

# Inclusion of a dry surface layer and modifications to the transpiration and canopy evaporation partitioning in the Canadian Land Surface Scheme Including biogeochemical Cycles (CLASSIC)

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November 22, 2022

## Abstract

Land surface/Earth System models depend upon accurate simulation of evapotranspiration (ET) to avoid excessive biases in simulated energy, water, and carbon cycles. The Canadian Land Surface Scheme including biogeochemical Cycles (CLASSIC), the land surface scheme of the Canadian Earth System Model (CanESM) shows reasonable ET fluxes globally, but CLASSIC's partitioning into evaporation ( $E$ ) and transpiration ( $T$ ) can be improved. Specifically, CLASSIC exhibited a high soil evaporation ( $E_s$ ) bias in sparsely vegetated areas during wet periods, which can deplete soil water and decrease photosynthesis and  $T$  later in the year.

A dry surface layer (DSL) parameterization was implemented to address biases in  $E_s$  through an increased surface resistance to water vapour and heat fluxes. In arid/semi-arid regions, the DSL decreased  $E_s$ , leading to improved seasonality of ET and increased gross primary productivity (GPP) due to an increase in soil moisture. The DSL simulations significantly (t-test,  $p < 0.01$ ) increased  $T/ET$  from 0.25 in baseline CLASSIC to 0.30 in the DSL simulations.  $T/ET$  was further increased to 0.41 ( $p < 0.01$ ), comparable to the CMIP5 model mean, by allowing  $T$  to occur from the dry canopy fraction while water evaporates from the wet fraction. This mainly affected densely vegetated areas, where  $T$  and  $ET$  increased significantly ( $p < 0.01$ ) and canopy  $E$  was reduced ( $p < 0.01$ ). In seasonally dry tropical forests, higher  $T$  and  $ET$  reduced GPP. Despite increases in arid/semi-arid regions, the reduced GPP in tropical forests resulted in 1.6% lower global GPP ( $p = 0.018$ ) than baseline CLASSIC. Including these modifications in CanESM might reduce biases in climate.

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2        **the transpiration and canopy evaporation partitioning**  
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8        **Key Points:**

- 9        • Including a dry surface layer (DSL) parameterization reduced excessive soil evap-  
10        oration ( $E_s$ ) in CLASSIC globally, especially in dry regions  
11        • Evapotranspiration (ET) partitioning modifications increased photosynthesis in  
12        arid/semi-arid regions  
13        • Global transpiration ( $T$ ) to ET ratios were brought closer to observation-based  
14        estimates due to increased  $T$  and reduced  $E_s$  in dry regions

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**Plain Language Summary**

An important component of the global water cycle is the return of liquid water to the atmosphere from the land surface. Evaporation ( $E$ ) occurs on the surface of plants and the soil while transpiration ( $T$ ) is water that plants release through their stomata. We investigated how well  $E$  and  $T$  are simulated by the Canadian Land Surface Scheme including biogeochemical Cycles (CLASSIC). We found that the model simulated the total amount of water lost from the land surface reasonably well, but too much was lost via  $E$  at the expense of  $T$ . To improve this we changed how water evaporates from bare soil to more realistically capture the resistance to evaporating water as a thin dry layer forms on the soil surface. We additionally allowed the model to transpire water from plant leaves while other leaves on the plant were wet and evaporating, which was previously not the case. Our results improve the partitioning of  $E$  and  $T$  in CLASSIC resulting in more realistic simulated plant productivity in dry regions.

**1 Introduction**

Evapotranspiration (ET) is an important component of the global terrestrial water budget, with about 60% of precipitation over land returned to the atmosphere via ET (Jung et al., 2010). ET can be separated into its components soil evaporation ( $E_s$ ), canopy evaporation ( $E_c$ ) and transpiration ( $T$ ). These components draw water from different sources and their relative contributions to ET vary seasonally. While  $E_s$  mainly originates from the soil surface and shallow soil depths,  $T$  uses water accessible to plant roots and  $E_c$  occurs for a limited time following precipitation events, when water is on the vegetation canopy due to precipitation interception.  $E_s$ ,  $E_c$  and  $T$  then also show different sensitivities to environmental drivers of ET such as vapor pressure deficit (VPD) and solar radiation (K. Wang & Dickinson, 2012). Total ET and the contributions of the different ET components vary significantly between ecosystems and seasons, as vegetation cover, vegetation characteristics, i.e. leaf area index (LAI) and rooting depths, and soil characteristics differ (K. Wang & Dickinson, 2012).

Available measurements of ET or latent (LE) and sensible heat fluxes ( $H$ ) range temporally from half-hourly to annual and spatially from point-scale to hundreds of meters based upon techniques using lysimeters, eddy covariance or Bowen ratio methods and scintillometers (Barr et al., 1994; Baldocchi et al., 2001; Gavilán & Berengena, 2007; Malek & Bingham, 1993; Savage, 2009). On monthly to annual time scales over large regions, the surface water balance method can be used. It integrates measurements of precipitation, river discharges and estimates of changes in terrestrial water storage to determine LE on regional to global scales (L. Zhang et al., 2008; Güntner, 2008). Satellite remote sensing algorithms can estimate ET using empirical relationships with satellite-derived data (e.g., radiation, temperature, VPD) (Q. Mu et al., 2011). The relative contributions of the ET components can be estimated using several techniques. Canopy interception, which determines the amount of water available for  $E_c$ , can be estimated as the difference between total precipitation and the precipitation measured underneath the canopy (Herbst et al., 2008). For transpiration, sap flow methods can determine its value at the tree level, which can then be up-scaled to the stand or landscape level (Smith & Allen, 1996; Čermák et al., 2004; Warren et al., 2018). All three ET components,  $E_s$ ,  $E_c$  and  $T$ , can also be measured with stable isotope techniques whereby variations in the stable isotopic composition of water vapour measured near the surface in combination with measurements of the isotopic composition of water from the soil and within the plant can determine the transpiration and evaporation fractions (Sutanto et al., 2014). However, direct measurements of ET and its components are only available at small scales (e.g., plant level), and thus there are large uncertainties in global  $T/ET$  with estimates varying between 0.43 and 0.75 with a mean value of  $0.57 \pm 0.07$  (Wei et al., 2017). Most of global  $T$  originates from tropical forests (Good et al., 2015) with an estimated  $T/ET$  of  $0.70 \pm 0.14$  (Schlesinger & Jasechko, 2014), while shrublands and desert ecosystems tend to have the lowest  $T/ET$  with estimated values of  $0.47 \pm 0.10$  and  $0.54 \pm 0.18$ , respectively (Schlesinger & Jasechko, 2014).

Land surface models (LSMs) are used to simulate water and energy fluxes, including the different components of ET, for historical simulations and future projections. Despite challenges validating the different ET components on large scales, it is important for LSMs to correctly partition ET as it affects the water, energy and carbon (C) cycles (Swenson & Lawrence, 2014). Poorly simulated ET also has implications for the simulated climate in Earth System Models (ESMs). Dong et al. (2022) attributed a warm bias in 2 m air temperatures occurring in the central United States in models contributing to the Climate Model Intercomparison Project Phase 6 (CMIP6) to an underestimated ET and a low  $T/ET$ . In the CMIP6 models, ET is highly dependent on shallow soil moisture and water intercepted by the canopy while less dependent on root zone soil moisture. This leads to an underestimated contribution of  $T$  to ET, as well as a low total ET. Summertime ET in CMIP6 models was underestimated which was suggested to be a result of an overestimation of water stress, as the ESMs were not able to adequately simulate the ability of plants to access soil moisture in deeper layers, which can sustain  $T$ , and thereby were overly dependent on precipitation to supply near-surface soil moisture (Dong et al., 2022). While simulated ET partitioning varies between models, e.g.  $T/ET$  ranged from 0.20 to 0.57 (Lian et al., 2018), on average, the CMIP5 models underestimated  $T$  with an ensemble mean  $T/ET$  of  $0.41 \pm 0.11$  (Lian et al., 2018) compared to the estimated  $0.57 \pm 0.07$  (Wei et al., 2017) derived from upscaling site measurements using ecosystem-specific LAI regressions and LAI and canopy interception estimates from remote sensing and land surface models. As the underestimation of  $T/ET$  in ESMs leads to underestimations of summertime ET as well as overestimations of air temperature, improving ET partitioning in LSMs is important for future projections of the water and C cycles (Dong et al., 2022). Dong et al. (2020) suggest that  $E_s$  stress functions, commonly used in LSMs where they rely upon simple relationships with soil texture, cause biases in soil moisture-ET coupling in LSMs. Especially in bare soil areas or regions with sparse vegetation canopies, LSMs tend to overestimate ET due to an overestimation of  $E_s$  during periods of high soil moisture (Swenson & Lawrence, 2014). Over the past decade,

119 studies have shown that simulated  $E_s$  can be improved by different means including re-  
 120 sistance to  $E$  due to water vapour diffusion through a dry layer developing at the soil  
 121 surface (Swenson & Lawrence, 2014), a viscous sublayer (Haghighi & Or, 2015; Decker  
 122 et al., 2017) or a litter layer (Decker et al., 2017; M. Mu et al., 2021). Biases in simu-  
 123 lated ET and its component fluxes were also shown to be reduced by an improved rep-  
 124 resentation of the effects of soil texture in the  $E_s$  stress function, which decreased soil  
 125 moisture-ET coupling strength biases in the Noah land surface model with multiparam-  
 126 eterization options (Noah-MP version 3.6) (Dong et al., 2020).

127 In this study, we investigate ET and its component fluxes in the Canadian Land  
 128 Surface Scheme Including biogeochemical Cycles (CLASSIC). In order to improve mod-  
 129 elled  $E_s$  and  $T$ , a process-based ground evaporation efficiency parameterization, in which  
 130  $E_s$  is determined by water vapour diffusion through a thin dry surface layer (DSL) fol-  
 131 lowing Swenson and Lawrence (2014) was implemented. The partitioning into  $E_c$  and  
 132  $T$  was also modified such that the dry fraction of the canopy can transpire while  $E_c$  oc-  
 133 curs from the wet canopy fraction following Fan et al. (2019). We compare the modi-  
 134 fied and baseline CLASSIC versions at the site-level as well as globally, evaluating them  
 135 using eddy covariance or satellite-based observations of  $\text{CO}_2$  and energy fluxes. Section  
 136 2 describes CLASSIC as well as the modifications made to its partitioning of ET. Sec-  
 137 tion 3 shows the site-level and global water and carbon fluxes using the original CLAS-  
 138 SIFIC, CLASSIC including the DSL parameterization and CLASSIC including the DSL  
 139 as well as a modified partitioning into  $E_c$  and  $T$ . Differences between the carbon and  
 140 water fluxes of the three CLASSIC versions, how they compare with other LSMs and pos-  
 141 sible future improvements are discussed in Section 4.

## 142 2 Methods

### 143 2.1 Model description

144 CLASSIC is the land surface component of the Canadian Earth System Model (CanESM)  
 145 (Swart et al., 2019) and the open-source community model successor to CLASS-CTEM  
 146 (Melton et al., 2020), which couples the physics (the Canadian Land Surface Scheme;  
 147 CLASS; Verseghy (2017)) and biogeochemistry (the Canadian Terrestrial Ecosystem Model;  
 148 CTEM; Arora (2003); Melton and Arora (2016)) sub-modules. The exchange of energy,  
 149 water, momentum, and C at the land surface is represented in CLASSIC (for details see  
 150 Melton et al. (2020)). The model is driven by seven atmospheric variables: air temper-  
 151 ature ( $T_a$ ), precipitation rate, air pressure, specific humidity ( $q$ ), wind speed, and incom-  
 152 ing shortwave and longwave radiation. Water and heat fluxes between the ground and  
 153 the atmosphere, as well as its transfer between soil layers and snow, when present, are  
 154 usually calculated on a half-hourly time step. The number of soil layers and their thick-  
 155 nesses can be chosen depending on the application, but typically CLASSIC uses 20 ground  
 156 layers of increasing thicknesses starting with 10 layers of equal 10 cm thickness giving  
 157 a maximum depth of 61.4 m. Heat transfer occurs within the whole ground column, in-  
 158 cluding both soil and bedrock layers. The movement of water, however, is limited to the  
 159 permeable soil layers. Canopy conductance and photosynthesis are calculated on the same  
 160 time step as the energy and water fluxes while vegetation (leaf, stem, root), litter and  
 161 soil C pools as well as respiratory fluxes are calculated on a daily time step. We prescribed  
 162 the vegetation cover which is represented by plant functional types (PFTs) and their per-  
 163 cent coverage (see Table 1). Vegetation biomass and height, LAI and rooting depths are  
 164 dynamically determined within the biogeochemistry sub-module based upon photosyn-  
 165 thesis and respiration, PFT-specific C allocation parameters and land surface charac-  
 166 teristics (e.g., soil temperatures, soil moisture and net radiation) obtained from the physics  
 167 sub-module. The physical land surface properties are calculated separately for up to four  
 168 subareas of each grid cell (bare ground, snow-covered bare ground, vegetation over soil  
 169 and vegetation over snow). In CLASSIC version 1.2 as used here, the vegetation, as seen  
 170 by the physics submodule, is composed of five broad categories of PFTs (i.e., needleleaf

171 trees, broadleaf trees, crops, grasses and shrubs). The biogeochemical calculations dif-  
 172 ferentiate between evergreen and deciduous (split into cold and drought deciduous) PFTs  
 173 and  $C_3$  and  $C_4$  photosynthetic pathways for crops and grasses, which results in 12 PFTs  
 174 for the biogeochemistry sub-module.

## 175 2.2 Evapotranspiration parameterization and partitioning

176 The calculation of ET in CLASSIC and modifications made to its partitioning are  
 177 described in detail in Appendix A. In short, ET is calculated as the sum of the compo-  
 178 nents  $E_s$ , which originates from bare soil and the soil underneath the vegetation canopy,  
 179  $E_c$  and  $T$ . The potential evaporation rate from the soil depends on the ground evapo-  
 180 ration efficiency ( $\beta$ ; unitless; Equation A20), which is determined using an empirical for-  
 181 mulation based on Lee and Pielke (1992) and Merlin et al. (2011). Based on simulations  
 182 at flux tower sites, we determined that CLASSIC overestimates  $E_s$  and thus ET in sparsely  
 183 vegetated arid or semi-arid regions (see Meyer et al. (2021);  $E_s$  and ET are not shown  
 184 here, but the ET bias is comparable to the bias in LE in Figure B3). During the wet sea-  
 185 son, CLASSIC simulates excessive amounts of  $E_s$ , limiting the amount of moisture within  
 186 the soil for the dry season, which causes a suppression of photosynthesis in these regions.  
 187 In order to avoid excessive  $E_s$  from bare soil surfaces in CLASSIC, we implemented a  
 188 process-based  $\beta$  parameterization building on the previously used empirical formulation  
 189 (Merlin et al., 2011; Meyer et al., 2021). In the new formulation,  $E_s$  is determined by  
 190 water vapour diffusion through a thin DSL whose thickness is calculated based on the  
 191 moisture content of the top soil layer following Swenson and Lawrence (2014). This ap-  
 192 proach was chosen, as it is likely to have a stronger effect on  $E_s$  than a litter layer for  
 193 example in sparsely-vegetated areas. The way the DSL parameterization was implemented  
 194 also works well with CLASSIC’s structure. In addition to the DSL’s effects on  $E_s$ , we  
 195 have implemented its effects on  $H$  and the ground heat flux ( $G$ ) via the thermal con-  
 196 ductivity, which depends on the degree of soil saturation (see Section A4 for details). Changes  
 197 in surface albedo, when the DSL is present, were also accounted for (see Section A5). Avail-  
 198 able observations, although uncertain, indicate that CLASSIC also underestimates the  
 199 global  $T/ET$ . To address this, we modified the partitioning of  $E_c$  and  $T$ , as in the origi-  
 200 nal CLASSIC formulation  $T$  could only occur when there was no water on the canopy.  
 201 The modified CLASSIC version allows the dry fraction of the canopy to transpire while  
 202  $E_c$  occurs from the wet canopy fraction following Fan et al. (2019) (see Section A6 for  
 203 details).

## 204 2.3 Simulations

205 We performed simulations using CLASSIC v.1.2 with the original CEVAP formu-  
 206 lation of soil evaporation (labelled ‘Baseline’), a model version including the DSL pa-  
 207 rameterization (labelled ‘DSL’; Section A3) and a version including both the DSL pa-  
 208 rameterization and a modified partitioning of  $E_c$  and  $T$  (labelled ‘DSL-EcT’; Section A6)  
 209 (see Table 1). Site-level simulations for a selection of sites (see Table 2) from the FLUXNET2015  
 210 dataset (Pastorello et al., 2020) were driven by observed meteorology at these sites. Site-  
 211 specific information such as vegetation cover and composition, soil texture and depth were  
 212 obtained from the literature (Melton et al., 2020). At these sites, CLASSIC was driven  
 213 by cycling through the meteorological measurements available and the atmospheric  $CO_2$   
 214 concentration from the first year of measurements at each site until the C pools reached  
 215 equilibrium (defined as annual  $NEP / NPP \leq 0.02$ , where NEP is the net ecosystem pro-  
 216 ductivity and NPP is the net primary productivity). Then, CLASSIC was run for the  
 217 years available at each site with transient atmospheric  $CO_2$  concentrations from Le Quéré  
 218 et al. (2018).

219 We also performed global simulations on the CanESM grid (approximately  $2.8^\circ$  by  
 220  $2.8^\circ$ ). In order to assess differences between the CLASSIC versions and account for un-  
 221 certainty in model forcing and geophysical inputs, simulations are driven by combina-

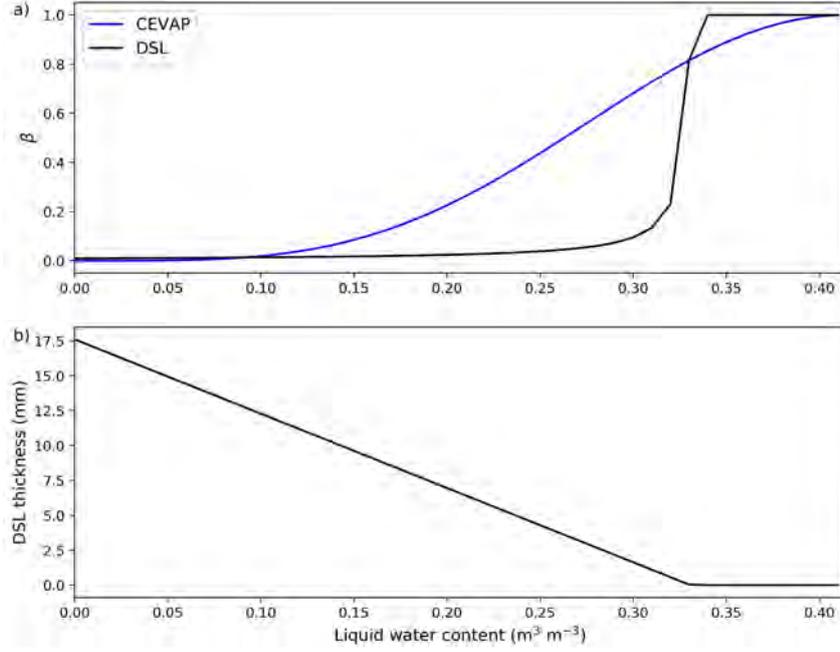


Figure 1: Soil evaporation efficiency ( $\beta$ ) determined using the original CEVAP parameterization (Equation A20; blue line) as well as the calculation using the resistance due to the DSL (first term on the right hand side of Equation A19; black line) for liquid water content values of the top soil layer ranging between 0 and the soil porosity (set to  $0.41 \text{ m}^3 \text{ m}^{-3}$  for this example) (a) and the thickness of the DSL for the respective liquid water content (b). This example was derived from annual average values of  $C_{DH} \times v_a$ ,  $\tau$ , and  $D_v$  from year 2005 at the US-Sta shrubland FLUXNET site (see Table 2).

