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1 **Impact of evapotranspiration on dry season climate in the**
2 **Amazon forest**

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ABSTRACT

7
8 Moisture recycling can be an important source of rainfall over the Amazon forest, but this
9 process relies heavily upon the ability of plants to access soil moisture. Evapotranspiration
10 (ET) in the Amazon is often maintained or even enhanced during the dry season, when net
11 radiation is high. However, ecosystem models often over predict the dry season water stress.
12 We removed unrealistic water stress in an ecosystem model (the Simple Biosphere model,
13 SiB3), and examined impacts of enhanced ET on the dry season climate when coupled to
14 a GCM. The "Stressed" model experiences dry season water stress and limitations on ET,
15 while the "Unstressed" model has enhanced root water access and exhibits strong drought
16 tolerance.

17 During the dry season in the southeastern Amazon, SiB3 Unstressed has significantly
18 higher latent heat flux (LH) and lower sensible heat flux (SH) than SiB3 Stressed. There
19 are two competing impacts on the climate in SiB3 Unstressed: cooling due to lower SH,
20 and moistening due to higher LH. During the average dry season, the cooling plays a larger
21 role and the atmosphere is more statically stable, resulting in less precipitation than in
22 SiB3 Stressed. During dry season droughts, significantly higher LH in SiB3 Unstressed is a
23 necessary but not sufficient condition for stronger precipitation. The moistening effect of LH
24 dominates when the Bowen ratio ($BR=SH/LH$) is >1.0 in SiB3 Stressed, and precipitation
25 is up to 26% higher in SiB3 Unstressed. An implication of this analysis is that forest
26 conservation could enable the Amazon to cope with drying conditions in the future.

27 1. Introduction

28 The Amazon forest stores huge amounts of carbon in its biomass (Saatchi et al. 2007,
29 2011), but its future is uncertain due to combined threats of climate change and deforestation
30 (Nepstad et al. 2008; Malhi et al. 2008). Recent Amazonian droughts have led to decreases
31 in biomass (Phillips et al. 2009; Lewis et al. 2011; Toomey et al. 2011) and increases in tree
32 mortality (Phillips et al. 2010), and several GCMs predict reduced dry season precipitation
33 throughout the 21st century (Malhi et al. 2008). Amazon droughts are linked to variability
34 in the tropical Atlantic and Pacific sea surface temperatures (Liebmann and Marengo 2001;
35 Marengo 2004; Chen et al. 2011). An anomalously warm tropical north Atlantic displaces the
36 Intertropical Convergence Zone (ITCZ) northward (Marengo et al. 2011), which weakens the
37 trade winds, reduces water vapor transport, and increases subsidence above the central and
38 southern Amazon (Espinoza et al. 2011). In effect, these meteorological changes lengthen
39 the dry season, and data from the Global Precipitation Climatology Centre suggest that
40 dry seasons have become longer since the 1990's (Marengo et al. 2011). El Niño events are
41 associated with drought in the northeastern Amazon (Ropelewski and Halpert 1987; Chen
42 et al. 2011) and large-scale subsidence due to a shift in the Walker Circulation (Malhi and
43 Wright 2004). Two recent severe droughts (in 2005 and 2010) attracted attention due to their
44 severity and ecological impacts. Both droughts were linked to an anomalously warm tropical
45 north Atlantic (Marengo et al. 2008b; Espinoza et al. 2011), a pattern that is predicted to
46 continue or perhaps increase (Cox et al. 2008). In 2005, drought coincided with the dry
47 season and resulted in significant biomass reductions (Aragao et al. 2008; Zeng et al. 2008;
48 Phillips et al. 2009) related to both heat and moisture stress (Toomey et al. 2011). The
49 severe drought in 2010 affected a larger area, negatively impacted biomass (Lewis et al.
50 2011), and resulted in widespread declines in vegetation greenness (Xu et al. 2011). This
51 drought was preceded by an El Niño, which limited wet season precipitation and contributed
52 to the severity of the drought (Marengo et al. 2011).

53 During a drought, plants may close their stomata to limit water loss (Fisher et al. 2006),

54 which can lead to mortality as the plants cease to assimilate carbon. Whether plant stomata
55 remain open or closed during a drought also impacts the latent heat flux (LH) between
56 the land and atmosphere. On average, moisture recycling due to evaporation from the
57 land surface contributes between one-quarter and one-third of the precipitation over the
58 Amazon, although this number varies in space and time (Eltahir and Bras 1994; Trenberth
59 1999). Evapotranspiration (ET) serves as a significant source of precipitation (Spracklen
60 et al. 2012), and loss of forest cover has been linked to drying in the southern Amazon
61 over the past 30 years (Lee et al. 2011). Moisture recycling can impact dry season climate
62 by affecting the timing of the wet season onset (Fu and Li 2004; Li and Fu 2004). Prior
63 to the transition from dry to wet season, surface LH increases the atmosphere’s convective
64 available potential energy (CAPE) and decreases convective inhibition energy (CINE). These
65 processes increase rainfall and initiate the transition period (Fu and Li 2004). When the
66 land surface is anomalously dry, LH is lower and sensible heat (SH) is higher than average.
67 CINE remains high and the wet season onset is delayed (Fu and Li 2004).

68 If stomatal conductance is severely limited during a drought, reduced ET could reduce
69 moisture recycling and reinforce drought conditions. This is a positive feedback on drought,
70 analogous to the delayed wet season onset described by Fu and Li (2004). Conversely, plants
71 can maintain or even increase transpiration during the dry season (Nepstad et al. 1994;
72 Oliveira et al. 2005; Lee et al. 2005; Hasler and Avissar 2007; da Rocha et al. 2009; Costa
73 et al. 2010). Due to the presence of deep roots and ample soil moisture, ET in the equatorial
74 Amazon is tightly coupled to net radiation, which is higher during the dry season (Hasler
75 and Avissar 2007). Moving south, the seasonality of precipitation increases, as does the
76 potential for water stress during the dry season. As a result, ET can be higher in the dry
77 season (Costa et al. 2010; Vourlitis et al. 2011) or the wet season (Costa et al. 2010; da Rocha
78 et al. 2009; Lathuilliere et al. 2012), depending on several factors such as vegetation type,
79 dry season intensity, and depth to the water table. For example, in the southern Amazon,
80 surface resistance can be twice as high during the dry season compared to the wet season

81 (Costa et al. 2010), contributing to lower dry season ET.

82 It is unknown how long into a drought the trees are able to maintain pre-drought pho-
83 tosynthesis and transpiration rates, but the immediate impacts of the 2005 drought possi-
84 bly included enhanced vegetation greenness (which implies increased transpiration) (Saleska
85 et al. 2007), although a subsequent study asserted that forest "green-up" did not occur
86 (Samanta et al. 2010). Satellite-based microwave retrievals based on improved algorithms
87 during the 2005 drought suggest a 3-month lag in forest response to water deficits in the
88 western Amazon, although the response was concurrent with the greatest water deficits in
89 the northeastern Amazon (Saatchi et al. 2013). If plants continue to transpire during a
90 drought, this could reduce the severity of the drought through moisture recycling. Evidence
91 of this phenomenon has been observed during the wet season onset, such that when the
92 land surface is wet, the enhanced LH can enable an earlier transition (although large-scale
93 circulation can counteract this) (Fu and Li 2004).

94 Modeling studies of Amazon climate are essential for preparing for possible future climate
95 and land cover scenarios, and it is equally important to accurately capture the impacts
96 of ET on dry season rainfall. Models that do not allow plants to access adequate soil
97 moisture during the dry season will overestimate the dry season Bowen ratio (SH/LH), and
98 are more likely to induce the positive feedback cycle above. Recent developments in the
99 Simple Biosphere model (SiB3) focused on accurately representing soil water stress in the
100 Amazon. Previously, SiB predicted water limitations on photosynthesis and transpiration
101 during the dry season. When coupled to a GCM (the BUGS model at Colorado State
102 University), reduced ET severely limited precipitation above the Amazon (Randall et al.
103 1996). Changes to SiB3's soil and roots enabled the trees to transpire through the dry
104 season, increased LH, reduced SH, and impacted local climate (Baker et al. 2008; Harper
105 et al. 2010). SiB3 can realistically simulate seasonal cycles of LH, SH, and net ecosystem
106 exchange at a handful of sites in the Amazon (Tapajos K83 in Baker et al. (2008); Manaus,
107 Tapajos K67, K83, Reserva Jaru, and Pe de Gigante in Baker et al. (2013); and Tapajos

108 K67, K83, and Caxiuana in Harper et al. (in prep.)).

109 The aim of the present study is to examine the impacts of increased ET on the dry season
110 climate in the Amazon, using SiB3 coupled to the BUGS5 GCM. The standard version of
111 SiB3 represents a strongly drought resistant forest, as it includes processes documented as
112 important for soil moisture access in the tropical forests of South America (Baker et al. 2008,
113 2013; Harper et al. in prep.). We use a second version of SiB3 that does not include these
114 adaptations and produces unrealistic dry season water stress. The standard version is called
115 SiB3 Unstressed (or SiB3U for short), and the latter model is SiB3 Stressed (or SiB3S). We
116 hypothesize that SiB3 Stressed will produce the positive feedback addressed above (reduced
117 dry season ET reinforcing dry conditions and further reducing precipitation). The methods
118 of the study are outlined in Section 2, and the overall performance of the BUGS5 model is
119 discussed in Section 3, with special attention on South American climate. In Section 4, we
120 assess the impacts on dry season climate of two extreme representations of forest drought
121 resistance.

