

## Supplementary Material

### S1. Additional photosynthesis equations

In JULES, leaf-level photosynthesis (Collatz et al 1991:1992) is calculated based on the limiting factor of three potential photosynthesis rates.

1. A light-limited rate,  $W_l$ :

$$W_l = \alpha(1 - \omega)I_{par} \left( \frac{c_i - \Gamma}{c_i + 2\Gamma} \right) \quad \text{for C3 plants} \quad (\text{A.1})$$

$$W_l = \alpha(1 - \omega)I_{par} \quad \text{for C4 plants} \quad (\text{A.2})$$

where  $\alpha$  is the quantum efficiency of photosynthesis ( $\text{mol CO}_2 \text{ mol PAR}^{-1}$ ) and  $\omega$  is the leaf scattering coefficient for PAR.  $I_{par}$  is the photosynthetically active radiation hitting the leaf ( $\text{mol m}^{-2} \text{ s}^{-1}$ ),  $\Gamma$  is the  $\text{CO}_2$  compensation point in the absence of mitochondrial respiration (Pa), and  $c_i$  is the internal  $\text{CO}_2$  concentration (Pa).

2. A Rubisco-limited rate,  $W_c$ :

$$W_c = V_{cmax} \left( \frac{c_i - \Gamma}{c_i + K_c (1 + O_a / K_o)} \right) \quad \text{for C3 plants} \quad (\text{A.3})$$

$$W_c = V_{cmax} \quad \text{for C4 plants} \quad (\text{A.4})$$

where  $K_O$  and  $K_C$  are the Michaelis-Menten parameters for  $\text{O}_2$  and  $\text{CO}_2$ , respectively, and  $V_{cmax}$  is the maximum rate of carboxylation of Rubisco ( $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ ).

3. A rate of transport of photosynthetic products for C3 plants, and PEPCarboxylase limitation for C4 plants,  $W_e$ :

$$W_e = 0.5V_{cmax} \quad \text{for C3 plants} \quad (\text{A.5})$$

$$W_e = 2 \times 10^4 V_{cmax} \frac{c_i}{P_*} \quad \text{for C4 plants} \quad (\text{A.6})$$

where  $P_*$  is the surface air pressure (Pa).

**S2. Data for N<sub>mass</sub>, LMA, and LL.** The table shows the data sources for the TRY data used in this study. For each source, the number of measurements for each source is provided for N<sub>mass</sub>/specific leaf area (SLA) pairs and for leaf lifespan.

Ref.	Contact	N <sub>mass</sub> + SLA	Leaf lifespan
Atkin et al., 1997; Campbell et al., 2007	Owen Atkin	218	
Xu and Baldocchi, 2003	Dennis Baldocchi	468	
Cavender-Bares et al., 2006	Jeannine Cavender-Bares	x	
unpublished	F. Stuart III Chapin	50	48
Cornelissen et al., 2004; Cornelissen et al., 2003, 1996; Diaz et al., 2004; Quested et al., 2003	Johannes Cornelissen	690	161
	Will Cornwell (+David Ackerly)	53	
Díaz et al., 2004 (maybe didn't use); Diaz et al. 2010 (definitely used)	Sandra Díaz	70	
Han et al., 2005; He et al., 2006, 2007	Jingyun Fang	148	
Freschet and Cornelissen, 2010; Freschet et al., 2010	Gregoire Freschet (+Hans Cornelissen)	40	
	Eric Garnier (+ Sandra Lavorel)	966	
Kattge et al., 2009	Jens Kattge*	1326	204
Kurokawa and Nakashizuka, 2008	Hiroko Kurokawa	399	89
	Daniel Laughlin	139	
Niinemets, 1999; Niinemets, 2001	Ülo Niinemets	264	33
Ordoñez et al., 2010; Ordoñez et al., 2010	Jenny Ordoñez (+Peter van Bodegom)	282	
Ogaya and Peñuelas, 2007a, 2007b, 2003; Ogaya, 2006; Sardans et al., 2008	Josep Peñuelas	808	
Poorter et al., 2009; 2006	Lourens Poorter		x
Reich et al., 2008, 2009	Peter Reich	720	199
Cornwell et al., 2007	Lawren Sack	30	

Shipley and Lechowicz 2000, Ecoscience, 7:183- 194	Bill Shipley	603
2. Meziane and Shipley, 1999, Plant Cell and Environment		
	Enio Sosinski	66
Soudzilovskaia et al, 2013, PNAS	Nadia Soudzilovskaia	155
	Peter van Bodegom	x x
Wright et al., 2011	S. Joseph (Joe) Wright	204
Wright et al., 2006, 2004	Ian Wright	1673 442

Data was analysed to calculate the parameters  $N_m$  and LMA for the new 9 PFTs (Table 2) and for the original 5 PFTs (Table S2). For JULES users who wish to run the model with the standard 5 PFTs but with updated parameters, the recommended parameters are given in Table S2. Additionally, it is important to set both ‘l\_trait\_phys’ and ‘l\_ht\_compete’ to True in the jules\_vegetation namelist. The former will switch to the trait-based physiology discussed in the main text, and the latter will allow for flexible number of PFTs to compete if the dynamic vegetation mode is used. Results of JULES with the new 9 PFTs and the height-based competition will be discussed in a follow up paper.

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### **S3. Energy and respiration results at the Fluxnet sites**

The TRY-based parameters give a lower  $N_{\text{root}}$  and  $N_{\text{stem}}$  for all PFTs, and a lower  $N_{\text{leaf}}$  for all PFTs except for C<sub>3</sub> grass and BET-Te (Fig. S2). For C<sub>3</sub> grass, the simulated  $N_{\text{leaf}}$  in JULES9 is higher during the winter than in JULES5, since a moderate LAI is maintained due to the new phenology.  $N_{\text{leaf}}$  is higher for BET-Te due to thicker leaves than previously. The respiration fluxes at BR-Ma2 provide a good example of the impacts of the new PFT

parameters (Fig. SM3). The lower  $N_{\text{stem}}$  and  $N_{\text{root}}$  in JULES9 compared to JULES5 (Fig. SM2) reduced simulated  $R_{\text{pm}}$  (Eqn. 8). As a result, the average increase in plant respiration ( $46 \text{ g C m}^{-2} \text{ yr}^{-1}$ ) was much smaller than the increase in GPP ( $377 \text{ g C m}^{-2} \text{ yr}^{-1}$ ), and NPP increased from  $867 \text{ g C m}^{-2} \text{ yr}^{-1}$  in JULES5 to  $1198 \text{ g C m}^{-2} \text{ yr}^{-1}$  in JULES9, which was higher than the observed value at Manaus of  $1011 \pm 140 \text{ g C m}^{-2} \text{ yr}^{-1}$  (Mahli et al., 2009). At Santarem, NPP also increased, but it was lower than the observed value of  $1440 \text{ g C m}^{-2} \text{ yr}^{-1}$ .

The higher ecosystem respiration in JULES9 compared to JULES5 that accompanied the increased GPP was less realistic (in terms of RMSE), with 3 exceptions (Fig. SM3), but the seasonal cycle of total respiration was improved at 8 sites. The RMSE decreased at ES-ES1, where lower GPP and R in the fall and winter were more realistic, at JP-Tom, where the switch from generic needle leaf to deciduous needle leaf improved all aspects of the simulation, and at FI-Kaa, where the new phenology of grass also improved the simulation.

In JULES, latent heat flux (LE) is due to evaporation from water stored on the canopy, evaporation of water from the top layer of soil, transpiration through the stomata, and sublimation of snow. The seasonal cycle of LE was improved at nine sites, however  $r$  decreased (by  $<0.03$ ) at BR-Sa1, ES-ES1, BR-Sa3, US-Bo1, and US-FPe, comparing JULES9<sub>ALL</sub> to JULES5. The RMSE increased by  $>4 \text{ W m}^{-2}$  in JULES9<sub>TRY</sub> compared to JULES5 at DE-Tha and FI-Hyy, and RMSE increased by a further  $4 \text{ W m}^{-2}$  at DE-Tha when the photosynthesis/respiration parameters were added due to the higher GPP and stomatal conductance. However, the correlation was  $>0.91$  for both sites. At some forest sites, simulated LE (SH) was too high (low) during the winter and spring (DE-Tha, US-Ha1, and US-MMS), however the LE component contributing to the high bias is site-dependent. For example, from Jan.-Mar. the largest source of LE is evaporation from snow/ice ( $E_i$ ) at

Harvard, canopy evaporation and  $E_i$  at Tharandt, and soil evaporation/transpiration at Morgan Monroe. These springtime errors were not affected by the new PFTs. Another consistent bias in the forests was high mid-summer LE (De-Tha, FI-Hyy, US-Ha1, and US-MMS), which in this case always results from the soil evaporation/transpiration. Because the new PFTs tend to increase GPP and stomatal conductance, the errors in summer LE are higher.