Table 1: Calculation of the surface evaporation efficiency ( $\beta$ ) and the canopy evaporation ( $E_c$ ) and transpiration ( $T$ ) components in the three CLASSIC versions (Baseline, DSL and DSL-EcT) used in this study as well as the meteorological forcing and land cover representations used in the simulations. The simulations, where the meteorological forcing and land cover are bold, are the ones shown in the geographic distributions and in Figure 5.

Simulation	Surface evaporation efficiency	$E_c - T$ partitioning	Meteorological forcing	Land cover
Baseline	CEVAP (Equation A20)	$T$ only occurs, when the whole canopy is dry	<b>CRUJRAv2.2</b> GSWP3W5E5	<b>ESACCI</b> ESACCI
DSL	determined using DSL (Equation A19)	$T$ only occurs, when the whole canopy is dry	<b>CRUJRAv2.2</b> GSWP3W5E5	<b>GLC2000</b> GLC2000
DSL-EcT	determined using DSL (Equation A19)	$E_c$ occurs from wet canopy fraction, $T$ from dry canopy fraction (Section A6)	<b>CRUJRAv2.2</b> GSWP3W5E5 <b>CRUJRAv2.2</b> GSWP3W5E5	<b>ESACCI</b> ESACCI <b>GLC2000</b> GLC2000

222 tions of two different meteorological forcing datasets and two different land cover rep-  
 223 resentations, resulting in four simulations for each CLASSIC version. Meteorological forc-  
 224 ing for the simulations was either provided by the Climate Research Unit Japanese 55-  
 225 year reanalysis version 2.2 (CRUJRAv2.2, 1901-2020; CRU-JRA (2021); Harris et al. (2014,  
 226 2020); Kobayashi et al. (2015)) or the Global Soil Wetness Project Phase 3 (GSWP3)  
 227 -WFDE5 over land merged with ERA5 over the ocean (W5E5) (GSWP3W5E5; 1901-  
 228 2016; Lange (2020)). The methodology of Melton and Arora (2016) was used to disag-

229 gregate the 6-hourly meteorological data to the half-hourly time step CLASSIC uses. Model  
 230 preparation for historical simulations included spinups that cycled through the meteo-  
 231 rological forcings from 1700-1725 using constant CO<sub>2</sub> concentrations from 1700 until an  
 232 equilibrium state was reached. Then, historical simulations with transient CO<sub>2</sub> concen-  
 233 trations, vegetation cover and composition, including the effects of land use change and  
 234 fire, were performed from 1700-2019 (for CRUJRAv2.2) or 1700-2016 (for GSWP3W5E5).  
 235 Two different land cover representations are used for the CLASSIC simulations, which  
 236 are based on the Global Land Cover 2000 (GLC2000) and the European Space Agency  
 237 Climate Change Initiative (ESACCI; ESA (2017)) datasets. As described in A. Wang  
 238 et al. (2006), these datasets are mapped onto CLASSIC's PFTs and a timeseries includ-  
 239 ing changes in crop area is created. While the site-level simulations can include all 12  
 240 PFTs described in 2.1, the global land cover representations used here do not include  
 241 shrub PFTs and sedges, so they only include nine PFTs for the biogeochemical calcu-  
 242 lations and four PFTs (i.e., needleleaf trees, broadleaf trees, crops and grasses) for the  
 243 physics.

#### 244 **2.4 Observation-based estimates of global $T/ET$**

245 Global average estimates of the  $T/ET$  ratio have been reported by other studies  
 246 and can be used to evaluate our model results. Techniques used to determine  $T/ET$  in-  
 247 clude isotopes, site measurements, remote sensing as well as a hybrid approach (Wei et  
 248 al., 2017). As the water remaining after  $E$  is enriched in the heavier oxygen (<sup>18</sup>O) and  
 249 hydrogen (<sup>2</sup>H) isotopes, whereas  $T$  does not change isotope ratios, measurements of oxy-  
 250 gen and hydrogen stable isotope ratios can be used to partition  $T$  and  $E$  (Jasechko et  
 251 al., 2013). Jasechko et al. (2013) used isotope measurements from Earth's large lakes and  
 252 calculated lake catchment  $T$  from stable isotope mass balances between water inputs and  
 253 losses. They then up-scaled their calculations to global  $T$  using a global freshwater stable  
 254 isotope mass balance resulting in  $T/ET$  of 0.80-0.90 (25th and 75th percentiles) (Jasechko  
 255 et al., 2013). Coenders-Gerrits et al. (2014), however, suggest that Jasechko et al. (2013)'s  
 256  $T/ET$  was overestimated and showed that using different inputs results in  $T/ET$  of 0.50-  
 257 0.80 with a median value around 0.65. Further tests with different inputs and increased  
 258 uncertainty estimates, decreased the median  $T/ET$  even further to 0.58 and increased  
 259 its uncertainty with 25th and 75th percentiles of 0.35 and 0.8 (Coenders-Gerrits et al.,  
 260 2014). Good et al. (2015) also comment that previous studies might have overestimated  
 261  $T/ET$ , as they neglected that  $E$  originates from multiple pools and did not account for  
 262 their connectivity. Good et al. (2015) estimate the fraction of surface water that is bound  
 263 in the soil and accessible by plants for  $T$  versus mobile water that quickly passes through  
 264 the soil through preferential flow paths and is assumed to have the same isotopic com-  
 265 position as precipitation, as it does not mix with soil water. Good et al. (2015) deter-  
 266 mined the global terrestrial isotope budget using an isotope mass balance approach, grid-  
 267 ded land-atmosphere water fluxes and an estimate of the soil water-surface water con-  
 268 nectivity resulting in  $T/ET$  between 0.56 and 0.74 (25th and 75th percentiles) and a mean  
 269 of 0.64. A compilation of site-level measurements of  $T$  and  $ET$  in different ecoregions  
 270 using a range of techniques (eddy covariance, sap flow or isotopic approaches in combi-  
 271 nation with biophysical models to partition  $ET$ ) found a global mean  $T/ET$  of  $0.61 \pm$   
 272  $0.15$  ( $\pm 1$  SD) (Schlesinger & Jasechko, 2014). L. Wang et al. (2014) showed that site-  
 273 level  $T/ET$  ranged from 0.38 to 0.77 (25th and 75th percentiles) and that 43% of the  
 274 variations in  $T/ET$  could be explained by differences in LAI and the growing stage of  
 275 the ecosystem. The remote sensing-based global  $T/ET$  estimates used to evaluate CLAS-  
 276 SIC (Section 3.2) were obtained from studies using remotely sensed datasets of meteo-  
 277 rological variables (e.g., radiation, air temperature, precipitation) and vegetation char-  
 278 acteristics to drive different  $ET$  algorithms, which included the Penman-Monteith model  
 279 (PM-MOD; Q. Mu et al. (2007, 2011)), the Global Land Evaporation Amsterdam Model  
 280 (GLEAM; Miralles et al. (2011)), the Priestley-Taylor Jet Propulsion Laboratory (PT-  
 281 JPL; Fisher et al. (2008)) model (Miralles et al., 2016) and the Penman-Monteith-Leuning

Table 2: FLUXNET sites from the FLUXNET2015 dataset used in this study.

Site ID	Site name	Latitude	Longitude	Elevation	Years	IGBP code	DOI
AU-Tum	Tumbarumba	35.6566	148.1517	1200	2001–2013	EBF	<a href="https://doi.org/10.18140/FLX/1440126">https://doi.org/10.18140/FLX/1440126</a>
BR-Sa1	Santarem-Km67-Primary Forest	2.8567	54.9589	88	2002–2011	EBF	<a href="https://doi.org/10.18140/FLX/1440032">https://doi.org/10.18140/FLX/1440032</a>
CA-DL1	Daring Lake - Mixed Tundra	64.8689	-111.5748	425	2004–2017	OSH	<a href="https://doi.org/10.5281/zenodo.4301133">https://doi.org/10.5281/zenodo.4301133</a>
CA-Qfo	Quebec – E. Boreal, Mature Black Spruce	49.6925	74.3421	382	2003–2010	ENF	<a href="https://doi.org/10.18140/FLX/1440045">https://doi.org/10.18140/FLX/1440045</a>
CA-TPD	Ontario – Turkey Point Mature Deciduous	42.6353	80.5577	260	2012–2014	DBF	<a href="https://doi.org/10.18140/FLX/1440112">https://doi.org/10.18140/FLX/1440112</a>
CG-Tch	Tchizalamou	4.2892	11.6564	82	2006–2009	SAV	<a href="https://doi.org/10.18140/FLX/1440142">https://doi.org/10.18140/FLX/1440142</a>
CN-Dan	Dangxiang	30.4978	91.0664	4313	2004–2005	GRA	<a href="https://doi.org/10.18140/FLX/1440138">https://doi.org/10.18140/FLX/1440138</a>
CZ-BK1	Bily Kriz forest	49.5021	18.5369	875	2004–2014	ENF	<a href="https://doi.org/10.18140/FLX/1440143">https://doi.org/10.18140/FLX/1440143</a>
DE-Kli	Klingenberg	50.8931	13.5224	478	2004–2014	CRO	<a href="https://doi.org/10.18140/FLX/1440149">https://doi.org/10.18140/FLX/1440149</a>
DE-Tha	Tharandt	50.9624	13.5652	385	1996–2014	ENF	<a href="https://doi.org/10.18140/FLX/1440152">https://doi.org/10.18140/FLX/1440152</a>
DK-Sor	Sorø	55.4859	11.6446	40	1996–2014	DBF	<a href="https://doi.org/10.18140/FLX/1440155">https://doi.org/10.18140/FLX/1440155</a>
DK-ZaH	Zackenbergh Heath	74.4733	-20.5545	38	2000–2014	GRA	<a href="https://doi.org/10.18140/FLX/1440224">https://doi.org/10.18140/FLX/1440224</a>
FL-Hyy	Hyytiälä	61.8474	24.2948	181	1996–2014	ENF	<a href="https://doi.org/10.18140/FLX/1440158">https://doi.org/10.18140/FLX/1440158</a>
FR-Fon	Fontainebleau-Barbeau	48.4764	2.7801	103	2005–2014	DBF	<a href="https://doi.org/10.18140/FLX/1440161">https://doi.org/10.18140/FLX/1440161</a>
FR-Pue	Puéchabon	43.7413	3.5957	270	2000–2014	EBF	<a href="https://doi.org/10.18140/FLX/1440164">https://doi.org/10.18140/FLX/1440164</a>
ES-Amo	Amoladeras	36.8336	-2.2523	58	2008–2012	OSH	<a href="https://doi.org/10.18140/FLX/1440156">https://doi.org/10.18140/FLX/1440156</a>
ES-LgS	Laguna Seca	37.0979	2.9658	2267	2007–2009	OSH	<a href="https://doi.org/10.18140/FLX/1440225">https://doi.org/10.18140/FLX/1440225</a>
ES-LJu	Llano de los Juanes	36.9266	-2.7521	1600	2004–2013	OSH	<a href="https://doi.org/10.18140/FLX/1440157">https://doi.org/10.18140/FLX/1440157</a>
GF-Guy	Guyanaflux (French Guiana)	5.2788	52.9249	48	2004–2014	EBF	<a href="https://doi.org/10.18140/FLX/1440165">https://doi.org/10.18140/FLX/1440165</a>
GH-Ank	Ankasa	5.2685	2.6942	124	2011–2014	EBF	<a href="https://doi.org/10.18140/FLX/1440229">https://doi.org/10.18140/FLX/1440229</a>
IT-Lav	Lavarone	45.9562	11.2813	1353	2003–2014	ENF	<a href="https://doi.org/10.18140/FLX/1440169">https://doi.org/10.18140/FLX/1440169</a>
IT-SRo	San Rossore	43.7279	10.2844	6	1999–2012	ENF	<a href="https://doi.org/10.18140/FLX/1440176">https://doi.org/10.18140/FLX/1440176</a>
IT-Tor	Torgnon	45.8444	7.5781	2160	1999–2012	GRA	<a href="https://doi.org/10.18140/FLX/1440237">https://doi.org/10.18140/FLX/1440237</a>
MY-PSO	Pasoh Forest Reserve	2.9730	102.3062	147	2003–2009	EBF	<a href="https://doi.org/10.18140/FLX/1440240">https://doi.org/10.18140/FLX/1440240</a>
NL-Loo	Loobos	52.1666	5.7436	25	1996–2014	ENF	<a href="https://doi.org/10.18140/FLX/1440178">https://doi.org/10.18140/FLX/1440178</a>
PA-SPs	Sardinilla-Pasture	9.3138	79.6314	68	2007–2009	GRA	<a href="https://doi.org/10.18140/FLX/1440179">https://doi.org/10.18140/FLX/1440179</a>
RU-Che	Cherski	68.6130	161.3414	6	2002–2005	WET	<a href="https://doi.org/10.18140/FLX/1440181">https://doi.org/10.18140/FLX/1440181</a>
RU-Fyo	Fyodorovskoye	56.4615	32.9221	265	1998–2014	ENF	<a href="https://doi.org/10.18140/FLX/1440183">https://doi.org/10.18140/FLX/1440183</a>
RU-Ha1	Hakasia steppe	54.7252	90.0022	446	2002–2004	GRA	<a href="https://doi.org/10.18140/FLX/1440184">https://doi.org/10.18140/FLX/1440184</a>
RU-SkP	Yakutsk Spasskaya Pad larch	62.2550	129.1680	246	2012–2014	DNF	<a href="https://doi.org/10.18140/FLX/1440243">https://doi.org/10.18140/FLX/1440243</a>
SD-Dem	Demokeya	13.2829	30.4783	500	2005–2009	SAV	<a href="https://doi.org/10.18140/FLX/1440186">https://doi.org/10.18140/FLX/1440186</a>
US-MMS	Morgan Monroe State Forest	39.3232	-86.4131	275	1999–2014	DBF	<a href="https://doi.org/10.18140/FLX/1440083">https://doi.org/10.18140/FLX/1440083</a>
US-SRC	Santa Rita Creosote	31.9083	-110.8395	950	2008–2014	OSH	<a href="https://doi.org/10.18140/FLX/1440098">https://doi.org/10.18140/FLX/1440098</a>
US-Sta	Saratoga	41.3966	-106.8024	2069	2005–2009	OSH	<a href="https://doi.org/10.18140/FLX/1440115">https://doi.org/10.18140/FLX/1440115</a>
US-WCr	Willow Creek	45.8059	90.0799	520	1999–2014	DBF	<a href="https://doi.org/10.18140/FLX/1440095">https://doi.org/10.18140/FLX/1440095</a>
US-Whs	Walnut Gulch Lucky Hills Shrub	31.7438	-110.0522	1370	2008–2014	OSH	<a href="https://doi.org/10.18140/FLX/1440097">https://doi.org/10.18140/FLX/1440097</a>
US-Wkg	Walnut Gulch Kendall grasslands	31.7365	109.9419	1531	2004–2014	GRA	<a href="https://doi.org/10.18140/FLX/1440096">https://doi.org/10.18140/FLX/1440096</a>
ZA-Kru	Skukuza	25.0197	31.4969	359	2000–2013	SAV	<a href="https://doi.org/10.18140/FLX/1440188">https://doi.org/10.18140/FLX/1440188</a>
ZM-Mon	Mongu	15.4378	23.2528	1053	2007–2009	DBF	<a href="https://doi.org/10.18140/FLX/1440189">https://doi.org/10.18140/FLX/1440189</a>

IGBP land classification abbreviations used include evergreen broadleaf forest (EBF), evergreen needleleaf forest (ENF), savanna (SAV), deciduous broadleaf forest (DBF), grassland (GRA), cropland (CRO), open shrubland (OSH), closed shrubland (CSH), permanent wetland (WET), and deciduous needleleaf forest (DNF).

282 (PML; Y. Zhang et al. (2016)) model. Of the remote-sensing-based global ET models,  
 283 PM-MOD is one of the most widely used (for example in the MODIS ET product MOD16).  
 284 Unlike other Penman-Monteith-based models, PM-MOD determines the surface and aero-  
 285 dynamic resistances without using soil moisture or wind speed inputs. Its resistance pa-  
 286 rameters have, however, been calibrated using EC towers, which is not required for GLEAM  
 287 and PT-JPL (Miralles et al., 2016). GLEAM determines the ET components based on  
 288 Priestley and Taylor (1972) apart from interception losses, which use Gash (1979)’s an-  
 289 alytical model driven by precipitation observations (Miralles et al., 2016). Despite us-  
 290 ing the same forcing datasets to drive PM-MOD, GLEAM and PT-JPL, where input re-  
 291 quirements overlapped, differences in modelled ET and its component fluxes were large  
 292 (Miralles et al., 2016). Compared to the other two models and data from ERA-Interim  
 293 reanalysis (Dee et al., 2011) and the model tree ensemble (MTE; Jung et al. (2009, 2010))  
 294 product, which uses a machine-learning algorithm trained on FLUXNET data, PM-MOD  
 295 tends to underestimate ET, especially in the tropics and dry subtropical regions, apart  
 296 from in the Northern high-latitudes (Miralles et al., 2016). In high latitudes, GLEAM  
 297 and PT-JPL had lower ET than PM-MOD likely due to deficiencies in the Priestley-Taylor  
 298 approach when available energy is low (Miralles et al., 2016). Partitioning of ET into  
 299  $E_s$ ,  $E_c$  and  $T$  largely differs between the models with  $T$  being much lower in PM-MOD  
 300 and  $E_s$  and  $E_c$  higher than in the other two models (Miralles et al., 2016). PM-MOD’s  
 301  $T/ET$  of 0.24 is an outlier compared with the other observation-based estimates (see Sec-  
 302 tion 3.2), which might in part be due to its underestimation of ET in the tropics and dry  
 303 sub-tropics, which tend to contribute the most to global  $T$  (Schlesinger & Jasechko, 2014).  
 304 PM-MOD also shows relatively high  $E_s$  in tropical regions, where GLEAM and PT-JPL  
 305 show very little  $E_s$ .  $E_c$ ’s contribution to ET is also much larger on average in PM-MOD  
 306 with 24% compared to 18% in PT-JPL and 10% in GLEAM (Miralles et al., 2016) and  
 307 PML (Y. Zhang et al., 2016).

### 308 3 Results

#### 309 3.1 Site-level results

310 Baseline CLASSIC simulations at a selection of FLUXNET sites (Table 2) showed  
 311 an overestimation of LE compared with eddy covariance measurements at sparsely veg-  
 312 etated sites such as open shrublands (e.g., ES-Amo, ES-LJu, US-Sta) during wet peri-  
 313 ods and an underestimation of LE during the peak growing season or drier periods at  
 314 these sites (Figure 2 and B3). During wet periods, when LE was overestimated,  $H$  was  
 315 underestimated (Figure B4). During dry periods, on the other hand,  $H$  was overestimated  
 316 compared to observations. The ground heat flux ( $G$ ; Figure B5) also tended to be over-  
 317 estimated at these sites, especially during summer. The overestimation of LE led to a  
 318 strong reduction in soil moisture in the top layer (Figure B6). Thus, GPP was reduced  
 319 and underestimated during the peak growing season at several open shrubland or grass-  
 320 land sites (e.g., ES-LJu, US-SRC, ES-Amo; Figure 2 and B2).