122 2. Methods

123 a. *SiB3*

124 SiB3 simulates biophysical processes and ecosystem metabolism (Sellers et al. 1986; Den-
125 ning et al. 1996; Sellers et al. 1996b,a; Baker et al. 2008). Carbon assimilation accounts
126 for enzyme kinetics (Farquhar et al. 1980) and is linked to stomatal conductance (Collatz
127 et al. 1991, 1992). The model simulates the turbulent exchange of CO₂, moisture, heat,
128 and momentum between the free atmosphere and a prognostic canopy air space (Vidale and
129 Stockli 2005). The surface hydrology scheme consists of water intercepted by the canopy,
130 the ground and a ten-layer soil model. Vertical movement of soil moisture is governed by
131 Darcy’s law, and the model has 10 soil layers which become thicker with depth. Runoff can
132 occur due to sub-surface drainage out of the lowest layer, or due to excess overland flow when

133 incoming rainfall cannot infiltrate the top layer. SiB3's evapotranspiration (ET) is the sum
134 of canopy transpiration and evaporation from puddles, the top soil layer, and the canopy.
135 The modifications to SiB3 Stressed alter the stomatal conductance, therefore differences in
136 the canopy transpiration dominate the model differences in ET. In the coupling with the
137 GCM, ET is converted to latent heat flux from the canopy air space to the mixed layer
138 (which is the lowest GCM level).

139 Leaf area index (LAI) and fraction of photosynthetically active radiation (fPAR) are
140 calculated from the Normalized Difference Vegetation Index (NDVI) from the Advanced
141 Very High Resolution Radiometer 4-km global area coverage data (Tucker et al. 2005). Each
142 grid cell is assigned one biome type for the entire simulation period (Sellers et al. 1996a),
143 therefore there is no land use change in the experiments. Grid cells in the Amazon forest are
144 designated as tropical broadleaf evergreen forest, and NDVI is held constant at the maximum
145 value during the measurement period. The use of a constant NDVI avoids known errors in
146 the remotely sensed vegetation index due to cloud and aerosol contamination (Los et al.
147 2000; Hilker et al. 2012; Samanta et al. 2012). The parameter fPAR is a strong determinant
148 of model potential photosynthesis and transpiration rates, and it saturates above an LAI
149 of $4 \text{ m}^2 \text{ m}^{-2}$. Therefore, the constant NDVI introduces only minor errors in regions with
150 high LAI, as is the case in much of the Amazon basin (Myneni et al. 2007; Malhado et al.
151 2009; Miller et al. 2004). However, semideciduous forests are common in the transition zone
152 between the evergreen tropical forests and savannas, and in these regions LAI can display
153 strong seasonality. For example, LAI varies from $2\text{-}2.5 \text{ m}^2 \text{ m}^{-2}$ during the dry season to
154 $5\text{-}6 \text{ m}^2 \text{ m}^{-2}$ during the wet season at a site northeast of Sinop, Mato Grosso ($11^\circ 24.75'S$,
155 $55^\circ 19.50'W$) (Vourlitis et al. 2011). In these regions, SiB3 will likely overestimate dry season
156 ET due to the constant LAI and fPAR.

157 SiB3 constrains the net ecosystem exchange (NEE) of CO_2 to be roughly zero each
158 year, since the model does not include dynamic vegetation or biomass storage and cannot
159 accumulate or lose carbon. NEE is not exactly zero because the respiration is based on the

160 previous year’s assimilation. Soil texture is based on maps from the International Geosphere-
161 Biosphere Programme (IGBP: Global Soil Data Task Group, 2000).

162 Potential photosynthesis in SiB3 is linearly weighted by three stress factors to give the
163 actual photosynthetic rate. The three factors range from 0.1 (maximum stress) to 1 (no
164 stress), and parameterize the impacts of less than optimal temperature, humidity, and soil
165 moisture on the gross carbon assimilation (Sellers et al. 1992, 1996b). The modifications in
166 SiB3 Stressed relate to the soil moisture stress, which at strong water deficits can induce
167 stomatal closure and reduce transpiration and photosynthesis. There are three differences
168 between SiB3 Stressed and Unstressed (Table 1). First, the soil is 3.5 meters deep in SiB3S
169 and 10 meters deep in SiB3U. Root depths vary by biome and density decreases exponentially
170 with depth (Jackson et al. 1996). Roots extend through the entire soil column in both
171 versions in the tropical broadleaf evergreen biome. Second, roots in SiB3U are able to access
172 soil moisture wherever it is in the soil column, regardless of root biomass (Baker et al. 2008).
173 This emphasizes the role of deep roots in efficiently accessing soil moisture. In SiB3S, root
174 water extraction is weighted by biomass, which emphasizes the shallow soil layers over the
175 deep layers. Third, the dependence of soil moisture stress on the volumetric water content
176 is revised, such that SiB3U experiences less stress at moderate soil moisture reductions. For
177 further details of these changes see Baker et al. (2008).

178 *b. BUGS5*

179 BUGS5 has evolved from the UCLA GCM to include a geodesic grid and modified sigma
180 coordinate (Suarez et al. 1983; Randall et al. 1985; Ringler et al. 2000) (<http://kiwi.atmos.colorado.edu/BUGS/BUGSoverview.html>). The planetary boundary layer (PBL) depth changes
181 due to horizontal mass flux divergence, entrainment, and convective mass flux. The en-
182 trainment rate is predicted by integrating the turbulent kinetic energy (TKE) conservation
183 equation over the depth of the PBL (Denning et al. 2008). BUGS5 uses a modified Arakawa-
184 Schubert cumulus parameterization with a prognostic cumulus kinetic energy (Ding and
185

186 Randall 1998; Pan and Randall 1998), which relaxes the quasi-equilibrium closure of the
187 models original Arakawa-Schubert parameterization. The stratiform parameterization in-
188 cludes prognostic variables for cloud water, cloud ice, rain, snow, and water vapor (Fowler
189 et al. 1996), and is directly coupled to the cumulus parameterization. The microphysical
190 parameterization follows Fowler and Randall (2002). The radiation scheme is adopted from
191 NCARs Community Atmosphere Model (CAM), which uses a 2-stream method for calculat-
192 ing broadband and heating rates in the shortwave and longwave, and accounts for infrared
193 scattering (Gabriel et al. 2001; Stephens et al. 2001).

194 The dynamical core is based on a spherical geodesic grid (Ringler et al. 2000), which solves
195 the vorticity and divergence equations with second-order accuracy. The model resolution is
196 10,242 grid cells, which yields an average cell area of $4.98 \times 10^{-4} \text{km}^2$ (for comparison a 2.5°
197 $\times 2.5^\circ$ grid has 10,368 grid cells). BUGS and SiB2 were initially coupled in the early 90's
198 (Randall et al. 1996; Denning et al. 1996), and SiB3 was tested in a single column version
199 of BUGS5 (Harper et al. 2010). In the present study, we ran SiB3U and SiB3S coupled to
200 BUGS5 with observed SSTs from 1997-2006. The SSTs are from the Program for Climate
201 Model Diagnosis and Intercomparison (PCMDI) as part of AMIP Phase II (Taylor et al.
202 2000; Hurrell et al. 2008). We ran five 10-year ensembles of each version of the model, each
203 initialized with a restart file from a previous, spun-up AMIP-style run created on the first
204 five days of 1997 (e.g. Ensemble 1 begins with the Jan. 1 restart, Ensemble 2 begins with
205 the Jan. 2 restart, and so on). The biome maps for SiB3 are identical for the two runs,
206 therefore the only differences are the changes in Table 1.

207 *c. Datasets and Analysis*

208 A number of datasets are used for comparison with model results. First, NCEP/DOE
209 Reanalysis version 2 (hereafter "NCEP2") (Kalnay et al. 1996) was provided by the NOAA
210 Earth System Research Laboratory's Physical Sciences Division, from their Web site at
211 <http://www.esrl.noaa.gov/psd/>. We limit our use of NCEP2 to the observation-based vari-

212 ables air temperature, relative humidity, vertical velocity (ω), and geopotential height. Pre-
213 cipitation is from the Global Precipitation Climatology Project (GPCP) version 2.1 (Adler
214 et al. 2003), and outgoing longwave radiation (OLR) is from the Earth’s Radiation Budget
215 Experiment (ERBE), which based OLR on observations from the Earth’s Radiation Budget
216 Satellite and the NOAA9 and NOAA10 satellites from February 1985 to April 1989. ERBE
217 data was accessed at <http://www2.cgd.ucar.edu>. The NCAR Command Language (NCL
218 2013) was used for much of the analysis (e.g. significance testing) and plotting. Statisti-
219 cal significance of differences between the models are determined with two-tailed Student’s
220 t-test. If the returned probability is less than 0.05 we reject the null hypothesis that the
221 means are from the same population, hence the differences are significant.