**Table SM1.** List of parameters and symbols in the text.

Symbol	Units	Equation	Description	Default Value <sup>a</sup>
$A_l$	$\text{kg C m}^{-2} \text{s}^{-1}$	5	Leaf-level photosynthesis	
$a_{wl}$	$\text{kg C m}^{-2}$	24	Allometric coefficient	
$a_{ws}$	--	24	Ratio of total to respiring stem carbon	
$b_{wl}$	--	24	Allometric exponent	1.667
$C_i$	Pa	6	Internal leaf $\text{CO}_2$ concentration	
$C_{mass}$	$\text{kg C [kg biomass]}^{-1}$	23	Leaf carbon concentration per unit mass	0.5 for this study
$C_s$	Pa	6	Leaf surface $\text{CO}_2$ concentration	
$D_{crit}$	$\text{kg kg}^{-1}$	7	Critical humidity deficit	
$d_T$	--	16	Rate of change of leaf turnover with temperature	
$f_0$	--	7	Stomatal conductance parameter	
$f_d$	--	4	Leaf dark respiration coefficient	
$g_s$	$\text{m s}^{-1}$	6	Leaf-level stomatal conductance	
$i_v$	$\mu\text{mol CO}_2 \text{ m}^{-2} \text{s}^{-1}$	19	Intercept for relationship between $N_{area}$ and $V_{cmax,25}$	
$k_n$	--	3, 20	Extinction coefficient for nitrogen	0.78
$h$	m	13, 23, 24	Canopy height	
$L_{bal}$	$\text{m}^2 \text{m}^{-2}$	12, 13, 22-24	Balanced leaf area index (maximum LAI given the plant's height)	
$L_{max}$	$\text{m}^2 \text{m}^{-2}$		Maximum LAI	
$L_{min}$	$\text{m}^2 \text{m}^{-2}$		Minimum LAI	
$LMA$	$\text{kg m}^{-2}$	18, 21, 22	Leaf mass per unit area (new parameter)	
$N_a$	$\text{kg N m}^{-2}$	18	Leaf nitrogen per unit area	
$n_{eff}$	$\text{mol CO}_2 \text{ m}^{-2} \text{s}^{-1}$ $\text{kg C [kg N]}^{-1}$	3	Constant relating leaf nitrogen to Rubisco carboxylation capacity	
$N_{lo}$	$\text{kg N [kg C]}^{-1}$	3	Top leaf nitrogen concentration (old parameter, mass basis)	
$N_m$	$\text{kg N kg}^{-1}$	18, 21-23	Top leaf nitrogen concentration (new parameter)	
$N_l$	$\text{kg N m}^{-2}$	11, 21	Total leaf nitrogen concentration	
$N_r$	$\text{kg N m}^{-2}$	12, 22	Total root nitrogen concentration	
$N_s$	$\text{kg N m}^{-2}$	13, 23	Total stem nitrogen concentration	
$p$	--	17	Phenological state (LAI/ $L_{bal}$ )	
$Q_{10,leaf}$	--	2	Constant for exponential term in temperature function of $V_{cmax}$	2
$R_a$	$\text{kg C m}^{-2} \text{s}^{-1}$	8	Total plant autotrophic respiration	
$R_d$	$\text{kg C m}^{-2} \text{s}^{-1}$	4, 5	Leaf dark respiration	
$r_g$	--	10	Growth respiration coefficient	0.25
$rootd$	m		e-folding root depth	
$s_v$	$\mu\text{mol CO}_2 \text{ g N}^{-1} \text{ s}^{-1}$	19	Slope between $N_{area}$ and $V_{cmax,25}$	
$T_{low}$	°C	1	Upper temperature parameter for $V_{cmax}$	
$T_{off}$	°C	16	Threshold temperature for phenology	
$T_{opt}^b$	°C		Optimal temperature for $V_{cmax}$	
$T_{upp}$	°C	1	Upper temperature parameter for $V_{cmax}$	
$V_{cmax,25}$	$\mu\text{mol m}^{-2} \text{s}^{-1}$	1, 9	The maximum rate of carboxylation of Rubisco at 25°C	
$W$	$\text{kg C m}^{-2} \text{s}^{-1}$	5	Smoothed minimum of the potential limiting rates of photosynthesis	
$\alpha$	$\text{mol CO}_2 [\text{mol PAR photons}]^{-1}$		Quantum efficiency	

$\beta$	--	5	Soil moisture stress factor	
$F^*$	Pa	7	$\text{CO}_2$ compensation point	
$\gamma_0$	[360 days] <sup>-1</sup>	16	Minimum leaf turnover rate	
$\gamma_{lm}$	[360 days] <sup>-1</sup>	16	Leaf turnover rate	
$\gamma_p$	[360 days] <sup>-1</sup>	17	Leaf growth rate	20
$\mu_{rl}$	--	12, 22	Ratio of nitrogen concentration in roots and leaves	
$\mu_{sl}$	--	13, 23	Ratio of nitrogen concentration in stems and leaves	
$\eta_{sl}$	kg C m <sup>-2</sup> LAI <sup>-1</sup>	13, 23	Live stemwood coefficient	0.01
$\sigma_L$	kg C m <sup>-2</sup> LAI <sup>-1</sup>	11, 12	Specific leaf density (old parameter)	

<sup>a</sup>Default values only provided for non-PFT-dependent parameters.

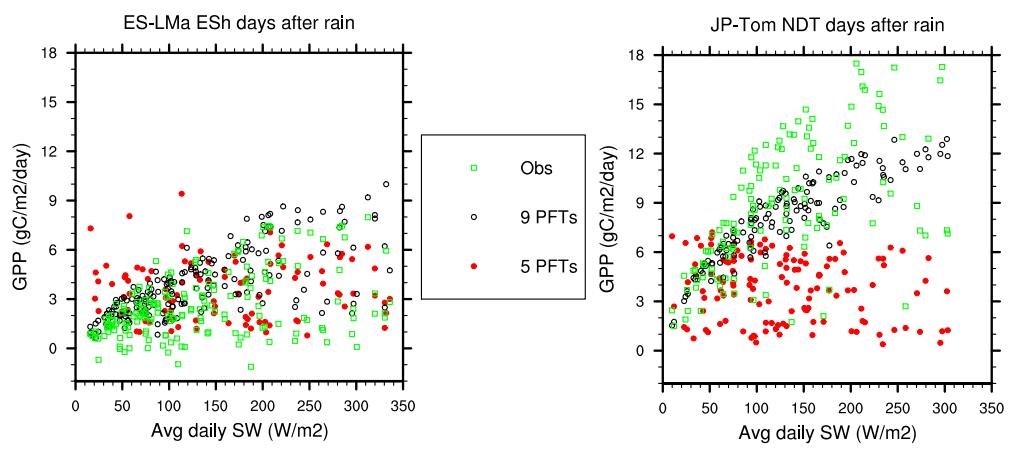
**Table SM2.** New trait-based parameters for 5 PFTs that are consistent with TRY data and updated parameters used in this study.