321 The DSL simulations mainly affected LE with statistically significant (t-test,  $p <$   
 322 0.01) differences between the bias in the DSL simulation and the bias in the Baseline sim-  
 323 ulation (Figure B8) at sparsely vegetated sites such as open shrubland sites with a large  
 324 bare ground area (e.g., ES-Amo, ES-LgS, ES-LJu, US-Sta, US-SRC, US-Whs), where  
 325 the DSL parameterization reduced LE during wet periods and increased LE during dry  
 326 periods (Figure 2 and B3). Thus, the DSL simulations eliminated or reduced overesti-  
 327 mation of LE during wet times and the simulated seasonal cycle of LE more closely rep-  
 328 resented observations at these sites. The reduction in LE in the wet season led to an in-  
 329 crease in soil moisture of the top layer (Figure B6) and thus to higher GPP later in the  
 330 year (Figure 2 and B2). At the majority of sites,  $G$  was only minimally affected (Fig-  
 331 ure B5).

332 The DSL-EcT modifications affected LE at both sparsely vegetated sites as well  
333 as sites with high LAI (e.g., evergreen forest sites; Figures 2, B3, B7 and B8). As in the  
334 DSL simulations, LE at the sparsely vegetated sites was reduced during wet periods and  
335 increased during dry periods due to the DSL parameterization (Figure 2). Effects of the  
336  $E_c$  and  $T$  partitioning modifications on LE at the open shrubland sites were minor, while  
337 LE at densely vegetated sites such as tropical forests in Brazil (BR-Sa1) or French Guiana  
338 (GF-Guy) was more strongly impacted (Figure B3). At the sites where LE increased in  
339 the DSL-EcT simulations (mainly denser forest sites),  $H$  decreased (Figure B4). For the  
340 more sparsely-vegetated sites, on the other hand,  $H$  slightly increased, especially dur-  
341 ing wetter periods. The effect on  $G$  at each site was minimal (Figure B5). The  $E_c$  and  
342  $T$  partitioning modifications resulted in slightly lower GPP than for the DSL simulation,  
343 as LE increased and the liquid water content of the top soil layer ( $\theta_{l,1}$ ) decreased.

344 Biases in simulated daily LE and GPP compared to observations for the FLUXNET  
345 sites were reduced in the DSL and DSL-EcT simulations compared to the Baseline sim-  
346 ulations at several of the sites (Figure 3 and 4). Averaged over all the sites, the mean  
347 absolute error (MAE) and root mean square error (RMSE) in LE were reduced in the  
348 DSL-EcT simulations compared to the Baseline with a decrease in mean MAE of 5% and  
349 in mean RMSE of 4% (Figure 3). T-tests showed that the majority of sites showed sta-  
350 tistically significant differences ( $p < 0.01$ ) between daily simulated and observed LE and  
351 GPP for all model configurations (Figure 3 and 4). The magnitude of reduction in MAE  
352 and RMSE varied between sites.

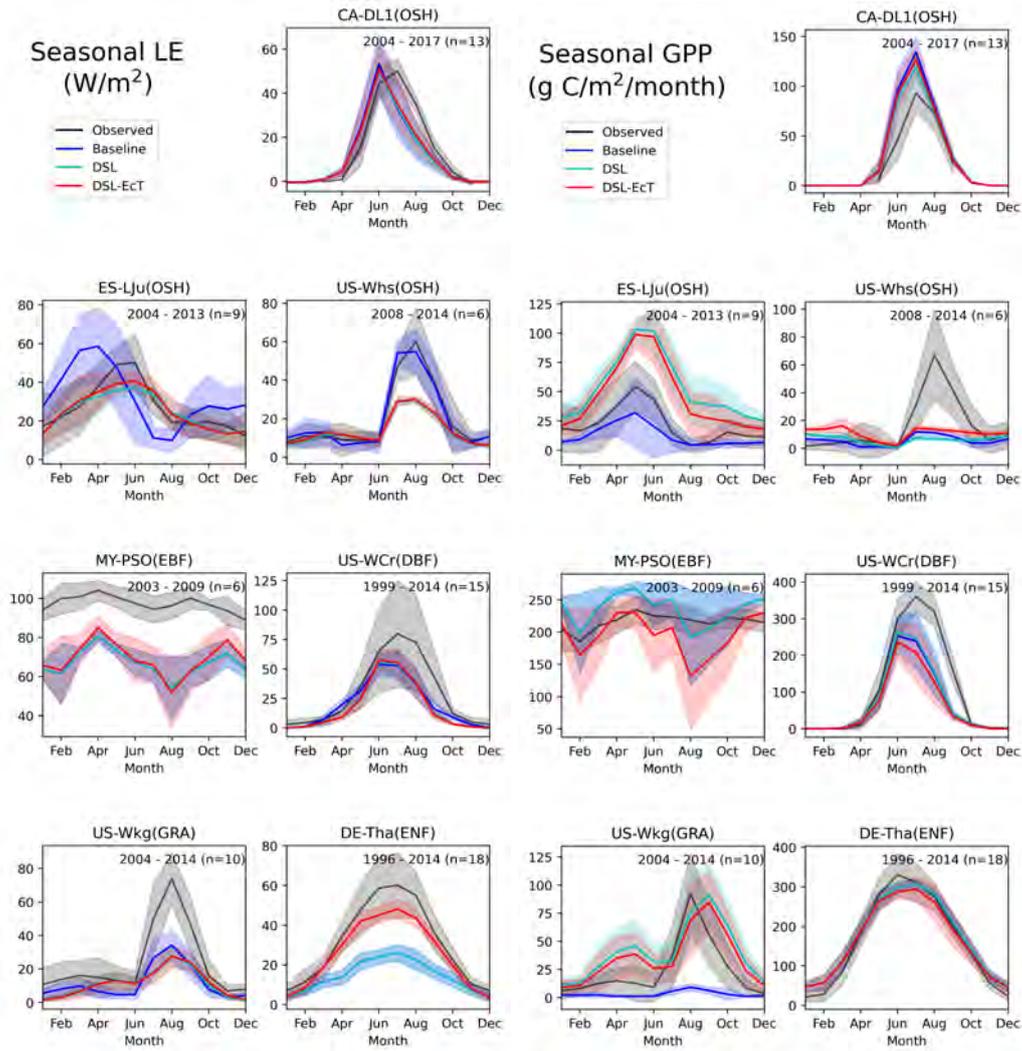


Figure 2: Monthly mean observed and simulated latent heat flux (LE) and gross primary productivity (GPP) for a selection of FLUXNET sites (showing different biomes and climates) for the Baseline, DSL and DSL-EcT simulations (Table 1). All sites are shown in Figures B2 and B3. The shading shows the standard deviation over the available years. Site names, their biomes and years of measurements used are listed for each site (for more details see Table 2). For some sites, the results from the different simulations and observations are overlapping and lines may be difficult to distinguish.

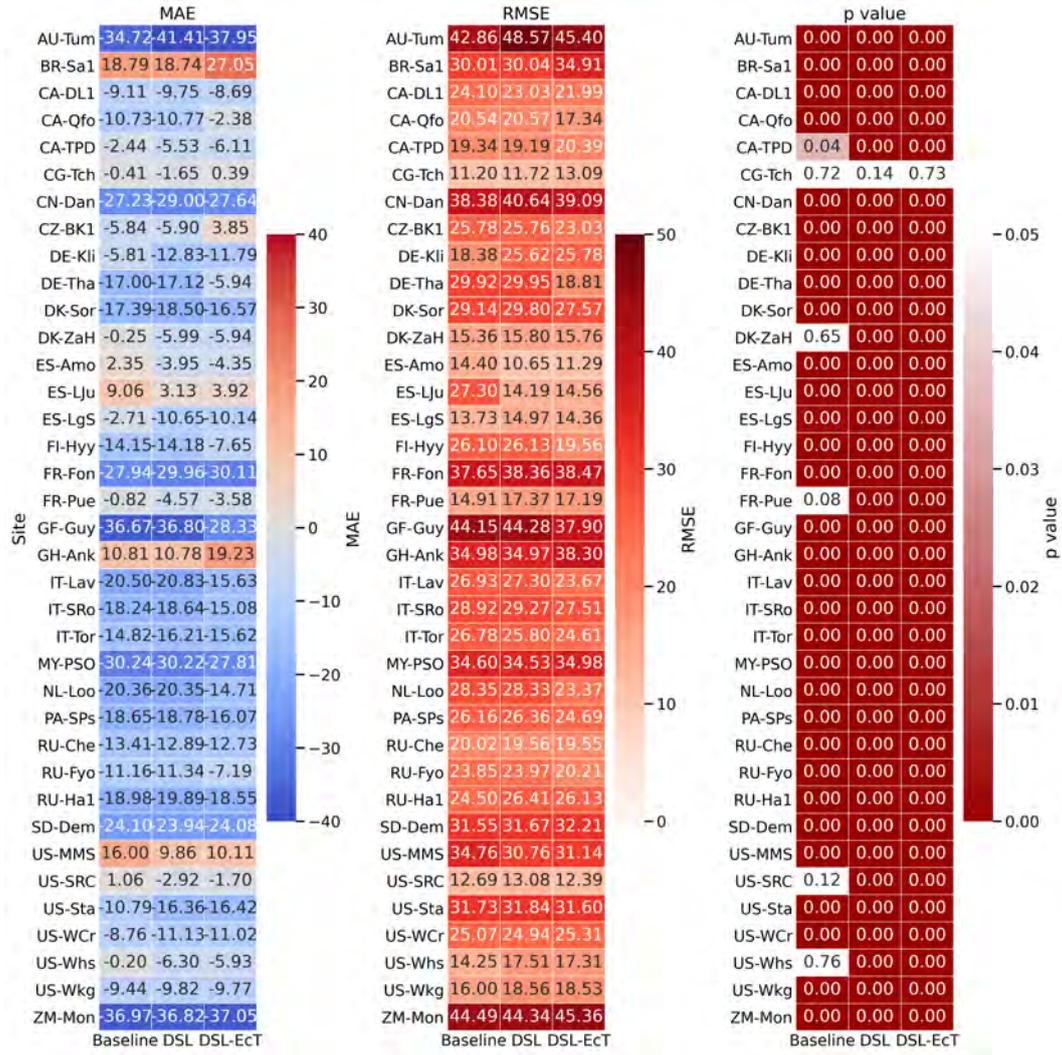


Figure 3: Mean absolute error (MAE,  $W m^{-2}$ ), root mean square error (RMSE,  $W m^{-2}$ ) and p value (determined using an independent two-sided t-test) between the observed and simulated daily latent heat flux (LE) for the FLUXNET sites (Table 2) for the Baseline, DSL and DSL-EcT simulations (Table 1). For MAE and RMSE, values closer to zero indicate better model performance.

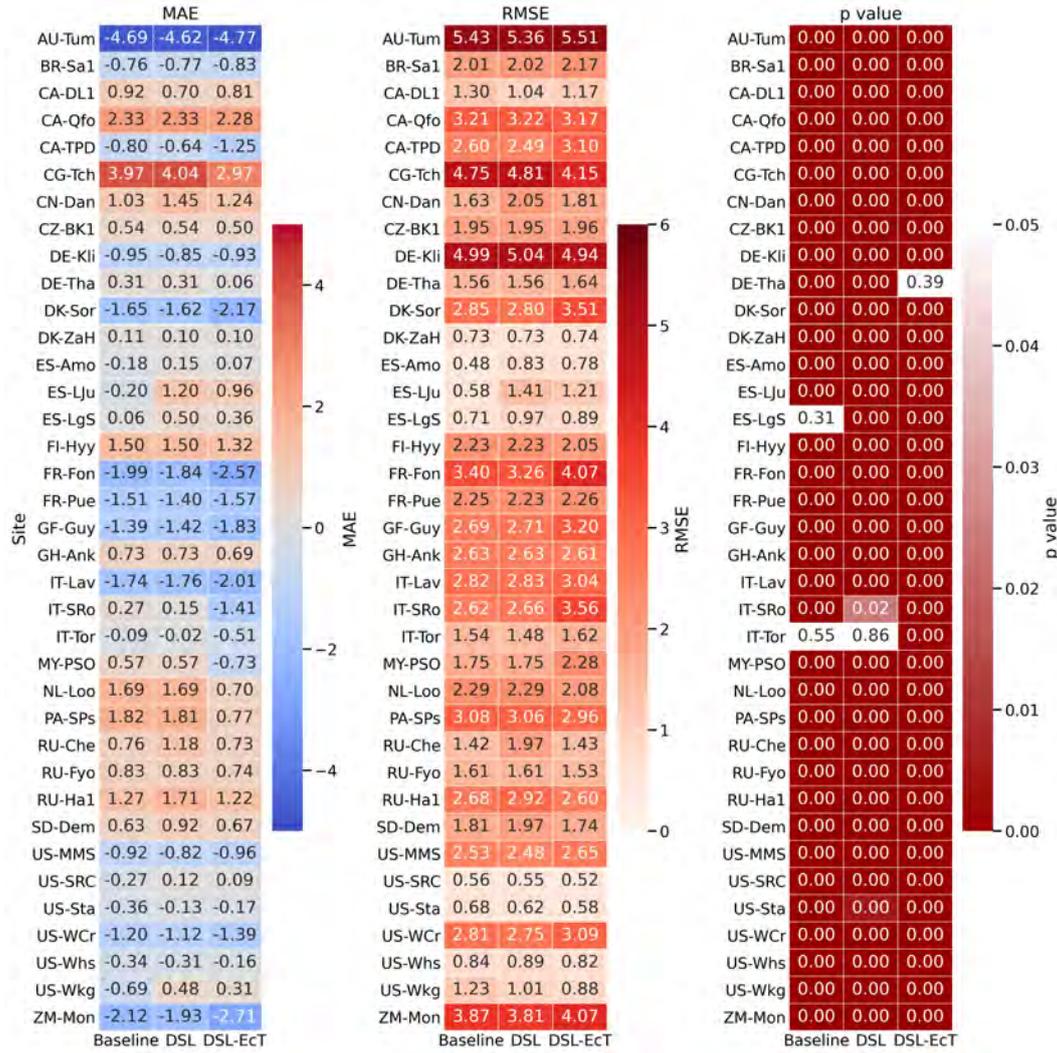


Figure 4: Mean absolute error (MAE,  $\text{g C m}^{-2} \text{ day}^{-1}$ ), root mean square error (RMSE,  $\text{g C m}^{-2} \text{ day}^{-1}$ ) and p value (determined using an independent two-sided t-test) between the observed and simulated daily gross primary productivity (GPP) for the FLUXNET sites (Table 2) for the Baseline, DSL and DSL-EcT simulations (Table 1). For MAE and RMSE, values closer to zero indicate better model performance.

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### 3.2 Global results

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For the baseline CLASSIC simulations, the largest contribution to global ET comes from  $E_s$  (39.5%), followed by  $E_c$  (34.6%) and  $T$  (25.9%) (Figure 5). Results shown in Figure 5 focus on simulations using one land cover and one meteorological forcing dataset as the results were similar across the simulations using different land cover and meteorological forcing. The ET partitioning in baseline CLASSIC results in a lower  $T/ET$  ratio than estimates from isotope, remote sensing (apart from the PM-MOD algorithm, which is an outlier compared to other remote sensing-based estimates as discussed in Section 2.4) or site measurements as well as several other process-based models suggest (Figure 6). Total global GPP, LE and  $H$  are within the uncertainty bounds of observation-based estimates (Figure C1a-f). Low productivity regions such as the southwestern United

364 States, southern Australia, southern South America, however, show very low to zero GPP.  
 365 Due to the low GPP,  $E_s$  exceeds  $E_c$  and  $T$  in these regions (not shown).

366 Implementing the DSL parameterization changes CLASSIC's ET partitioning by  
 367 significantly reducing  $E_s$  (t-test,  $p < 0.01$ ) and increasing  $E_c$  (t-test,  $p = 0.013$ ) and  $T$   
 368 (t-test,  $p < 0.01$ ) compared to baseline CLASSIC (Figure 5). As  $T$  increased and ET  
 369 decreased, global  $T/ET$  significantly increased (t-test,  $p < 0.01$ ) from  $\sim 0.25$  on average  
 370 (taken across the simulations listed in Table 1) in baseline CLASSIC to  $\sim 0.30$  in the DSL  
 371 simulations (Figure 6). Slightly increased  $T$  in the DSL simulations was due to an in-  
 372 crease in GPP, especially in arid and semi-arid regions, where  $E_s$  was reduced and soil  
 373 moisture available to the vegetation increased.

374 Combining the DSL parameterization with the modifications to  $E_c$  and  $T$  parti-  
 375 tioning resulted in slightly increased ET (t-test,  $p < 0.01$ ), a reduction in  $E_s$  (t-test,  $p$   
 376  $< 0.01$ ), a slight decrease in  $E_c$  (t-test,  $p < 0.01$ ) and an increase in  $T$  (t-test,  $p < 0.01$ )  
 377 (Figure B9). Thus, the  $T/ET$  ratio is significantly higher (t-test,  $p < 0.01$ ) for the DSL-  
 378 EcT simulations than the CEVAP simulations by 0.15 - 0.17 depending on the land cover  
 379 and meteorological forcing (Table B1 and Figure 5). The DSL-EcT modifications decreased  
 380 both  $E_s$  and  $E_c$  while increasing  $T$  (all statistically significant, t-test,  $p < 0.01$ ). Despite  
 381 differences in the water fluxes between simulations using the different forcings (see Fig-  
 382 ure B9 and Section 2.3), the modified CLASSIC versions changed ET and its partition-  
 383 ing more than the different forcing datasets did (Figure 5 and Table B1). The  $T/ET$  for  
 384 the DSL-EcT simulations remained lower than several observation-based estimates us-  
 385 ing isotopes, other site measurements or remote sensing algorithms (except PM-MOD)  
 386 suggest, but was closer to estimates from other models and is close to the CMIP5 en-  
 387 semble mean value of 0.41 (Lian et al., 2018).

388 As CLASSIC's  $T/ET$  remained low compared to observations and some other LSMs,  
 389 we considered further options to improve its ET partitioning. Lian et al. (2018) suggested  
 390 that the simulation of  $T/ET$  in ESMs could be improved by taking into account the dif-  
 391 fuse fraction of incoming radiation, as it would affect the photosynthetic activity of shaded  
 392 leaves and likely increase  $T$ . Including the diffuse radiation fraction using a 2-leaf pho-  
 393 tosynthesis scheme instead of the big-leaf scheme in CLASSIC (Arora, 2003), however,  
 394 only had minor impacts on  $T$  and  $T/ET$ . Both increased by  $\sim 2\%$ , respectively, compared  
 395 to the big-leaf scheme without the diffuse radiation fraction in CLASSIC resulting in  $T/ET$   
 396 of 0.427 driven with CRUJRA and ESACCI.

