commonly used flux partitioning algorithms. The correlation analysis showed that the linear correlation between the different temperature observations and nighttime NEE was similar. However, the correlation with air temperature was slightly but significantly higher across FLUXNET
15 sites. For specifically selected sites soil temperature can explain more of the nighttime NEE variation. The highest explained variance was achieved with an averaged temperature with optimised weights, in spite of the cost of having an additional parameter. We recommend site-specific evaluations to determine which temperature explains most of the variability of the ecosystem respiration (nighttime NEE). As surface temperature
20 showed a good correlation with nighttime NEE, observations of longwave radiation or surface temperature could be a useful extension of the measurements at flux sites.

The differences in R_{eco} and GPP estimates using the Reichstein et al. (2005) method are generally the same because GPP is simply the difference between R_{eco} and NEE. On the annual time scale they show an average difference of 26 g C m^{-2} yr^{-1}. The
25 median of deviations for GPP based on Lasslop et al. (2010) is slightly smaller, and for R_{eco} it is close to zero. The reason could be revealed by analysing the mean diurnal cycles: for the R_{eco} of the Lasslop et al. (2010) method nighttime differences are cancelled

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out by differences during daytime. An estimate of the potential bias, i.e. the difference owing to the choice of temperature, for the global upscaling of eddy-covariance GPP is 3.2 Pg C per year assuming a constant difference of 25 g C m^{-2} yr^{-1}.

The intersite pattern of annual differences can be partly explained by the vegetation structure (e.g. vegetation types, lag between temperatures). The time lag between soil and air temperature was the best indicator for the difference between the estimate using soil or air temperature respectively.

The strongest influence of the choice of the temperature observations for the ecosystem parameter values was on the temperature sensitivity, but all parameters showed a significant difference across sites.

At our current level of understanding of the magnitude of uncertainties in flux partitioning algorithms, we recommend using both soil and air temperature as drivers for partitioning algorithms and including both datasets to account for the uncertainty due to the choice of temperature.

