

Water table depth modulates productivity and biomass across Amazonian forests

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**Water table depth modulates productivity and biomass
across Amazonian forests**

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Title: Water table depth modulates productivity and biomass across Amazonian forests

Abstract

Aim Water availability is the major driver of tropical forest structure and dynamics. While most research has focused on the impacts of climatic water availability, remarkably little is known about the influence of water table depth and excess soil water on forest processes. Nevertheless, since plants take-up water from the soil, the impacts of climatic water supply on plants are likely to be modulated by soil water conditions.

Location Lowland Amazonian forests

Time period 1971 to 2019

Methods We use 344 long-term inventory plots distributed across Amazonia to analyse the effects of long-term climatic and edaphic water supply on forest functioning. We modelled forest structure and dynamics as a function of climatic, soil-water, and edaphic properties.

Results Water supplied by both precipitation and groundwater affect forest structure and dynamics, but in different ways. Forests with shallow water table (depth < 5 m) had 18% less above-ground-woody productivity and 23% less biomass stock than deep water table, while forests in drier climates (maximum cumulative water deficit < -160 mm) had 21% less productivity and 24% less biomass than those in wetter climates. Productivity was affected by the interaction between climatic water deficit and water table depth: on average, in drier climates shallow water table forests had lower productivity than deep water table forests, with this difference decreasing within wet climates where lower productivity is confined to very shallow water table.

Main conclusions We show that the two opposites of "water availability" (excess and deficit) both reduce productivity in Amazon upland (*terra-firme*) forests. Biomass and productivity across Amazonia respond not simply to regional climate but rather to its interaction with highly locally differentiated water table conditions. Our study disentangles the relative contribution of those factors, helping to improve understanding of tropical-ecosystem functioning and how they are likely to respond to climate change.

Keywords: groundwater, tropical ecology, seasonality, forest dynamics, above-ground biomass, carbon

Introduction

Tropical forests hold a disproportionate share of the Earth's biodiversity and carbon stocks, providing environmental services of global importance through their hydrological and carbon cycles (Fauset et al., 2015; Fearnside, 2008; Pokhrel et al., 2014; ter Steege et al., 2013). Amazonia represents the largest of all tropical forests, and plays a fundamental role as a long-term carbon sink, mostly due to the carbon accumulated in woody plants (Pan et al., 2011; Phillips & Brien, 2017). Therefore there is great interest in understanding underlying controls on biomass productivity and dynamics of the Amazonian forests, and how climate change is and will affect them (Llopart et al., 2018; Malhi et al., 2009; Zhao & Running, 2010). Amazonian climates are naturally characterized by spatial and temporal variability in the distribution of

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3 45 rainfall, and recently both droughts and floods have become more frequent, probably
4 46 driven by anthropogenic climate change (Gloor *et al.*, 2013, 2015; Marengo &
5 47 Espinoza, 2016). In this context, it is essential to understand the impact of water
6 48 availability on forest functioning. While this has been studied from the perspective of
7 49 changes in precipitation seasonality and climatic water deficits (e.g., Phillips *et al.*,
8 50 2009; Toledo *et al.*, 2011b; Álvarez-Dávila *et al.*, 2017) there has been much less
9 51 attention paid to the role of water availability in the soil, as regulated by groundwater
10 52 (but see Nobre *et al.*, 2011; Ivanov *et al.*, 2012; Esteban *et al.*, 2020; Chitra-Tarak *et al.*,
11 53 2021), and no account of how groundwater affects forest productivity and biomass
12 54 measured on the ground currently exists.

