

Criteria of selecting publications

1. New publications were searched on web of science (www.webofknowledge.com), google scholar (<https://scholar.google.com/>) and www.cnki.net (Chinese publications).

2. Carbon use efficiency (CUE = NPP/GPP) is calculated from the observational values or directly taken from publications. NPP and GPP is based on an annual scale, or growing season for Tundra or boreal area assuming very low GPP and NPP in non-growing season.
3. If NPP is not reported, but Ra (autotrophic respiration), CUE is calculated as:

$$CUE = \frac{GPP - Ra}{GPP} = 1 - \frac{Ra}{GPP}$$

4. GPP is based on eddy covariance or component based (Malhi et al., 2015; Xu et al., 2016).

5. NPP is biometric or harvesting (grass/crop) based estimates (Zanotelli et al., 2013), rather than any model based estimate.

6. Sites with multi-year measurements, an observation of each year was taken as independent value (Campioli et al., 2011). If only the average is given, the mean value will be taken or writing to corresponding author for original data.

7. Only eddy covariance derived-GPP is used if multiple approaches report GPP for the same study.

8. If NPP and GPP are graphically presented, (1) writing to corresponding author for original data; (2) if not, grab tool, such as WebPlotDigitizer (<http://arohatgi.info/WebPlotDigitizer/>), will be used, however, there might be a small random error.

20 **Plausibility check**

GPP is separated into different components, shown in Figure S1, suggested by(Van Oijen et al., 2010):

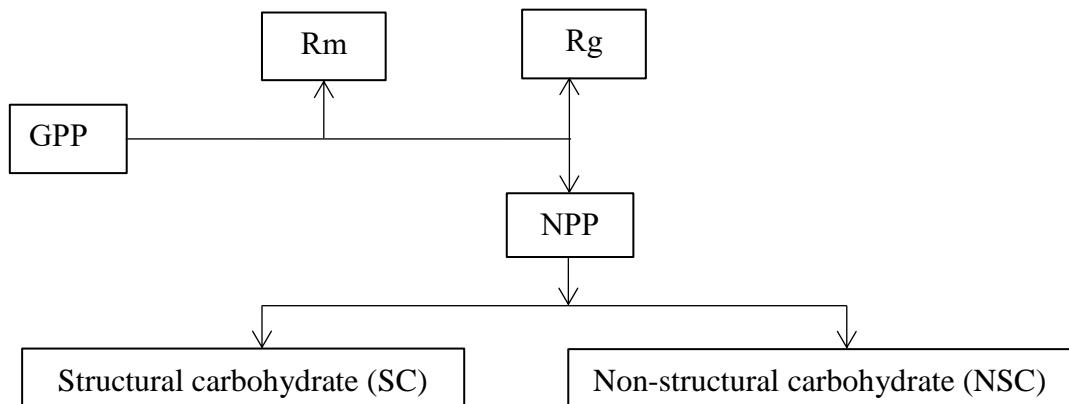


Figure S1. Ecosystem carbon flow in terms of maintenance respiration, growth respiration, allocation of NPP to structural carbohydrate (SC) and non-structural carbohydrate (NSC)

25 The total GPP can be expressed as the sum of respiration and net primary production (NPP):

$$GPP = Rg + Rm + NPP = Rg + Rm + SC + NSC \quad (1)$$

$$NPP = SC + NSC \quad (2)$$

$$CUE = NPP/GPP \quad (3)$$

Assumption: sugars and mineral nutrients are required to produce biomass. Therefore, the growth yield
30 can be expressed by:

$$Yg = \frac{SC}{Rg+SC} \leq 1 \quad \Rightarrow \quad SC = \frac{Rg \cdot Yg}{1-Yg} \quad (4)$$

Yg can be measured in the terrestrial ecosystems; i.e. the amount of structural biomass formed per unit of photosynthates. Yg is independent of environmental conditions(De Vries et al., 1974).

Combining (2), (3) and (4) to introduce the carbon use efficiency (CUE) term:

$$35 \quad NPP = SC + NSC = Yg \cdot \frac{Rg}{1-Yg} + NSC = GPP \cdot CUE \quad (5)$$

Solving Rg:

$$Rg = \frac{(GPP \cdot CUE - NSC)(1-Yg)}{Yg} \quad (6)$$

Dividing GPP:

$$\frac{Rg}{GPP} = \frac{(CUE - NSC/GPP)(1-Yg)}{Yg} \geq 0 \quad (7)$$

40 Defining a unitless term of β :

$$\beta = \frac{NSC}{GPP} \quad (8)$$

If the β term is known, we know how the photosynthesis (GPP) is allocated to NSC.

Since $Yg=[0,1]$

$$0 \leq \frac{(1-Yg)}{Yg} \quad \text{with (7)} \rightarrow CUE - \beta \geq 0 \rightarrow$$

45 $\beta \leq CUE \quad (9)$

$$Rm = GPP - GPP \cdot CUE(NPP) - Rg = GPP(1 - CUE) - \frac{(GPP \cdot CUE - S)(1 - Yg)}{Yg} \quad (10)$$

$$\frac{Rm}{GPP} = 1 - CUE - \frac{(CUE - \beta)(1 - Yg)}{Yg} \geq 0 \quad (11)$$

From the different publications, measured $Rg = 0.8$ is taken as the most favorable value: (Schnapp et al., 1991) (Rambal et al., 2004) (Cannell and Thornley, 2000) (Gifford, 1995) (Lavigne and Ryan, 1997)

50 (Thornley and Cannell, 2000)

Thus, equation (11) can be further simplified:

$$1 - CUE - \frac{(CUE - \beta)}{4} \geq 0 \rightarrow CUE \leq \frac{4 + \beta}{5} \quad (12)$$

Combining (9) and (12):

$$\beta \leq CUE \leq \frac{4 + \beta}{5} \quad (13)$$

55 β , the allocation of GPP to NSC, determines CUE.