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Table A1. List of FLUXNET sites and years used, reference, values of LAI and SOC included in the analysis, column LW indicates whether longwave radiation was available for the specific site, and d is the measurement depth of the soil temperature in m.

sites	years	reference	LAI	SOC	LW	d
AT-Neu	2003	Wohlfahrt et al. (2008)	6.5	4.25		0.05
AU-Fog	2006	–	–	–	x	
AU-Tum	2003	van Gorsel et al. (2008)	2.4	–		0.02
BE-Lon	2005, 2006	Moureaux et al. (2006)	5.4	3.7		
BE-Vie	97-8,00-03,05-06	Aubinet et al. (2001)	5.1	3.82		
BR-Sa2	2001	Sakai et al. (2004)	4.91	–	x	
BW-Ma1	2000	Veenendaal et al. (2004)	1.3	0.5	x	
CA-Ca1	1999,2002	Humphreys et al. (2006)	8.4	–	x	
CA-Ca2	2001–05	Humphreys et al. (2006)	2.2	–		
CA-Ca3	2002–05	Humphreys et al. (2006)	6.7	–	x	
CA-Let	1999–05	Flanagan et al. (2002)	0.8	–		
CA-Mer	1999–05	Lafleur et al. (2003)	1.3	–	x	0.02
CA-Oas	1997–00,02–05	Black et al. (2000)	2.1	1.63	x	
CA-Obs	2000–03,05	Bergeron et al. (2007)	3.8	–	x	
CA-Ojp	2000–03,05	Howard et al. (2004)	2.6	1.58	x	
CA-Qcu	2002–03,05,06	Giasson et al. (2006)	0.82	–	x	
CA-Qfo	2004	Bergeron et al. (2007)	3.7	3.5	x	
CA-SJ3	2005	Zha et al. (2009)	2.9	–	x	
CA-TP4	2004	Arain and Restrepo-Coupe (2005)	8	3.7	x	0.02
CA-WP1	2004–5	Syed et al. (2006)	2.61	–	x	
CN-Do1	2005	Yan et al. (2008)	5.1	–	x	
CN-Do2	2005	Yan et al. (2008)	3.78	–	x	
CN-Do3	2005	Yan et al. (2008)	1.63	–	x	
DE-Bay	1998–99	Staudt and Foken (2007)	5.3	17.02		0.02
DE-Geb	2004–06	Anthoni et al. (2004)	–	–	x	0.02
DE-Gri	2006	Prescher et al. (2010)	4.93	–	x	
DE-Hai	2001,03	Knohl et al. (2003)	6.08	12.2		0.05
DE-Har	2005–06	Schindler et al. (2006)	–	–	x	
DE-Kli	2005	Prescher et al. (2010)	4.7	9.7	x	
DE-Tha	1998–00,03–06	Grünwald and Bernhofer (2007)	8	16	x	
DE-Wet	2004,06	Rebmann et al. (2010)	4.78	–	x	0.02
DK-Sor	1997–02,05–06	Pilegaard et al. (2003)	5	–	x	0.02
ES-ES1	1999–00,05–06	Reichstein et al. (2005)	2.63	–		
ES-ES2	2006	–	3.015	–	x	
ES-LMa	2004–05	–	2	3.32	x	
ES-VDA	2004	Gilmanov et al. (2007)	1.35			
FI-Hyy	97,99,01–02,04	Suni et al. (2003b)	6.7	5.6		0.01–0.05
FI-Sii	2005	Aurela et al. (2007)	0.4	–		
FI-Sod	2001,05–06	Suni et al. (2003a)	1.2	3.14		
FR-Fon	2006	–	5.055	10.2		
FR-Hes	1998,03–06	Granier et al. (2000)	6.7	7.17	x	0.05

Table A1. Continued.

sites	years	reference	LAI	SOC	LW	d
HU-Bug	2006	Nagy et al. (2007)	1.5	–		
IT-BCi	2005	Reichstein et al. (2003a)	5.8	–		
IT-Cpz	1997,03	Garbulsky et al. (2008)	3.5	4.31		
IT-Ro1	2002–05	Rey et al. (2002)	3.03,2.42	11.3	x	
IT-Ro2	2002–03,06	Tedeschi et al. (2006)	3.9	11.3		
IT-SRo	2001–03,06	Chiesi et al. (2005)	4.2	2.15	x	