222 To diagnose behavior during dry season droughts, we first computed an area-averaged
223 time series of precipitation from 50-60°W and 9-14°S, considering tropical forest points only
224 (Fig. S1; see box in Fig. 3). This region contains the largest differences in surface fluxes
225 between SiB3 Stressed and Unstressed, and it encompasses the transition between the humid
226 tropical forests and the more arid savannas. Within this region, the dry season lasts from
227 May through September. The average seasonal cycle was removed to avoid a seasonal bias
228 when determining drought months, and we applied a 5-month running mean to remove
229 short-lived precipitation anomalies. The resultant anomaly timeseries is shown in Fig. S2,
230 and we defined dry season droughts for each ensemble as austral winter months (JJA) with
231 precipitation anomalies < -1 . Composites of drought conditions in each ensemble were then
232 averaged together for analysis of land-atmosphere interactions.

233 We also determined drought months using two well-known drought indices: the Stan-
234 dardized Precipitation Index (SPI) (McKee et al. 1993; Taylor et al. 2012) and the Soil
235 Moisture Anomaly (SMA) (e.g. Burke and Brown (2008)). The SPI was calculated by fit-
236 ting a gamma distribution to the precipitation time series in Fig. S1, and standardizing
237 the resultant time series. By definition, the SPI3 is based on anomalies from the preceding
238 three months and highlights short-term droughts, while the SPI6 is based on the previous

239 six months and identifies longer-term (but sub-annual) droughts. We calculated the SPI for
240 each ensemble and determined drought months (during JJA only) as summarized in Table
241 2. The SMA is directly related to the soil moisture stress felt by the model:

$$SMA = SM - SM_c \quad (1)$$

242 where SM is the soil moisture content for the entire rooting profile averaged over the pre-
243 ceding 12 months, and SM_c is the soil moisture climatology for the individual ensemble. The
244 SMA was standardized, and we defined droughts as months (during JJA) when the SMA <
245 -1.

246 3. Evaluation of BUGS5 Climatology

247 a. *Global climatology*

248 The overall patterns of modeled climate agree well with observations, and the global
249 climate is roughly similar in BUGS5 with SiB3U and SiB3S (Fig. S3). In general, the
250 model tends to produce over-vigorous precipitation at the expense of growing high clouds,
251 as indicated by globally high biases in precipitation and outgoing longwave radiation (OLR).
252 During July, BUGS5 captures observed patterns of global precipitation, but the global mean
253 is too high due to overestimations in tropical convergence zones (Fig. S3). Modeled OLR
254 is higher than the global observed average, indicating an underestimation of cloud cover,
255 especially high clouds. Previous work with BUGS showed sensitivity of tropical rainfall to
256 the parameter alpha in the cumulus parameterization (Lin et al. 2000). The cloud mass flux
257 is inversely proportional to the square root of alpha (Pan and Randall 1998). This study
258 uses the default value of $\alpha = 10^8$ but a larger value might yield more realistic precipitation
259 throughout the Tropics (Lin et al. 2000). Precipitable water is also too high in most of
260 the tropics (not shown). Global mean temperature is slightly higher in BUGS5 than in
261 the NCEP2 Reanalysis, mostly due to overestimation in subtropical dry zones (such as

262 the Sahara and Arabian Peninsula), and in the mid-latitudes (recall that SSTs are set by
263 observed values). Many of the same biases are seen in the January climatology (Fig. S4),
264 and the model performance during January is discussed in the Supplementary Material.

265 *b. Tropical South America climate*

266 Observed annual precipitation has a maximum in the northwestern Amazon (Fig. 1), and
267 high annual rainfall extends to the southeast through the South Atlantic convergence zone.
268 The models capture the mean pattern of high annual rainfall in the northwest and lower
269 rainfall in the southeast. There is too much rainfall in the Intertropical convergence zone
270 (ITCZ) and over high topography, such as the Andes and above southern Brazil. The average
271 precipitation for all tropical forest points in South America is 6.64 and 6.74 mm day⁻¹ in
272 SiB3S and SiB3U, respectively. This is high compared to GPCP (P=5.2 mm day⁻¹), but
273 within the range of measurements compiled by Marengo (2006) (5.2-8.6 mm day⁻¹).

274 For the majority of the Amazon, the wet season occurs during DJF and the dry season
275 occurs during JJA (Fig. 1). This seasonal cycle is reversed north of the Equator. A
276 predominant feature of the lower atmospheric circulation is the Trade Winds, which transport
277 low-level moisture from the tropical Atlantic Ocean and Caribbean Sea, across the continent
278 and toward the Andes (Fig. S5a,c). The highest rainfall occurs in the western Amazon
279 basin, when the Andes force the air southward. During July, winds south of the Equator are
280 southeasterly which limits moisture transport into the southern Amazon compared to the
281 wet season (Fig. S5c). At 850 hPa, there is anticyclonic flow off the southern coast of Brazil,
282 and at 200 hPa westerlies dominate the circulation (Fig. S5d). BUGS5 captures these mean
283 circulation patterns well with a few exceptions: exaggeration of the anticyclone at low levels
284 in July (related to the coarsely resolved SE Brazilian highlands) and underestimation of the
285 southerly component of winds in July in the central Amazon.

286 Average precipitation for South American tropical forests has a similar seasonal cycle in
287 both versions of the model. These are controlled by the large-scale circulation patterns de-

288 scribed above. To better understand precipitation biases in the model, we averaged seasonal
 289 rainfall over two regions used in an analysis of models from the fifth Coupled Model Intercom-
 290 parison Project (CMIP5) (Yin et al. 2012): the southern Amazon (SAma: 5-15°S, 50-70°W)
 291 and northern Amazon (NAma: 5°N-5°S, 55-70°W) (Fig. S6). In the SAma, the models
 292 produce dry biases of 2.6 and 2.4 mm day⁻¹ (in SiB3 Stressed and Unstressed, respectively)
 293 during the wet season (DJF), and wet biases of 1.6 and 1.5 mm day⁻¹ during the dry season
 294 (JJA) (Fig. S6b). The JJA wet bias is in contrast to the majority of the CMIP5 models,
 295 which mostly produce a dry bias (Yin et al. 2012). Compared to the ECMWF ERA-Interim
 296 reanalysis, the majority of CMIP5 models overestimate dry season moisture divergence in
 297 the SAma, possibly related to an over-active ITCZ and strong subsidence over the Amazon.
 298 The two models without a dry season dry bias (HadGEM2-CC and HadGEM2-ES) com-
 299 pensate for high moisture divergence by also having high ET. Following the methodology in
 300 (Yin et al. 2012), we calculated moisture convergence as:

$$MC = P - ET + \Delta TWV \quad (2)$$

301 where ΔTWV is the monthly change in atmospheric total water vapor. In the SAma, the
 302 BUGS5 model demonstrates a similar trade-off between ET and MC as the two Hadley
 303 Centre models (Fig. S7). In SiB3 Stressed, ET is low and MC is near 0, meaning very small
 304 moisture divergence. In SiB3 Unstressed, ET is higher by 0.69 mm day⁻¹, but P is slightly
 305 lower (by 0.04 mm day⁻¹). The excess water vapor originating from ET is transported away
 306 from the Amazon, and MC is more negative by 0.73 mm day⁻¹. A similar result was found
 307 using SiB3 Stressed and Unstressed coupled to a single column version of BUGS5 (Harper
 308 et al. 2010).

309 In the NAma, observed rainfall is relatively high year-round, but the driest (wettest)
 310 months are SON (MAM) (Fig. S6). The BUGS5 modeled seasonal cycle does not match
 311 observations: the driest (wettest) months occur during DJF (SON) in BUGS5. Due to these
 312 high biases in the northern Amazon, the focus of the remaining analysis is on the southern
 313 Amazon. However, since the mean state of the dry season in the southern Amazon is too

314 wet, drought intensities and responses might be dampened in these experiments.

315 4. Results

316 a. *Impact of water stress on dry season fluxes and precipitation*

317 Due to the changes in root zone biophysics (see Section 2a; Table 1), SiB3 Unstressed
318 avoids moisture-related stress during the dry season. For illustrative purposes, Fig. 2 shows
319 daily averages from the two versions of the model during 1999 at a point in the southern
320 Amazon. During the dry season, the stomatal resistance is lower in SiB3 Unstressed, leading
321 to higher rates of photosynthesis and transpiration, and therefore higher LH. Because the
322 net surface energy must be balanced, the higher LH results in lower SH. During JJA, there
323 are significant model differences in surface fluxes along the southern edge of the forest (Fig.
324 3).