13 55 Water is essential to life and, together with temperature, a key determinant of
14 56 global patterns of plant distribution and productivity (Ellison *et al.*, 2017; Law *et al.*,
15 57 2002; Webb *et al.*, 1978; Whittaker, 1975). Although variation in precipitation is
16 58 associated with large-scale variation in forest structure and dynamics, soil-water
17 59 availability to plants is the result of the fine-scale interplay of precipitation and terrain
18 60 properties at landscape scales. The major landscape factors affecting the redistribution
19 61 of water entering the system as rainfall are topography and soil texture (Fan, 2015; Fan
20 62 & Miguez-Macho, 2011; Moeslund *et al.*, 2013). Topography affects the water flow to
21 63 groundwater, and groundwater movement to lower gravitational positions (lower
22 64 relative elevation in the landscape) creates gradients of increasing water availability
23 65 from uplands towards valleys (Fan, 2015; Nobre *et al.*, 2011; Rennó *et al.*, 2008). The
24 66 retention of water depends on soil texture, decreasing with soil particle size, so that it is
25 67 greater in clays than in sands (Costa *et al.*, 2013; Hillel, 1998; Parahyba *et al.*, 2019).
26 68 The dynamics of water drainage and retention in the soil supply the groundwater,
27 69 influencing seasonal and interannual fluctuations in the water table (Hodnett *et al.*,
28 70 1997; Miguez-Macho & Fan, 2012), and also affects soil-water conditions in the rooting
29 71 zone.

30 72 Water table depth (WTD) can be used as a proxy for the accessibility of
31 73 groundwater to plants, mediated by root depth, which is highly constrained by WTD
32 74 (Fan *et al.*, 2017), and soil density (Emilio *et al.*, 2013; Quesada *et al.*, 2012). In
33 75 Amazon non-flooded (*terra-firme*) forests, at low topographic positions roots are in
34 76 direct contact with the superficial water tables or capillary fringe year-round or during
35 77 the wet season, but roots become progressively decoupled from the groundwater with
36 78 increasing ground elevation relative to the local water table (Fan, 2015; Fan *et al.*,
37 79 2017). During normal dry seasons, the water table level drops and the soil surface
38 80 becomes drier, but the intensity of this effect depends not simply on climate but also on
39 81 the soil retention properties and subsidy of groundwater flowing from higher
40 82 topographic positions (Tanco & Kruse, 2001; Tomasella *et al.*, 2008). Understanding
41 83 this process is especially important because a considerable portion (~ 50%) of
42 84 Amazonian forest have a relatively superficial water table of 5m depth or less (Costa *et al.*,
43 85 2022; Fan & Miguez-Macho, 2010).

44 86 Water table depth is expected to play a key role in the regional patterns of plant
45 87 growth and mortality (Costa *et al.*, 2022). Easier access to groundwater in shallow water
46 88 table forests is likely to reduce the effects of precipitation water deficit during the dry

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89 season, thus promoting greater productivity in these environments than in sites in the
90 same climate where the water table is deep. However, excess water in shallow water
91 table conditions during the wet season leads to anoxic stress, which may result in
92 reduced plant growth. Water excess inhibits oxygen flow to the roots and limits plant
93 growth, since alternative anaerobic routes of energy production are much less efficient
94 than aerobic respiration (Gibbs & Greenway, 2003; Parolin, 2012). Thus, optimal
95 conditions for growth may be restricted to a short window of time, limiting the potential
96 for biomass accumulation. Additionally, to avoid anoxic conditions, tree roots are
97 typically superficial in shallow water table environments (Canadell et al., 1996; Fan et
98 al., 2017; Jackson et al., 1996). The resulting poor anchorage, in combination with the
99 loose aggregation of soil particles in waterlogged conditions, increases the risk of
100 treefall (Gale & Barfod, 1999; Gale & Hall, 2001; Ferry *et al.*, 2010). Together, these
101 constraints lead to the expectation that where water tables are shallow, low soil oxygen
102 will lead to low productivity, and weak root anchorage will lead to higher mortality
103 rates, and reduced stand biomass. While some local studies have documented these
104 patterns, major uncertainties remain, in part because forests with shallow water tables
105 tend to be understudied, but also because in some local contexts shallow water table
106 forests may not have lower biomass productivity than nearby deep water table forests
107 under the same climatic conditions (Damasco et al., 2013; Grogan & Galvão, 2006).