JP-Mas	2002–03	Saito et al. (2005)	5.45	–		
JP-Tom	2001–03	Hirano et al. (2003)	9.2	–	x	
NL-Loo	2005–06	Elbers et al. (2011)	2.03	2.4	x	0.03
RU-Fyo	1999–00,03–06	Kurbatova et al. (2008)	2.8	–	x	0.05
SE-Nor	1996–97	Lagergren et al. (2008)	4.53			
UK-Gri	1998	Rebmann et al. (2005)	7	15		
US-ARM	2003–05	Fischer et al. (2007)	2.9,1.6	–	x	
US-Bkg	2005–06	Gilmanov et al. (2005)	3	–	x	
US-Bo1	97–98,00,05–06	Meyers and Hollinger (2004)	5.5	–	x	
US-Bo2	2006	Meyers and Hollinger (2004)	–	–	x	
US-FPe	2004–06	–	2.5	–	x	
US-Goo	2006	–	2	–	x	
US-Ha1	1995,98,01	Urbanski et al. (2007)	4.5	8.8		
US-IB1	2006	Allison et al. (2005)	1.75	6.3		
US-IB2	2005–06	Allison et al. (2005)	3.96,3.03	6.3		
US-KS2	2002,04–05	Powell et al. (2006)	2.5	3.6		
US-MMS	2003–05	Schmid et al. (2000)	4.7,4.8,4.4	6.6	x	0.05
US-MOz	2006	Gu et al. (2006)	4.01	–	x	
US-Me3	2004–05	Vickers et al. (2009)	0.52	10		
US-NR1	2000,02–03	Monson et al. (2002)	5.6	16	x	
US-Ne1	2003	Verma et al. (2005)	2.8	18.4		0.02
US-Ne2	2004	Verma et al. (2005)	4.4	21.1		0.02
US-Ne3	2003–04	Verma et al. (2005)	4.27	–		0.02
US-PFa	1999	Davis et al. (2003)	4.05	20.2		
US-SO2	2004–06	Lipson et al. (2005)	3	0.87		
US-SO3	2005–06	Lipson et al. (2005)	1.1	–		
US-SO4	2004–06	–	3	–		
US-SP1	2005	Powell et al. (2008)	5.5	8		0.1
US-SP2	2001,04	Bracho et al. (2012)	1.05,6.85	–		0.1
US-UMB	2000,02–03	Nave et al. (2011)	4.59,4.26,4.47	3.6		
US-Var	2001–03,06	Xu and Baldocchi (2004)	2.5	–		
US-WBW	1995,99	Wilson and Baldocchi (2001)	5.75	–		
US-WCr	2000,02	Cook et al. (2004)	5.4	9.47	x	

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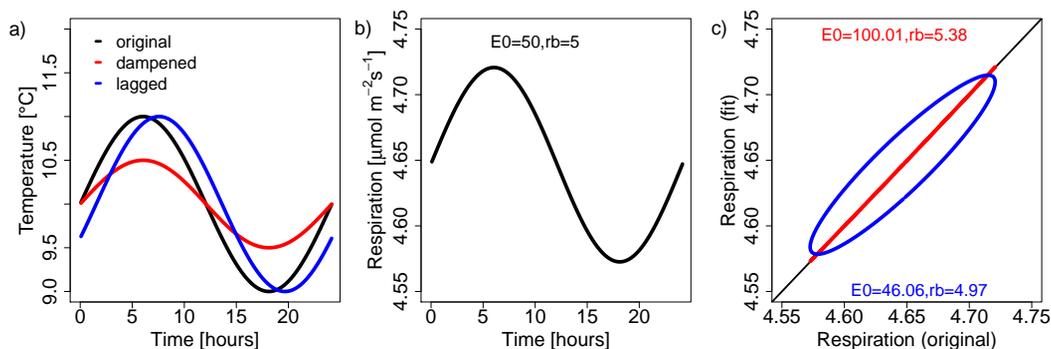
Fig. 1. Illustration of the potential effects of using temperatures that are lagged or dampened on the fit with the respiration model. **(a)** Artificial temperature time series, b: respiration model output [µmol C m^{-2} s^{-1}] with original (see **a**) temperature time series and parameter values used, and **(c)** model output of **(b)** versus the respiration model outputs optimised for the model output of **(b** using the lagged and dampened time series of **(a)** and optimised parameter values.

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Fig. 2. Histograms of lags **(a)** and dampening **(b)** between air and soil temperature (1-slope of the linear regression between the two). Blue vertical lines indicate the median of the distribution. Both lag and dampening was computed using the whole growing season time series. Negative values of the dampening mean the soil temperature has the higher amplitude.
