

Response to Reviewer #2

[Response] We thank the Reviewer for the helpful comments and suggestions, which improved our manuscript significantly. We have addressed all the suggestions and comments in our revision. Please find below the Reviewer's comments (italics), followed by our responses (roman font), with red color indicating relevant changes in the manuscript. We hope that the revised version addresses all the issues raised by the Reviewer.

In this manuscript, Zhang et al. compare the atmosphere-vegetation decoupling factor ω simulated by the land surface model ORCHIDEE with observation-based estimates derived from 106 eddy covariance sites across PFTs. The motivation of this study is valuable as it attempts to evaluate the model performance of simulating an ecosystem property rather than just the simulation of fluxes themselves. This is useful as it allows deeper insights into the functioning of the model compared to more common evaluation approaches. However, I found one critical point that needs to be corrected before the article can be published. In Eq. 3, z should be the reference height (or sensor/measurement height for the eddy covariance towers), not the vegetation height. See e.g. the Monteith and Unsworth 2013 textbook (4th edition, Eq. 17.6) or Liu et al. 2007. To be consistent with the tower observations, z must be set to the height of the sensor at the flux sites in the model. If this is not done a series of biases are introduced which affect the rest of the results. For example, the interpretation of a measured wind speed at a flux site depends on the height where it's measured (logarithmic increase with height), but in the model the assumption is made that the wind speed was always measured at vegetation height, which is incorrect and which causes biases in G_a that affect different PFTs to a different extent.

[Response] Thanks for pointing out this problem in our method. To address this problem, we checked out the FLUXNET dataset and re-run all our simulations at site where the measurement height is available (Table S1). In the new simulations, we kept the distance between canopy top and the measurement height consistent with the observations (Eq. 3). All the results in the manuscript are updated accordingly, using the new simulation outputs. The description of the new simulations has been added to

the revised manuscript, Lines 266-269: “In all the simulations, we kept the distance between measurement height and canopy height consistent with the observations, to ensure unbiased estimates of aerodynamic conductance in the model.”

Our updated results show that the biases in current ORCHIDEE model are not from the biases in measurement height, but from model processes.

A second major point is that more care must be taken in how ‘observations’ of omega are used in such an evaluation exercise. The omega values here are modelled products as well (observation-based at best) which come with a range of assumptions that will affect its interpretation. For example, the canopy boundary layer conductance in de Kauwe et al. 2017 from Thom et al. 1972 was derived for a bean crop but applied to all PFTs in that study. That will inevitably lead to biases in the flux-derived omega values affecting different PFTs to a different extent. Similar issues arise from a non-closure of the energy balance that lead to negative biases in Gs (probably also different in different biomes/PFTs). The authors point to these issues in the discussion, but it would be most useful to the reader if the consequences of these potential biases on the results were elaborated. Just describing that an issue causes a bias is not very useful. It would be much better to show what factors are likely to have major impacts on the comparison by conducting e.g. sensitivity analyses. Knowing that a model-data mismatch could be caused by either the observations or the model is absolutely crucial for such an analysis.

[Response] Thanks for this suggestion. Following the reviewer’s suggestion, we tested how biases in G_a and G_s affect our results (Fig R2).

First, we tested how increasing and decreasing observation-based G_a by 30% affects G_s and omega. We found that perturbing G_a does not result in large changes in G_s . However, using a reference G_a that is 30% smaller than current observation-based estimates, we obtained a better consistency between G_a and omega from ORCHIDEE Ctrl simulation and the reference in forest PFTs. Whereas in short PFTs, decreasing reference G_a to the level of model output (-30%) results in even larger biases in omega, indicating that the large biases in model vegetation coupling strength in short vegetation is not due to uncertainties in the observation-based estimates.

Then we tested how large the energy imbalance affects our evaluation. Using the same method as in De Kauwe et al. (2017), we corrected the energy imbalance and recalculated Gs and omega. The model shows larger Gs biases across all PFTs when