In the publications, the NSC is often expressed by percentage, which is:

$$NSC\% = \frac{NSC}{Biomass} \rightarrow NSC = Biomass \times NSC\% \quad (14)$$

For any time t (year, for consistency to GPP),

$$NSC_t = Biomass_t \times NSC\%_t \quad (15)$$

60 And for one year after: $t+1$

$$NSC_{t+1} = Biomass_{t+1} \times NSC\%_{t+1} \quad (16)$$

Therefore, the changes of NSC as a result of photosynthesis (GPP) could be:

$$NSC_{t+1} - NSC_t = Biomass_{t+1} \times NSC\%_{t+1} - Biomass_t \times NSC\%_t \quad (17)$$

It is assumed that NSC% between two near years is the same or similar, thus eq. (17) is further simplified:

$$\Delta \text{NSC} = \text{NSC}_{t+1} - \text{NSC}_t = (\text{Biomass}_{t+1} - \text{Biomass}_t) \times \text{NSC\%}_t = \text{NPP} \times \text{NSC\%} \quad (18)$$

From eq. (8), based on one year scale:

$$\beta = \frac{\text{NSC}}{\text{GPP}} = \frac{\Delta \text{NSC}}{\text{GPP}} = \frac{(\text{Biomass}_{t+1} - \text{Biomass}_t) \times \text{NSC\%}}{\text{GPP}} \quad (19)$$

In view of ecosystem scale, biomass should include different carbon components, such as overstory, understory, roots and litter(Roxburgh et al., 2005). Therefore, since the loss of carbon from other fluxes, such as volatile organic C compounds and root exudations, is difficult to measure and only account a small amount of carbon pool, $\text{Biomass}_{t+1} - \text{Biomass}_t \approx \text{NPP}$. Thus, Eq. 19 can be further simplified:

$$\beta = \frac{\text{NPP} \times \text{NSC\%}}{\text{GPP}} \quad (20)$$

From the physiological view, NPP*NSC% is considered as the amount of GPP to non-structural carbohydrate.

Combining eq. 13,

$$\text{CUE}_{max} = f(\beta_{max}) = f(\text{NSC\%}_{max}) \quad (21)$$

$$\text{CUE}_{min} = f(\beta_{min}) = f(\text{NSC\%}_{min}) \quad (22)$$

From the measured data by averaging for different seasons for different species, NSC% = [2%, 23%], detailed as follows:

15% of the total net radiation for maize and 7% for soybean(Meyers, 2004)

Stem NSC of 85 wood species: S/P=2.4-23% (Poorter and Kitajima, 2007)

Total tree NSC of ten species: 5%-14% (O'Brien et al., 2014)

For all organs of NSC of four species: 5%-20%(Newell et al., 2002)

NSC for shrubs:16%-18.2% (Marquis et al., 1997)

NSC of Douglas-fir trees at different heights: 4%-15%(Woodruff and Meinzer, 2011)

Three Mediterranean species: 2%-15%(Rosas et al., 2013)

Based NPP and GPP dataset of 434 observations,

$$\beta_{max} = 0.2146$$

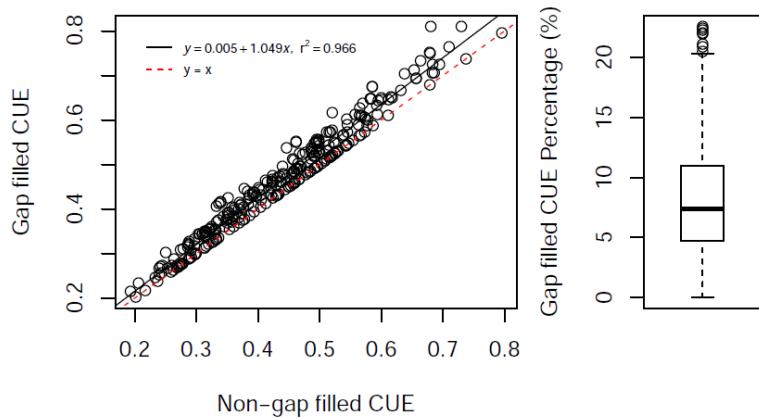
$$\beta_{min} = 0.0038$$

Therefore, $0 \leq \text{CUE} \leq 0.843$

Table S1. Variables used for predicting temporal and spatial CUE.