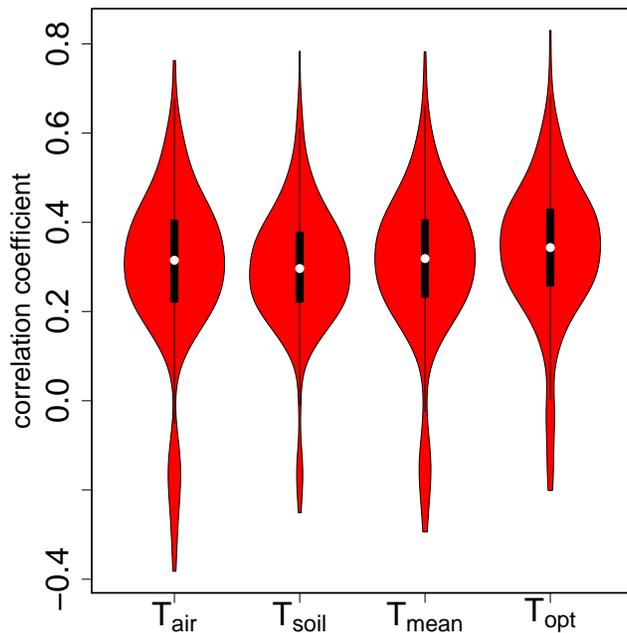
Fig. 3. Correlation with nighttime NEE with T_{air}, T_{soil}, the average of the two, T_{mean}, and an weighted average with optimised weights, T_{opt}, is computed for 12 days time windows. The violin plot shows the distributions of the median of significant correlations for the growing season of each year across FLUXNET sites, the extent in x-direction indicates the frequency of the y-axis value. The point indicates the median, the thick black line is the interquartile range.
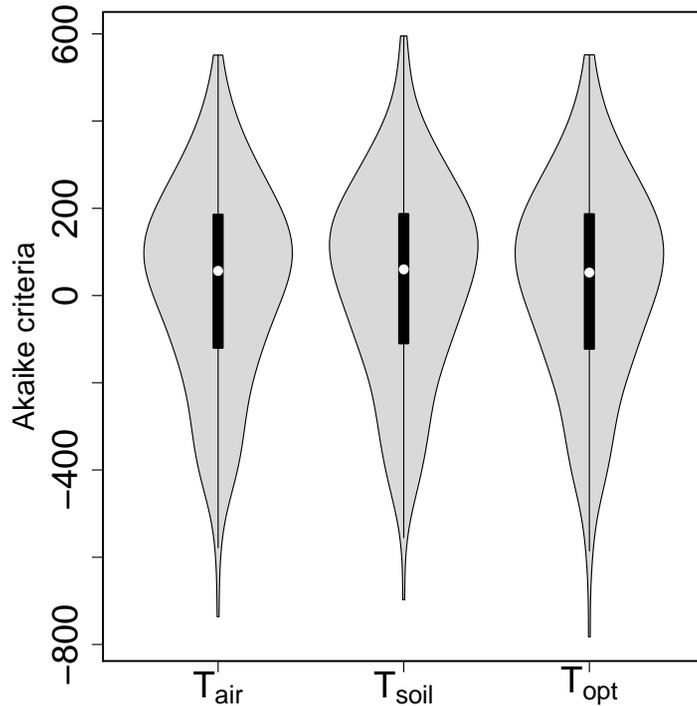
Fig. 4. Median of the Akaike information criterion for 12-days windows with significant correlation with nighttime NEE for all included annual datasets. The point indicates the median, the thick black line is the interquartile range.
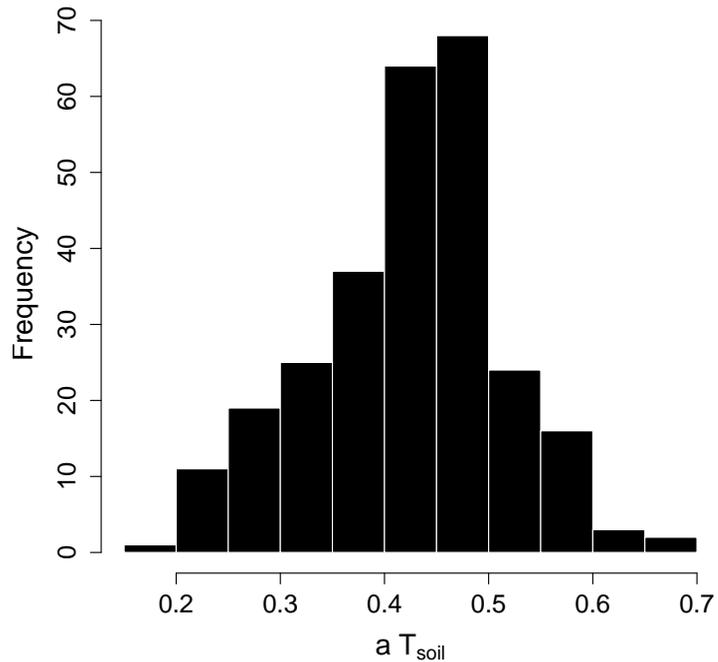
Fig. 5. Median weights of the optimised temperature for the soil part across site and year.