397 GPP significantly increased (t-test,  $p < 0.01$ ) in arid and semi-arid regions such  
 398 as the southwestern United States and Australia due to the modifications in ET parti-  
 399 tioning while some densely vegetated areas (especially in the Tropics) showed a reduc-  
 400 tion in GPP (Figure 7b). The majority of the regions where GPP increased in the DSL-  
 401 EcT simulations showed an increase in  $H$  and a small reduction in LE (Figure 7d and  
 402 f) or ET, as the decrease in  $E_s$  (Figure 8b) exceeded the increases in  $E_c$  and  $T$  (Figure  
 403 8d and f). In areas with reduced GPP,  $H$  decreased and LE (Figure 7b, d and f) and  
 404 ET tended to increase, as  $E_s$  increased or decreased only slightly,  $E_c$  decreased moder-  
 405 ately and  $T$  increased statistically significantly (t-test,  $p < 0.01$ ; Figure 8d and f). In  
 406 tropical forests (here defined as areas between  $25^\circ\text{S}$  and  $25^\circ\text{N}$  with a leaf area index  $>$   
 407  $3 \text{ m}^2 \text{ m}^{-2}$ ), the DSL-EcT simulation mainly affects  $E_c$  and  $T$  while the change in  $E_s$  is  
 408 relatively small. In semi-arid regions, on the other hand,  $E_s$  and  $T$  are more strongly af-  
 409 fected than  $E_c$ .

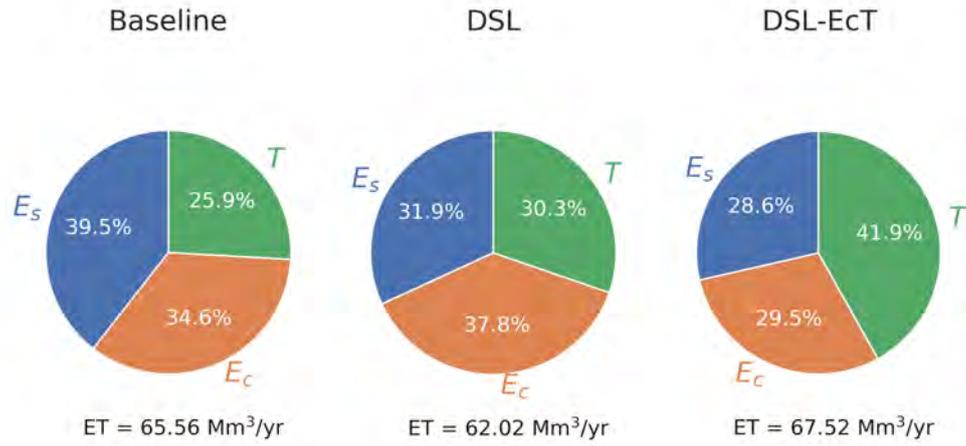


Figure 5: Partitioning of evapotranspiration (ET) into its components soil evaporation ( $E_s$ ), canopy evaporation ( $E_c$ ) and transpiration ( $T$ ) for the three different CLASSIC versions (see Section 2.3; Table 1). Percentages of  $E_s$ ,  $E_c$  and  $T$  are global averages over 1997-2016 for simulations using the CRUJRA meteorological forcing and the ESACCI land cover. Total ET values for the different CLASSIC versions are shown below the pie charts.

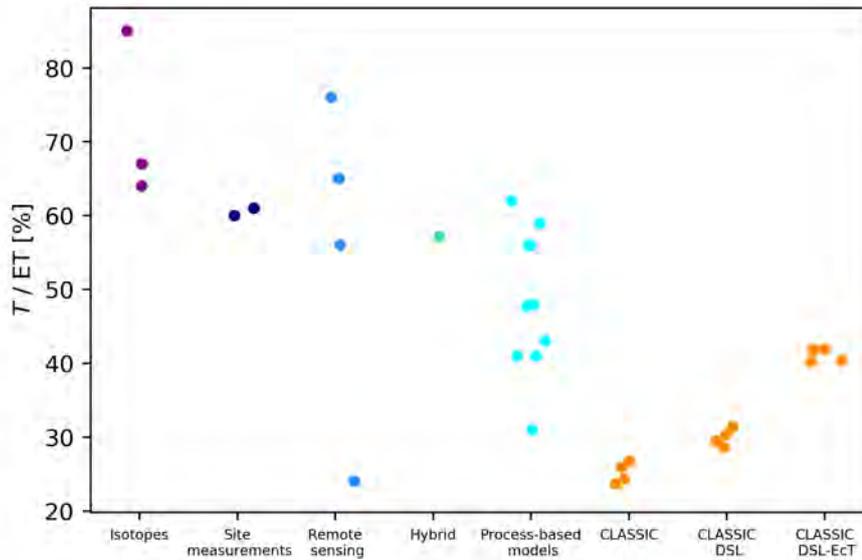


Figure 6: Globally averaged  $T/ET$  from observation-based datasets (see Section 2.4) and different land surface or Earth System models (adapted from Wei et al. (2017)) alongside the versions of CLASSIC tested in our study (see Section 2.3). The “Hybrid” reference dataset uses site measurements, satellite-based observations, as well as land surface model (a complex physically based model i.e., the Community Land Model 4.5, the remote sensing-based Global Land Evaporation Amsterdam Model (GLEAM) and a simple biophysical model i.e., Penman-Monteith-Leuning Model; PML) outputs, to upscale site-level measurements of the ET components (Wei et al., 2017). For each of the CLASSIC versions (Table 1), the four points represent the results using a combination of the two different meteorological forcing datasets and the two land cover representations (Section 2.3). The horizontal displacement of the dots is just to allow each one to be visible.

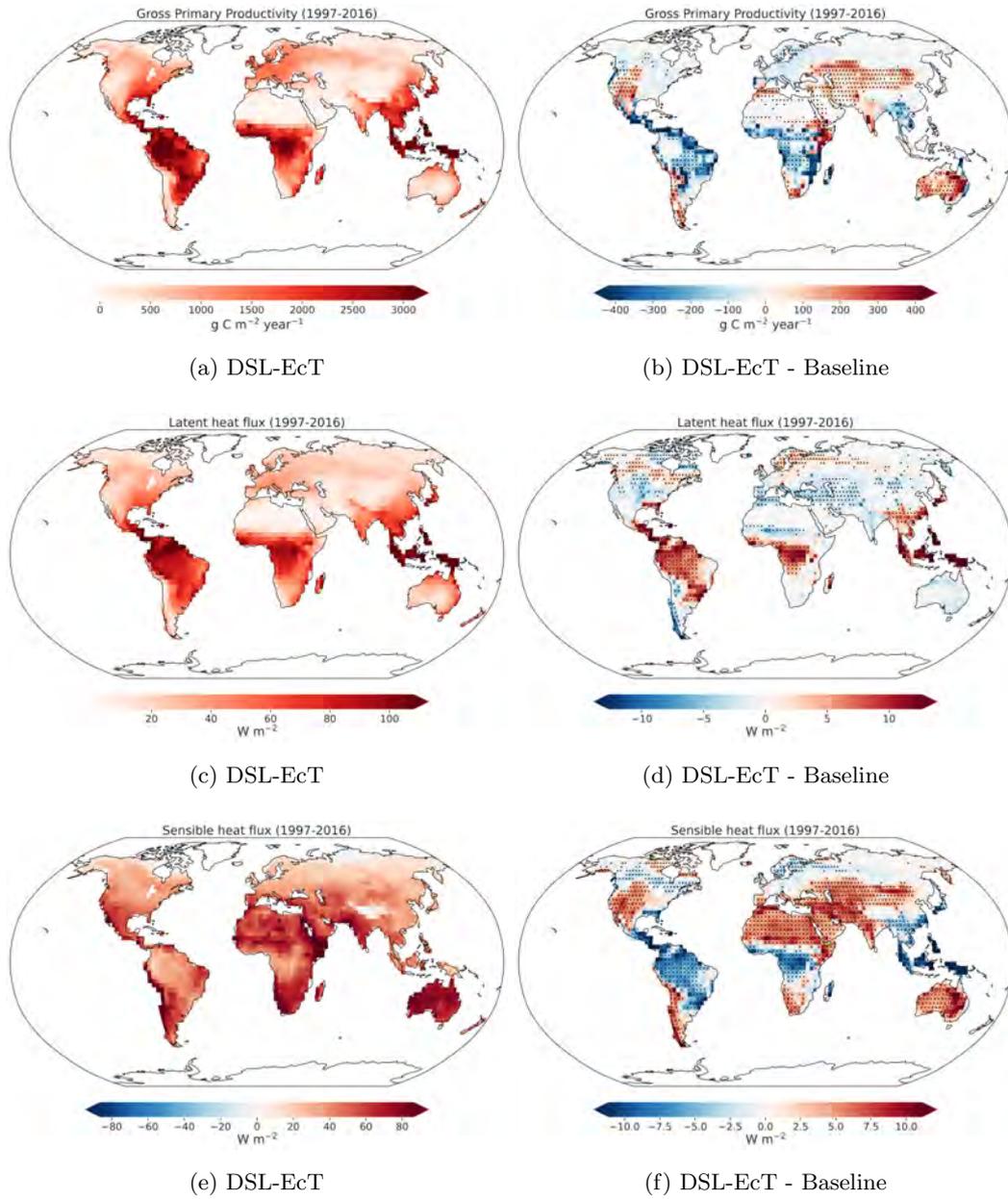


Figure 7: Geographic distribution of annual gross primary productivity (GPP), latent (LE) and sensible ( $H$ ) heat flux averaged over 1997-2016 for the DSL-EcT simulation (a, c, e) and the difference between the DSL-EcT and Baseline simulations (b, d, f) using the CRUJRA meteorological forcing and the ESACCI land cover. Grid cells with dots indicate that differences are statistically significant (independent two-sample t-test  $p$  level < 0.01).

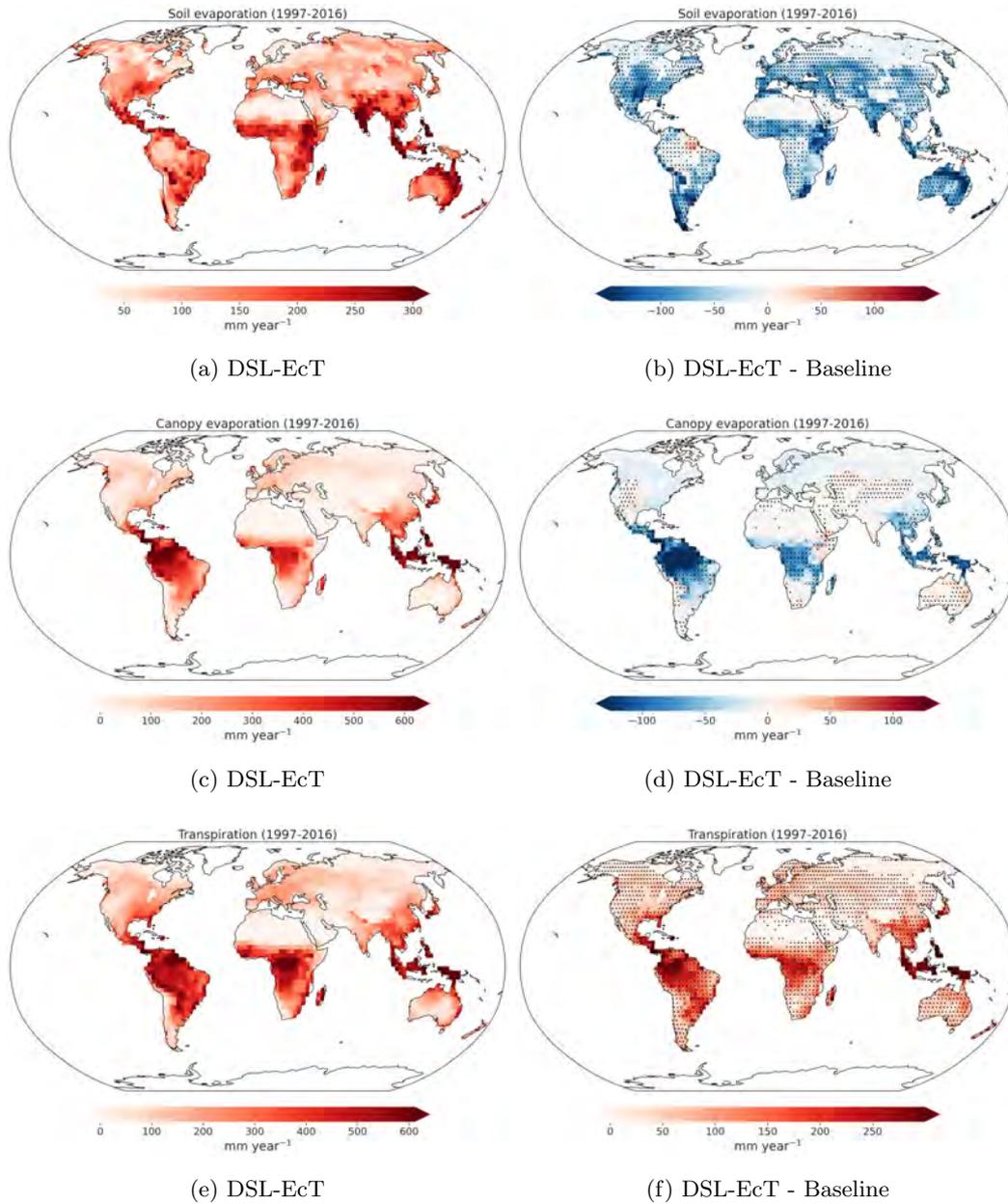


Figure 8: Geographic distribution of soil evaporation ( $E_s$ ), canopy evaporation ( $E_c$ ) and transpiration ( $T$ ) averaged over 1997-2016 for the DSL-EcT simulation (a, c, e) and the difference between the DSL-EcT and Baseline simulations (b, d, f) using the CRUJRA meteorological forcing and the ESACCI land cover. Grid cells with dots indicate that differences are statistically significant (independent two-sample t-test  $p$  level  $< 0.01$ )

410 Parameter values which determine the DSL thickness ( $z_{max}$  and  $K$ ; Equations A12  
 411 and A13) and the interception capacity of the canopy (the maximum storage of liquid  
 412 water;  $p_l$ , Equation A31) are uncertain. To investigate how the chosen parameter val-  
 413 ues impact simulated energy fluxes and GPP, we conducted a sensitivity analysis with  
 414 global simulations. The simulations demonstrated that LE has the opposite response to  
 415  $H$  and GPP as  $K$ ,  $z_{max}$  and  $p_l$  are changed (Figure 9). Changing  $K$  or  $p_l$  affected LE,

416  $H$  and GPP more than changes in  $z_{max}$ . Of the ET components,  $E_s$  is most affected by  
 417 changes in  $K$  and  $z_{max}$ , while changes in  $p_l$  affected  $E_c$  the most. Changes in ET par-  
 418 titioning due to a modified  $p_l$  were relatively small, however. A reduction in  $p_l$  by 50%  
 419 from  $0.2 \text{ kg m}^{-2}$  to  $0.1 \text{ kg m}^{-2}$  reduced  $E_c/ET$  only by 15% and increased  $E_s/ET$  and  
 420  $T/ET$  by 7% and 6%, respectively. Thus, even with a significantly reduced interception  
 421 capacity, CLASSIC simulated an  $E_c$  that still contributed 25% of ET, which is higher  
 422 than GSWP-2 (16%) (Dirmeyer et al., 2006) and CLM4 (20%) (D. M. Lawrence et al.,  
 423 2011), while  $T$  was 44% of ET.

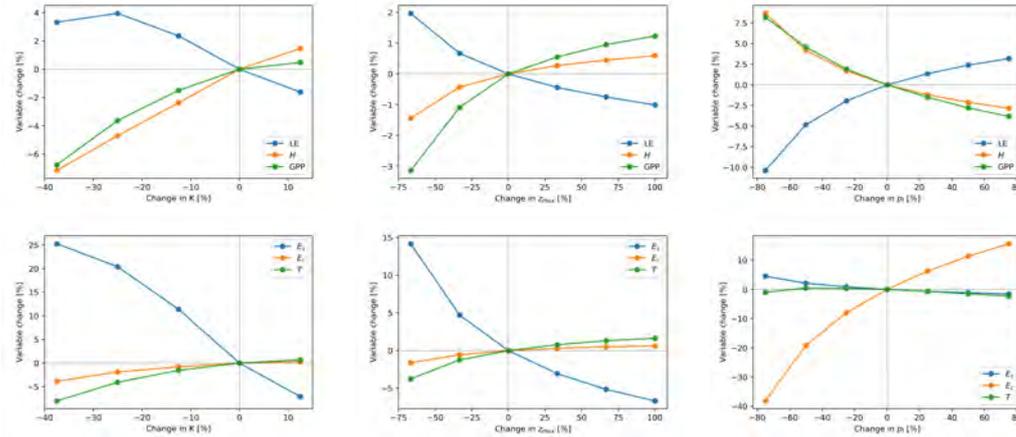


Figure 9: Percentage change in LE,  $H$  and GPP (top) and in  $E_s$ ,  $E_c$  and  $T$  (bottom) with change in the parameters  $K$ , maximum DSL thickness ( $z_{max}$ ) and the maximum storage of liquid water ( $p_l$ ) in the sensitivity simulations performed using CLASSIC with the DSL-EcT configuration.