	BT	NT	C3	C4	SH
$N_m$	0.0185	0.0117	0.0240	0.0113	0.0175
<b>LMA</b>	0.1012	0.2240	0.0495	0.1370	0.1023
$s_v$	25.48	18.15	40.96	20.48	23.15
$i_v$	6.12	6.32	6.42	0.00	14.71
$V_{\text{cmax},25}$	53.84	53.88	55.08	31.71	56.15
$T_{\text{off}}$	5	-40	5	5	-40
$d_T$	9	9	0	0	9
$\gamma_0$	0.25	0.25	3.0	3.0	0.66
$\gamma_p$	20	15	20	20	15
$L_{\min}$	1	1	1	1	1
$L_{\max}$	9	7	3	3	4
$D_{\text{crit}}$	0.09	0.06	0.051	0.075	0.037
$f_0$	0.875	0.875	0.931	0.800	0.950
$f_d$	0.010	0.015	0.019	0.019	0.015
$rootd$	3	2	0.5	0.5	1
$T_{\text{low}}$	5	0	10	13	0
$T_{\text{opt}}$	39	32	28	41	32
$T_{\text{upp}}$	43	36	32	45	36
$\alpha$	0.08	0.08	0.06	0.04	0.08
$\mu_{rl}$	0.67	0.67	0.72	0.72	0.67

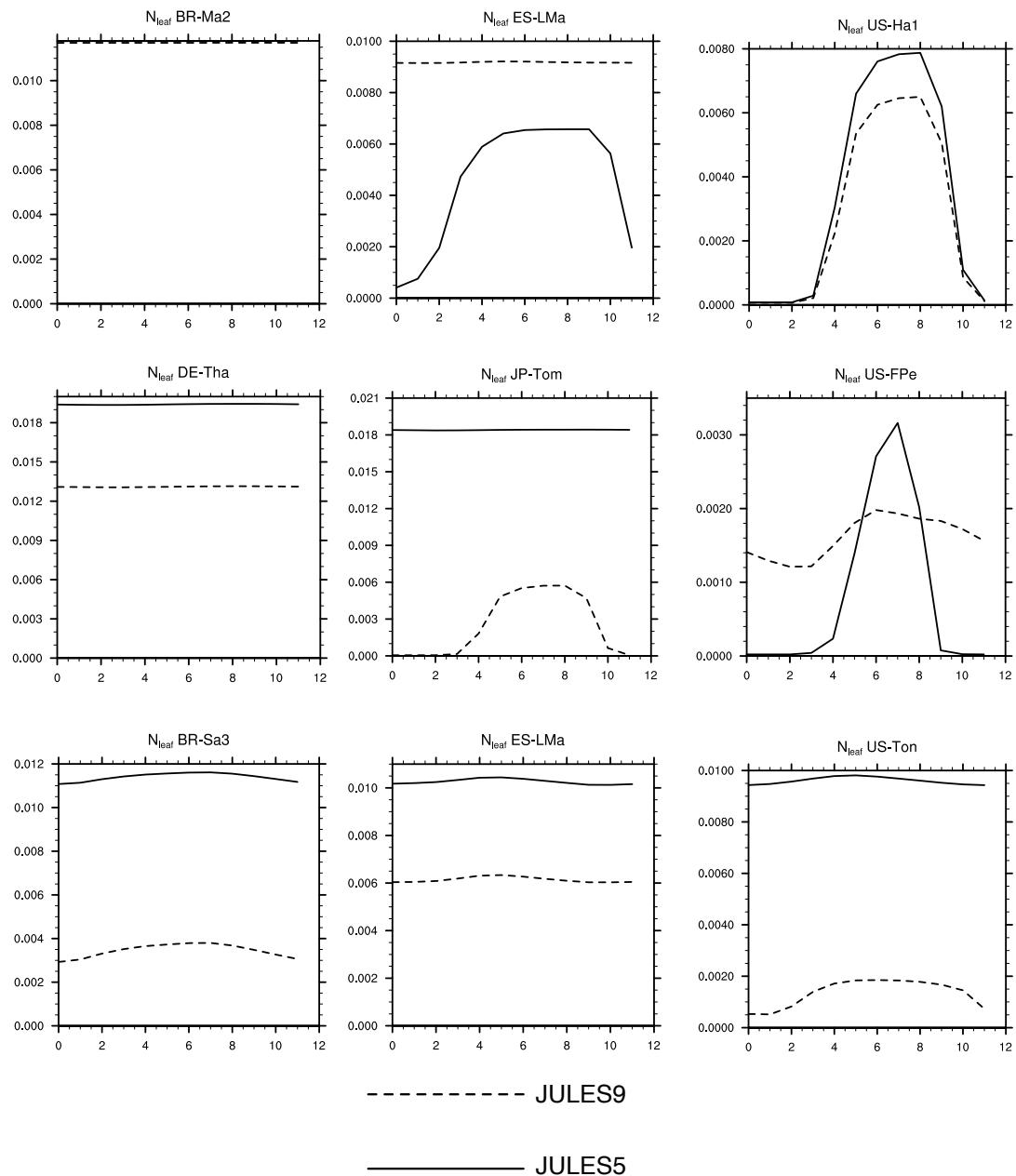
**Table SM3.** Relationship between WWF ecoregions and the eight biomes used in this study.

WWF ecoregion	Biome for this study
Tropical & Subtropical Moist Broadleaf Forests	Tropical forests
Tropical & Subtropical Dry Broadleaf Forests	Tropical forests
Tropical & Subtropical Coniferous Forests	Tropical forests
Temperate Broadleaf & Mixed Forests	Extratropical mixed forests
Temperate Conifer Forests	Boreal and coniferous forests
Boreal Forests/Taiga	Boreal and coniferous forests
Tropical & Subtropical Grasslands, Savannas & Shrublands	Tropical savannas
Temperate Grasslands, Savannas & Shrublands	Temperate grasslands
Flooded Grasslands & Savannas	Temperate grasslands
Montane Grasslands & Shrublands	Temperate grasslands
Tundra	Tundra
Mediterranean Forests, Woodlands & Scrub	Mediterranean woodlands
Deserts & Xeric Shrublands	Desert
Mangroves	Tropical forests