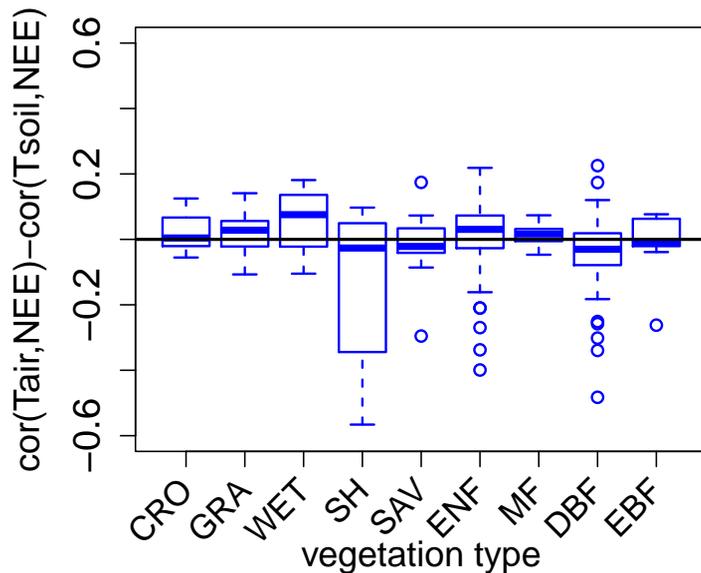
Fig. 6. Differences between the median of significant correlations with air and soil temperature with nighttime NEE for the growing season of one year across FLUXNET sites for different vegetation types. CRO: crops, DBF: deciduous broadleaf forest, EBF: evergreen broadleaf forest, ENF: evergreen needleleaf forest, GRA: grassland, MF: mixed forest, SAV: savanna, SH: shrubs, WET: wetland.
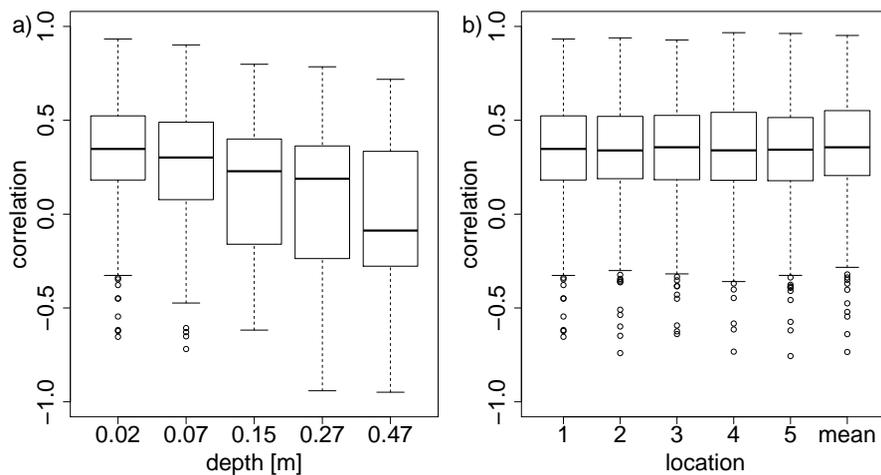
Fig. 7. Correlations between temperature and nighttime NEE at the Finish station Hyytiälä, based on temperature measurements in 5 depths **(a)** and the top measurement (0.02 m) at 5 locations **(b)**.