325 We define the region with the largest model differences as the southeastern (SE) Amazon
326 for the purpose of further analysis (50-60°W, 9-14°S, tropical forest points only: See box in
327 Fig. 3). (Note that this region is different from the SAma region in Section 3.) The dry
328 season is relatively long and dry, lasting from May-September with average precipitation of
329 1.3-1.6 mm day⁻¹ in the models, therefore the impact of enhanced soil water access is greater
330 than in regions with a less pronounced dry season. Sensible heat flux in SiB3 Stressed is
331 more than twice that from SiB3 Unstressed (56 W m⁻² compared to 27 W m⁻²) (Table 3),
332 and LH is on average 40% higher in SiB3 Unstressed, with the largest difference of 48 W
333 m⁻². As mentioned in Section 2a, LAI and fPAR are prescribed at a constant value in SiB3
334 and are likely overestimated during the dry season in this region. High LAI could lead to
335 overestimated ET, but this bias is present in both versions of the model.

336 Enhanced LH could increase rainfall through moistening and destabilizing the lower at-
337 mosphere (Fu and Li 2004). The model differences in precipitation are small but significant
338 (p<0.05) (Fig. 3). On average, precipitation in the South American tropical forests is

339 marginally greater in SiB3U (by 0.1 mm day^{-1}). Within the SE region, however, precipita-
340 tion is greater in SiB3S (JJA average 1.95 mm day^{-1} compared to 1.61 mm day^{-1} in SiB3U),
341 with the largest difference of 0.8 mm day^{-1} .

342 Convective activity in the model depends on the atmospheric static stability. In SiB3U,
343 enhanced surface humidity decreases the static stability (due to more latent heat in the low
344 atmosphere), while cooler surface temperatures from reduced SH increase stability. During
345 the average dry season, the latter effect is greater, and overall the static stability is greater
346 in SiB3U, resulting in reduced convectively available potential energy (CAPE) and rainfall
347 compared to SiB3S. We examine these differences in more detail in Fig. 4, which shows both
348 latitude-height cross-sections (averages from $50\text{-}60^\circ\text{W}$ along $0\text{-}15^\circ\text{S}$) and an average vertical
349 profile (from $50\text{-}60^\circ\text{W}$ and $11\text{-}13^\circ\text{S}$). The largest differences in temperature, relative humidity
350 (RH), and moist static energy (h) occur near 12°S , where differences in surface fluxes are
351 the greatest. In the lower atmosphere, SiB3S is roughly 1.5 K warmer than SiB3U, and
352 SiB3U has slightly higher surface relative humidity (Table 3). However, less of this moist
353 air is transported upward due to large-scale subsidence in SiB3U. The air is slightly warmer
354 and more moist in SiB3S above 700 hPa . The combined effect is higher h in SiB3S, with
355 significant differences between the models below 600 hPa and south of 5°S .

356 SiB3S has rising air in the low atmosphere, while SiB3U has subsiding air throughout the
357 profile. In comparison, this region is characterized by subsidence in the NCEP2 reanalysis
358 (bearing in mind that in data-sparse regions such as the Amazon, reanalysis products rely
359 heavily upon a model to fill in gaps between observations). The NCEP2 profile is slightly
360 cooler and drier than the modeled profiles (and RH is much lower), resulting in lower h . This
361 is consistent with the result of BUGS5 overestimating dry season precipitation. Compared
362 to the observations, the tendency toward subsidence and lower precipitation rates in SiB3U
363 is an improvement, although a large bias in atmospheric RH still exists.

364 *b. Land-atmosphere interactions during dry season droughts*

365 The following discussion focuses on the previously defined SE Amazon region (50-60°W,
366 9-14°S), where dry season differences in surface fluxes are most pronounced. Over this
367 region, we defined dry season droughts as explained in Table 2, first exploring composites
368 based on the simple anomaly time series. During an average dry season, winds at 850 hPa
369 are predominantly easterly above the central Amazon basin in the NCEP2 Reanalysis, and
370 there is anticyclonic rotation above southern Brazil (Fig. S5). During modeled dry season
371 droughts, this flow is reversed: the 850 hPa winds are anomalously southeasterly above
372 southern Brazil, and westerly near the Equator (Fig. 5). Instead of moist air flowing onto
373 the continent from the tropical Atlantic, drier air is advected into the Amazon region from
374 the south/southeast. The atmosphere is warmer and drier than during an average dry season,
375 and the differences in surface fluxes between the models are enhanced (Fig. 6, Table 3).

376 Contrary to the results for an average dry season, the enhanced ET in SiB3U can play
377 an important role in moisture recycling during dry season droughts. Precipitation in the SE
378 Amazon region is approximately 20% higher in SiB3U (Fig. 6, Table 3). The Unstressed
379 atmosphere is cooler (by up to 1.8 K at 850 hPa) and more moist (by on average 1 g kg^{-1}).
380 The result is slightly higher h between the surface and 850 hPa, although the differences are
381 not significant (Fig. 7). The moist static energy is higher near the surface and lower near
382 700 hPa, and there is slightly more CAPE in SiB3U. Above 700 hPa, subsidence is stronger
383 in SiB3S.

384 Although rainfall during droughts is higher in SiB3U, the number of drought months
385 is also higher. This is likely a result of the selection criteria. During the dry season, the
386 modeled rainfall is infrequent but heavy rains are possible, leading to a bimodal probability
387 distribution for dry season rain rates in both versions of the model. In SiB3U this distribution
388 is spread out: the very dry and very wet months are more frequent at the expense of 'average'
389 months (Fig. S8). Therefore, it is more likely for this model to encounter dry season months
390 with low rain rates. To test the dependence of results on the definition of drought months, we

391 composite dry season droughts based on the Standardized Precipitation Index (SPI) and Soil
392 Moisture Anomaly (SMA) (Table 2). There are more "drought" months during JJA using
393 these two indices than with the original index. The timing of droughts in the original index is
394 most similar to the 3-month SPI. The annual SMA best captures the long-term nature of the
395 2005 drought in the SE Amazon region, which was particularly severe because it followed an
396 anomalously dry wet season (Marengo et al. 2008a). The model simulates drought conditions
397 during the austral winter of 2005 in 3 of the 5 ensembles for SiB3 Stressed, and in 4 of the
398 5 ensembles in SiB3 Unstressed.

399 During droughts based on the SPI, the average precipitation over the SE Amazon region
400 is higher in the model with higher RH throughout the profile and higher vertically integrated
401 h (Figs. 8 and S9). An additional factor is the Bowen ratio ($BR=SH/LH$) in SiB3 Stressed
402 (Table 4). The BR for SiB3 Unstressed is always between 0.25 and 0.35, and it is always
403 higher in SiB3 Stressed (0.8-1.3). When $BR>1$ in SiB3 Stressed, the model differences in
404 lower atmospheric moisture are substantial, leading to stronger moisture recycling in SiB3
405 Unstressed. For example, during Extreme and Moderate droughts as defined with the 6-
406 month SPI, $BR>1$ in SiB3S, and lower atmospheric RH and h are both higher in SiB3U
407 (Fig. 8). As a result, precipitation is stronger in SiB3U (Table 4). Conversely, $BR<1$
408 in SiB3S during Severe droughts and abnormally dry periods, the lower atmosphere's h is
409 higher, and precipitation is higher than in SiB3U. Similar relationships between surface fluxes
410 in SiB3 Stressed and precipitation are seen with the 3-month SPI (Supplemental Material).
411 Droughts defined with the SMA are generally less severe than the SPI droughts. The vertical
412 structures of model differences in RH, h , and ω are similar to those during an average JJA
413 (Fig. S10). There is more moist static energy in the lower atmosphere in SiB3S, and stronger
414 precipitation in this model (Table 4).

416 Next we analyze the dry season land-atmosphere interactions at 9°S, 50°W, which is a
417 tropical forest point near the forest/savanna transition (See Fig. 3). While the previous
418 analysis focused on large-scale patterns, this allows more detailed investigation of weekly
419 variability in surface energy fluxes and their impact on the overlying circulation. We pur-
420 posefully use an area near the forest/savanna transition because it is impacted by more arid
421 air, which increases evaporative demand and intensifies the difference in LH between the
422 models. During the average dry season, plants are able to draw on stored soil moisture in
423 both models. Modeled ET rates decline during July, similar to the observed seasonal cycle
424 from a nearby tower (Javaes: da Rocha et al. (2009)). However there are two complica-
425 tions with directly comparing the model results to in-situ observations. First, the wet bias
426 during JJA results in low moisture stress in both versions of the model. Average modeled
427 dry season precipitation was 100-106 mm month⁻¹, while observed precipitation near Sinop
428 (11°24.75'S, 55°19.50W) was 50-100 mm month⁻¹ (Vourlitis et al. 2011). Second, the SiB3
429 input data classifies the region as "tropical broadleaf evergreen" biome, while the true vege-
430 tation coverage is a semideciduous forest (Vourlitis et al. 2011; Costa et al. 2010; Lathuilliere
431 et al. 2012). The semideciduous forests typically have a dry season decrease in ET, due to
432 both phenology and stomatal control (Costa et al. 2010; Vourlitis et al. 2011). Although
433 SiB3 Unstressed simulates a dry season increase in aerodynamic resistance, the stomatal
434 resistance shows almost no seasonal cycle (Fig. 2). Therefore, it is likely that the Unstressed
435 model overestimates dry season ET at this point, and the model differences in this section
436 can be viewed as representing two extremes in land-atmosphere interactions.