108 In summary, the impacts of water on forests depend on much more than simply
109 how much rain falls. Although soil moisture is difficult to measure and characterize
110 over the relevant scales of individual trees and plots across the Amazon, some key
111 determinants of the local hydrological conditions in non-flooded upland forests -
112 precipitation, water table depth and soil texture (Fan et al., 2017; Freeze & Cherry,
113 1979; Zipper et al., 2015) - can be estimated. The effects of those hydrological
114 components on plant responses are not expected to be simple linear and additive effects,
115 but rather involve complex interactions, as different combinations may give rise to
116 water deficit, excess of water or mesic conditions.

117 Here, we use a unique, extensive long-term forest-monitoring dataset across
118 Amazonia, resulting from the efforts of hundreds of researchers and field assistants
119 working for decades (ForestPlots.net *et al.*, 2021), to address two central questions: (1)
120 How do the structure and dynamics of Amazonian forests vary with water table depth
121 and the long-term average climatic water deficit?, and (2) How does water table depth
122 interact with climatic water deficit and soil properties to influence Amazonian forest
123 structure and dynamics? There are reasons to expect that above-ground-biomass
124 productivity and above-ground-biomass stock are lower, and mortality higher, with both
125 water deficit and with water excess. Considering the challenges imposed on plant
126 growth by saturated soils, we predict that the combination of a wet climate and a
127 shallow water table leads to the lowest productivity and highest mortality, while shallow
128 water table within a dry climate mitigates the climatic water deficit, allowing higher
129 productivity than in deep water table settings. Soil texture is expected to further
130 modulate those responses, as soils with low-water-retention capacity could reverse the
131 positive interaction of shallow water tables and dry climates.

133 **Materials and methods**

134 **Vegetation data**

135 To address our questions, we analyzed plot-level data from long-term ground-
 136 based monitoring of Amazon forests, using available records from intact old-growth
 137 forests in lowland (125 ± 115 m altitude) Amazonia that are not seasonally or permanently
 138 flooded, i.e. *terra-firme* forests. We used data from 344 plots monitoring Amazon
 139 vegetation from the RAINFOR and PPBio networks (Lopez-Gonzalez et al., 2011;
 140 Magnusson et al., 2013) (see Table S1 for plot details). Only plots with two or more
 141 censuses were included in this study. The vegetation monitoring followed standardized
 142 measurement protocols. In RAINFOR plots, all trees and palms with a diameter (D) at
 143 1.3 m (or above buttress) ≥ 10 cm were tagged and measured (196 plots in this dataset)
 144 (Phillips et al., 2010). In PPBio plots all stems with $D \geq 30$ cm are sampled in the full 1
 145 ha per plot, stems with $10 \text{ cm} \leq D < 30 \text{ cm}$ were measured in a subplot of 0.5 ha per plot
 146 (148 plots in this dataset) (Magnusson et al., 2005). Field data were curated and accessed
 147 via the ForestPlots.net database (Lopez-Gonzalez et al., 2011), and subject to strict quality
 148 control to identify possible measurement or annotation errors, as described in Brienen et
 149 al. (2015).

150 To evaluate the forest structure and dynamics, we estimated the plot-based above-
 151 ground biomass stock (AGB) and above-ground woody productivity (AGWP) of trees
 152 and palms per hectare, in each plot. AGB was calculated for each census (Mg ha^{-1}), and
 153 AGWP for each census interval ($\text{Mg ha}^{-1} \text{ yr}^{-1}$), and then a time-weighted mean was taken
 154 to give one value per plot. Tree biomass was estimated based on the diameter (D), wood
 155 density (ρ) and height (H), using the pantropical equation developed by Chave et al.
 156 (2014):

$$157 \text{AGB}_{\text{trees}} = 0.0673 \times (\rho D^2 H)^{0.976}$$

158 Species wood density was obtained from the global wood-density database (Chave
 159 et al., 2009; Zanne et al., 2009). A 3-parameter regional height-diameter Weibull equation
 160 was adjusted using the BiomasaFP R package (Lopez-Gonzalez et al., 2015) to estimate
 161 heights.