#### 4 Discussion

424

425 The original CLASSIC version had unreasonably low  $T/ET$  compared to most observation-  
 426 based estimates (see Section 2.4 and Figure 6) and other LSMs with a global mean value  
 427 of  $0.25 \pm 0.01$  (mean  $\pm$  standard deviation of the ensemble of four simulations using two  
 428 meteorological forcings and two land cover representations; Table B1). Similar to results  
 429 of Swenson and Lawrence (2014) using the Community Land Model (CLM), implement-  
 430 ing a DSL parameterization in CLASSIC increased the resistance to  $E_s$ , improved simu-  
 431 lated ET and increased productivity in arid / semi-arid regions (Figures 2 and 7b). In  
 432 areas, where the original  $\beta$  formulation (Equation A20) simulated high  $E_s$ , which reduced  
 433 soil moisture availability later in the growing season, the DSL parameterization gener-  
 434 ally improved simulated LE and increased GPP, as there was more water available in the  
 435 root zone (Figures 2 and B6). This agrees with findings in other studies using different  
 436 models such as Swenson and Lawrence (2014) (CLM4.5) and Decker et al. (2017) (CA-  
 437 BLE). Inclusion of a DSL parameterization alone increased CLASSIC's global  $T/ET$  to  
 438  $0.30 \pm 0.01$  from  $0.25 \pm 0.01$  in baseline CLASSIC. A further modification which influ-  
 439 ences the canopy fluxes,  $E_c$  and  $T$ , allowed  $T$  to occur from the dry portion of the canopy  
 440 while intercepted water is evaporating from the wet canopy fraction. This change increased  
 441 CLASSIC's global  $T/ET$  further to a value of  $0.41 \pm 0.01$ . This value is lower than observa-  
 442 tion-based global estimates of  $0.57 \pm 0.07$  (Wei et al., 2017), but it equals the CMIP5 ensem-  
 443 ble mean (Lian et al., 2018) and is 0.16 higher than the baseline CLASSIC simulations.  
 444 Other models have been working on improving their  $T/ET$  ratio. For example, imple-  
 445 mentation of a bare soil resistance term for dry soils in the ORganizing Carbon and Hy-

446 hydrology in Dynamic EcosystEms (ORCHIDEE) LSM increased simulated  $T/ET$  at semi-  
 447 arid shrub, grass and forest sites in the Southwestern US (MacBean et al., 2020). In the  
 448 Community Atmosphere Biosphere Land Exchange (CABLE) model,  $T/ET$  at several  
 449 FLUXNET sites was increased from an average value of 0.28 (ranging from 0.08 to 0.71  
 450 depending on the site) to 0.70 (ranging from 0.29 to 0.84) by implementing pore-scale-  
 451 based resistance formulations of  $E_s$ , which reduced overestimation of  $E_s$  (Decker et al.,  
 452 2017). In CLM, global  $T/ET$  has been increased from 0.43 in CLM3.5, to 0.48 in CLM4,  
 453 which rises to 0.56 when the nitrogen (N) cycle is explicitly simulated, as ground evap-  
 454 oration decreased from 39% to 32% and 23%, respectively (D. M. Lawrence et al., 2011).  
 455 To accomplish these increases in  $T/ET$  from CLM3.5 to CLM4 new litter resistance and  
 456 canopy turbulence functions were added, which increased the resistance to ground  $E$  when  
 457 litter was present and turbulent exchange was reduced under dense canopies. The dif-  
 458 ferences in  $T/ET$  for the two versions of CLM4 (with and without the N cycle) are likely  
 459 due to a higher LAI with the N cycle explicitly simulated, which increased  $T$  and  $E_c$  and  
 460 reduced  $E_s$  (D. M. Lawrence et al., 2011). Our simulations with the modified DSL-EcT  
 461 CLASSIC, which had higher  $T$  and  $T/ET$  than the baseline CLASSIC, changed LAI re-  
 462 gionally, showing a statistically significant (t-test, p value < 0.01) increase in LAI in arid/semi-  
 463 arid regions and a decrease in some tropical forests. Global mean LAI, however, did not  
 464 appreciably change compared to the baseline CLASSIC and is higher than AVHRR and  
 465 MODIS observations suggest in both simulations (Figure C1g and h). Comparing the  
 466 CLASSIC DSL-EcT against CLM4 (no N cycle) (D. M. Lawrence et al., 2011) show CLAS-  
 467 SIC DSL-EcT to have lower  $T$  (48% CLM4, 41% CLASSIC DSL-EcT), similar  $E_s$  (32%  
 468 CLM4, 30% CLASSIC DSL-EcT) and higher  $E_c$  (20% CLM4, 29% CLASSIC DSL-EcT).  
 469 The Global Soil Wetness Project Phase 2 (GSWP-2) multi-model mean (including thir-  
 470 teen land surface models; see Dirmeyer et al. (2006)) contributions to ET were 48%  $T$ ,  
 471 36%  $E_s$  and 16%  $E_c$  (Dirmeyer et al., 2006). Variability between global estimates of the  
 472 ET components from CLM4, GSWP-2 and other models such as GLEAM, PT-JPL and  
 473 PM-MOD (Miralles et al., 2016) is large and uncertainties are high (see Section 2.4). Com-  
 474 pared against observation-based estimates and other models, however,  $E_c$  remains too  
 475 high in CLASSIC DSL-EcT while  $T$  is too low. As our parameter sensitivity tests (Sec-  
 476 tion 3.2 and Figure 9) showed, the higher  $E_c$  is in part due to a higher maximum stor-  
 477 age of liquid water ( $p_l$ ) compared with CLM4 and a lower  $p_l$  of 0.1 kg m<sup>-2</sup> would reduce  
 478 CLASSIC's  $E_c$  to ~25% of ET from 29%. Measurements of maximum water storage per  
 479 leaf area index show large variability depending on the ecosystem, vegetation species and  
 480 stand age with values ranging from 0.14 to 0.88 mm (Hadiwijaya et al., 2021), which sug-  
 481 gest that a  $p_l$  of 0.1 kg m<sup>-2</sup> could be too low.

482 In order to improve the simulation of the different ET components and especially  
 483  $T/ET$  in LSMs or ESMs, further processes have been highlighted as potentially impor-  
 484 tant in other studies using different models. Chang et al. (2018) found that simulated  
 485  $T/ET$  of a subhumid, mountainous catchment improved when an empirical resistance  
 486 formulation to  $E$  was replaced by a process-based soil surface resistance parameteriza-  
 487 tion, and lateral flow, redistributing precipitation in mountainous terrain, was included  
 488 in a process-based ecohydrological model. Here, we have included a process-based sur-  
 489 face resistance parameterization through the simulation of the DSL, however, terrain-  
 490 driven lateral flow is not included in CLASSIC. Its inclusion could improve  $T/ET$  fur-  
 491 ther, as lateral flow affects soil moisture along hillslopes resulting in drier surfaces on up-  
 492 per slopes suppressing  $E$  more than  $T$  (Chang et al., 2018). Water redistribution in semi-  
 493 arid ecosystems, however, is complex and in addition to lateral flow, local microtopog-  
 494 raphy and biocrusts forming on bare soils can affect runoff and channel water to vege-  
 495 tated patches, where it infiltrates more easily and increases productivity (Chen et al.,  
 496 2013; Rodríguez-Caballero et al., 2018). Another issue observed in LSMs is that root growth  
 497 and distribution and interactions between soil moisture and root dynamics are often not  
 498 adequately represented (Chang et al., 2018; P. Wang et al., 2018). P. Wang et al. (2018)  
 499 showed that a dynamic root scheme combined with the simulation of the ground water  
 500 table implemented in the Noah LSM, where root dynamics depend on fluctuating ground-

501 water levels, improves simulation of root water uptake and latent heat fluxes in arid or  
 502 semi-arid regions. During growing season periods when the water table declines, roots  
 503 extract water from the saturated zone or directly from groundwater. In forests with deep  
 504 roots, for example, ground water dynamics can impact energy, water and carbon fluxes  
 505 as well as simulated soil moisture (De Pue et al., 2022; MacBean et al., 2020; Decharme  
 506 et al., 2019). Including groundwater recharge from an aquifer in Niu et al. (2007) was  
 507 shown to increase soil moisture and ET especially in transition areas from arid to wet  
 508 regions (e.g., riparian zones in arid regions). The simulation of seasonal drought effects  
 509 in LSMs or ESMs was found to be improved by combining the representation of ground-  
 510 water replenishment from an aquifer with lateral flow and dynamic root distributions  
 511 instead of commonly used static, prescribed root profiles (P. Wang et al., 2018). Uncer-  
 512 tainties in pedotransfer functions, which are used to determine soil physical properties,  
 513 also affect the ability of LSMs to adequately represent soil moisture (De Pue et al., 2022).  
 514 Simulated soil moisture and infiltration might be improved by incorporating improved  
 515 pedotransfer functions (Gupta et al., 2021; Pinnington et al., 2021), which depend on  
 516 climatology and land use in addition to soil texture (Fatichi et al., 2020; Vereecken et  
 517 al., 2019). Simulated drought response can also be improved by implementing a plant  
 518 hydraulics scheme, which determines  $g_c$  based on xylem hydraulics instead of using an  
 519 empirical moisture stress function (Eller et al., 2018). Especially under extreme climatic  
 520 conditions or a changing climate, process-based models of  $g_c$  can improve simulated wa-  
 521 ter fluxes during droughts. Eller et al. (2018) showed that their hydraulics-based  $g_c$  model  
 522 was able to better represent effects of drought on  $T$  of tropical forests during El Niño  
 523 events than an empirical drought scheme.

524 Future work, which would likely improve ET partitioning in CLASSIC and sim-  
 525 ulated  $T/ET$ , could include the representation of terrain-dependent lateral flow, plant  
 526 hydraulics and possibly modifications to canopy interception such as inclusion of wind-  
 527 driven loss of intercepted water or snow which increases throughfall (Véliz-Chávez et al.,  
 528 2014). As Dong et al. (2022) attributed a warm bias in the central US in CMIP6 mod-  
 529 els, which CanESM exhibits as well, to low ET and  $T/ET$ , we are also planning to in-  
 530 vestigate the effects of the DSL and  $E_c-T$  partitioning modifications in the ESM CanESM  
 531 to determine their effects on land C and water fluxes as well as the climate, when the  
 532 land and the atmosphere interact.

## 533 5 Conclusions

534 LSMs often show poor ET partitioning with positive biases in  $E$  and negative bi-  
 535 ases in  $T$ , resulting in an underestimation of  $T/ET$  (Chang et al., 2018; Lian et al., 2018).  
 536 These biases impact the simulation of C cycle processes. For example, we found that over-  
 537 estimation of  $E_s$  during periods of high soil moisture in sparsely vegetated areas such  
 538 as low-latitude shrublands resulted in excessive plant water stress during the growing  
 539 season and depressed GPP in CLASSIC simulations. To address CLASSIC’s bias in  $E_s$ ,  
 540 we implemented a dry surface layer (DSL) parameterization that increases the surface  
 541 resistance to water vapour and heat fluxes. To further improve simulated  $T$ ,  $T$  is now  
 542 allowed to occur from the dry fraction of the plant canopy at the same time as water evap-  
 543 orates from the wet fraction, which previously did not allow  $T$  when a canopy was even a  
 544 small fraction wet. After these modifications, in arid and semi-arid regions  $E_s$  and ET  
 545 were reduced during wet periods leading to improved seasonality of ET and an increase  
 546 in GPP. However, the impact of our modifications globally was for GPP to decrease slightly  
 547 ( $\sim 1.6\%$ ) compared to the baseline CLASSIC simulations as a result of increased  $T$  and  
 548 ET and drier soils in other biomes including seasonally dry tropical forests. Globally, the  
 549 proportion of  $T$  relative to ET was improved compared to observations with an increase  
 550 from  $\sim 25\%$  in baseline CLASSIC to  $\sim 41\%$  in the DSL-EcT simulations. As the simu-  
 551 lated global  $T/ET$  of 0.41 remains lower than observation-based estimates of  $0.57 \pm 0.07$   
 552 (Wei et al., 2017), possible future improvements to CLASSIC include implementing terrain-

553 driven lateral flow redistributing water, and including a plant hydraulics-based  $g_c$  scheme  
 554 instead of an empirical moisture stress function to improve the representation of plant  
 555 water use and the vegetation's response to drought stress. Improvements in ET parti-  
 556 tioning in LSMs and ESMs are important to simulate carbon and water fluxes well in  
 557 historical and especially future simulations, as warmer climates are expected to enhance  
 558 water cycles and impact ESM climate simulations.

## 559 Appendix A

### 560 A1 Evapotranspiration parameterization

ET is the sum of  $E_s$ ,  $E_c$  and  $T$ .  $E_s$  consists of  $E$  originating from bare soil and from soil underneath the vegetation canopy. The potential evaporation rate from bare soil,  $E(0)$  ( $\text{mm s}^{-1}$ ), is calculated as

$$E(0) = \rho_a C_{DH} v_a (q(0) - q_a), \quad (\text{A1})$$

where  $\rho_a$  is the air density ( $\text{kg m}^{-3}$ ),  $C_{DH}$  the stability-dependent surface drag coefficient for heat (unitless),  $v_a$  the wind speed at the reference height ( $\text{m s}^{-1}$ ),  $q(0)$  the specific humidity at the surface ( $\text{kg kg}^{-1}$ ) and  $q_a$  the specific humidity at the reference height ( $\text{kg kg}^{-1}$ ) (Verseghy, 2017). The saturated surface specific humidity,  $q(0)_{sat}$  ( $\text{kg kg}^{-1}$ ),  $q_a$ , and the surface evaporation efficiency ( $\beta$ ; unitless; Equation A20) are used to determine  $q(0)$  as

$$q(0) = \beta q(0)_{sat} + (1 - \beta) q_a. \quad (\text{A2})$$

The surface evaporation rate is limited to a maximum value,  $E(0)_{max}$  ( $\text{mm s}^{-1}$ ) determined by the water content of the top soil layer ( $\theta_1$ ;  $\text{m}^3 \text{m}^{-3}$ ) and the depth of water ponded on the surface ( $Z_p$ ,  $\text{m}$ ) as

$$E(0)_{max} = \rho_w [Z_p + (\theta_1 - \theta_{min}) \Delta Z_1] / \Delta t, \quad (\text{A3})$$

with the density of water  $\rho_w$  ( $\text{kg m}^{-3}$ ), the depth of the top soil layer  $\Delta Z_1$  (e.g. 0.10  $\text{m}$ ) and the time interval  $\Delta t$  ( $\text{s}$ ) (typically 900-1800  $\text{s}$  for CLASSIC) (Verseghy, 2017).  $\theta_{min}$  ( $\text{m}^3 \text{m}^{-3}$ ) is the residual soil liquid water content remaining after freezing or evaporation. This is set to 0.04  $\text{m}^3 \text{m}^{-3}$  for mineral and fibric organic soils and 0.15 and 0.22  $\text{m}^3 \text{m}^{-3}$  for hemic and sapric organic soils, respectively. Underneath the vegetation, the maximum surface evaporation rate,  $E(0)_{max,c}$  ( $\text{mm s}^{-1}$ ), is determined as

$$E(0)_{max,c} = \rho_w (\theta_1 - \theta_{min}) \Delta Z_1 / \Delta t. \quad (\text{A4})$$

The potential evaporation rate from soil under the vegetation,  $E(0)_c$  ( $\text{mm s}^{-1}$ ), is calculated as

$$E(0)_c = \frac{\rho_a}{r_{a,g}} (q(0) - q_{a,c}), \quad (\text{A5})$$

where  $q_{a,c}$  is the specific humidity of the canopy air ( $\text{kg kg}^{-1}$ ) and  $r_{a,g}$  ( $\text{s m}^{-1}$ ) is the surface resistance, whose inverse is derived from Deardorff (1972) as

$$\frac{1}{r_{a,g}} = 1.9 \times 10^{-3} (T(0)_v - T_{ac,v})^{1/3}, \quad (\text{A6})$$

with the virtual potential temperature at the surface ( $T(0)_v$ ;  $\text{K}$ ) and of the canopy air ( $T_{ac,v}$ ;  $\text{K}$ ) and the constant  $1.9 \times 10^{-3}$  in  $\text{m s}^{-1} \text{K}^{-1/3}$ . The evapotranspiration rate from the vegetation ( $\text{ET}_c$ ;  $\text{mm s}^{-1}$ ), i.e., the sum of  $E_c$  and  $T$ , which is equivalent to the latent heat flux from the vegetation canopy divided by the latent heat of vaporization, is calculated as

$$\text{ET}_c = \rho_a \frac{q_c - q_{a,c}}{r_b + r_c}, \quad (\text{A7})$$

where  $q_c$  is the saturated specific humidity at the canopy temperature ( $\text{kg kg}^{-1}$ ),  $r_b$  the leaf boundary layer resistance ( $\text{s m}^{-1}$ ) and  $r_c$  the stomatal resistance ( $\text{s m}^{-1}$ ). The relative contributions from  $E$  or  $T$  differ depending on the circumstances in the model. If

there is snow on the canopy,  $ET_c$  is limited to the intercepted snow amount and is in the form of  $E_c$  through sublimation. If instead, the canopy has liquid water upon it, the calculated  $ET_c$  is first drawn from the amount of liquid water stored on the canopy ( $W_l$ ;  $\text{kg m}^{-2}$ ). If that amount is insufficient to satisfy the calculated  $ET_c$ ,  $T$  is possible after checking there is enough soil water available in the root zone.  $E_c$  is then set to  $W_l$  and the remainder of  $ET_c$  is allocated to  $T$ . Thus,  $T$  only occurs, when there is no water available on the canopy and enough soil water is available, i.e., the liquid water content ( $\theta_l$ ;  $\text{m}^3 \text{m}^{-3}$ ) exceeds  $\theta_{min}$  for the respective soil layer. Based on Bonan (1996); McNaughton and Van Den Hurk (1995), the inverse of  $r_b$  is calculated as

$$1/r_b = v_{ac}^{1/2} \sigma f_i \gamma_i \text{PAI}^{1/2} / 0.75 [1 - \exp(-0.75 \text{PAI}^{1/2})] \quad (\text{A8})$$

with the wind speed in the canopy air space  $v_{ac}$ , the fractional coverage of each PFT  $f_i$ , the PFT-dependent parameter describing leaf dimension  $\gamma_i$  (unitless), and the plant area index (PAI). By default, CLASSIC uses Leuning (1995)'s stomatal conductance ( $g_c$ ;  $\text{mol CO}_2 \text{m}^{-2} \text{s}^{-1}$ ) formulation (details in Arora (2003); Melton and Arora (2016)) and  $g_c$  is calculated as

$$g_c = m \frac{G_{canopy,net} p}{(c_s - \Gamma)} \frac{1}{(1 + \text{VPD}/V_o)} + b \text{LAI}, \quad (\text{A9})$$

where  $G_{canopy,net}$  is the net canopy photosynthesis rate ( $\text{mol CO}_2 \text{m}^{-2} \text{s}^{-1}$ ),  $p$  is the surface atmospheric pressure (Pa) and  $\Gamma$  is the  $\text{CO}_2$  compensation point (Pa). The parameter  $m$  (unitless) is 9.0 for needle-leaf trees, 12.0 for other  $C_3$  plants and 6.0 for  $C_4$  plants,  $b$  is set to  $0.01 \text{mol m}^{-2} \text{s}^{-1}$  for  $C_3$  and  $0.04 \text{mol m}^{-2} \text{s}^{-1}$  for  $C_4$  plants. The parameter  $V_o$  has values of 2000 Pa for trees and shrubs and 1500 Pa for crops and grasses. The partial pressure of  $\text{CO}_2$  at the leaf surface,  $c_s$  (Pa), depends on the atmospheric  $\text{CO}_2$  partial pressure  $c_{ap}$  (Pa),  $G_{canopy,net}$  and the aerodynamic conductance  $g_b$  ( $\text{mol CO}_2 \text{m}^{-2} \text{s}^{-1}$ ) and is defined as

$$c_s = c_{ap} - \frac{1.37 G_{canopy,net} p}{g_b}. \quad (\text{A10})$$

The units of  $g_c$  and  $g_b$  can be converted from  $\text{mol CO}_2 \text{m}^{-2} \text{s}^{-1}$  to  $\text{m s}^{-1}$  using

$$g_c(\text{m s}^{-1}) = 0.0224 \frac{T_c p_0}{T_f p} g_c(\text{mol CO}_2 \text{m}^{-2} \text{s}^{-1}), \quad (\text{A11})$$

561 with the standard atmospheric pressure  $p_0 = 101\,325$  Pa and the freezing temperature  
562  $T_f = 273.16$  K.