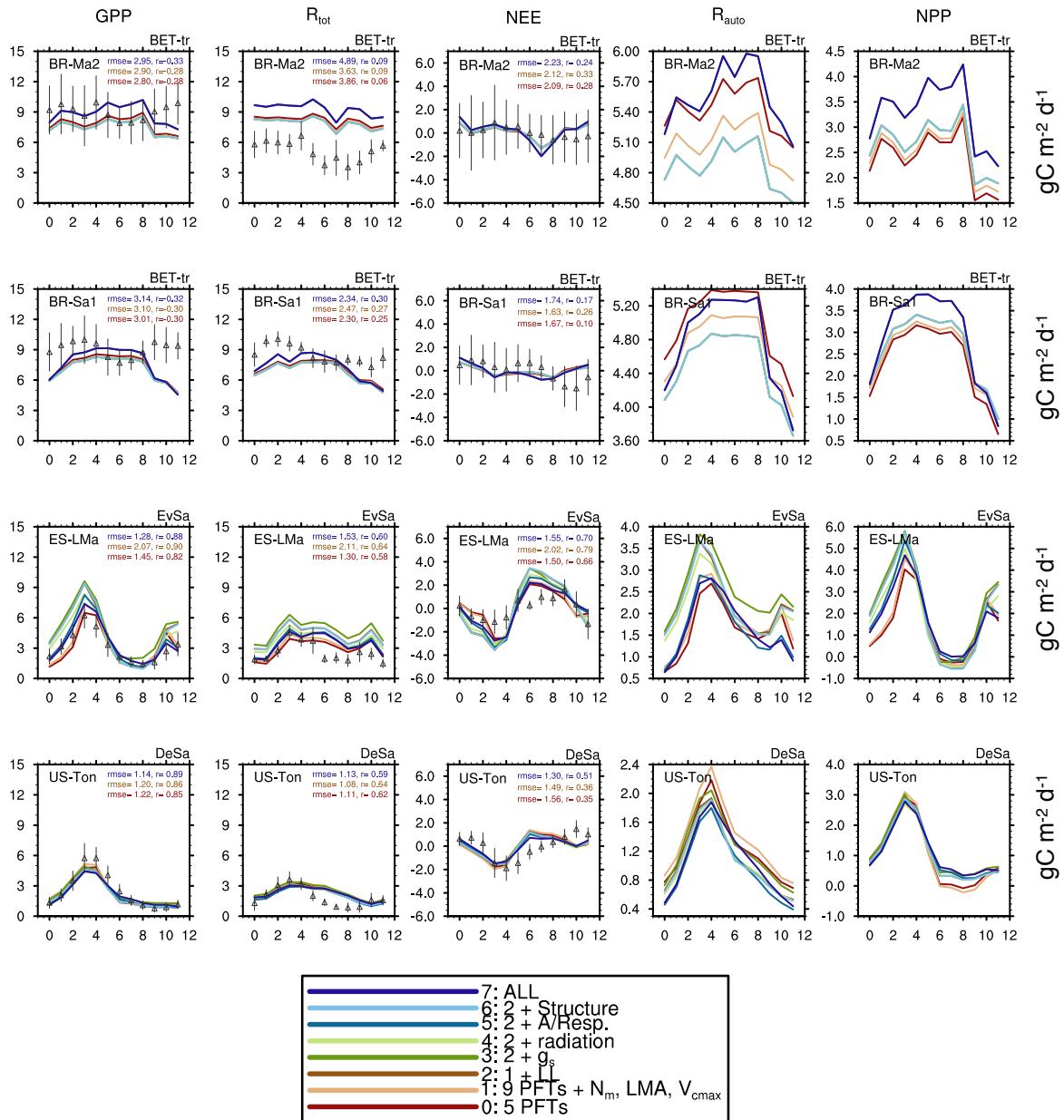
**Supplemental Material:  
Figures**



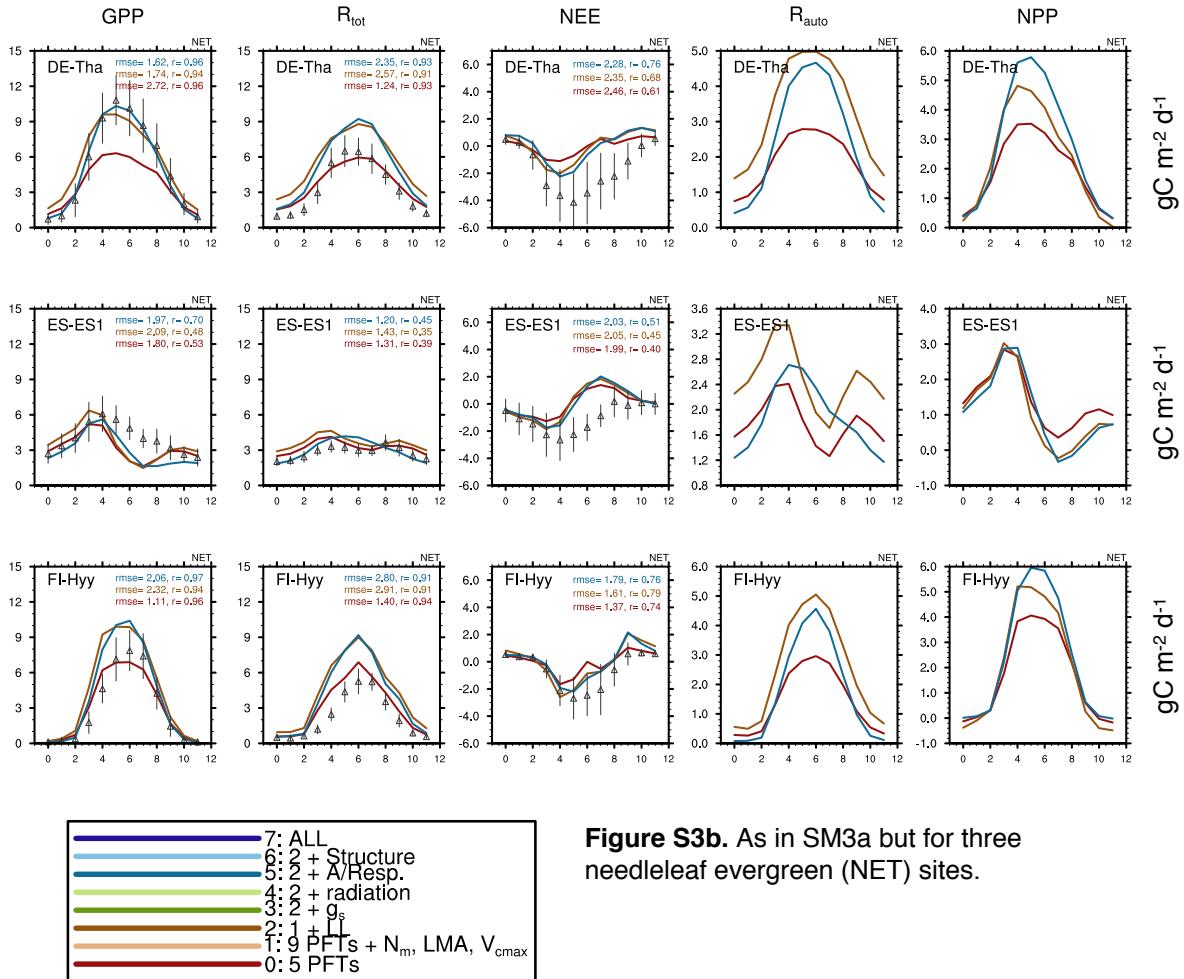
**Figure S1.** Daily average GPP versus shortwave radiation on days following rainfall when leaf area index is at or near its seasonal maximum.