162 The biomass of palms (Arecaceae family) was calculated from the allometric
 163 equation developed by Goodman et al. (2013), based on diameter (D):

$$164 \ln(\text{AGB}_{\text{palm}}) = -3.3488 + 2.7483 \cdot \ln(D)$$

165 Palm trees were excluded from the productivity calculations as variations in
 166 diameter are closely related to fluctuation in water content, and most growth of palm trees
 167 occurs through increases in height (Tomlinson, 1990; Stahl *et al.*, 2010).

168 AGWP was calculated from the sum of biomass growth of surviving trees and
 169 trees that recruited. Biomass-productivity estimates are affected by several factors,
 170 including census length, unobserved growth, recruitment, and mortality within each
 171 census interval; we corrected these using the method proposed by Talbot et al., (2014).

172 To assess biomass mortality, we first estimated the above-ground woody loss over
 173 time, in units of $\text{Mg hr}^{-1} \text{ yr}^{-1}$. We also estimated the ‘biomass mortality rate’, as
 174 $\text{AGB}_{\text{mortality}}/\text{AGB}$, in units of $\text{hr}^{-1} \text{ yr}^{-1}$. This standardization was performed in order to be
 175 able to compare the proportional rate of biomass loss among plots with different standing
 176 biomass stock.

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3 177 We also calculated stem mortality, measured as mean annual mortality rate (λ) as:
4 $\lambda = \frac{[\ln(N_0) - \ln(N_s)]}{t}$, where N_0 and N_s are the number of stems counted of the initial
5 178 population, and the number of stems surviving to time t , respectively (Sheil *et al.*, 1995).
6 179 For each site we also calculated annual recruitment rates (μ) as:
7 180 $\mu = [\ln(N_f/N_s)]/t$, where N_f is the final number of stems, N_s is the original number of
8 181 stems surviving to final inventory and t is the number of years between inventories.
9 182 Mortality and recruitment rates were calculated for each census interval (% yr⁻¹), and
10 183 then a time-weighted mean based on the census-interval lengths was taken to give one
11 184 value per plot. With these results we derived the stem turnover rate, defined as the
12 185 mean of recruitment and mortality (Phillips *et al.*, 1994). The length of the census
13 186 intervals can affect rate estimates, with long intervals between censuses more likely to
14 187 underestimate rates due to unobserved mortality and recruitment (Lewis *et al.*, 2004).
15 188 To account for potential impacts of varying census intervals on the rate estimates, we
16 189 applied the correction factor proposed by Lewis *et al.* (2004).
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192 Environmental data

193 We modelled forest structure and dynamics as a function of climatic, soil-water,
194 and edaphic properties. Maximum cumulative water deficit (MCWD) was used as an
195 inverse proxy to the climatic water supply, water table depth (WTD) was used as a proxy
196 for local soil-water supply, and soil texture was used as a proxy for soil-water-retention
197 capacity. Maximum temperature and soil fertility were also included in the multiple
198 models in order to control for their known effects on Amazon ecosystem functions (Baker
199 *et al.*, 2003; Malhi *et al.*, 2004; Quesada *et al.*, 2012; Sullivan *et al.*, 2020), thus making
200 it possible to assess the role of hydrological variables, our focus in this manuscript, more
201 clearly.

202 We calculated MCWD based on the long-term average of annual MCWD of each
203 plot, from 1971 to 2019, thus reflecting the climatic conditions experienced by each plot
204 over time and corresponding to the time window of our dataset. MCWD corresponded to
205 the maximum value of the monthly accumulated climatic water deficit reached in each
206 location, i.e., the difference between precipitation and evapotranspiration within each
207 hydrological year (Esquivel-Muelbert *et al.*, 2019). This metric represents the sum of
208 water-deficit values over consecutive months when evapotranspiration is greater than
209 precipitation (Aragão *et al.*, 2007). Precipitation data were extracted from the
210 TerraClimate data set (Abatzoglou *et al.*, 2018), at ~4 km (1/24th degree) spatial
211 resolution from 1971 to 2019. Monthly evapotranspiration was assumed as fixed at 100
212 mm month⁻¹, considering that Amazonian forest canopies have a nearly constant
213 evapotranspiration rate (Shuttleworth, 1988; Rocha *et al.*, 2004).