## 563 **A2 Determination of the dry surface layer thickness**

To avoid numerical instabilities due to thin soil layers, CLASSIC uses a 10 cm thick top soil layer. In reality, soil moisture can vary strongly within the top 10 cm of soil, especially during extended dry periods where a thin layer at the top of the soil surface gets very dry while the soil below stays moist (Goss & Madliger, 2007; Kurc & Small, 2004; Li et al., 2020). To approximate the effects of this thin dry layer on surface water and energy fluxes, a DSL parameterization is implemented in CLASSIC following Swenson and Lawrence (2014). Their DSL parameterization determines when a DSL is present, its thickness, and the resulting surface resistance to evaporation. The formation of a DSL is initiated when the soil moisture of the top soil layer falls below a defined moisture threshold,  $\theta_{DSL0}$  ( $\text{m}^3 \text{m}^{-3}$ ), which is determined as

$$\theta_{DSL0} = K \theta_{p,1}, \quad (\text{A12})$$

where  $\theta_{p,1}$  is the porosity of the top soil layer ( $\text{m}^3 \text{m}^{-3}$ ) and  $K$  is a constant (unitless), here with a value of 0.8 following Swenson and Lawrence (2014). The thickness of the DSL (m) is calculated as

$$DSL = \begin{cases} z_{max} \frac{\theta_{DSL0} - (\theta_{l,1} + \theta_{ice,1})}{\theta_{DSL0} - \theta_{air}} & \text{for } \theta_{l,1} + \theta_{ice,1} < \theta_{DSL0} \\ 0 & \text{for } \theta_{l,1} + \theta_{ice,1} \geq \theta_{DSL0}, \end{cases} \quad (\text{A13})$$

where  $z_{max}$  is the maximum DSL thickness (m), here set to 0.015 m. As the liquid ( $\theta_l$ ) and frozen ( $\theta_{ice}$ ) water contents of the bare ground and ground under canopy subareas can differ in CLASSIC, the DSL thickness and the resistance to evaporation are calculated separately for these two subareas. The ‘‘air-dry’’ soil moisture value ( $\theta_{air}$ ) was determined following Dingman (2002) as

$$\theta_{air} = \theta_{p,1} \left( \frac{\Psi_{sat,1}}{\Psi_{air}} \right)^{\frac{1}{b}} \quad (\text{A14})$$

with the saturated soil matric potential  $\Psi_{sat}$  (m), the air-dry matric potential  $\Psi_{air} = 10^4$  m (Swenson & Lawrence, 2014) and the Clapp and Hornberger empirical soil water characteristic ‘‘b’’ parameter (unitless). The soil resistance to evaporation from bare ground or the ground under the canopy  $R_{soil}$  ( $\text{s m}^{-1}$ ) is determined as

$$R_{soil} = \frac{DSL}{\tau D_v}, \quad (\text{A15})$$

where  $D_v$  ( $\text{m}^2 \text{s}^{-1}$ ) is the molecular diffusivity of water vapour in the air and calculated as (D. M. Lawrence et al., 2020)

$$D_v = 2.12 \times 10^{-5} \left( \frac{T_1}{273.15} \right)^{1.75}, \quad (\text{A16})$$

where  $T_1$  is the temperature of the top soil layer (K).  $\tau$  ( $\text{m}^3 \text{m}^{-3}$ ) in Equation A15 is the tortuosity of the vapour flow paths through the soil and determined following Moldrup et al. (2003) as

$$\tau = \Phi_{air}^2 \left( \frac{\Phi_{air}}{\theta_{p,1}} \right)^{3/b} \quad (\text{A17})$$

with the air-filled pore space  $\Phi_{air}$  ( $\text{m}^3 \text{m}^{-3}$ ) calculated as

$$\Phi_{air} = \theta_{p,1} - \theta_{air}. \quad (\text{A18})$$

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### A3 DSL effect on surface evaporation

An increasing thickness of the DSL acts to decrease surface evaporation and thus the latent heat flux in CLASSIC through a decrease in the surface evaporation efficiency ( $\beta$ ; unitless).  $\beta$  has a value between 0 and 1, where a value of 1 means that the specific humidity at the surface equals the saturated surface specific humidity and does not limit  $E$ , i.e. a DSL thickness of 0, whereas a  $\beta$  value of 0 means no surface evaporation can occur.  $\beta$  is calculated as the minimum, more limiting value, between the soil evaporation efficiency ( $R_{soil}$ ; Equation A15) derived from the DSL thickness and that calculated by using CLASSIC’s original soil evaporation efficiency (Meyer et al., 2021; Merlin et al., 2011) (CEVAP), which limits  $\beta$  values below 1 except when soils are fully saturated when the value can be 1.

$$\beta = \min \left( \frac{1}{C_{DH} v_a R_{soil} + 1}, \text{CEVAP} \right). \quad (\text{A19})$$

CEVAP is defined as

$$\text{CEVAP} = \begin{cases} 0 & \text{for } \theta_{l,1} < \theta_{min} \\ 0.25(1 - \cos(\pi\theta_{l,1}/\theta_{p,1}))^2 & \text{for } \theta_{min} < \theta_{l,1} \leq \theta_{p,1}. \end{cases} \quad (\text{A20})$$

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When there is snow or ponded water on the surface,  $\beta$  is set to 1 and  $q(0)$  is set to  $q(0)_{sat}$ . Equation A19 gives a  $\beta$  that is constrained to CEVAP, when the soil is too moist for a DSL to develop. Figure 1 shows an example of the soil evaporation efficiency determined using the original CEVAP parameterization as well as the calculation using the resistance due to the DSL and the minimum of the two parameterizations for a range of liquid water content values of the top soil layer.

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#### A4 DSL effect on the thermal conductivity

In CLASSIC, the soil thermal conductivity ( $\lambda_{soil}$ ;  $\text{W m}^{-1} \text{K}^{-1}$ ) is determined from the saturated thermal conductivity ( $\lambda_{sat}$ ;  $\text{W m}^{-1} \text{K}^{-1}$ ), the dry thermal conductivity ( $\lambda_{dry}$ ;  $\text{W m}^{-1} \text{K}^{-1}$ ) and a relative thermal conductivity ( $\lambda_r$ ; unitless; ranging from 0 for dry soils to 1 for saturated soils) following Côté and Konrad (2005) (Verseghy, 2017)

$$\lambda_{soil} = \lambda_r(\lambda_{sat} - \lambda_{dry}) + \lambda_{dry}. \quad (\text{A21})$$

An empirical coefficient  $\kappa$  (unitless), which differs for frozen ( $\kappa = 1.2, 0.85,$  and  $0.25$  for coarse mineral, fine mineral, and organic soils, respectively) and unfrozen ( $\kappa = 4.0, 1.9,$  and  $0.6$  for coarse mineral, fine mineral, and organic soils, respectively) soils, and the degree of saturation ( $S_r$ ; unitless) determine  $\lambda_r$

$$\lambda_r = \frac{\kappa S_r}{1 + S_r(\kappa - 1)}, \quad (\text{A22})$$

where

$$S_r = \frac{\theta_{l+ice}}{\theta_p}. \quad (\text{A23})$$

$\lambda_{dry}$  depends on  $\theta_p$  and is calculated as

$$\lambda_{dry} = \begin{cases} 0.75 \exp(-2.76 \theta_p) & \text{for mineral soil} \\ 0.30 \exp(-2.00 \theta_p) & \text{for organic soil.} \end{cases} \quad (\text{A24})$$

The thermal conductivities of liquid water ( $\lambda_l = 0.57 \text{ W m}^{-1} \text{K}^{-1}$ ), ice ( $\lambda_{ice} = 2.24 \text{ W m}^{-1} \text{K}^{-1}$ ) and the soil particles ( $\lambda_s$ ;  $\text{W m}^{-1} \text{K}^{-1}$ ; values are  $2.5 \text{ W m}^{-1} \text{K}^{-1}$  for sand and clay and  $0.25 \text{ W m}^{-1} \text{K}^{-1}$  for organic matter) determine  $\lambda_{sat}$  following De Vries (1963) as

$$\lambda_{sat} = \begin{cases} \lambda_l \theta_p + \lambda_s(1 - \theta_p) & \text{for unfrozen soil} \\ \lambda_{ice} \theta_p + \lambda_s(1 - \theta_p) & \text{for frozen soil.} \end{cases} \quad (\text{A25})$$

Similar to the latent heat flux (Section A3), the sensible heat flux should be limited by the DSL because the thermal properties, i.e., the thermal conductivity and heat capacity, which is influenced by changes in soil moisture, of the DSL differ from those of the top soil layer, as the DSL is drier and has more air filled-pore space. When a DSL is present, for mineral soils and organic soils in uplands, the thermal conductivity at the top of the first soil layer ( $\lambda$ ;  $\text{W m}^{-1} \text{K}^{-1}$ ) is linearly interpolated between the “dry” ( $\lambda_{dry}$ ) and calculated top soil layer thermal conductivity ( $\lambda_{soil}$ ) values depending on the DSL thickness

$$\lambda = \lambda_{soil} - \frac{DSL}{z_{max}}(\lambda_{soil} - \lambda_{dry}). \quad (\text{A26})$$

572

#### A5 DSL effect on the ground albedo

In CLASSIC, the visible and near-infrared ground albedos ( $\alpha_g$ ; unitless) are soil moisture dependent. As the top of the soil wets from a liquid water content value of 0.22 to  $0.26 \text{ m}^3 \text{ m}^{-3}$ , the albedo values follow a linear relationship between the “dry” ( $\alpha_{g,dry}$ ; unitless) and “wet” albedo values ( $\alpha_{g,wet}$ ; unitless) of the respective soil colour index (P. J. Lawrence & Chase, 2007). Outside of this range of liquid water content, the model adopts either the dry or wet albedo value accordingly. With the DSL formulation, if a DSL exists, a DSL-dependent  $\alpha_g$  is calculated as

$$\alpha_g = \alpha_{g,wet} - \frac{DSL}{z_{max}}(\alpha_{g,wet} - \alpha_{g,dry}) \quad (\text{A27})$$

573

and the  $\alpha_g$  value used by the model is set to the higher value of the original CLASSIC calculation and the value determined in Equation A27.

574

## A6 Modifications to canopy evaporation and transpiration in CLASSIC

In the CLASSIC v.1.2 formulation for snow-covered canopies the evaporative flux from the canopy ( $ET_c$ ; Equation A7) is first assigned to sublimation, as liquid water, if present, is assumed to be within or underneath the snow and no  $T$  is expected to occur. When there is only liquid water present on the canopy, we modified the original formulation so that  $T$  is allowed to occur from a partially-wet canopy instead of from a completely dry canopy only. To separate the calculated amount of evapotranspired water ( $ET_c$ ; see Equation A7) into  $E_c$  and  $T$ , we determine the wet ( $f_{wet}$ ) and dry ( $f_{dry}$ ) fractions of the canopy similar to Fan et al. (2019) as

$$f_{wet} = \begin{cases} F_l & \text{for } F_l \geq 0.01 \text{ and } F_l \leq 0.99 \\ 0 & \text{for } F_l < 0.01 \\ 1 & \text{for } F_l > 0.99 \end{cases} \quad (\text{A28})$$

$$f_{dry} = \begin{cases} (1 - f_{wet}) \frac{LAI}{PAI} & \text{for } f_{wet} \geq 0.01 \text{ and } f_{wet} \leq 0.99 \\ 1 & \text{for } f_{wet} < 0.01 \\ 0 & \text{for } f_{wet} > 0.99 \end{cases} \quad (\text{A29})$$

In general, only the leaves and not stems of a canopy can transpire,  $f_{dry}$  is adjusted by the LAI to PAI ratio.  $F_l$  is the fractional coverage of the canopy covered by liquid water (unitless) determined as

$$F_l = \begin{cases} \min(W_l / W_{l,max}, 1) & \text{for } W_{l,max} > 0 \\ 0 & \text{for } W_{l,max} = 0 \end{cases} \quad (\text{A30})$$

where  $W_l$  ( $\text{kg m}^{-2}$ ) is the amount of liquid water stored on the canopy and  $W_{l,max}$  ( $\text{kg m}^{-2}$ ) is the storage capacity of the canopy for liquid water, which is calculated as

$$W_{l,max} = p_l \times PAI \quad (\text{A31})$$

with the maximum storage of liquid water  $p_l$  set as  $0.20 \text{ kg m}^{-2}$  (Bartlett et al., 2006).  $W_l$  is calculated as the sum of  $W_l$  of the previous time step and the rainfall intercepted by the canopy during the current time step

$$W_{l,t} = \min(W_{l,t-1} + \Delta t \rho_w (P - \chi P), W_{l,max}), \quad (\text{A32})$$

where  $P$  is the rainfall rate ( $\text{m s}^{-1}$ ),  $\chi$  is the canopy gap fraction (unitless),  $\Delta t$  is the model physics timestep (s) and  $\rho_w$  ( $\text{kg m}^{-3}$ ) the density of liquid water. To determine the canopy fractional coverage of liquid water exposed to the air,  $F_l$  is decreased by the fractional snow coverage ( $F_s$ ).

$$F_l = \max(0, \min(F_l - F_s, 1)) \quad (\text{A33})$$

and, similar to  $F_l$ ,  $F_s$  is found by

$$F_s = \begin{cases} \min(W_f / W_{f,max}, 1) & \text{for } W_{f,max} > 0 \\ 0 & \text{for } W_{f,max} = 0, \end{cases} \quad (\text{A34})$$

576 where  $W_f$  ( $\text{kg m}^{-2}$ ) is the amount of frozen water stored on the canopy and  $W_{f,max}$  ( $\text{kg}$   
577  $\text{m}^{-2}$ ) is the storage capacity of the canopy for frozen water. If there is no plant available  
578 water in the root zone, the wet canopy fraction is set to 1, as  $T$  is not allowed to occur.

The predicted mass of water evapotranspired from the canopy ( $W_E$ ;  $\text{kg m}^{-2}$ ) is calculated as

$$W_E = ET_c \times \rho_w \Delta t, \quad (\text{A35})$$

where  $ET_c$  is the evapotranspiration rate from the canopy ( $\text{m s}^{-1}$ ; see Equation A7). The wet and dry canopy fractions as well as  $F_{RbRc}$  determine the fractions of  $W_E$  coming from

$E_c$  and  $T$ , respectively. The amount of  $W_l$  (see Equation A32) is adjusted by the amount of water evaporated from the wet canopy fraction as

$$W_l = W_l - (1 - f_{dry}) (1 - F_{RbRc}) W_E \text{ for } W_E (1 - f_{dry})(1 - F_{RbRc}) \leq W_l \quad (\text{A36})$$

and  $W_E$  is reduced by the amount being evaporated

$$W_E = \begin{cases} W_E (F_{RbRc} + f_{dry} - f_{dry} F_{RbRc}) & \text{for } W_E (1 - f_{dry}) (1 - F_{RbRc}) \leq W_l \\ W_E - W_l & \text{for } W_E (1 - f_{dry}) (1 - F_{RbRc}) > W_l \end{cases} \quad (\text{A37})$$

The contribution of the leaf boundary layer resistance ( $r_b$ ;  $\text{s m}^{-1}$ ; Equation A8) to the total resistance, the sum of  $r_b$  and the stomatal resistance ( $r_c$  or  $1/g_c$ ;  $\text{s m}^{-1}$ ; Equation A9), is calculated as a proportion of the total resistance from the leaf boundary layer and stomata to determine when canopy evaporation is dominant and when  $T$  can occur, as

$$F_{RbRc} = r_b / (r_b + r_c). \quad (\text{A38})$$

579 In the second case of Equation A37, where  $W_l$  could not meet the calculated amount of  
 580 water to be evaporated,  $W_l$  is then set to zero. If the predicted mass of water evapotran-  
 581 spirated from the vegetation ( $W_E$ ) after considering evaporation from wet leaves is greater  
 582 than zero, it is treated as  $T$ . If there is enough water available in the root zone and  $T$   
 583 can occur, the soil water content removed by  $T$  and the  $T$  flux are calculated for each  
 584 soil layer and the liquid water content of each soil layer containing roots is updated.

## 585 Appendix B

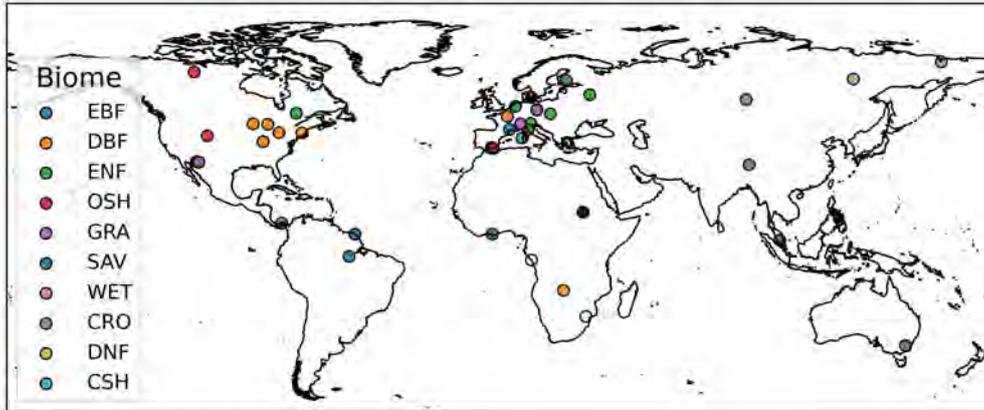


Figure B1: Map of the FLUXNET sites used in this study (including their biomes).

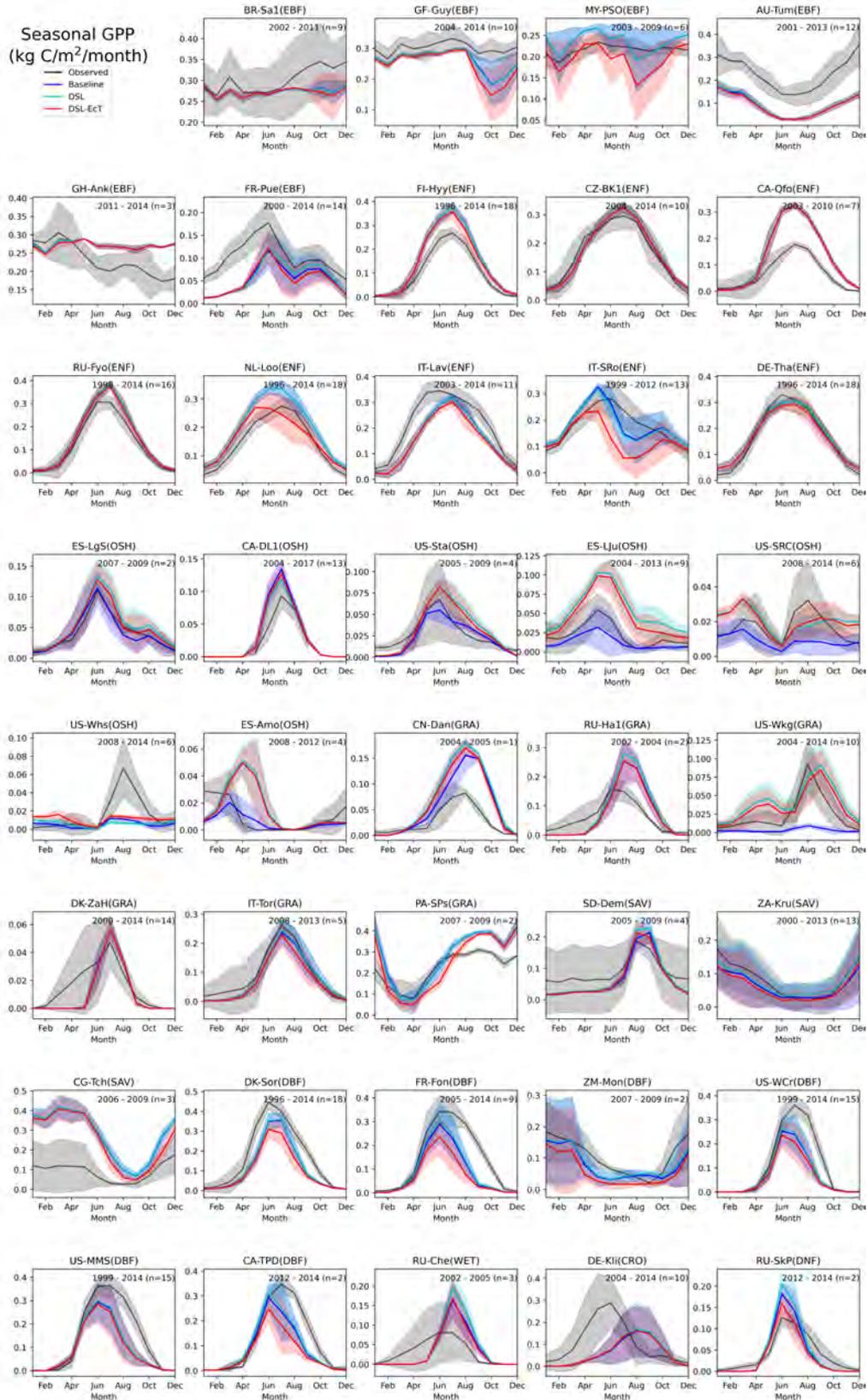


Figure B2: Monthly mean observed and simulated gross primary productivity (GPP) for the FLUXNET sites for the Baseline, DSL and DSL-EcT simulations (Table 1). The shading shows the standard deviation over the available years. Site names, their biomes and years of measurements used are listed for each site in Table 2.