	Variables	Abbreviation	Type	Type of variability	Sources
Climate	Annual mean temperature	AMT	Split	Yearly	https://crudata.uea.ac.uk/cru/data/hrg/cru_ts_4.01/ , (Harris et al., 2014)
	Annual mean precipitation	AMP	Split	Yearly	
	Potential evapotranspiration	PET	Split	Yearly	
	Diurnal temperature range	DTR	Split	Yearly	
	Vapour-pressure deficit	VPD	Split	Yearly	
	Nitrogen deposition	Ndep	Split	Yearly	
	Sensible heat flux	Heat	Split	Yearly	
Satellite	Palmer Drought Severity Index	PDSI	Split	Yearly	https://www.esrl.noaa.gov/psd/data/gridded/data.pdsi.html (Dai et al., 2004)
	GIMMS leaf area index	GimmsLAI	Split	Yearly	http://sites.bu.edu/cliveg/datacodes/ (Zhu et al., 2013)
	GIMMS normalized difference vegetation index	GimmsNDVI	Split	Yearly	
Soil	GIMMS Fraction of Absorbed Photosynthetically Active Radiation	GimmsfPAR	Split	Yearly	
	Soil carbon stock (0-30 cm)	SoilST	-	Static	https://soilgrids.org/#!/?layer=TAXNWRB_250m (Hengl et al., 2017)
Site	Management	Manag	-	Static	Publications
	Climate zone	CliZ	-	Static	
	Ecosystem type	EcoT	-	Static	

Figures



100 **Figure S2.** Carbon use efficiency (CUE) change after gap-filling in forest ecosystem. The gap-filling process followed Vicca et al. (2012). There was a strong correlation between the gap filled CUE and non-gap filled CUE ($R^2=0.967$). After the gap-filling, the mean CUE increased by 7% and there is no significant impact on CUE trend.

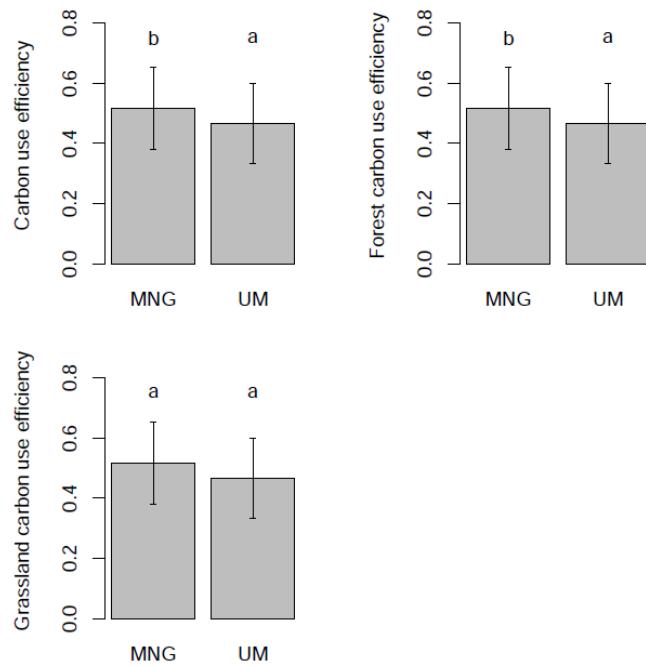
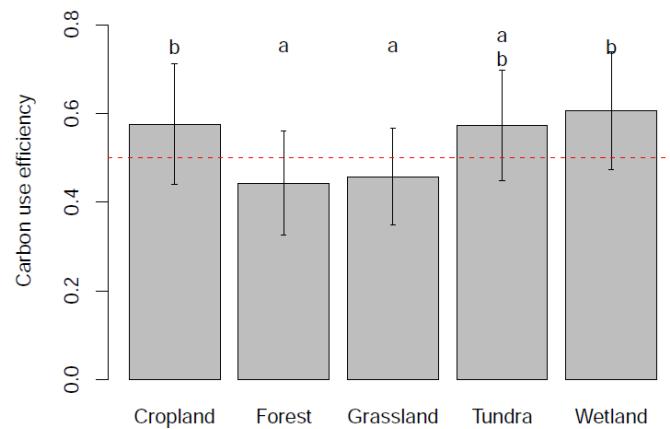
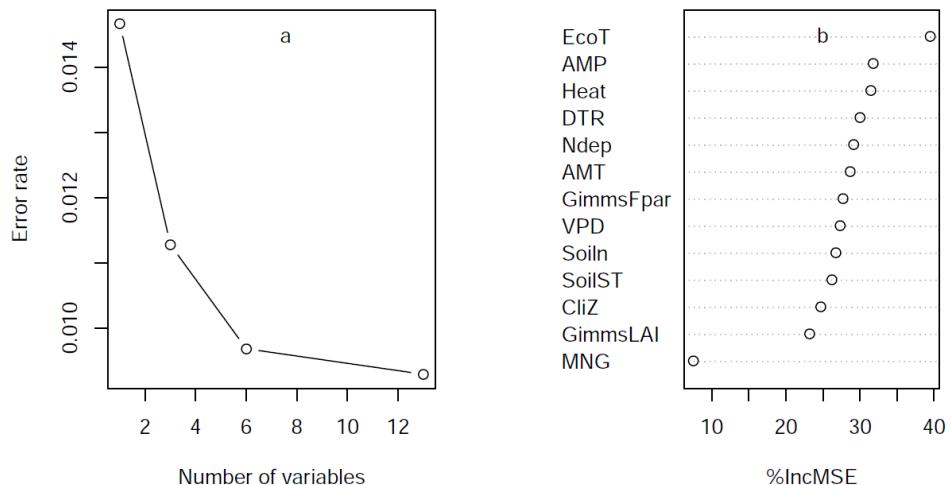


Figure S3. Effects of site management on CUE of all ecosystem types, forest and grass ecosystem. 105 Different letters above the plots indicate significant difference at $p < 0.05$. MNG = managed, UM = unmanaged. Values are mean \pm 1 standard deviation. This result indicated that site management significantly increased CUE in forest ecosystems, but not in grass ecosystems. Therefore, managed forest sites were excluded for spatial modelling (details in Methods).