437 In SiB3S, canopy transpiration declines throughout the dry season, and the PBL is
438 warmer and drier. During a simulated dry season drought (1999), transpiration is below
439 average in both models, but LH is lower by up to 60 W m⁻² in SiB3S (Fig. 9). The LH
440 is closely linked to rainfall events in SiB3S, while SiB3U produces a higher flux with less
441 variability. As a result, the atmospheric precipitable water (PRW) remains relatively low in

442 SiB3S through mid-August, while both PRW and h increase about a month earlier in SiB3U.
443 Although both versions of the model simulate a drought, the anomalously dry conditions
444 last longer in SiB3S.

445 Figure 9 also shows results from two ensembles. In Ensemble 2 (dashed line), SiB3U's LH
446 is fairly steady through June and isolated rainfall events keep the monthly mean precipitation
447 high. Precipitable water is similar in the two models until mid-July, when continued dry
448 conditions result in low LH in SiB3S. Latent heat flux reaches a minimum by mid-July, and
449 PRW, h , and CAPE also experience strong reductions. The average rainfall during July is
450 0.39 ± 0.26 mm day⁻¹ in SiB3S, compared to 1.62 ± 1.08 mm day⁻¹ in SiB3U. Alternatively,
451 SiB3U produces less rainfall than SiB3S during June-July 1999 in Ensemble 3. In this case,
452 large-scale dry conditions overshadow the higher LH in SiB3U and result in low CAPE,
453 PRW, and h . This represents a limit on the moisture recycling capacity of the forest.

454 The differences in soil moisture access between the models also have implications for the
455 carbon cycle. The net CO₂ flux from the canopy to the atmosphere is the difference between
456 uptake by the forest through photosynthesis and efflux due to respiration (the convention is
457 Fig. 9 is such that a negative flux is uptake). SiB3U generally simulates the land as a carbon
458 sink during the dry season of 1999, due to higher rates of photosynthesis than respiration,
459 while the opposite is true for SiB3S. Localized precipitation events can temporarily switch
460 the carbon sink to a source. For example, in Ensemble 3, large pulses of soil respiration
461 following heavy rains in late August temporarily convert the forest to a carbon source in
462 SiB3U.

463 5. Conclusion

464 Access to deep soil moisture by efficient rooting systems is important for drought sur-
465 vival (Nepstad et al. 1994, 2007; Jipp et al. 1998). The current study investigates how the
466 avoidance of dry season water stress can increase moisture recycling and mitigate drought

467 intensity. Accounting for drought tolerance mechanisms in SiB3U enables a more realistic
468 simulation of the average dry season in the southern Amazon. Increased LH and reduced
469 SH in SiB3U cool the lower atmosphere, thereby increasing the static stability and reducing
470 convection. The result is somewhat unexpected, since SiB3U has significantly higher ET
471 than SiB3S and a more moist lower atmosphere. There is no evidence for a positive feedback
472 between low precipitation and reduced ET during the average dry season, since SiB3S has
473 lower ET and yet more precipitation.

474 During a dry season drought, maintained ET has the potential to dampen the drought's
475 intensity if the moistening effect of higher LH is stronger than the cooling effect of lower SH.
476 Precipitation is higher in SiB3U during dry season droughts when the atmospheric conditions
477 are amenable to convection, as indicated by high relative humidity and moist static energy
478 relative to SiB3S. Additionally, when sensible heat flux is higher than latent heat flux in
479 SiB3S, the hot and dry lower atmosphere limits precipitation relative to SiB3U.

480 Two factors could limit moisture recycling during drought. First, enhanced moisture
481 availability cannot override a strongly statically stable atmosphere, as was shown to be
482 the case in the example from Ensemble 3 (Fig. 9). In this case, the moist static energy,
483 CAPE, and precipitable water vapor all are anomalously low in SiB3U (even compared to
484 the average during a drought). This is a similar result to the observed impacts of LH on wet
485 season onset from Fu and Li (2004). In that study, which was based on ECMWF Reanalysis,
486 an anomalously wet surface was shown to be a necessary but not sufficient condition for early
487 transition from wet to dry season.

488 A second factor affecting moisture recycling during drought is the diversity of plant
489 response to drought. Given the high species diversity of the Amazon forest, its trees likely
490 employ a variety of mechanisms for drought tolerance and avoidance of hydraulic failure.
491 In addition, the modeling study does not account for semideciduous trees in the southern
492 Amazon, nor land use change. Due to these limitations, the response from SiB3 Unstressed
493 can be interpreted as an upper limit to the ability of the forest to recycle precipitation.

494 Rainfall exclusion studies have illuminated drought responses in two equatorial Amazon
495 sites, but modeling the subtleties of these responses presents many challenges (e.g. Powell
496 et al. (2013)). The current study does not incorporate spatial heterogeneity in drought
497 response but future model development in Amazonia should account for gradients in plant
498 and soil hydraulic and physiological responses to drought, which could be a function of
499 soil type, rainfall variability, and/or nutrient availability. Continued observations of forest
500 response to drought are essential for such work to move forward.

501 During particularly strong and/or long droughts, trees reach a limit in their ability to
502 access and use soil moisture. Observational evidence suggests that such a threshold has
503 been reached during droughts in the past decade (Phillips et al. 2010; Lewis et al. 2011).
504 In terms of moisture recycling, it appears there is a threshold in the model which occurs
505 when the Bowen ratio is greater than one. Although this study did not directly address land
506 use change, an important implication is that forest preservation is essential for enabling the
507 Amazon forest to withstand a potentially drier climate. Pasture and secondary forests do
508 not have the extensively developed rooting systems present in primary forest, and loss of
509 vegetation coverage increases runoff during heavy rain. Pasture is more likely to experience
510 dry season water stress and seasonal reductions in ET, particularly in the southern Amazon
511 (von Randow et al. 2012), and deforestation can reduce moisture recycling and down-wind
512 precipitation (Spracklen et al. 2012). In addition, forests that border pasture or savanna are
513 more prone to desiccation and fire impacts (Malhi et al. 2008). Large areas of undisturbed
514 forest are more likely to maintain ET during dry periods and recycle rainfall.

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TABLE 1. Differences in the model formulation between SiB3 Stressed and Unstressed.

	SiB3 Stressed	SiB3 Unstressed
Number of soil layers	10	10
Soil depth (m)	3.5	10
Treatment of root water extraction	Extraction weighted by root biomass	Extraction weighted by soil moisture in the layer
Soil moisture stress function	Stress increases linearly with decreasing soil moisture	Stress increases gradually (slower) with decreasing soil moisture

TABLE 2. Methods for defining dry season drought months. In each case the time series (TS) refers to the area-averaged precipitation over the SE Amazon region, as shown in Fig. 3.

Method	Description
Anomaly Timeseries	Standardized anomalies from the deseasonalized TS with 5-month running mean.
SPI3	Standardized precipitation index based on the previous 3-months.
SPI6	Standardized precipitation index based on the previous 6-months.
SPI: Very Extreme	$-2.00 > \text{SPI}$
SPI: Extreme	$-1.60 > \text{SPI} > -1.99$
SPI: Severe	$-1.30 > \text{SPI} > -1.59$
SPI: Moderate	$-0.80 > \text{SPI} > -1.29$
SPI: Abnormally dry	$-0.51 > \text{SPI} > 0.79$
SMA	Soil moisture anomalies based on the annual soil moisture.

TABLE 3. Average latent heat (LH: W m^{-2}), sensible heat (SH: W m^{-2}), Bowen ratio (BR=SH/LH), precipitation (P: mm day^{-1}), temperature at 850 hPa (T_{850} : K), and specific humidity at 850 hPa (q_{850}): kg kg^{-1}) during an average JJA, and during dry season droughts

	<i>SiB3S(JJA)</i>	<i>SiB3U(JJA)</i>	<i>SiB3S(Drought)</i>	<i>SiB3U(Drought)</i>
LH	77	114	63	112
SH	56	27	65	32
BR	0.72	0.24	1.03	0.29
P	1.95	1.61	1.29	1.57
T_{850}	292.1	290.8	292.0	290.8
q_{850}	0.0122	0.0125	0.0108	0.0122

TABLE 4. Model differences during an average dry season, and during dry season droughts as defined by the original Anomaly Time Series, the Soil Moisture Anomalies (SMA), and the Standardized Precipitation Index (SPI). Bowen ratio (BR) is shown as the value for SiB3 Stressed only. Only drought composites are shown for which >2 months of drought occurred in each model. $\Delta P = \frac{P_{SiB3U} - P_{SiB3S}}{P_{SiB3S}}$, $\Delta q = q_{850,SiB3U} - q_{850,SiB3S}$, and $\Delta T = T_{850,SiB3U} - T_{850,SiB3S}$.