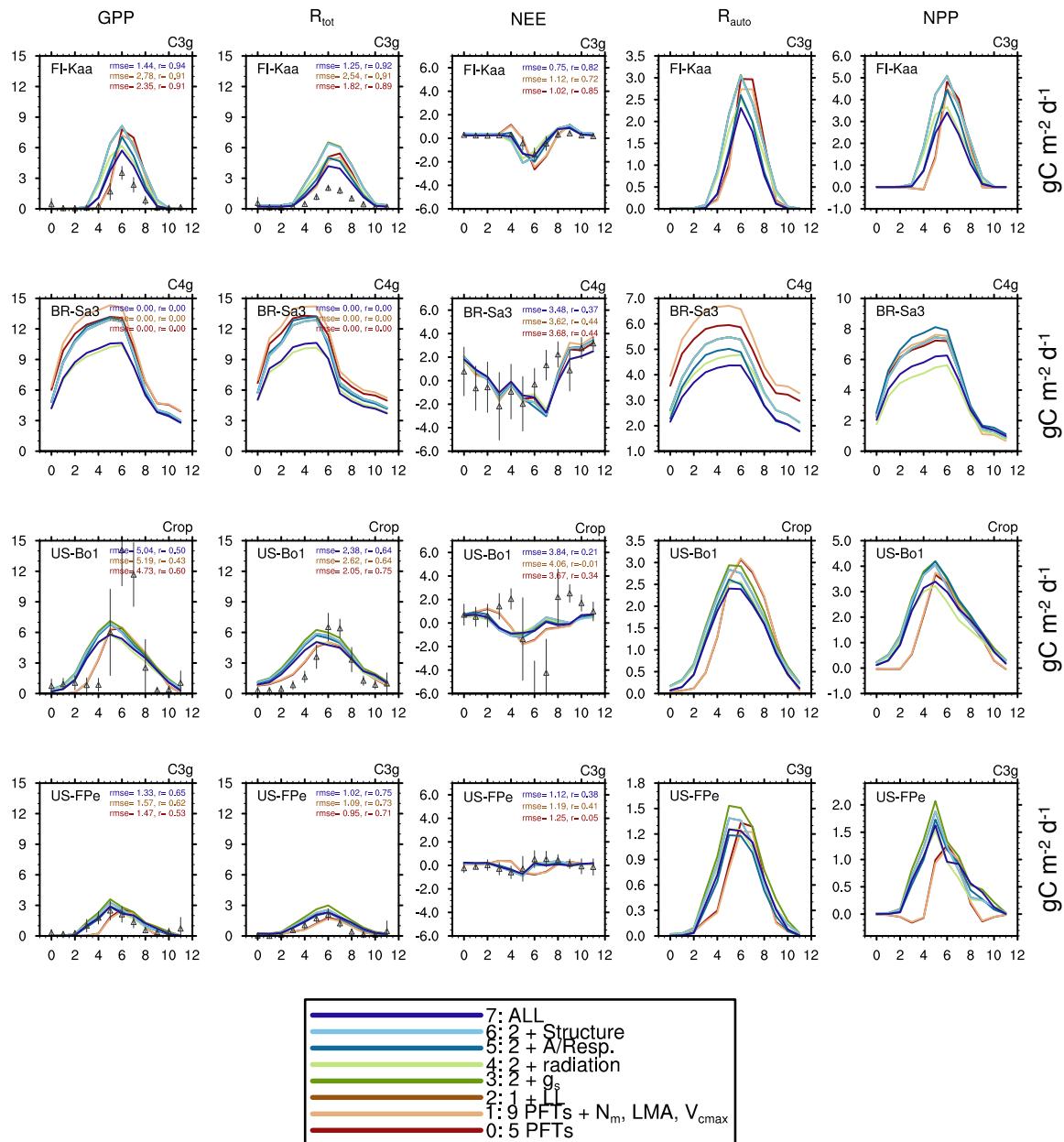


**Figure S2.** Monthly mean leaf nitrogen content (scaled by LAI) at nine sites representative of each of the new PFTs.

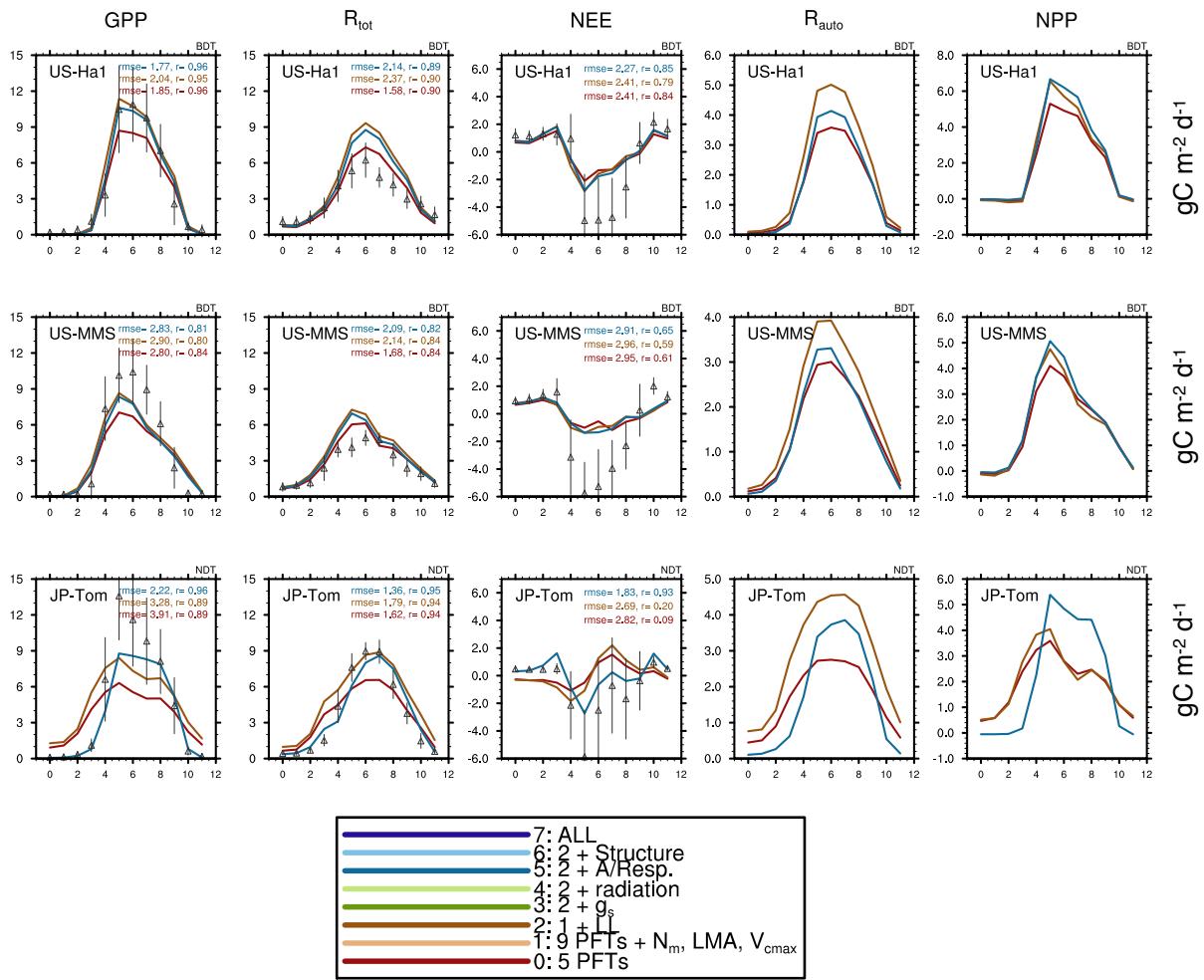


**Figure S3a.** Monthly mean fluxes of GPP, total ecosystem respiration, NEE, autotrophic respiration, and NPP at two tropical forest sites (BET-tr) and two savannah sites (EvSa=Evergreen Savannah, DeSa=Deciduous Savannah). Observations  $\pm$  standard deviation from Fluxnet are shown with triangles and vertical lines. All simulations in Table 4 in the main text are shown. Also shown are the daily root mean square error (rmse) based on daily fluxes and the correlation coefficient ( $r$ ) based on monthly mean fluxes for all years of the simulations. All units are in gC m<sup>-2</sup> d<sup>-1</sup>.