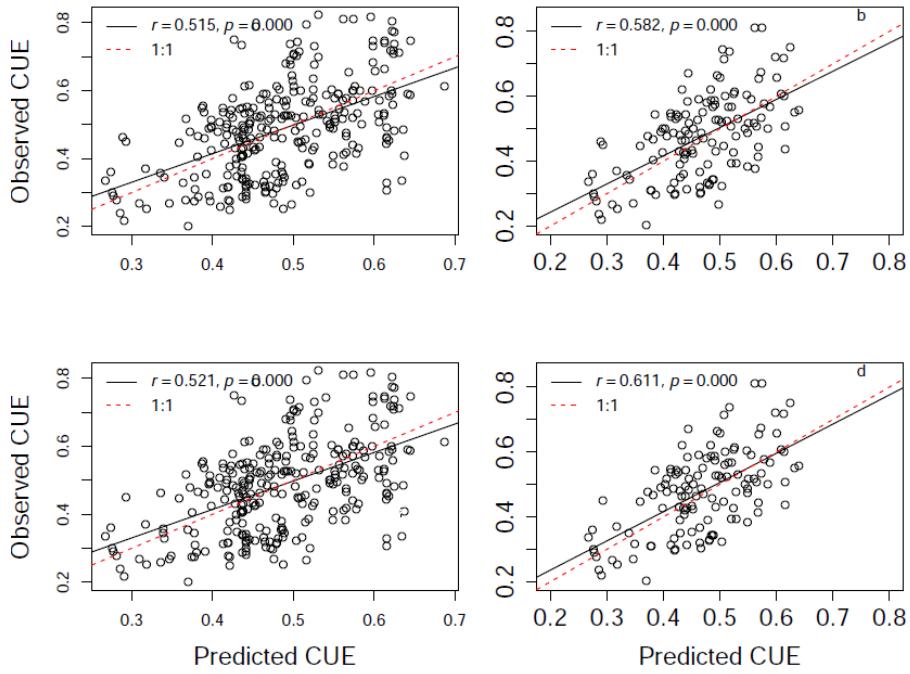
110 **Figure S4.** One-way ANOVA analysis of CUE after removing managed forest sites. Different letters above the plots indicate significant difference at $p < 0.05$. Values are mean ± 1 standard deviation. This results indicated that after removing the managed forest site, the significance of CUE among different ecosystem types did not change (Fig. 1) and removing the managed forest sites for modelling CUE is appropriate.



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Figure S5. (a) The error rate along with the number of the variables used to predict CUE using Random Forest by 10-fold cross-validation, (b) importance values of variables estimated by Random forest expressed by the mean decrease of model accuracy. These selected variables can explain about 51% variance of CUE and the ecosystem type was the most important variables with highest importance value. The higher of %IncMSE is, the higher the variable importance is. After the variables selection, the first six variables with higher importance values were selected to predict CUE and these variables can explain 49% variance of CUE. Noticeably, although management increased CUE, it had little power to explain the variation of CUE analysed by random forest. GIMSS NDVI and PET were removed for modelling due to multi-collinearity.



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Figure S6. (a) Leave-one-site-out, (b) mean-site, (c) leave-one-latitude-out and (d) mean-latitude cross-validation of CUE. Leave-one-site-out cross-validation (leave-one-latitude-out) means leaving all year observations out within the same site (latitude) and predicted by the rest observations using RF for each site (latitude). Mean-site cross-validation (mean-latitude) means that building a RF model using all site-year observations and validating mean-site (mean-latitude) CUE as a new dataset (one mean value for each site (latitude) regardless of the number of the observational years).

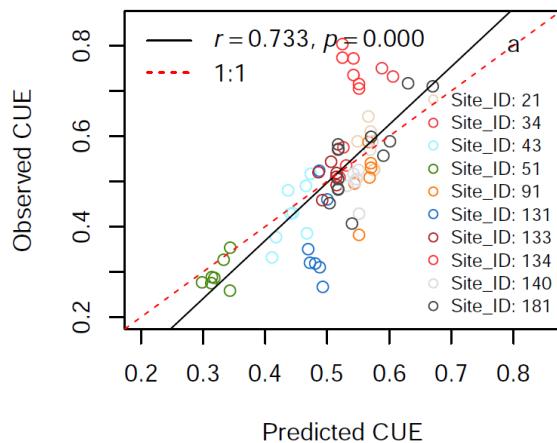


Figure S7. The relationship between CUE extracted from the predicted CUE map and CUE observations for sites with multi-year (longer than four years) observations.