	<i>Average</i>	<i>AnomalyTS</i>	<i>SMA</i>	<i>3 – monthSPI</i>			
				<i>extreme</i>	<i>severe</i>	<i>moderate</i>	<i>abnormal</i>
BR	0.72	1.03	0.8	1.29	0.79	0.94	0.89
ΔT (K)	-1.3	-1.2	-1.4	-1.2	-1.3	-1.7	-1.6
Δq (g kg ⁻¹)	0.3	1.4	0.6	1.0	0.4	0.8	0.3
ΔP	-17%	22%	-11%	22%	-8%	-31%	-1%
	<i>6 – monthSPI</i>				<i>12 – monthSPI</i>		
	<i>extreme</i>	<i>severe</i>	<i>moderate</i>	<i>abnormal</i>	<i>moderate</i>	<i>abnormal</i>	
<i>BR</i>	1.2	0.82	1.05	0.69	0.78	0.93	
ΔT (K)	-1.3	-1.6	-1.3	-1.4	-1.5	-1.3	
Δq (g kg ⁻¹)	1.3	0.0	1.2	0.3	0.6	1.0	
ΔP	26%	-7%	9%	-30%	-13%	1%	

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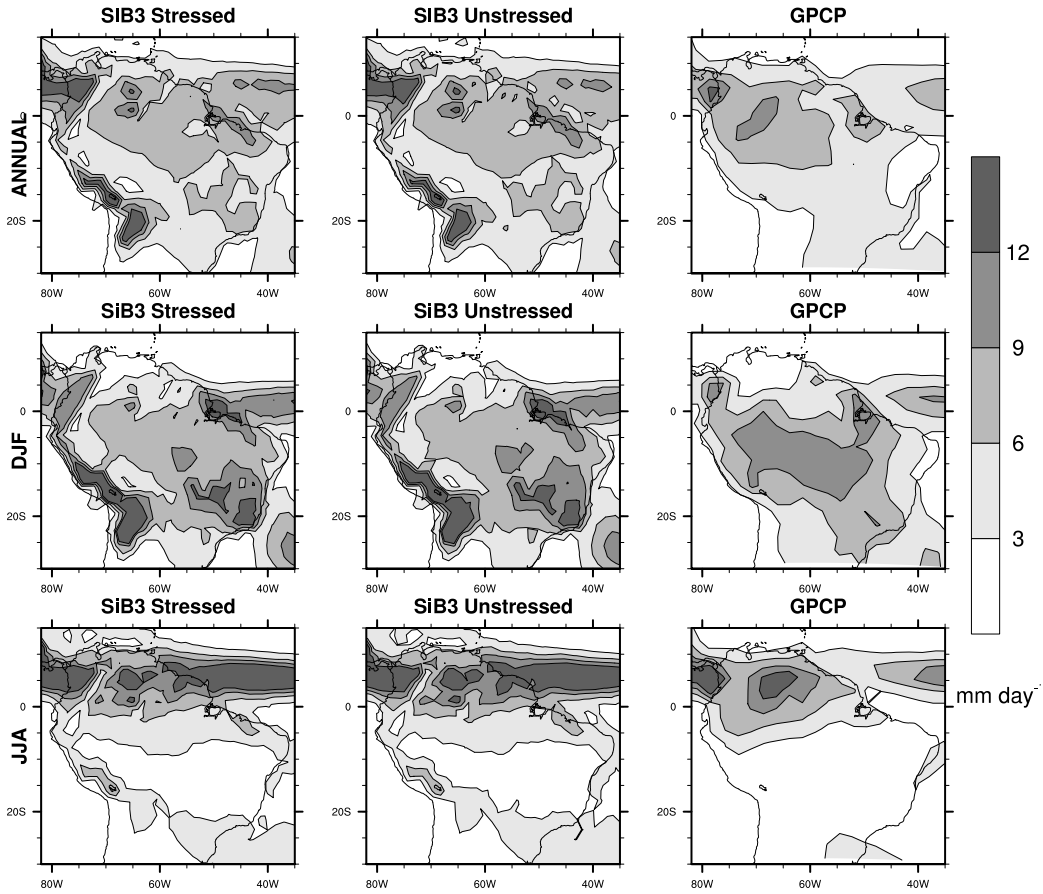


FIG. 1. Average precipitation annually, and during DJF and JJA in SiB3 Stressed, Unstressed, and GPCP. The models and observations are plotted in their native grid ($2.5^{\circ} \times 2.5^{\circ}$ for GPCP and roughly $2.5^{\circ} \times 2.5^{\circ}$ for the models). Time period for averages is 1997-2006.

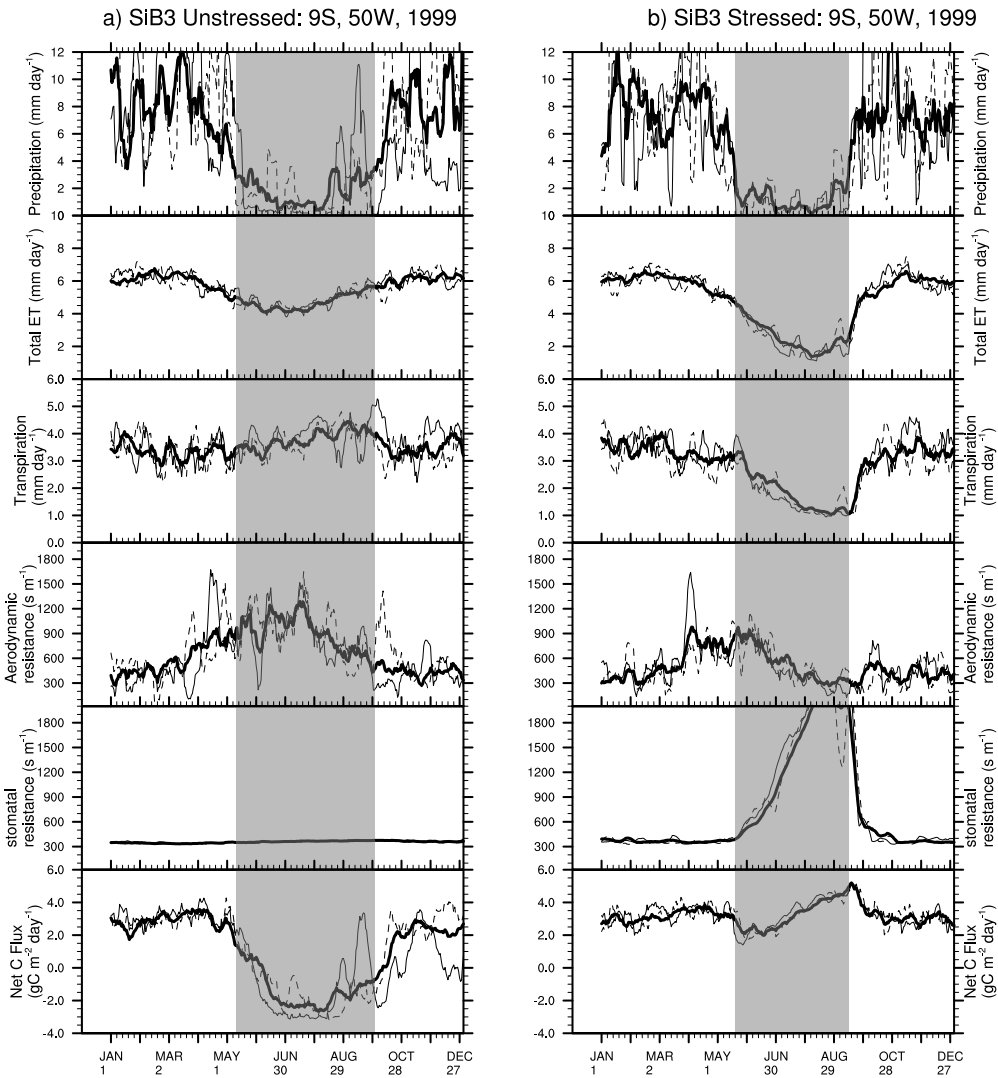


FIG. 2. Precipitation, ET, transpiration, aerodynamic resistance, stomatal resistance, and net C flux from the canopy. The dark line is the average of 5 ensembles. Patterned lines show the variability seen in individual ensembles: dashed line (Ensemble 2) and thin solid line (Ensemble 3). All time series have a 10-day running mean applied. Dark shading indicates the dry season in the ensemble average.