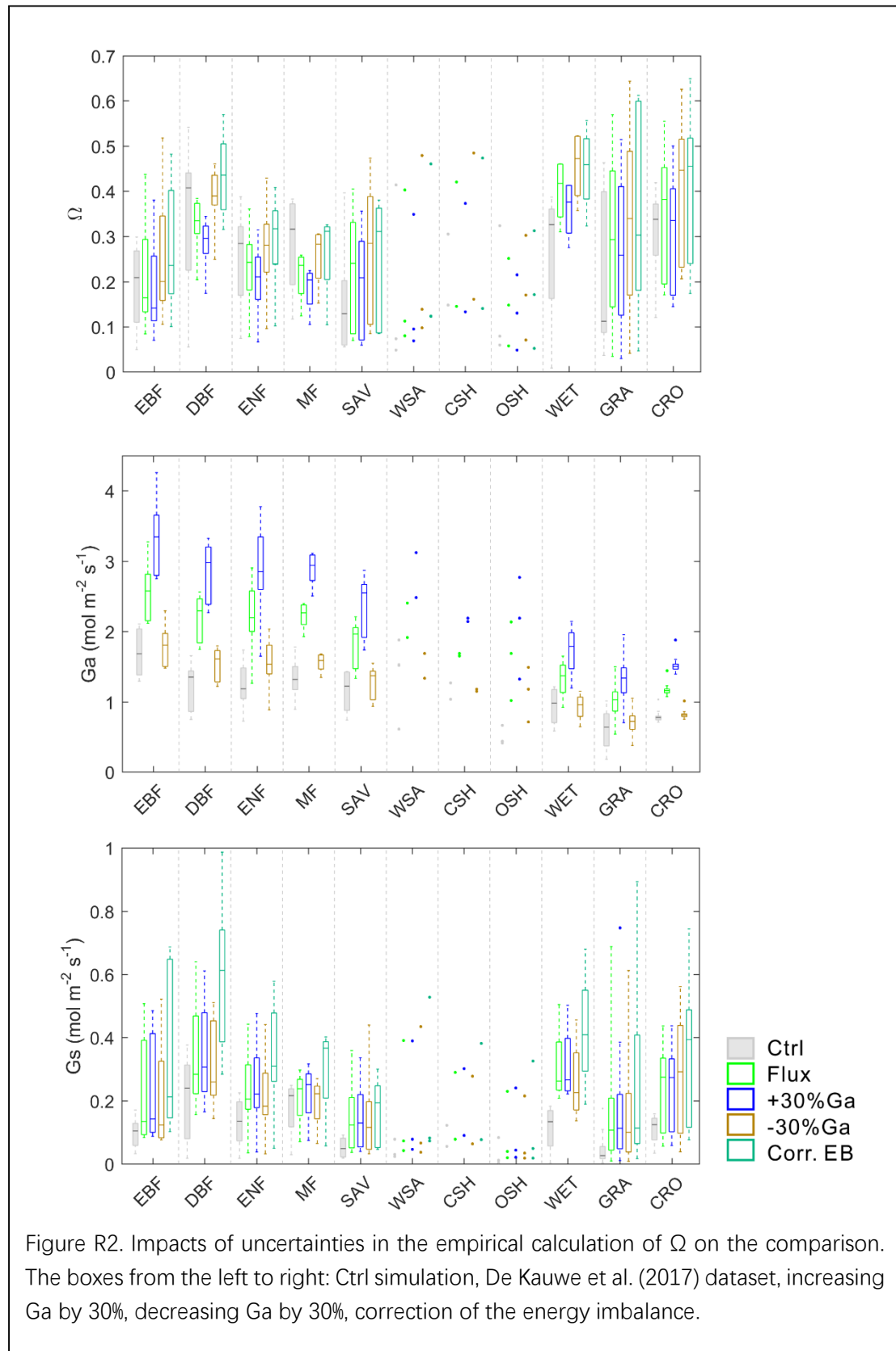


Figure R2. Impacts of uncertainties in the empirical calculation of Ω on the comparison. The boxes from the left to right: Ctrl simulation, De Kauwe et al. (2017) dataset, increasing G_a by 30%, decreasing G_a by 30%, correction of the energy imbalance.

compared with the estimates after correction. Although, the biases in G_s compensate to the existing biases in G_a and result in good performance of omega simulation in forest PFTs.

We have added the new results and discussion to the revised manuscript. Lines 625-634: “In the observation-based estimates, G_a was estimated using an empirical method from Thom et al. (1975), which was derived from a bean crop. G_a estimates from this method are found to be 81%-116% of the estimates of a more physically based method (Knauer et al., 2017) in 6 forest sites. To test how biased G_a affects our evaluation, we increased/decreased G_a by 30% and re-estimated G_s and Ω (Fig S6). We found that perturbing G_a does not result in large changes in G_s . However, when G_a is 30% smaller than current observation-based estimates, we obtained smaller biases in G_a and Ω in ORCHIDEE Ctrl simulation in forest PFTs. Whereas in short PFTs, decreasing the reference G_a results in even larger biases in Ω , indicating that the large biases in model vegetation coupling strength in short vegetation is not due to uncertainties in the observation-based estimates.”

Lines 641-645: “When the energy imbalance is corrected by adjusting the Bowen-ratio following De Kauwe et al. (2017), we obtained larger G_s estimates (Fig S6), resulting in even larger modeled G_s bias than in this study. The increased biases in the corrected G_s compensate for the existing biases in G_a , leading to a “good” performance of omega simulation in forest PFTs.”