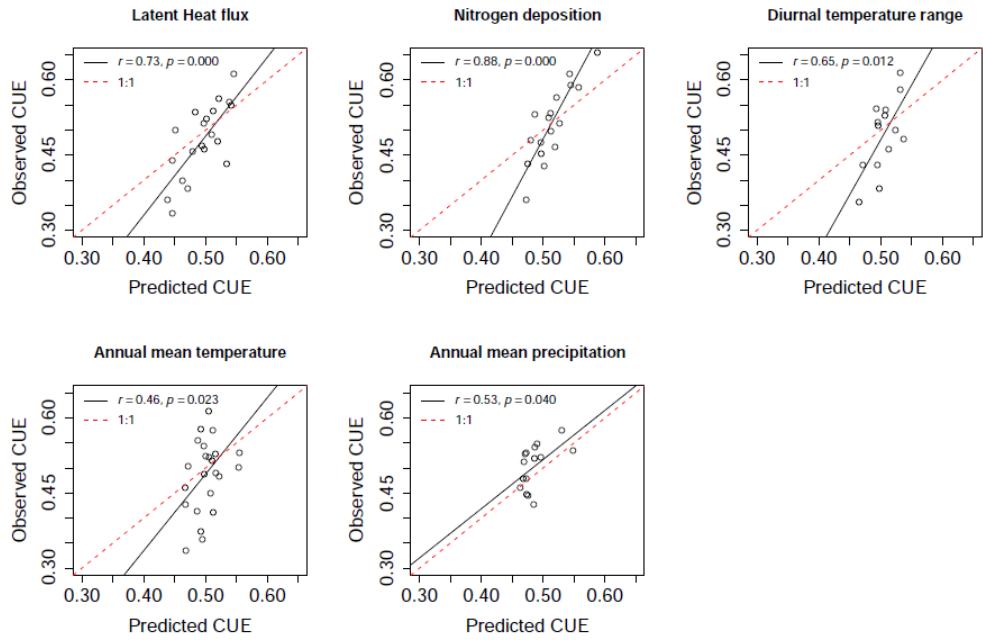
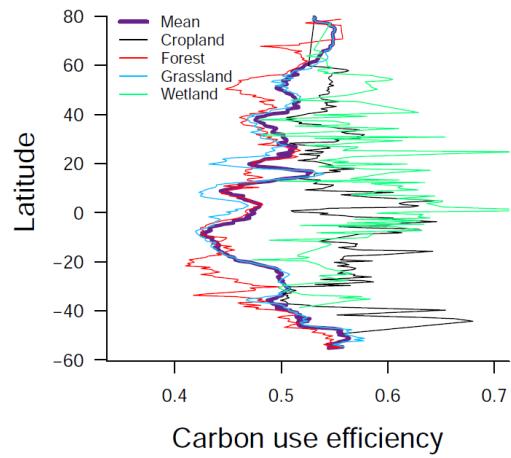


Figure S8. The relationships between CUE extracted from the predicted CUE map and CUE observations within the same selected range of the predictors (latent heat flux, annual mean precipitation, annual mean temperature, nitrogen deposition and diurnal temperature range). First, the values of predictors were divided into different ranges; second, the predicted CUE using RF was masked by those ranges; third, the mean masked predicted CUE and mean observed CUE within the same range were plotted.



145 **Figure S9.** Latitudinal distribution of CUE of forest (red), grass (blue), wetland (green), crop (black) and total mean (purple). Due to the limited number of grids of wetland, this figure may not reflect the “real” latitudinal distribution of wetland CUE.