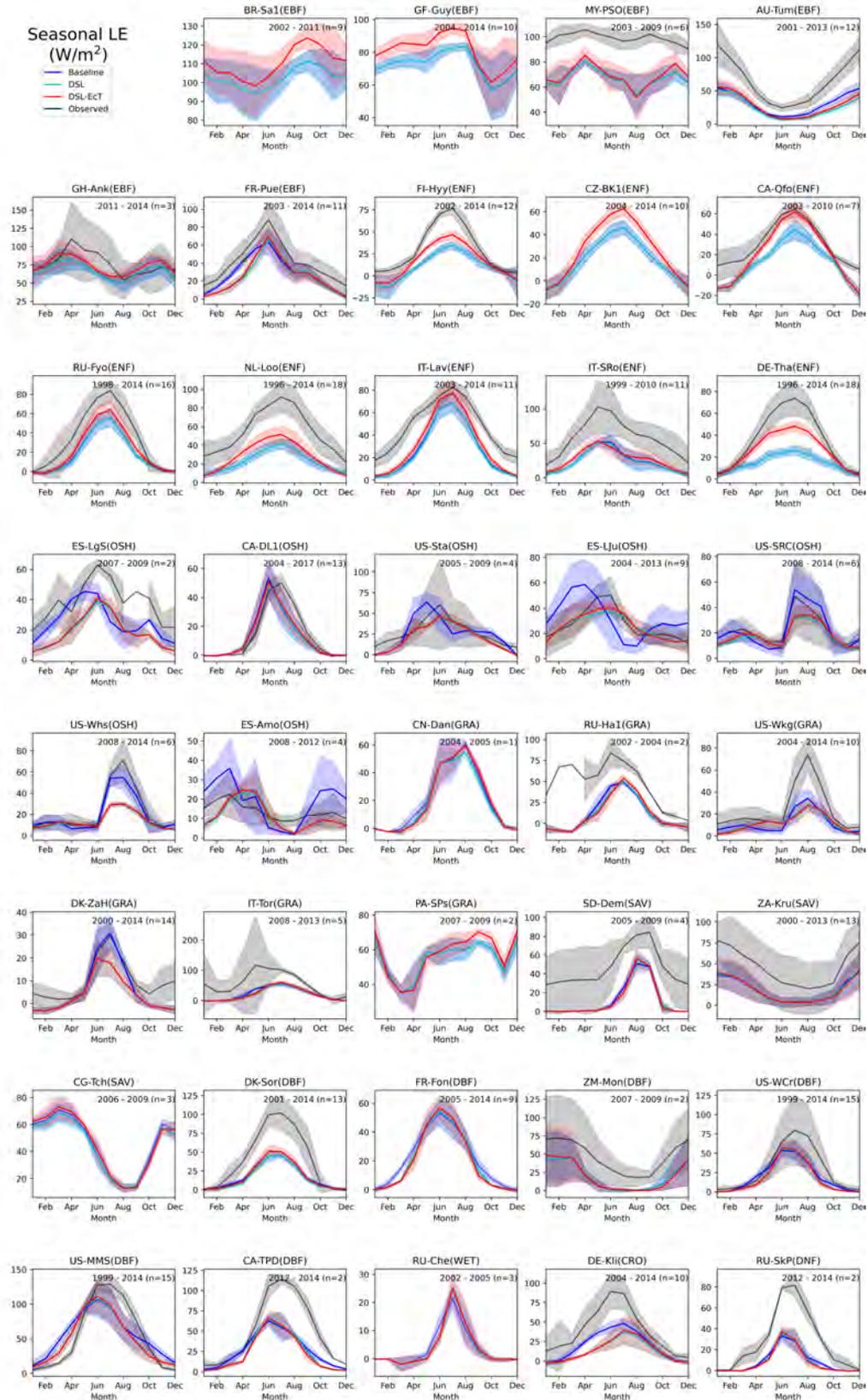


Figure B3: Monthly mean observed and simulated latent heat flux (LE) for the FLUXNET sites for the Baseline, DSL and DSL-EcT simulations (Table 1).

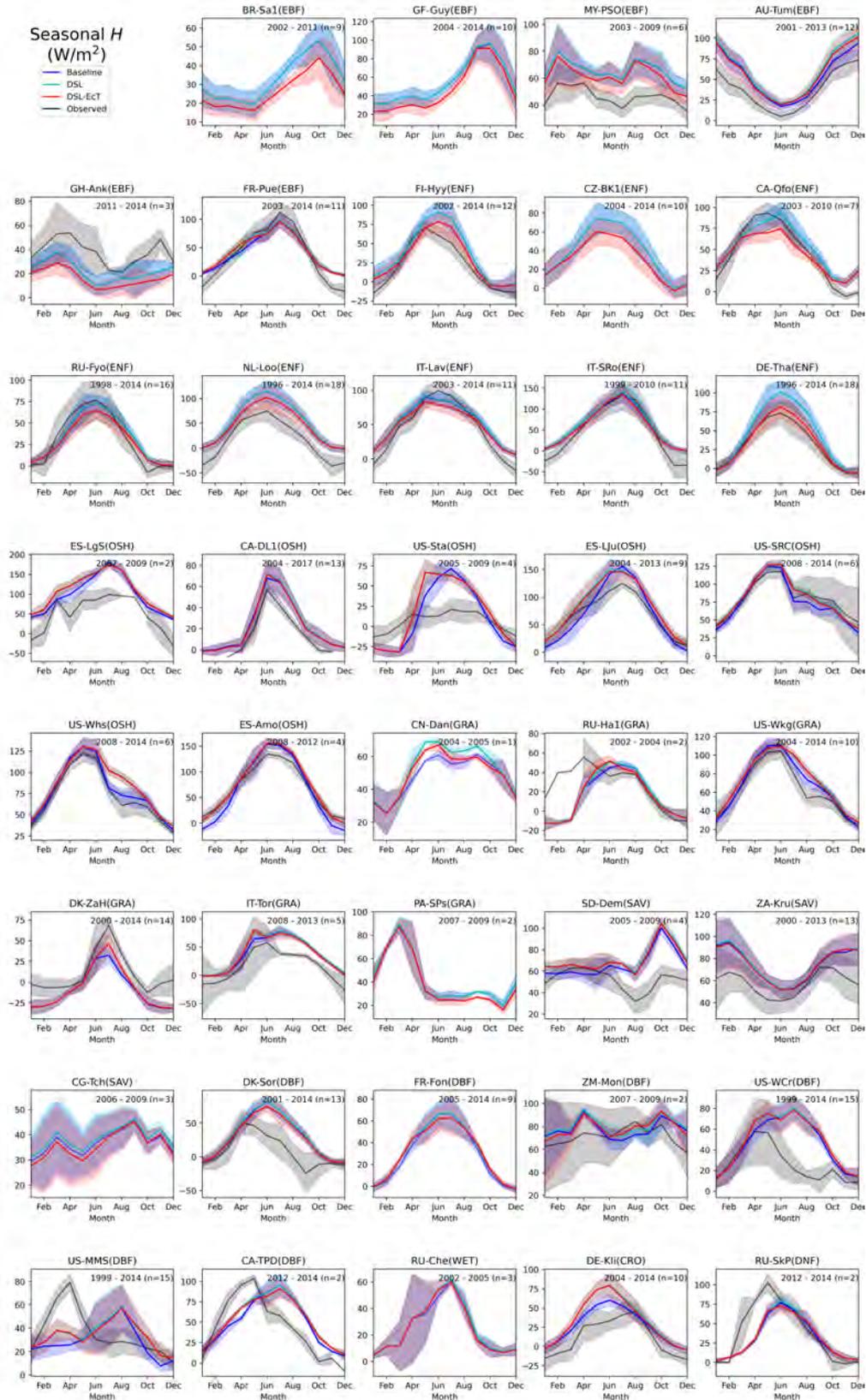


Figure B4: Monthly mean observed and simulated sensible heat flux ( $H$ ) for the FLUXNET sites for the Baseline, DSL and DSL-EcT simulations (Table 1).

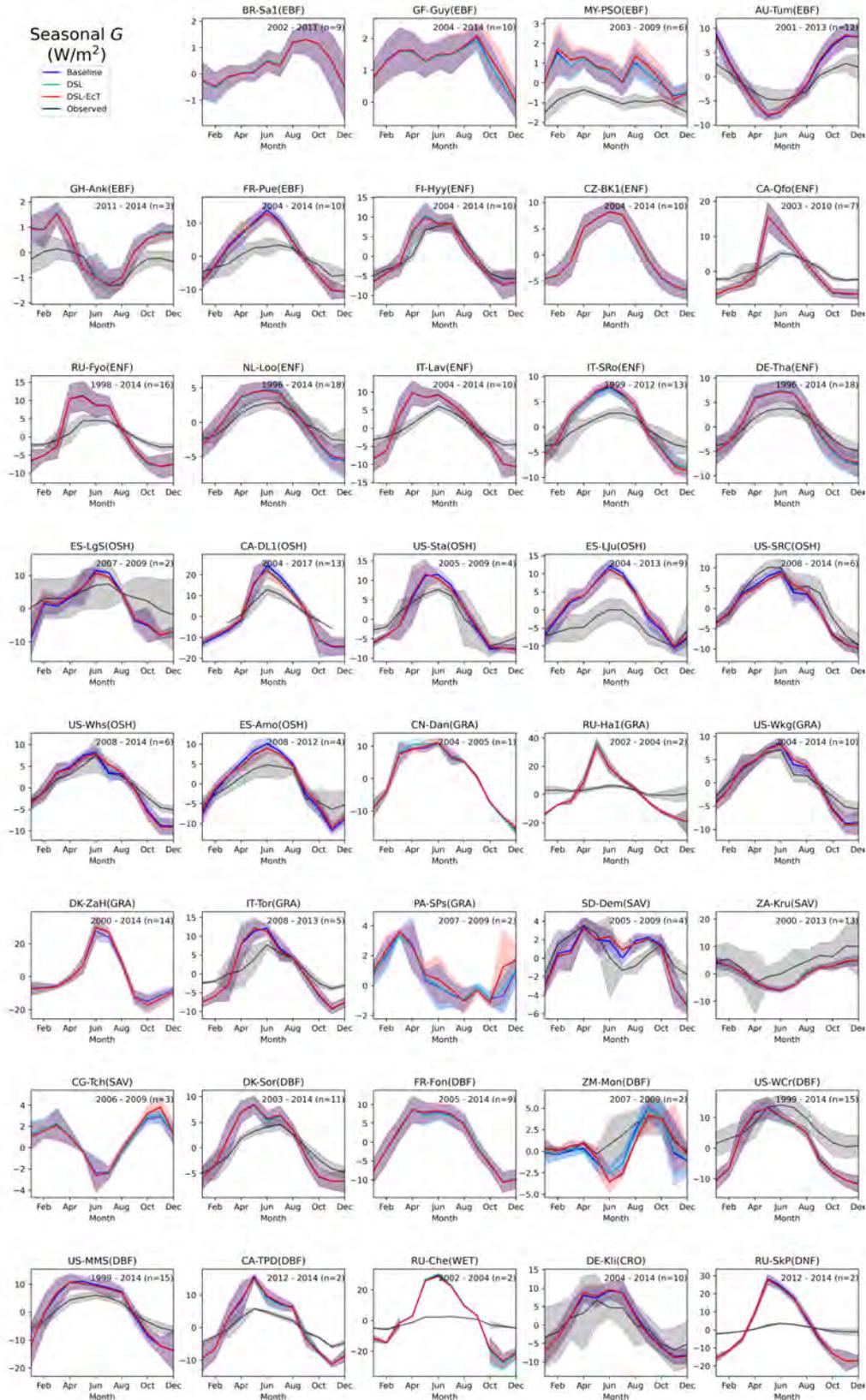


Figure B5: Monthly mean observed and simulated ground heat flux ( $G$ ) for the FLUXNET sites for the Baseline, DSL and DSL-EcT simulations (Table 1).

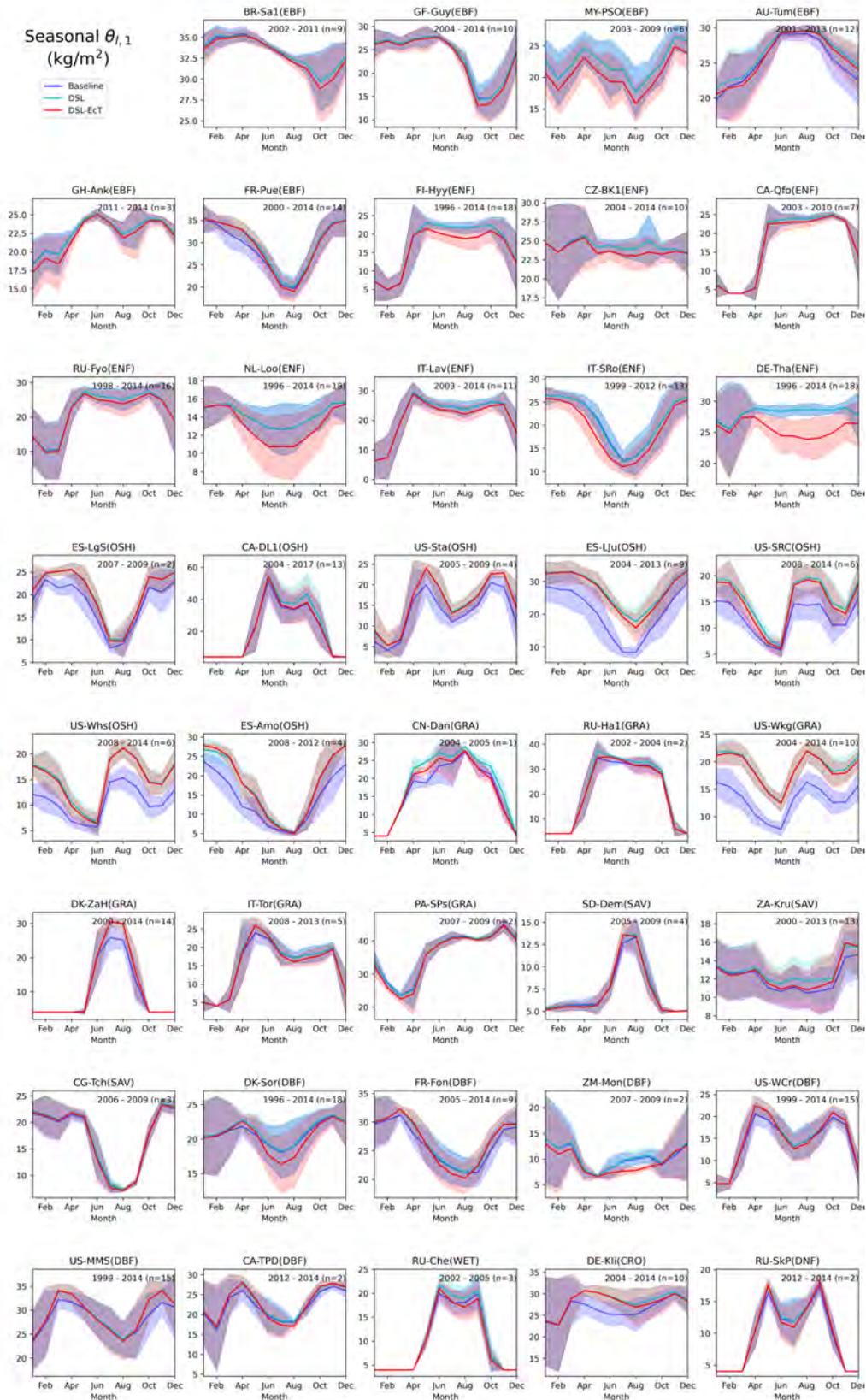


Figure B6: Monthly mean simulated liquid water content of the top soil layer (0-10 cm depth;  $\theta_{l,1}$ ) for the FLUXNET sites for the Baseline, DSL and DSL-EcT simulations (Table 1).

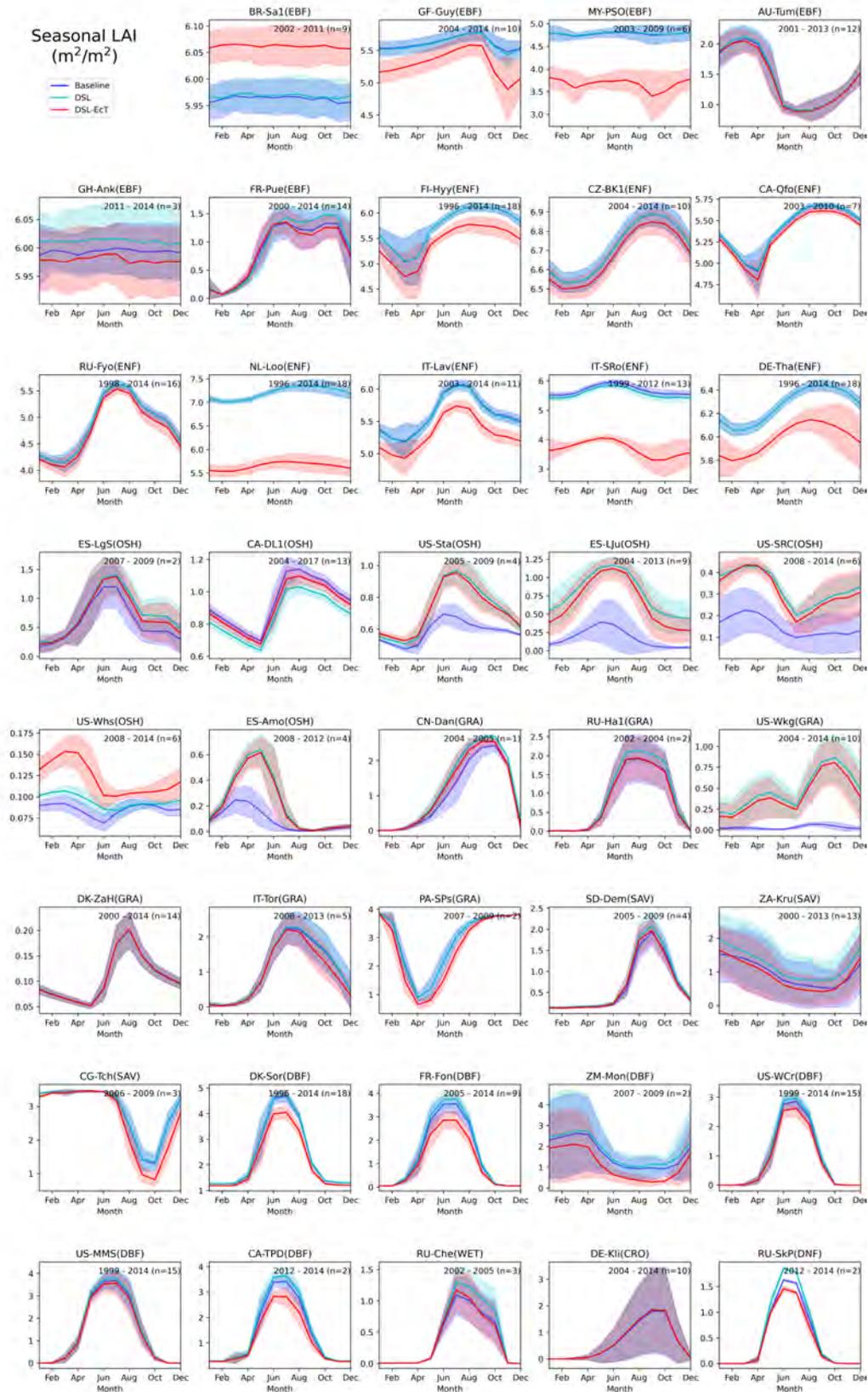


Figure B7: Monthly mean simulated leaf area index (LAI) for the FLUXNET sites for the Baseline, DSL and DSL-EcT simulations (Table 1).