Minor comments:

Introduction:

l. 43: add ‘of’ after simulation

l. 45: one cannot add conductances if they are in series, only resistances. Please rephrase.

[Response] Corrected

L. 52: surely there must be more references supporting this statement.

[Response] Thanks for this suggestion, we have added more references to the manuscript.

Methods:

L100: please describe how stomatal conductance (gs) is scaled to the canopy level.

[Response] The description of the stomatal conductance is added. Line 153-154: “The canopy level stomatal conductance is calculated by integrating gs across all leaves in the canopy.”

L 115: please give the equation for z0h here, not only the reference. The z0h to z0m ratio is relevant for the canopy boundary layer conductance and thus Ga. Does the equation imply differences in z0h/z0m across PFTs?

[Response] We have added the equations to the manuscript.

Lines 168-177: “z0h is estimated using z0m :

$$z_{0h} = \frac{z_{0m}}{e^{\kappa B^{-1}}} \quad (6)$$

B is the Stanton number. κB^{-1} is estimated following Su et al. (2001; 2002):

$$\kappa B^{-1} = \frac{\kappa C_d}{4C_t \eta (1 - e^{-\frac{nec}{2}})} f c^2 + 2f c f_s \frac{\kappa \eta \frac{z_{0m}}{z}}{C_t^*} + \kappa B_s^{-1} f_s^2 \quad (7)$$

Where Cd, Ct are drag and heat transfer coefficient of leaves, nec is within canopy wind profile extinction coefficient, calculated as $nec = CdLAI/(2\eta^2)$. fc, fs are the fraction of canopy and bare soil, C_t^* is the heat transfer coefficient of soil. Bs is the Stanton number for bare soil, with κB_s^{-1} estimated following Brutsaert (1999):

$$\kappa B_s^{-1} = 2.46 Re_*^{\frac{1}{4}} - \ln(7.4) \quad (8)$$

Where Re_* is the Reynolds number.”

The z_{0h}/z_{0m} is affected by LAI and canopy height, thus implicitly different across PFTs. In current ORCHIDEE version, the parameters to calculate z_{0h}/z_{0m} is the same for different PFTs.

L 120: add ‘radiation’ after short-wave down and long-wave down

[Response] Added accordingly.

Where in Su et al. 2001 can Eq. 4 be found?

[Response] Eq. 4 in this manuscript is derived from Eq. 10 in Su et al. (2001)

Section 2.2 is inaccurately named as there is basically no information on the FLUXNET data used. I argue that this section needs to be split up: one part giving more

information on the data used (e.g. what sites from what biomes), which could be combined with section 2.3, and then a separate section for the simulation setup.

[Response] Thanks for this suggestion, we have split this section as the reviewer suggested. Now Section 2.2 is “**FLUXNET data and empirical calculation of Ω** ”, and Section 2.3 is “**Simulation setup and modeled Ω calculation**”.

L 152ff: I do not understand why the integrated canopy level stomatal conductance was not used? Surely the canopy-integrated value needs to be used in order to be comparable with the flux-derived omega.

[Response] Here we used the Gs calculated from inverted Penman–Monteith equation because the latent heat flux simulated by ORCHIDEE includes bare soil evaporation and evaporation from the rainfall intercepted by the canopy. Also the observation-based estimates of Gs implicitly include these processes. To be consistent with the reference, we used the same way to calculate modeled Gs.

L. 158: sites

L. 209: observations

L. 214: compensation

[Response] Revised accordingly.

Discussion

L 327: what other factors? Please explain.

[Response] The diurnal change of radiation may strongly affect the coupling strength. We added this into the manuscript. Line 532-534: “**At a sub-daily time scale, this VPD- Ω relationship is not easily observed due to the strong impacts of other factors, such as radiation (Wullschleger et al., 2000; Zhang et al., 2018).**”

L. 370: Limitations

[Response] Corrected.

L. 380: I find this point a bit senseless. If conditions with wind speed don't occur across the data set, why should these conditions matter? They may be of interest from a theoretical point of view but not for LSM evaluation.

[Response] Thanks for this comment. Here we discuss this point because we think that our framework is not only helpful to understand the bias but also has the potential to

calibrate the model. However, there remains observation gaps. In the manuscript, we emphasized this point. Line 612-614: “New observation methods are needed to fill this gap so that future calibrations can ensure the models to correctly simulate vegetation under all different conditions.”

L. 389: what are the uncertainties and how would this affect observation-derived omega values?

[Response] Please see the response to the 2nd major point

L. 397: how and under which conditions would this bias the results? This needs to be explained better.

[Response] The bias is explained in the revised manuscript. Lines 649-652: “A higher surface than ambient air temperature (daytime) tends to overestimate Gs in the inverted Penman-Monteith equation with observed LE, which can further overestimate omega.”