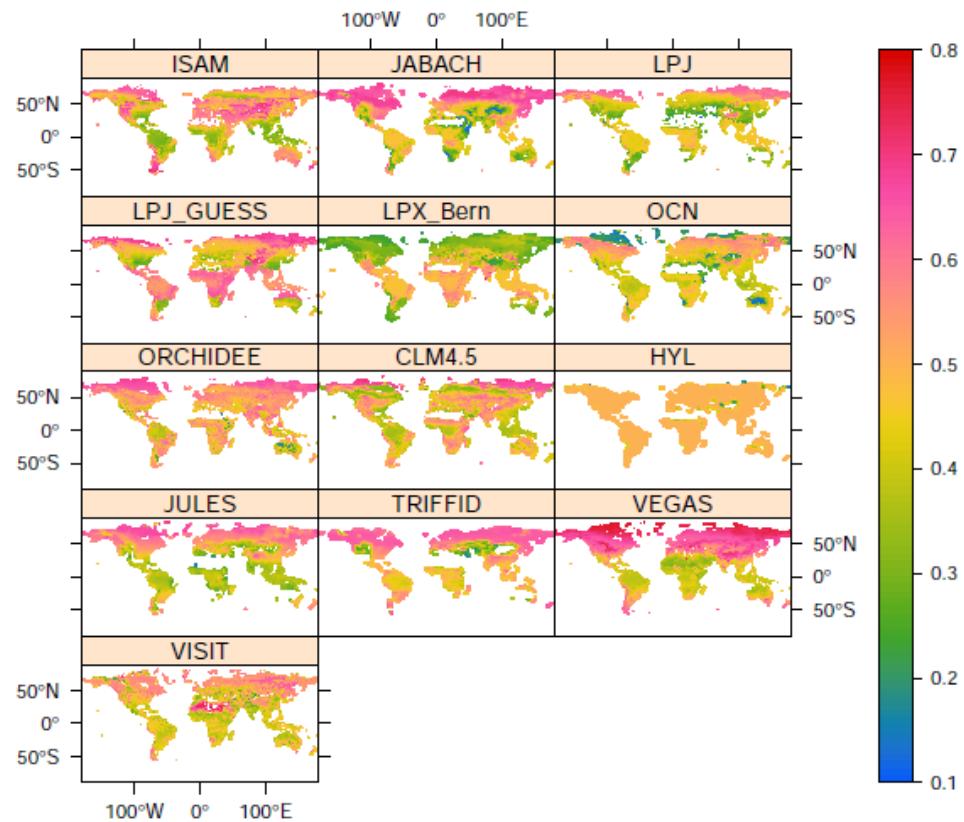


Figure S10. CUE of 13 TRENDY models

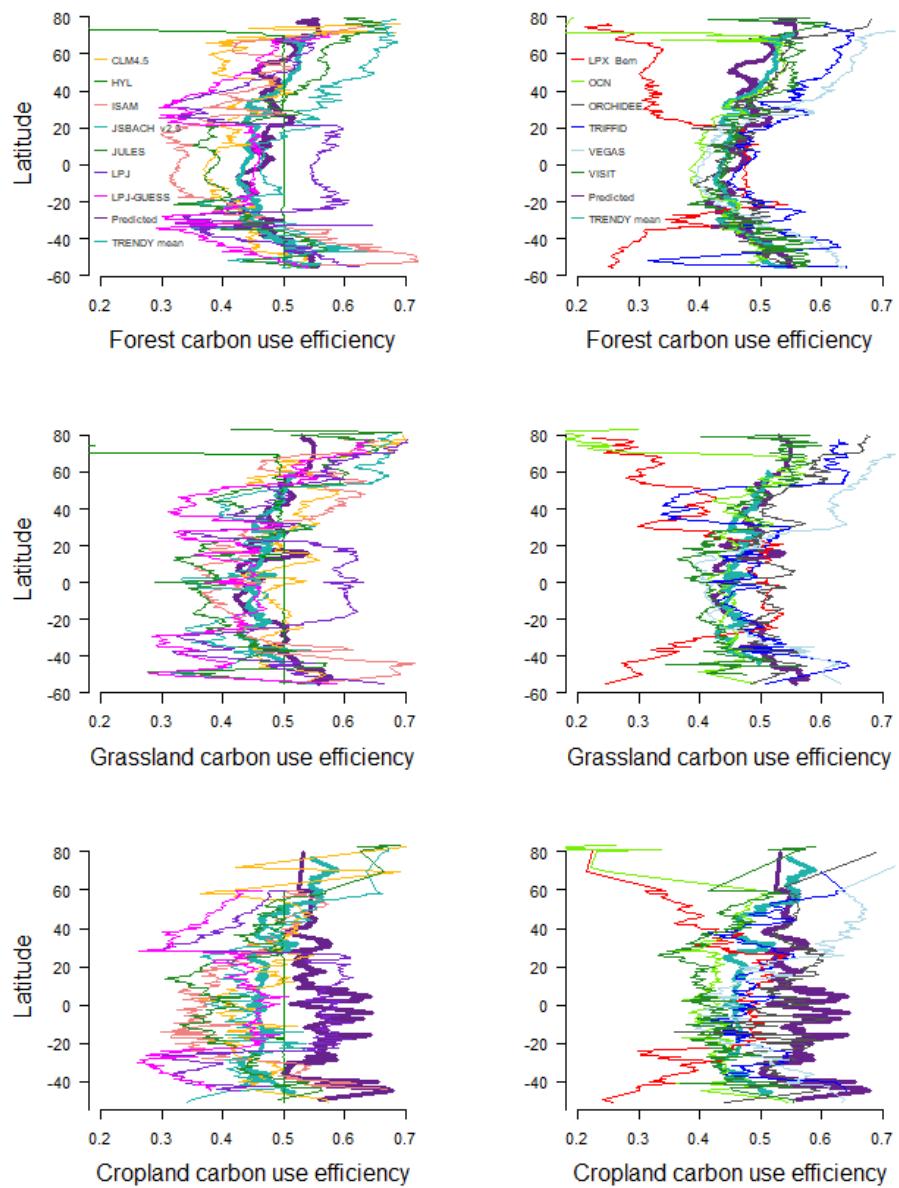
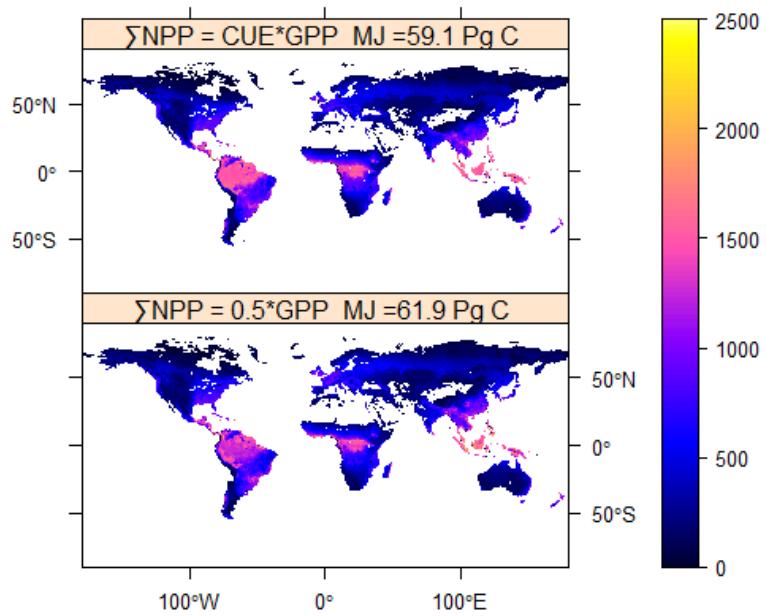


Figure S11. Latitudinal pattern of TRENDY-CUE for forest, grass and crop ecosystems. The bold purple line represents predicted CUE. Due to a limited number of grid cells for wetland, the latitudinal CUE was not presented.



155 **Figure S12.** Constant and variable CUE based NPP ($\text{g C m}^{-2} \text{ a}^{-1}$). Total constant CUE (0.5*GPP) based estimation of NPP was 61.9 Pg C a^{-1} globally, which was 2.8 Pg C a^{-1} higher than that of variable CUE (data-driven CUE*GPP) based estimation of NPP.