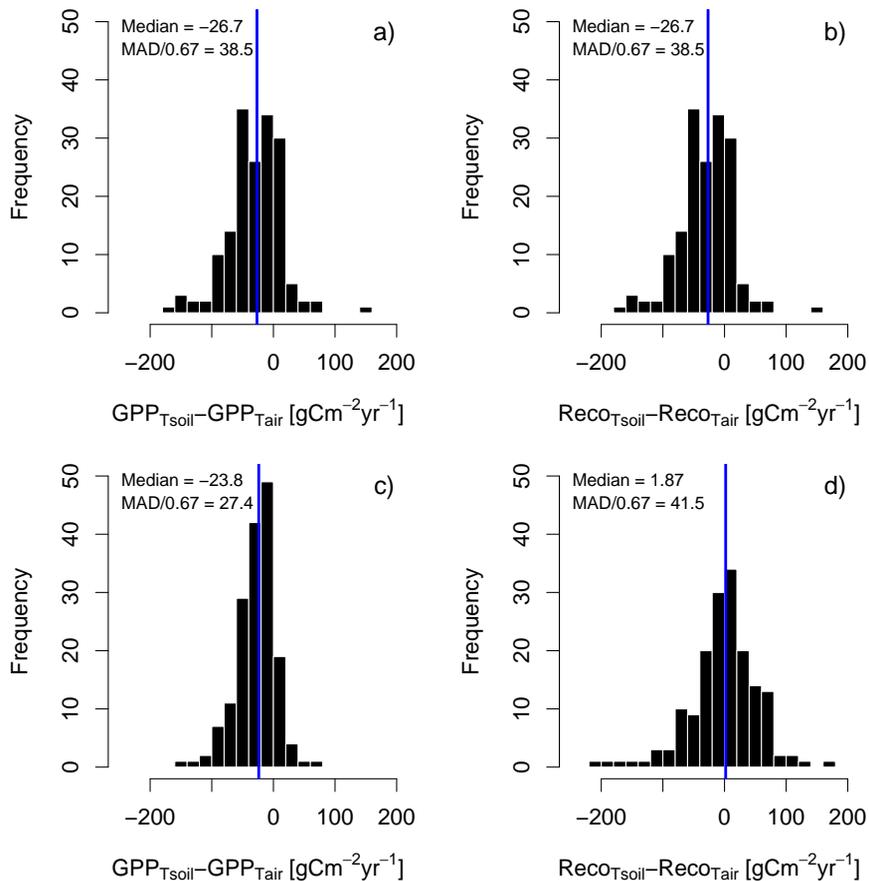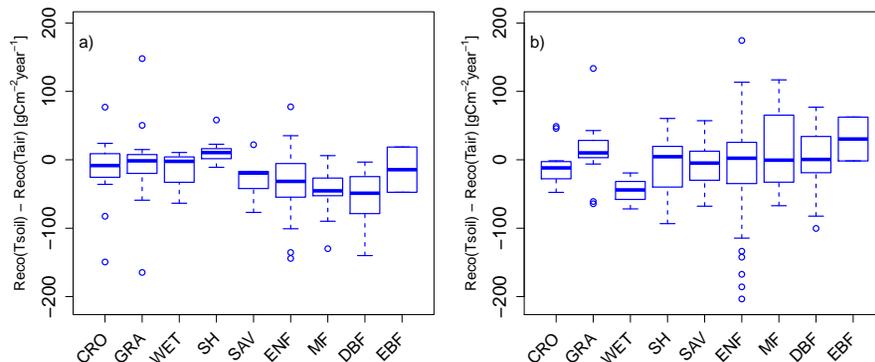
Fig. 8. Difference in annual estimates using the Reichstein et al. (2005) method **(a, b)** and the Lasslop et al. (2010) approach **(c, d)**, blue vertical line indicates the median value, MAD is the median absolute deviation. Note that **(a)** and **(b)** are identical due to the methodology.