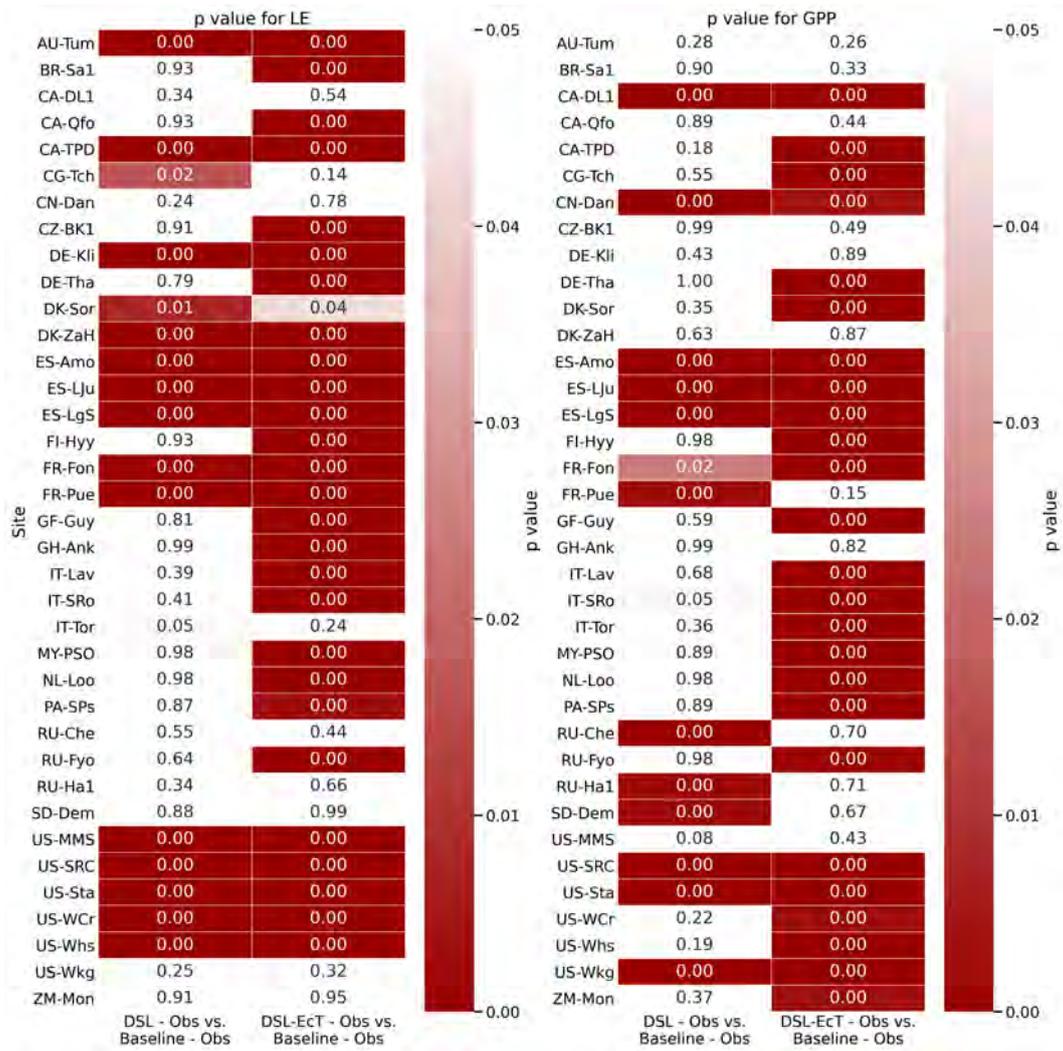


Figure B8: Two-sided t-test p value between the error in simulated daily gross primary productivity (GPP) and latent heat flux (LE) for the FLUXNET sites (Table 2) for DSL and DSL-EcT simulations compared to the Baseline simulation (Table 1).

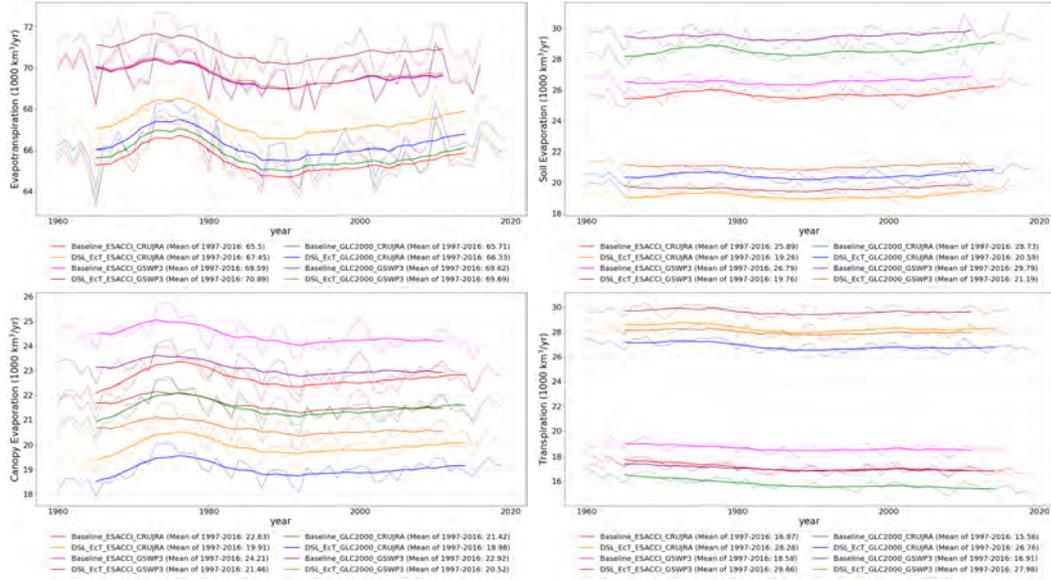


Figure B9: Annual ET, soil  $E$ , canopy  $E$  and  $T$  for 1960-2016 for the four Baseline and DSL-EcT simulations (Table 1), respectively.

Table B1: Transpiration ( $T$ ) to evapotranspiration (ET) ratios of the different CLASSIC simulations (Table 1) averaged over 1997-2016.

Forcing data	Baseline	DSL	DSL-EcT
ESACCI and CRUJRA	25.92%	30.33%	41.93%
ESACCI and GSWP3	26.72%	31.43%	41.85%
GLC2000 and CRUJRA	23.69%	28.62%	40.35%
GLC2000 and GSWP3	24.30%	29.48%	40.17%

## Appendix C

Zonal ensemble plots of Baseline and DSL-EcT simulations show slight differences in GPP with the DSL-EcT simulations having slightly lower GPP in the Tropics and slightly higher GPP in the higher latitudes than the Baseline simulations. Globally, GPP is lower in the DSL-EcT simulations and shows less variability between the four simulations (using two different meteorological forcings and two different land cover representations) (Figure C1). In the Tropics, LE tends to be higher in the DSL-EcT simulations.

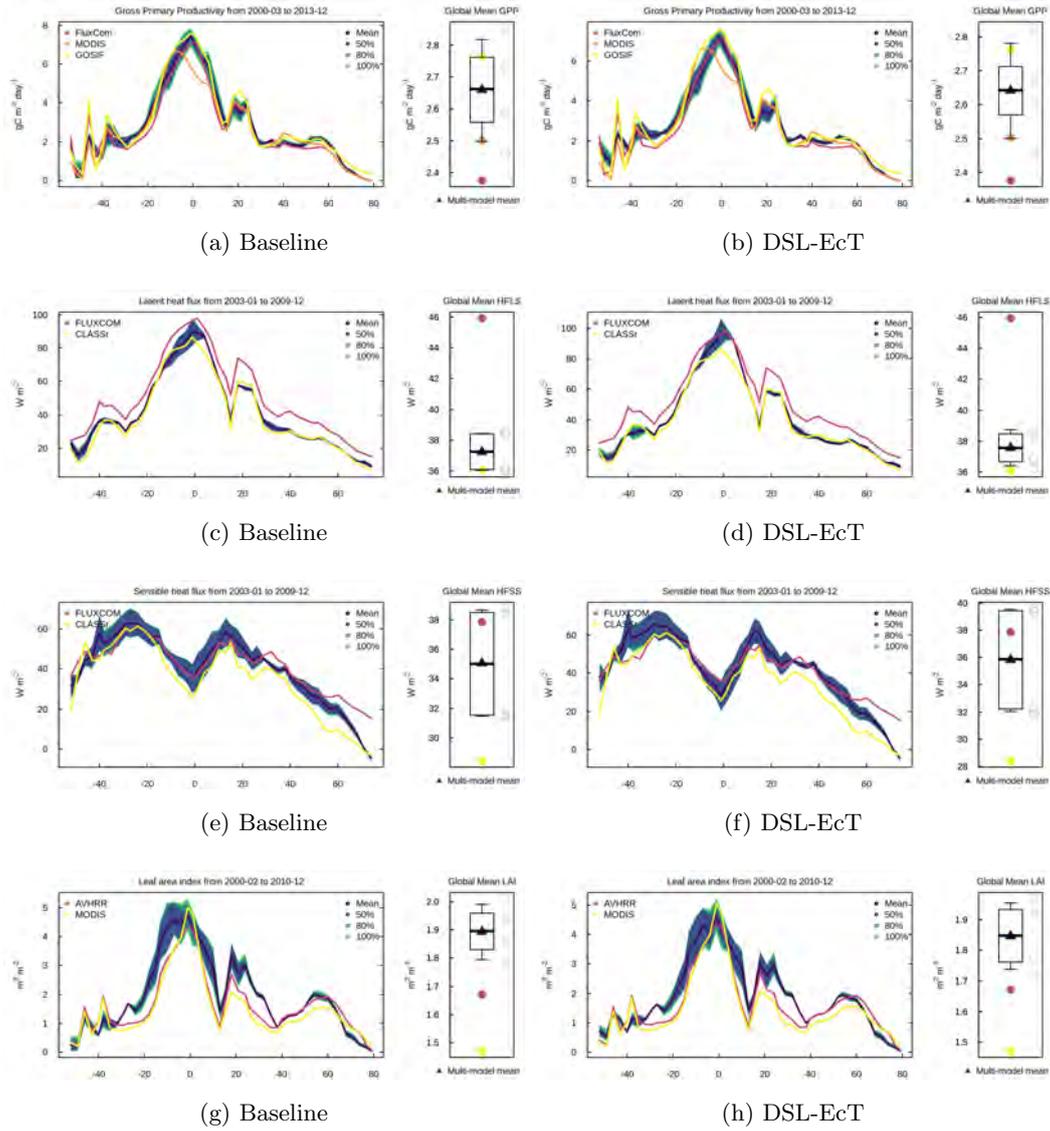


Figure C1: Zonally averaged and global mean GPP, LE,  $H$  and LAI over land for the Baseline and DSL-EcT simulations. The ensemble includes the four simulations using a combination of two different meteorological forcing datasets and two land cover representations (Table 1).

593 Seasonal averages for the 11 TRANSCOM regions show that the Baseline and DSL  
 594 simulations have the greatest differences in the North American Boreal, the South Amer-  
 595 ican Tropics, Eurasian Temperate and Australia. During the spring, the DSL simu-  
 596 lations tend to overestimate GPP in Australia (Figure C2), while they perform well dur-  
 597 ing the rest of the year.

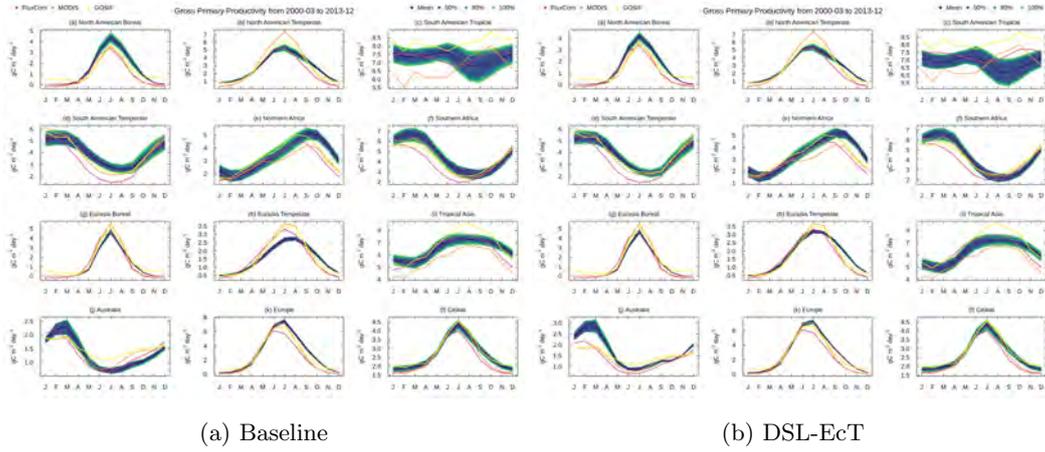


Figure C2: Monthly averaged GPP for 11 TRANSKOM regions and globally from March 2000 to December 2013.

598 In LE and  $H$  (Figure C3 and C4), the Baseline and DSL simulations show differ-  
 599 ences especially in the South American Tropical, Northern Africa and Tropical Asia. How-  
 600 ever, in all of the TRANSKOM regions as well as globally both simulations tend to lie  
 601 within the uncertainty bounds of observations.

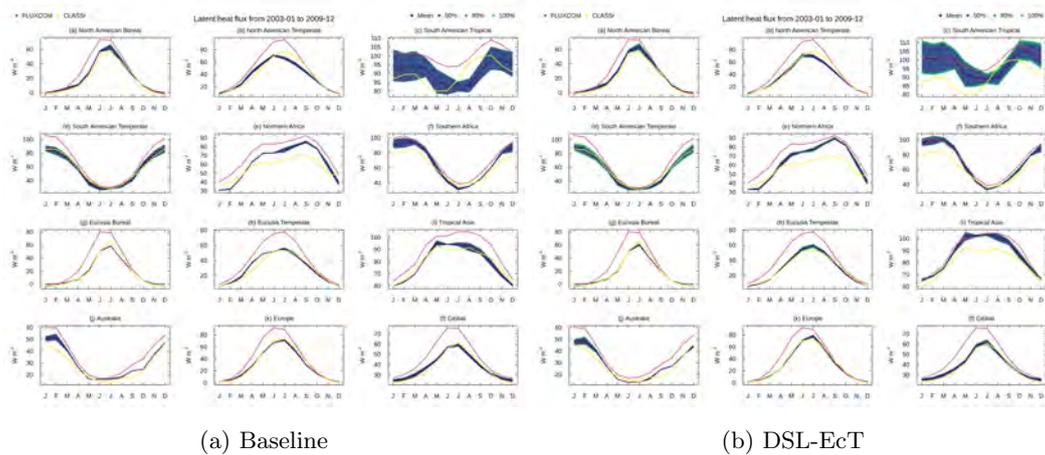


Figure C3: Monthly averaged LE for 11 TRANSKOM regions and globally from January 2003 to December 2009.

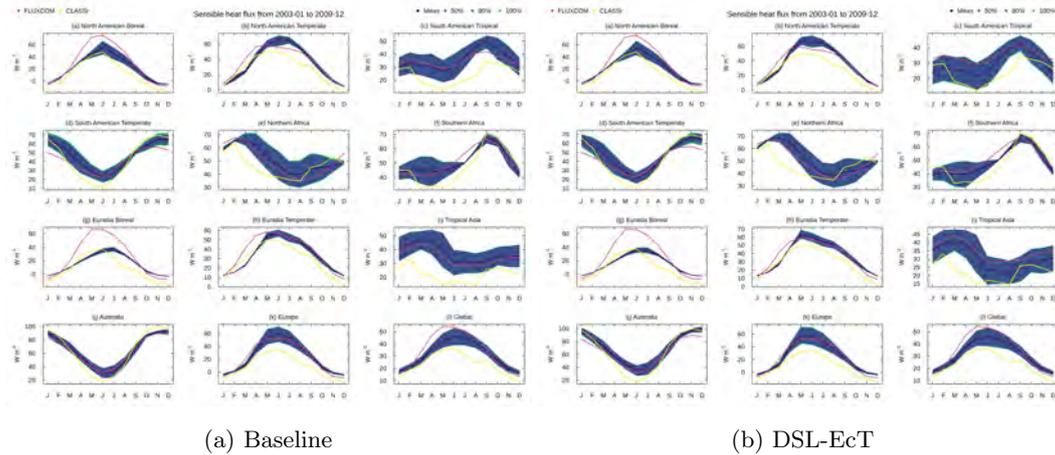


Figure C4: Monthly averaged  $H$  for 11 TRANSKOM regions and globally from January 2003 to December 2009.

## Appendix D Open Research

### D1 Data Availability Statement

The CLASSIC code versions (Baseline, DSL and DSL-EcT) and model outputs presented in our paper are archived on Zenodo (<https://doi.org/10.5281/zenodo.7015764>; Meyer et al. (2022)).

### Acknowledgments

This work used eddy covariance data acquired and shared by the FLUXNET community, including these networks: AmeriFlux, AfriFlux, AsiaFlux, CarboAfrica, CarboEuropeIP, CarboItaly, CarboMont, ChinaFlux, Fluxnet-Canada, GreenGrass, ICOS, KoFlux, LBA, NECC, OzFlux-TERN, TCOS-Siberia, and USCCC. The FLUXNET eddy covariance data processing and harmonization were carried out by the European Fluxes Database Cluster, AmeriFlux Management Project, and Fluxdata project of FLUXNET with the support of CDIAC and ICOS Ecosystem Thematic Center, and the OzFlux, ChinaFlux, and AsiaFlux offices. We thank Christian Seiler for providing comments on a pre-submission version of the manuscript.

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