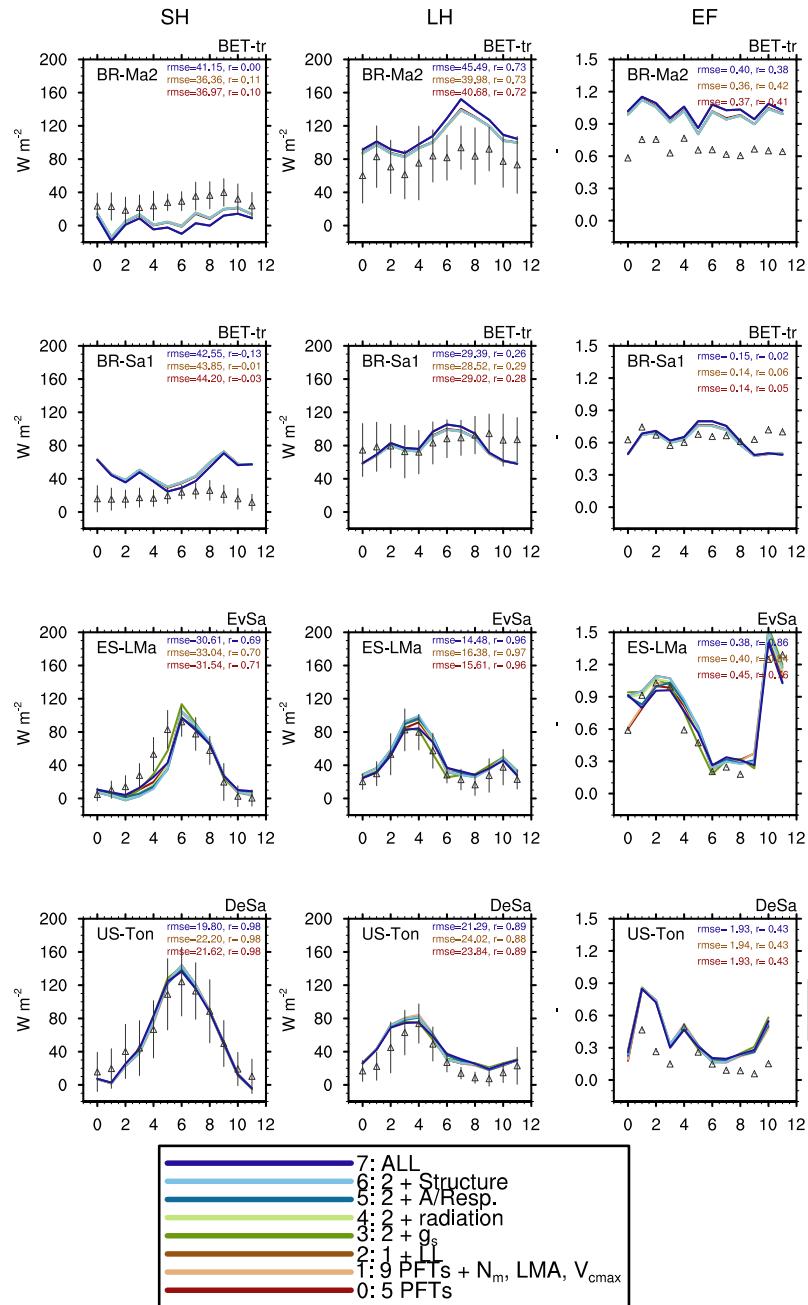




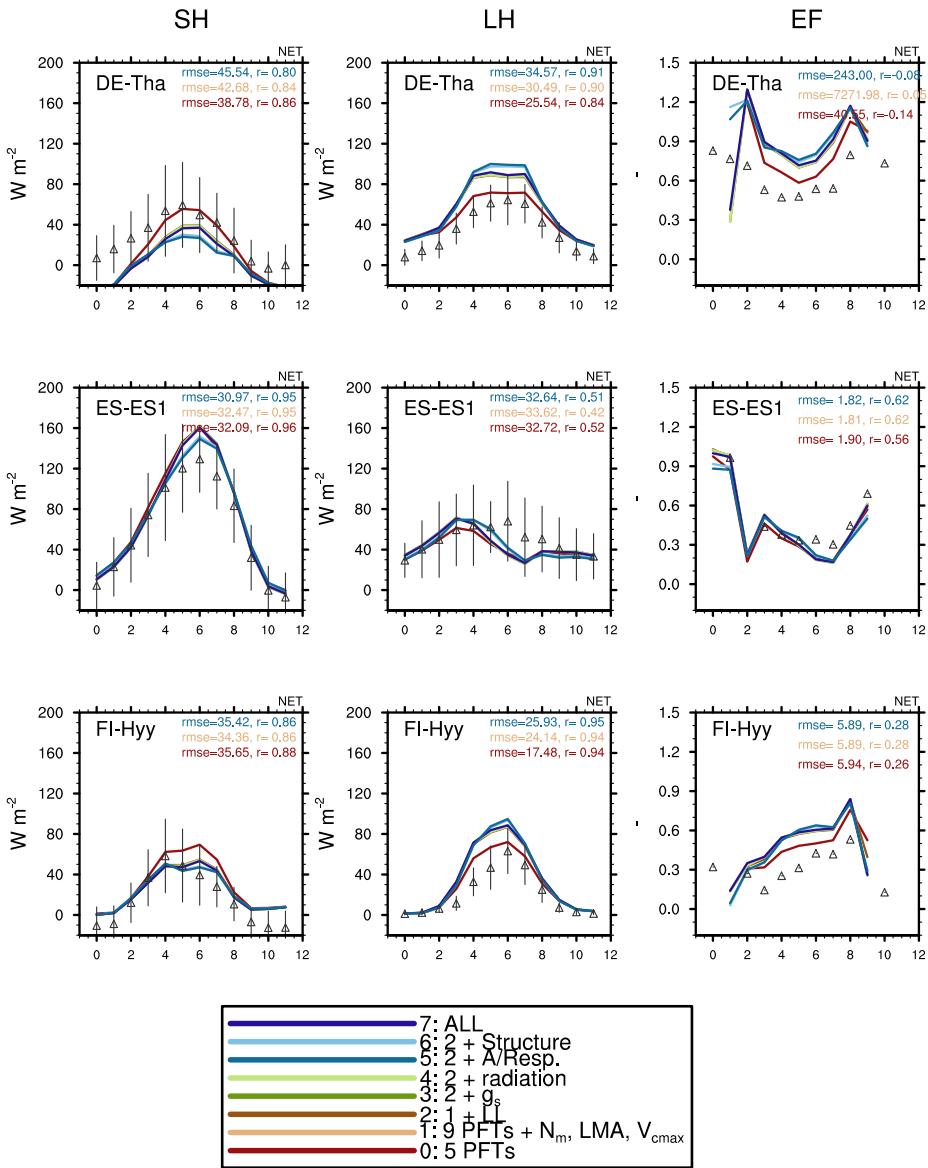
**Figure S3c.** As in SM3a but for four grass sites.



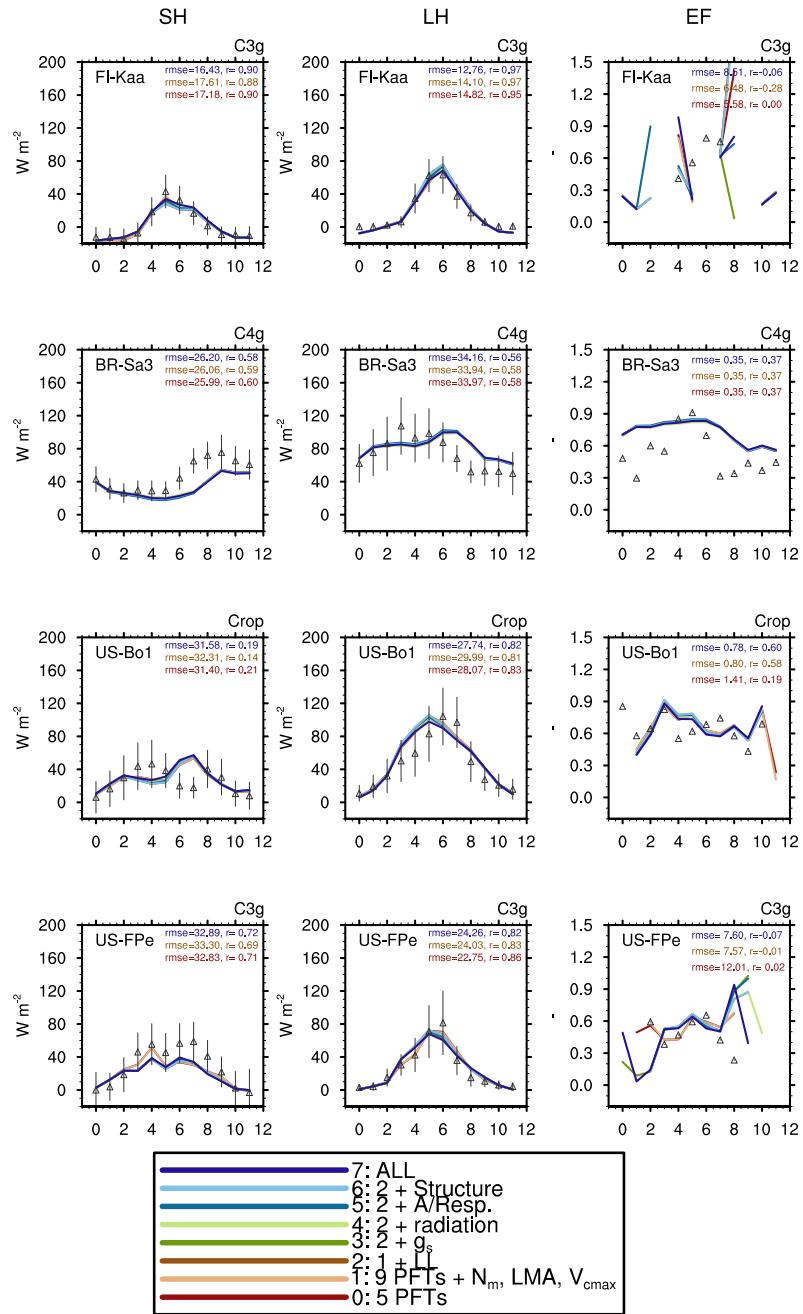
**Figure S3d.** As in SM3a but for two broadleaf deciduous (BDT) sites and one needleleaf deciduous (NDT) site.