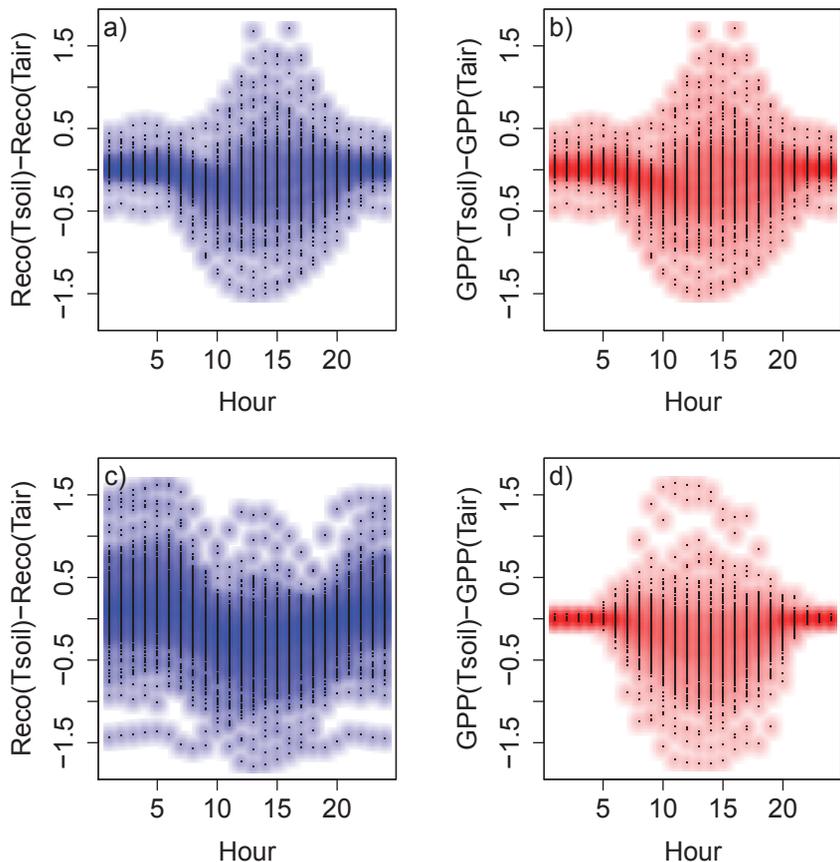
Fig. 9. Mean diurnal cycle of differences [μmol C m^{-2} s^{-1}] between the estimates using soil and air temperature, averaged over the growing season for the Reichstein et al. (2005) **(a, b)** and for the Lasslop et al. (2010) approach **(c, d)**.
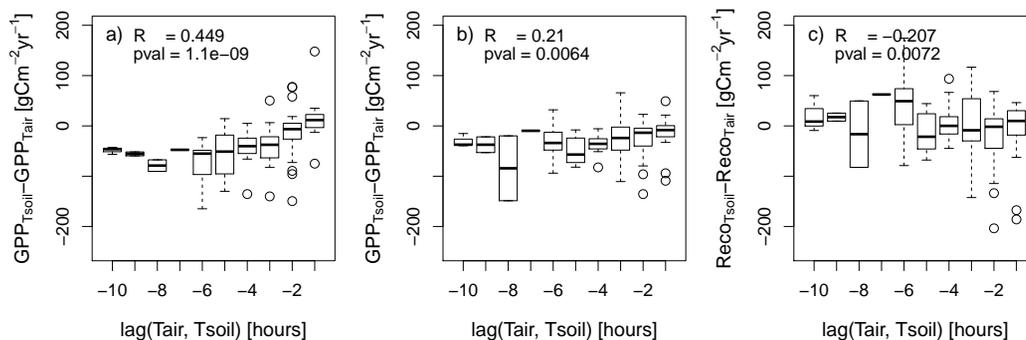
Fig. 10. Annual differences between the carbon fluxes estimated with soil and air temperature versus lag between air and soil temperature for the Reichstein et al. (2005) **(a)** and Lasslop et al. (2010) **(b, c)** approach. Note that for the Reichstein et al. (2005) approach results are the same for both components.