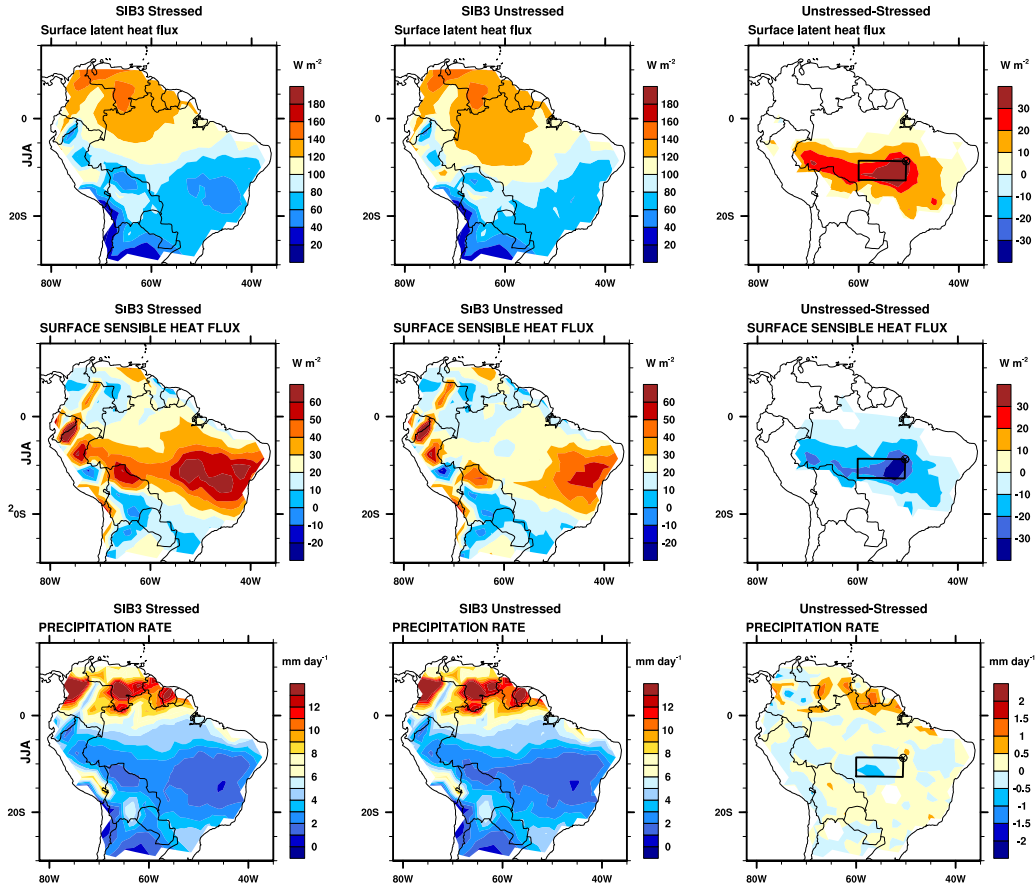


FIG. 3. Latent heat flux (W m^{-2}) during JJA in SiB3S, SiB3U, and the difference between the models (Unstressed - Stressed). Second and third rows are the same but for sensible heat flux (W m^{-2}) and precipitation (mm day^{-1}). Only significant differences are shown ($p < 0.05$). The box is the southeast Amazon (SE) region discussed in the text ($50\text{-}60^\circ\text{W}$, $9\text{-}14^\circ\text{S}$). The circle in the top-right of the box marks the location used in Figs. 2 and 9.

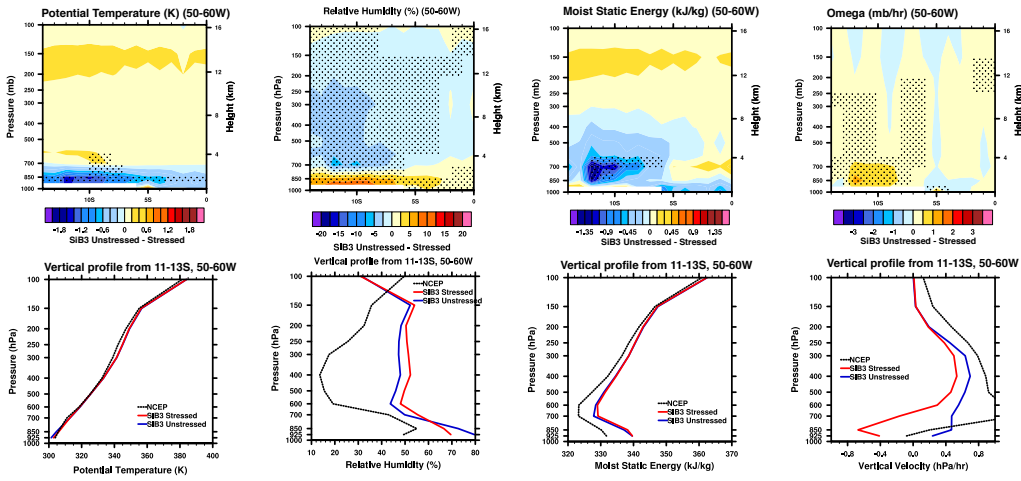


FIG. 4. (Top row) Latitude-height cross sections of model differences (SiB3U-SiB3S) of potential temperature (K), relative humidity (%), moist static energy h (kJ kg^{-1}), and the vertical velocity ω (hPa hr^{-1}) from 0-15°S (averaged from 50-60°W) during JJA. Stippling indicates regions of significant differences between the models ($p < 0.05$). (Bottom row) Vertical profiles averaged over 11-13°S and 50-60°W from the two models and NCEP2 Reanalysis.

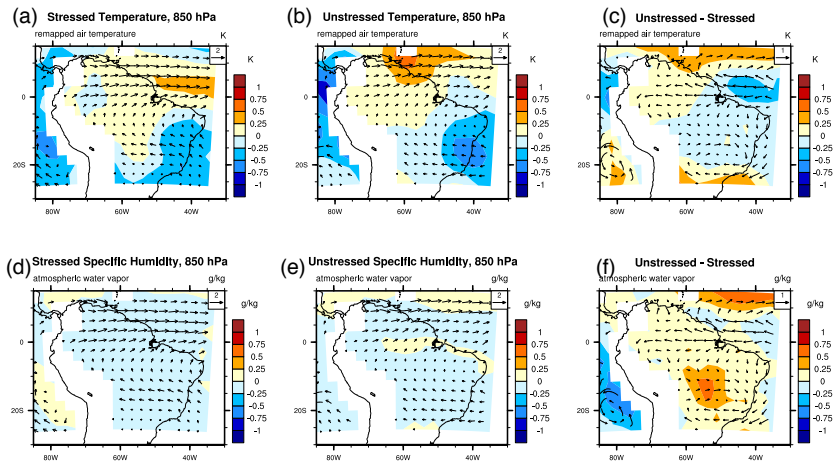


FIG. 5. Composites of anomalous 850 hPa winds, temperature, and specific humidity during JJA droughts (as defined with the anomaly time series) in SiB3 Stressed, Unstressed, and the difference between the two models.

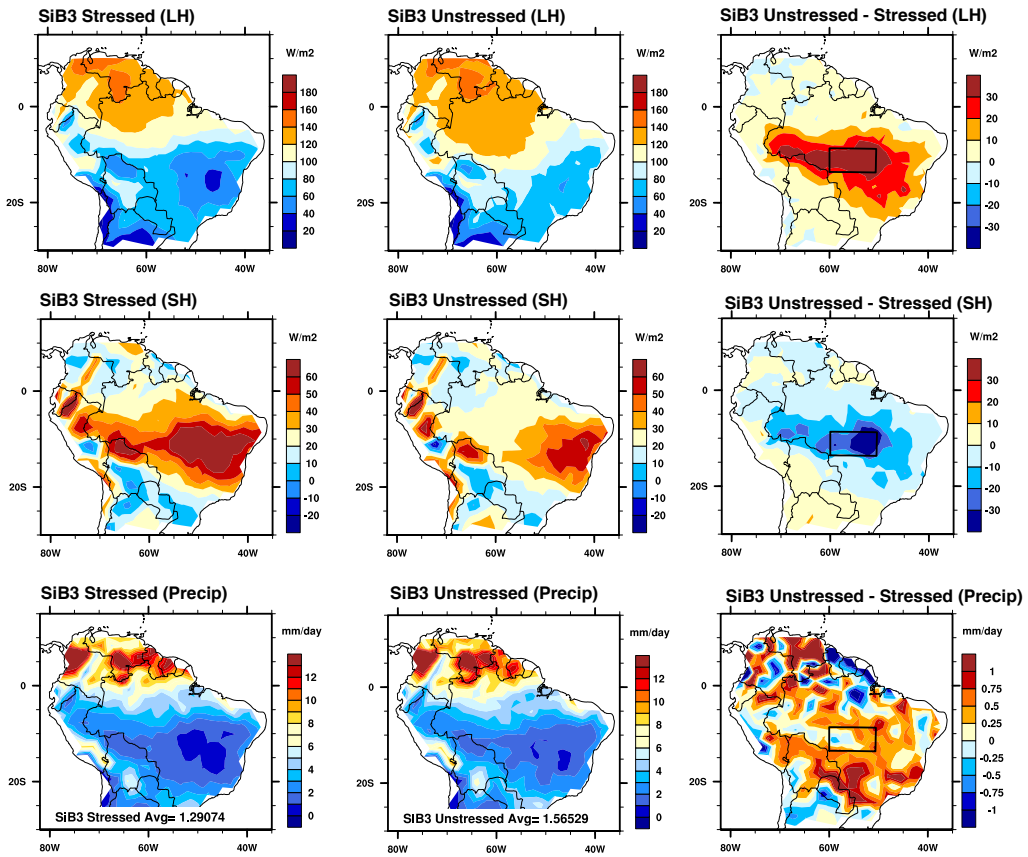


FIG. 6. As in Figure 3 but averages are during dry season drought months. Note that all differences are shown. LH=Latent heat flux; SH=Sensible heat flux.