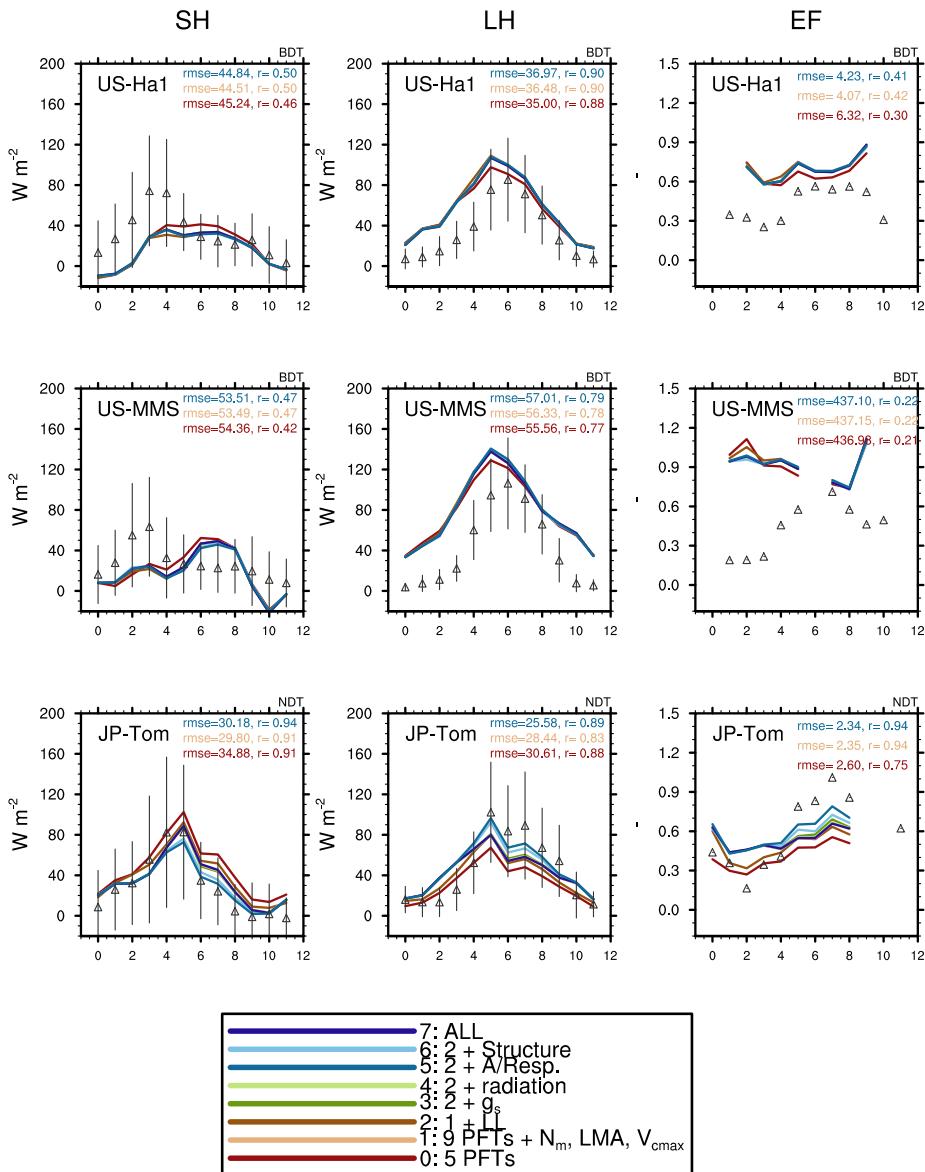
**Figure S4a.** Monthly mean fluxes of latent heat, sensible heat, and evaporative fraction (=LE/(SH +LE)) at two tropical forest sites (BET-tr) and two savannah sites (EvSa=Evergreen Savannah, DeSa=Deciduous Savannah). Observations  $\pm$  standard deviation from Fluxnet are shown with triangles and vertical lines.



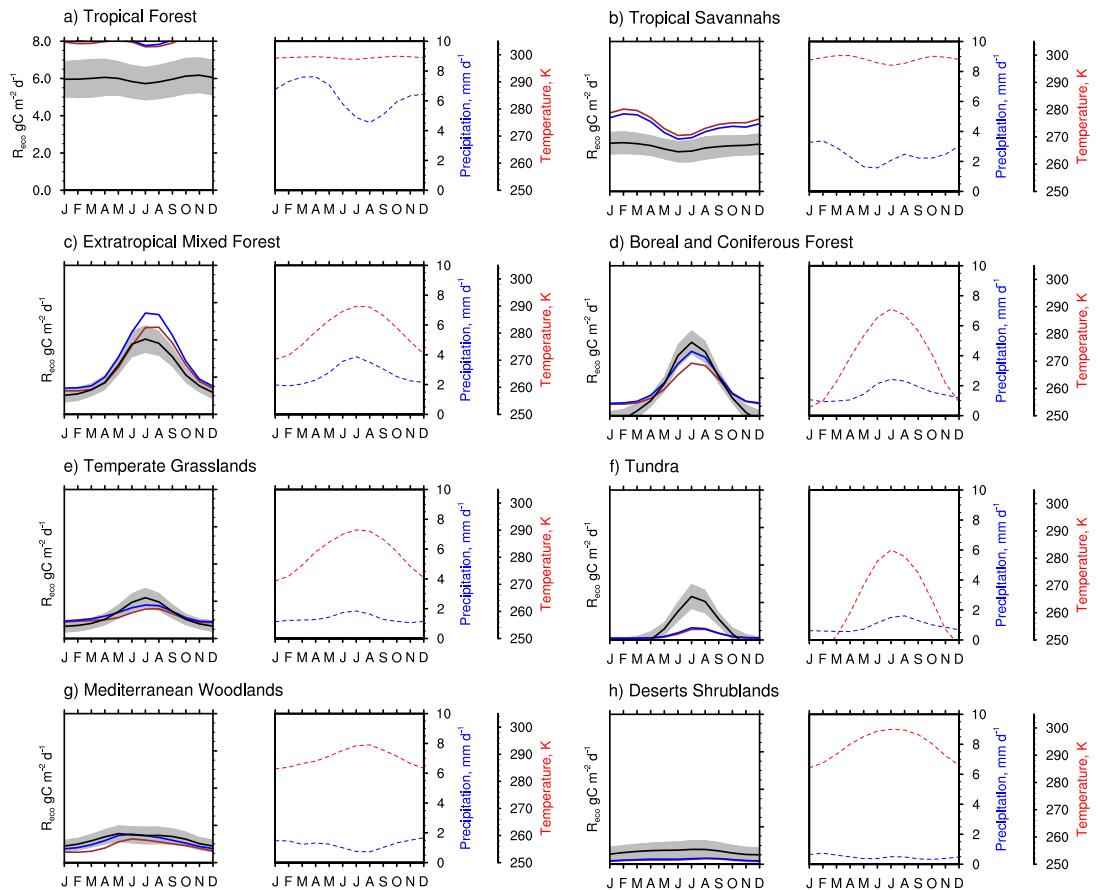
**Figure S4b.** As in Fig. S4a but for the NET sites.