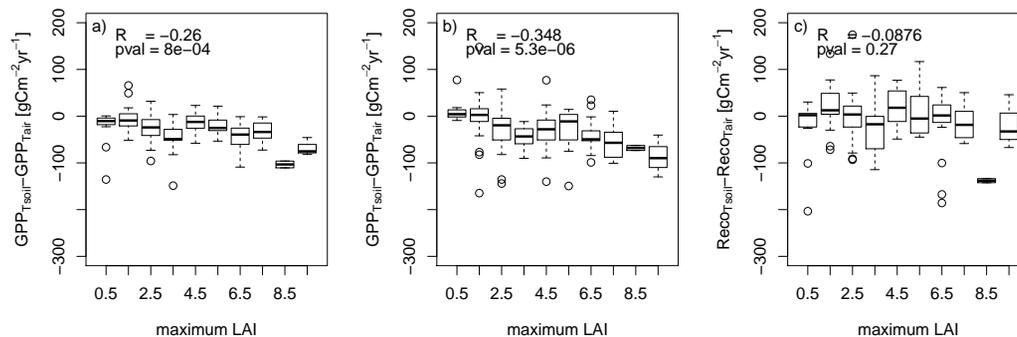
Fig. 11. Difference in annual GPP estimates using the Reichstein et al. (2005) approach **(a)** (note that for the R_{eco} estimates the plot would be exactly the same) and in annual GPP **(b)** and R_{eco} **(c)** estimates using the Lasslop et al. (2010) approach versus the maximum LAI measured at the site.

Fig. 12. Difference in annual R_{eco} estimates using the Reichstein et al. (2005) approach **(a)** and the Lasslop et al. (2010) approach **(b)**, distinguished according to vegetation types (see Fig. 5 for vegetation types definition).

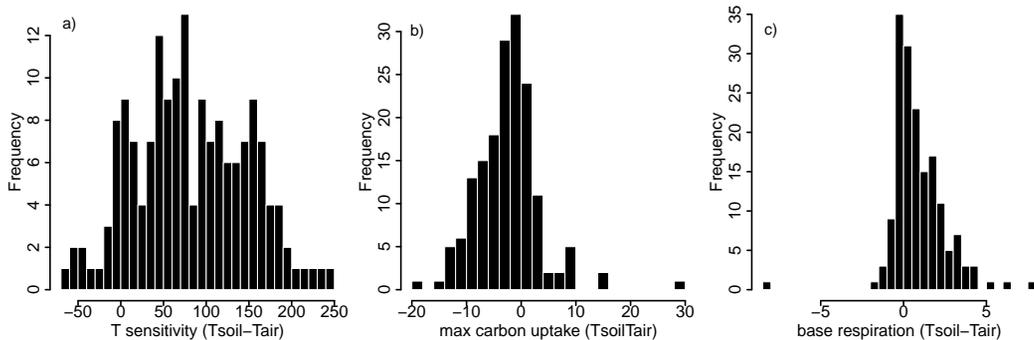**Fig. 13.** Differences in the estimates of the temperature sensitivity [°C] **(a)**, maximum carbon uptake rate [μmol C m^{-2} s^{-1}] **(b)** and base respiration [μmol C m^{-2} s^{-1}] **(c)** using soil and air temperature across FLUXNET sites. The histogram shows the differences of the parameters derived with air and soil temperature across FLUXNET sites.