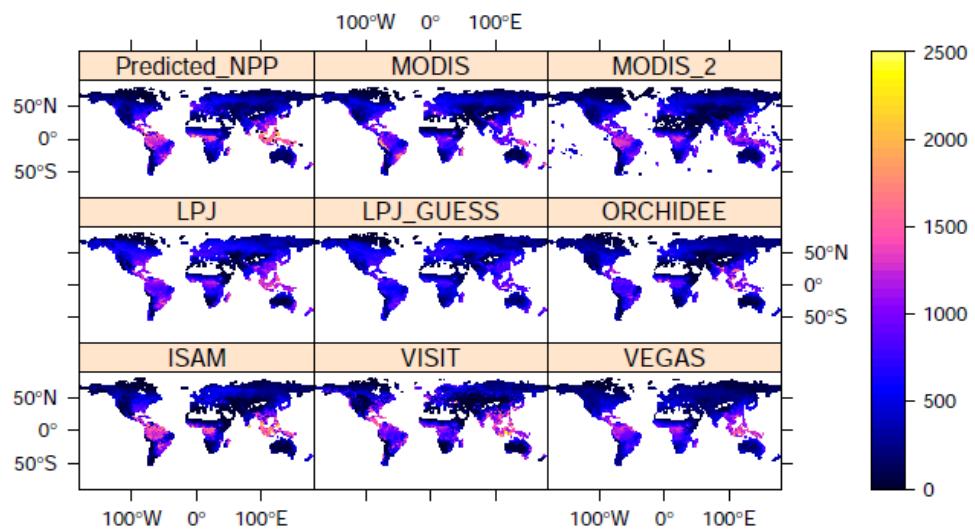


Figure S13. Global distribution of predicted NPP, MODIS NPP and TRENDY NPP ($\text{g C m}^{-2} \text{ a}^{-1}$).
 MODIS_2 NPP estimated from MODIS GPP multiplied by constant CUE (0.5).

References

Campioli, M., Gielen, B., Gockede, M., Papale, D., Bouriaud, O., and Granier, A.: Temporal variability of the NPP-GPP ratio at seasonal and interannual time scales in a temperate beech forest, 165 Biogeosciences, 8, 2481-2492, <http://dx.doi.org/10.5194/bg-8-2481-2011>, 2011.

Cannell, M., and Thornley, J.: Modelling the components of plant respiration: some guiding principles, Ann. Bot., 85, 45-54, 2000.

Dai, A., Trenberth, K. E., and Qian, T.: A Global Dataset of Palmer Drought Severity Index for 1870–2002: Relationship with Soil Moisture and Effects of Surface Warming, Journal of Hydrometeorology, 170 5, 1117-1130, 10.1175/JHM-386.1, 2004.

De Vries, F. W. T. P., Brunsting, A. H. M., and Van Laar, H. H.: Products, requirements and efficiency of biosynthesis a quantitative approach, J. Theor. Biol., 45, 339-377, 10.1016/0022-5193(74)90119-2, 1974.

Gifford, R. M.: Whole plant respiration and photosynthesis of wheat under increased CO₂ concentration and temperature: long-term vs. short-term distinctions for modelling, Glob. Chang. Biol., 175 1, 385-396, 1995.

Harris, I., Jones, P., Osborn, T., and Lister, D.: Updated high-resolution grids of monthly climatic observations—the CRU TS3. 10 Dataset, Int. J. Climatol., 34, 623-642, <http://dx.doi.org/10.1002/joc.3711>, 2014.

Hengl, T., Mendes de Jesus, J., Heuvelink, G. B., Ruiperez Gonzalez, M., Kilibarda, M., Blagotic, A., Shangguan, W., Wright, M. N., Geng, X., Bauer-Marschallinger, B., Guevara, M. A., Vargas, R., MacMillan, R. A., Batjes, N. H., Leenaars, J. G., Ribeiro, E., Wheeler, I., Mantel, S., and Kempen, B.: SoilGrids250m: Global gridded soil information based on machine learning, PLoS One, 12, e0169748, 180 <http://dx.doi.org/10.1371/journal.pone.0169748>, 2017.

Lamarque, J. F., Dentener, F., McConnell, J., Ro, C. U., Shaw, M., Vet, R., Bergmann, D., Cameron-Smith, P., Dalsoren, S., Doherty, R., Faluvegi, G., Ghan, S. J., Josse, B., Lee, Y. H., MacKenzie, I. A., Plummer, D., Shindell, D. T., Skeie, R. B., Stevenson, D. S., Strode, S., Zeng, G., Curran, M., Dahl-Jensen, D., Das, S., Fritzsche, D., and Nolan, M.: Multi-model mean nitrogen and sulfur deposition from the Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP): evaluation of historical and projected future changes, Atmos. Chem. Phys., 13, 7997-8018, 10.5194/acp-13-7997-2013, 2013.

Lavigne, M. B., and Ryan, M. G.: Growth and maintenance respiration rates of aspen, black spruce and jack pine stems at northern and southern BOREAS sites, Tree Phys., 17, 543-551, 10.1093/treephys/17.8-9.543, 1997.