**Figure S4c.** As in Fig. S4a but for the grass sites.

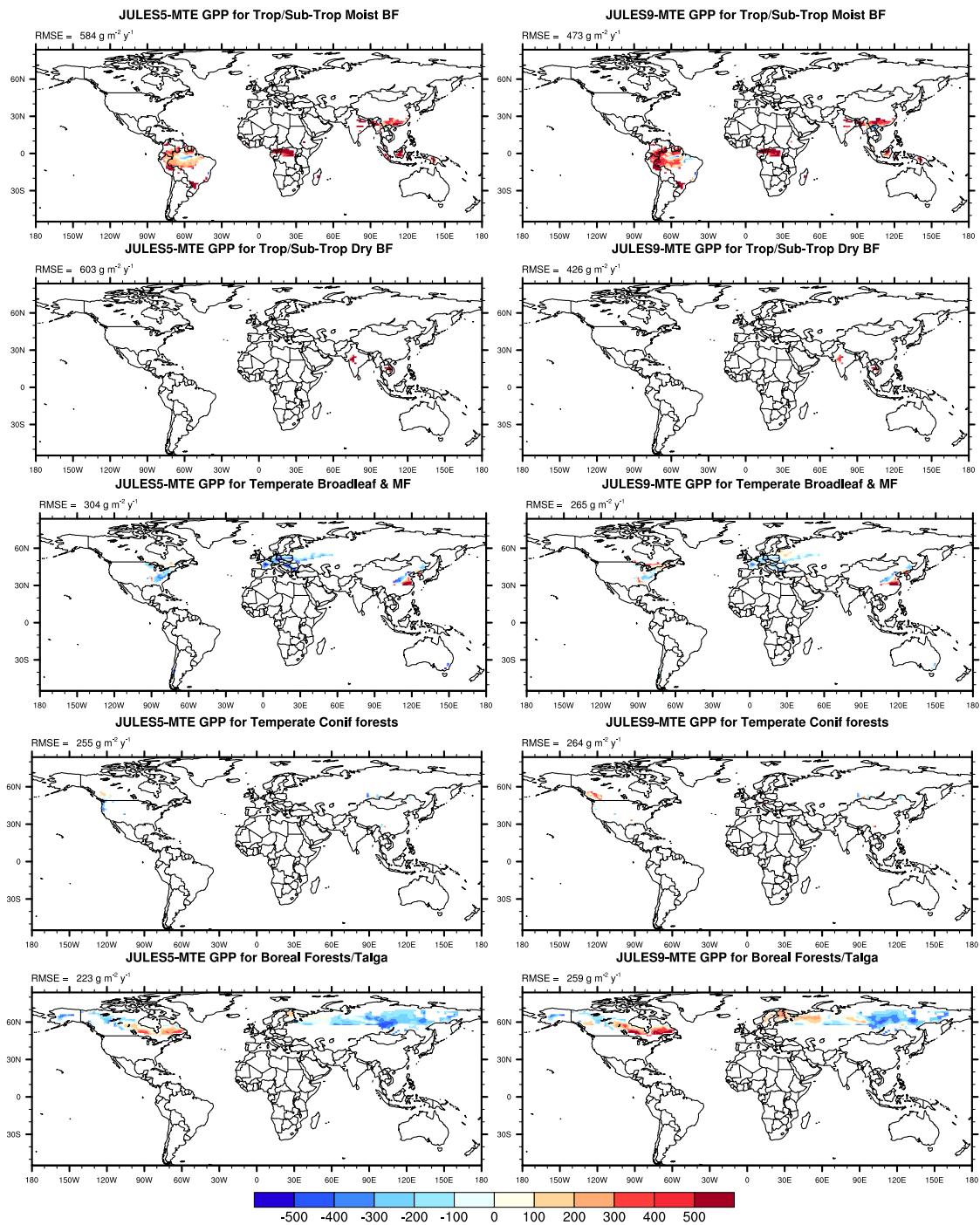


**Figure S4d.** As in Fig. S4a but for the deciduous tree sites.

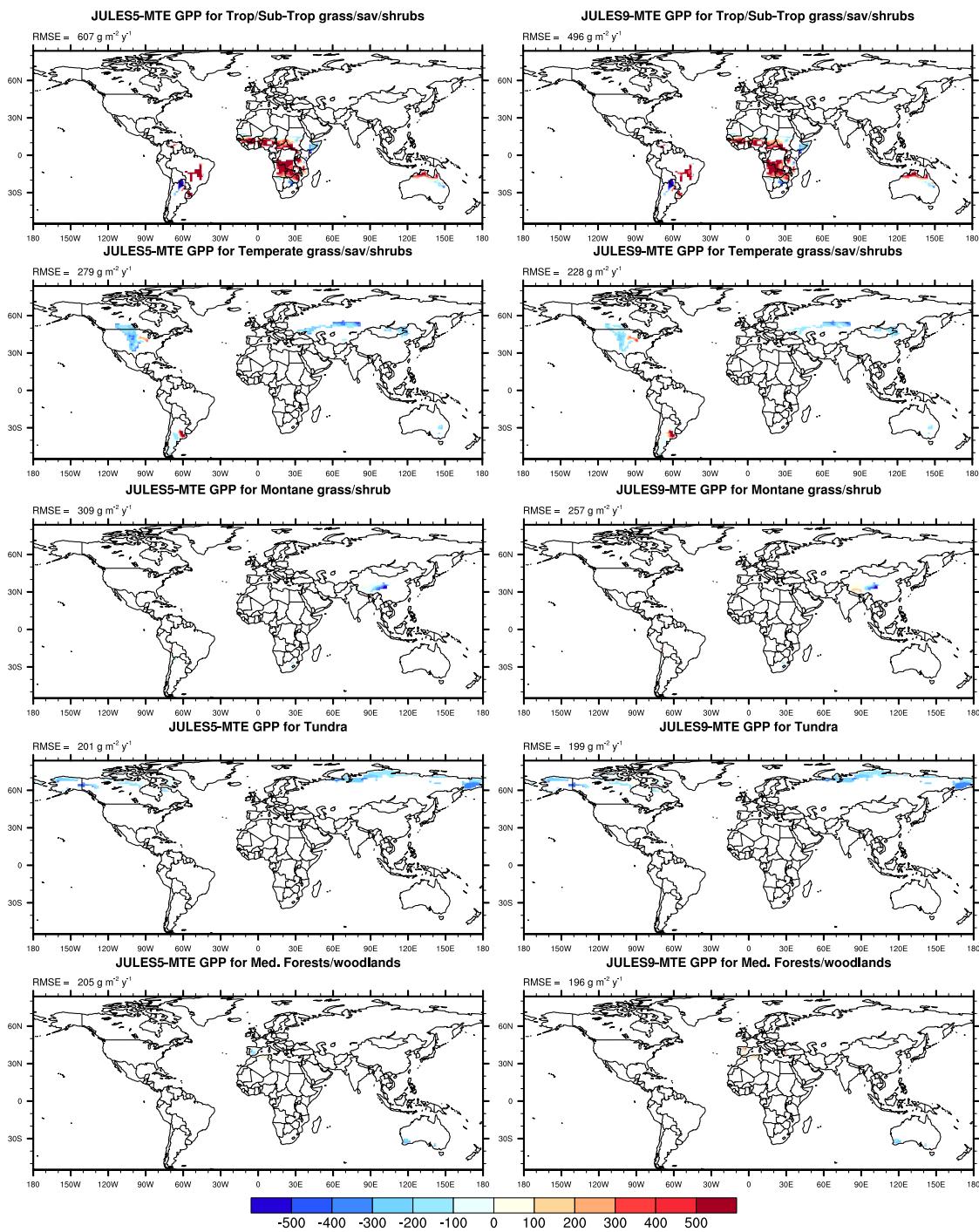


**Figure S5.** Seasonal cycles of Reco from the biomes shown in Fig. 3, comparing JULES5, JULES9, and the Jung et al. (2011) MTE. Also shown are the temperature and precipitation from the CRU-NCEP dataset used to force the JULES simulations.

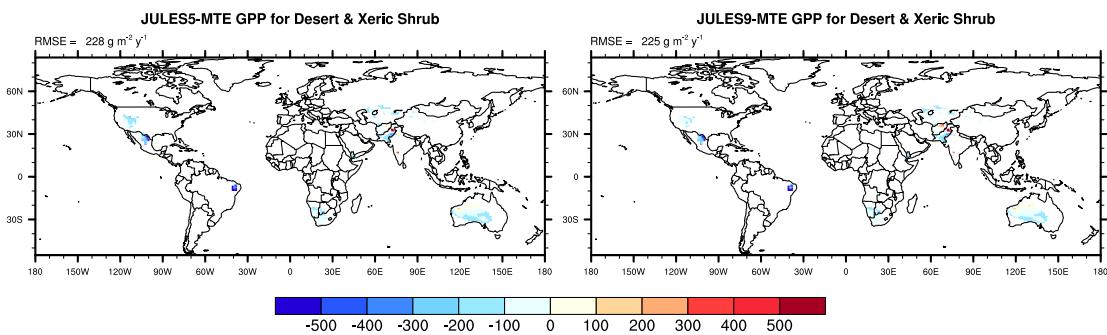
- MTE
- JULES9-ALL
- JULES5



**Figure S6a.** Differences between JULES5 and JULES9 and the MTE GPP for each of the 11 major biomes from the WWF database (biomes with area > 1,000 km<sup>2</sup>). The area-average root mean square error is given for each map.



**Figure S6b.** Differences between JULES5 and JULES9 and the MTE GPP for each of the 11 major biomes from the WWF database (biomes with area > 1,000  $\text{km}^2$ ). The area-average root mean square error is given for each map.



**Figure S6c.** Differences between JULES5 and JULES9 and the MTE GPP for each of the 11 major biomes from the WWF database (biomes with area > 1,000 km<sup>2</sup>). The area-average root mean square error is given for each map.