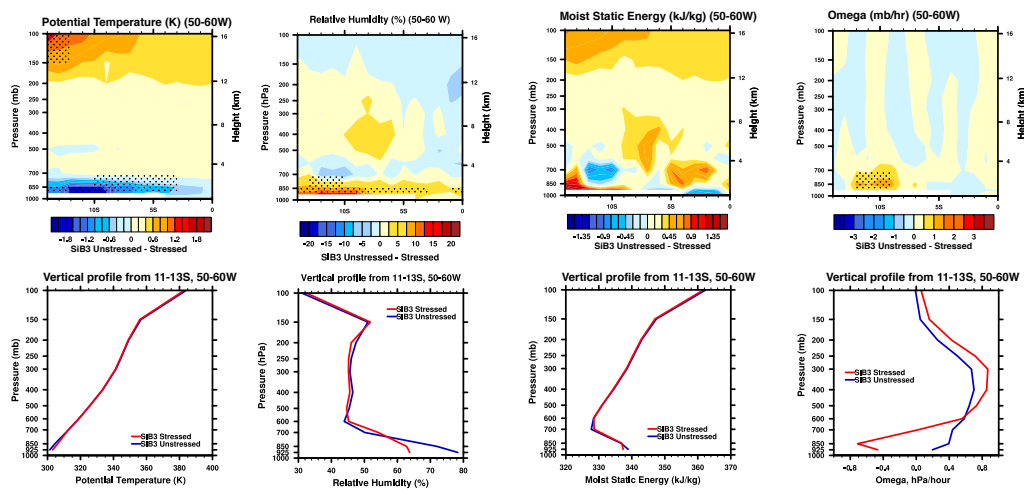


FIG. 7. As in Figure 4 but during JJA droughts.

JJA Droughts defined from SPI6 (differences = Unstressed - Stressed)

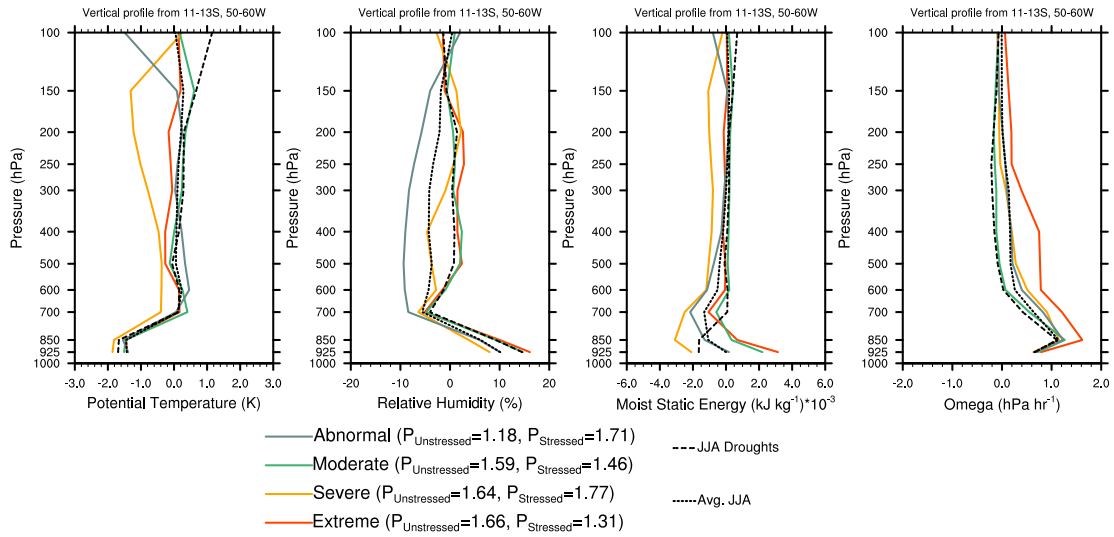


FIG. 8. Differences (SiB3 Unstressed–SiB3 Stressed) in the vertical profiles during droughts defined from the 6-month Standardized Precipitation Index. Profiles are averaged over 11-13°S and 50-60°W. Drought intensities are defined in Table 3. For reference the profiles are shown for an average JJA and a JJA drought defined with the Anomaly Time Series.

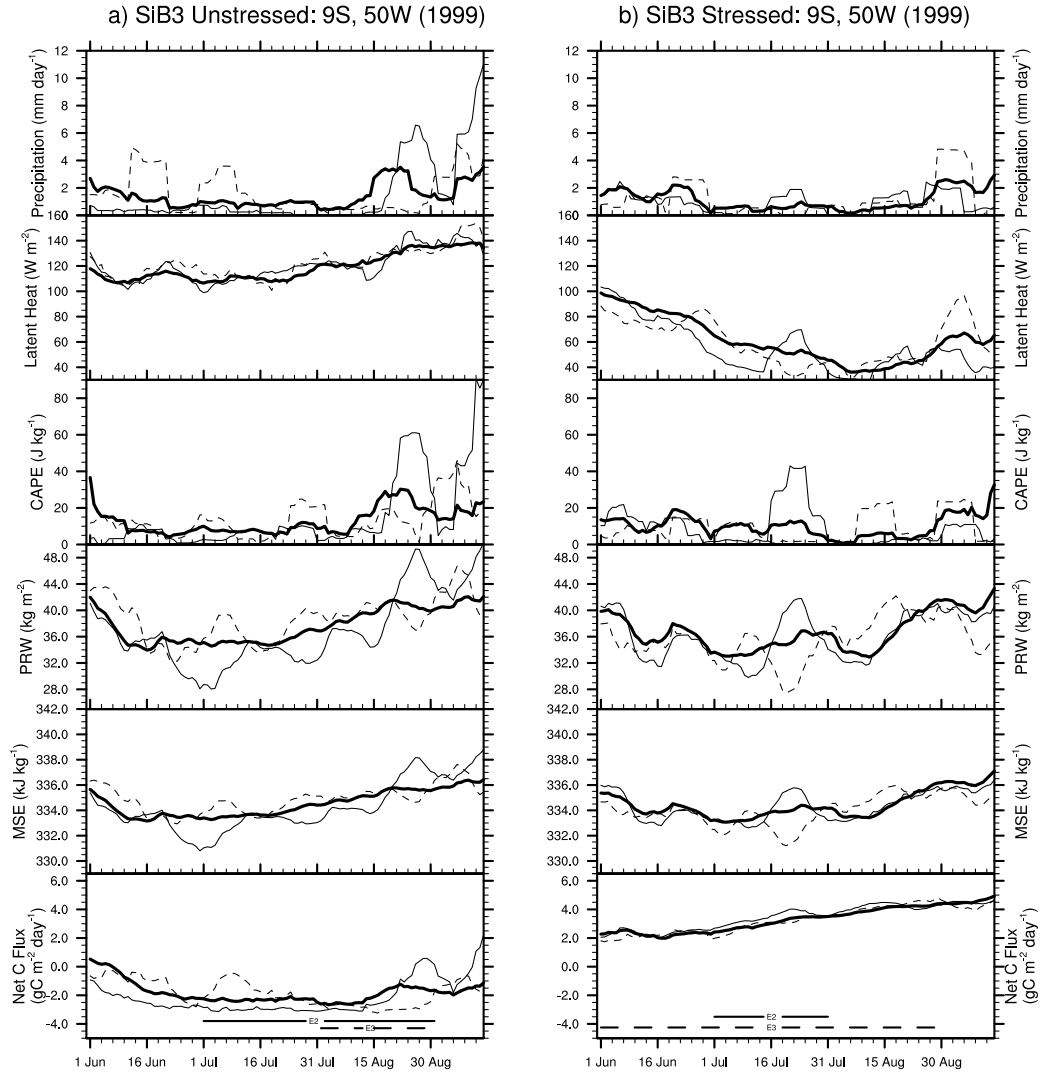


FIG. 9. Precipitation, latent heat flux, convectively available potential energy (CAPE), total column precipitable water (PRW), vertically integrated moist static energy (MSE: from 1000 to 100 hPa), and net C flux from the canopy during the dry season of 1999. The dark line is the average of 5 ensembles. Patterned lines show the variability seen in individual ensembles: dashed line (Ensemble 2) and thin solid line (Ensemble 3). All time series have a 10-day running mean applied. The climatological dry season is May-Sept. The months which qualified as droughts were: July-Aug. (SiB3U, Ensemble 2), Aug. (SiB3U, Ensemble 3), July (SiB3S, Ensemble 2), June-Aug. (SiB3S, Ensemble 3). The beginning and end of the droughts are denoted with the horizontal lines above the x-axis.