Malhi, Y., Doughty, C. E., Goldsmith, G. R., Metcalfe, D. B., Girardin, C. A., Marthews, T. R., Del Aguila-Pasquel, J., Aragao, L. E., Araujo-Murakami, A., Brando, P., da Costa, A. C., Silva-Espejo, J. E., Farfan Amezquita, F., Galbraith, D. R., Quesada, C. A., Rocha, W., Salinas-Revilla, N., Silverio, D., Meir, P., and Phillips, O. L.: The linkages between photosynthesis, productivity, growth and biomass in lowland Amazonian forests, Glob. Chang. Biol., 21, 2283-2295, 10.1111/gcb.12859, 2015.

Marquis, R., Newell, E., and Villegas, A.: Non-structural carbohydrate accumulation and use in an understorey rain-forest shrub and relevance for the impact of leaf herbivory, Funct. Ecol., 11, 636-643, 1997.

Meyers, T.: An assessment of storage terms in the surface energy balance of maize and soybean, Agric. For. Meteorol., 125, 105-115, 10.1016/j.agrformet.2004.03.001, 2004.

205 Newell, E., Mulkey, S., and Wright, J.: Seasonal patterns of carbohydrate storage in four tropical tree species, *Oecologia*, 131, 333-342, 10.1007/s00442-002-0888-6, 2002.

O'Brien, M. J., Leuzinger, S., Philipson, C. D., Tay, J., and Hector, A.: Drought survival of tropical tree seedlings enhanced by non-structural carbohydrate levels, *Nature Climate Change*, 4, 710-714, 10.1038/nclimate2281, 2014.

210 Poorter, L., and Kitajima, K.: Carbohydrate Storage and Light Requirements of Tropical Moist and Dry Forest Tree Species, *Ecology*, 88, 1000-1011, 2007.

Rambal, S., Joffre, R., Ourcival, J. M., Cavender-Bares, J., and Rocheteau, A.: The growth respiration component in eddy CO₂ flux from a *Quercus ilex* mediterranean forest, *Glob. Chang. Biol.*, 10, 1460-1469, 2004.

215 Rosas, T., Galiano, L., Ogaya, R., Peñuelas, J., and Martínez-Vilalta, J.: Dynamics of non-structural carbohydrates in three Mediterranean woody species following long-term experimental drought, *Frontiers in Plant Science*, 4, 10.3389/fpls.2013.00400, 2013.

Roxburgh, S. H., Berry, S. L., Buckley, T. N., Barnes, B., and Roderick, M. L.: What is NPP? Inconsistent accounting of respiratory fluxes in the definition of net primary production, *Funct. Ecol.*, 19, 378-382, 10.1111/j.1365-2435.2005.00983.x, 2005.

220 Schnapp, S. R., Curtis, W. R., Bressan, R. A., and Hasegawa, P. M.: Estimation of growth yield and maintenance coefficient of plant cell suspensions, *Biotechnol. Bioeng.*, 38, 1131-1136, 1991.

Sheffield, J., Goteti, G., and Wood, E. F.: Development of a 50-Year High-Resolution Global Dataset of Meteorological Forcings for Land Surface Modeling, *J. Clim.*, 19, 3088-3111, 10.1175/JCLI3790.1, 2006.

225 Thornley, J., and Cannell, M.: Modelling the components of plant respiration: representation and realism, *Ann. Bot.*, 85, 55-67, 2000.

Van Oijen, M., Schapendonk, A., and Hoglind, M.: On the relative magnitudes of photosynthesis, respiration, growth and carbon storage in vegetation, *Ann. Bot.*, 105, 793-797, 10.1093/aob/mcq039, 2010.

230 Vicca, S., Luyssaert, S., Penuelas, J., Campioli, M., Chapin, F. S., 3rd, Ciais, P., Heinemeyer, A., Hogberg, P., Kutsch, W. L., Law, B. E., Malhi, Y., Papale, D., Piao, S. L., Reichstein, M., Schulze, E. D., and Janssens, I. A.: Fertile forests produce biomass more efficiently, *Ecol. Lett.*, 15, 520-526, <http://dx.doi.org/10.1111/j.1461-0248.2012.01775.x>, 2012.

235 Woodruff, D. R., and Meinzer, F. C.: Water stress, shoot growth and storage of non-structural carbohydrates along a tree height gradient in a tall conifer, *Plant Cell Environ.*, 34, 1920-1930, 10.1111/j.1365-3040.2011.02388.x, 2011.

Xu, X., Shi, Z., Chen, X. C., Lin, Y., Niu, S. L., Jiang, L. F., Luo, R. S., and Luo, Y. Q.: Unchanged carbon balance driven by equivalent responses of production and respiration to climate change in a 240 mixed-grass prairie, *Glob. Chang. Biol.*, 22, 1857-1866, 10.1111/gcb.13192, 2016.

Zanotelli, D., Montagnani, L., Manca, G., and Tagliavini, M.: Net primary productivity, allocation pattern and carbon use efficiency in an apple orchard assessed by integrating eddy covariance, biometric and continuous soil chamber measurements, *Biogeosciences*, 10, 3089-3108, 10.5194/bg-10-3089-2013, 2013.

245 Zhu, Z., Bi, J., Pan, Y., Ganguly, S., Anav, A., Xu, L., Samanta, A., Piao, S., Nemani, R., and Myneni, R.: Global Data Sets of Vegetation Leaf Area Index (LAI)3g and Fraction of Photosynthetically Active Radiation (FPAR)3g Derived from Global Inventory Modeling and Mapping Studies (GIMMS)

Normalized Difference Vegetation Index (NDVI3g) for the Period 1981 to 2011, Remote Sensing, 5, 927, 2013.

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