214 Water table depth was extracted from a map developed for the entire Amazon (Fan
215 *et al.*, 2013; Fan & Miguez-Macho, 2010), at ~270 m spatial resolution, based on model
216 simulation constrained by over 1,000,000 direct well measurements from government
217 archives and publications. We extracted water table depth values for the geographic
218 coordinates for each plot and did not interpolate values of the surrounding pixels to avoid
219 degrading the already coarse resolution of the WTD data. Clay-content data were obtained

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3 220 from the SoilGrids database, at 250 m resolution (Hengl *et al.*, 2017). As a proxy for soil
4 221 fertility, we used the soil concentration of exchangeable base cations (Ca + Mg + K),
5 222 extracted from the Amazon-wide model of Zuquim *et al.* (2019), since this is the best
6 223 continuous layer of soil fertility available for the entire study area. SoilGrids has a layer
7 224 of cation exchange capacity (CEC) (Hengl *et al.*, 2017), but the correlation of measured
8 225 cations and the mapped CEC has been shown to be low, as CEC includes the
9 226 concentration of aluminium, which is not a nutrient (Moulatlet *et al.*, 2017). Although
10 227 phosphorus is widely considered as a key limiting nutrient for growth in tropical forests,
11 228 this variable is not available for all plots or as a continuous estimated layer. However, the
12 229 availability of exchangeable cations tends to be correlated well to the amount of
13 230 phosphorus (Quesada *et al.*, 2010, 2012) and also predicts forest growth well (Quesada *et*
14 231 *al.*, 2012). We estimated long-term maximum temperature, using a dataset from
15 232 TerraClimate, at ~4 km (1/24th degree) spatial resolution from 1971 to 2019.
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22 234 **Data analyses**

23 235 To achieve our goal of understanding the hydrological effects on forest
24 236 functioning, we used a spatial analysis of the influence of our proxies on the water
25 237 conditions of each site (water table depth, MCWD and soil texture), including their
26 238 potential interactions, on the metrics of forest structure and dynamics (biomass stock,
27 239 productivity and mortality; stem mortality, recruitment and turnover). To test these
28 240 effects, we ran multiple linear models considering in addition to hydrological variables
29 241 (MCWD, WTD and soil texture), soil fertility and air temperature, since they are
30 242 recognized as important determinants of structure and dynamics of Amazon forests. Our
31 243 models included interactions because we expected the effect of water table depth on the
32 244 forest dynamics to depend on the levels of water-deficit (MCWD) and soil texture (Table
33 245 S2). Before running the models, we tested for multicollinearity among predictors. The
34 246 Variance Inflation Factors (VIF) were estimated and only low multicollinearity was
35 247 detected (VIF < 5, Table S3). To detect if spatial aggregation of plots (which could induce
36 248 autocorrelation) interfered in our results, we ran generalized linear mixed models
37 249 (GLMM) with and without a random factor representing the clusters of plots within 50
38 250 km of each other, checked the model summaries and compared their Akaike's information
39 251 criterion (Table S4). Adding the random factor improved the models (smaller AIC
40 252 values), but did not qualitatively change the results, so we present here the models without
41 253 the random factor.

42 254 We weighted the plots in regression analyses when testing the effects of the
43 255 environmental predictors on forest dynamics and structure according to the plot size and
44 256 monitoring time, as larger plots and those monitored for longer periods are expected to
45 257 provide better estimates of local, long-term forest properties. To achieve this, following
46 258 Lewis *et al.* (2009), we plotted the residuals from linear models against plot area and
47 259 monitoring period, and selected the root transformations of plot area and monitoring
48 260 period that removed the nonlinear patterns in the residuals when applied as a weight.
49 261 These empirically-determined weights were: AGWP, $\text{Area}^{1/2}$; AGB, $\text{Area}^{1/3}$; AGB
50 262 mortality, $\text{Area}^{1/2} + \text{Monitoring length}^{1/4} - 1$; Mortality rate, $\text{Area}^{1/2} + \text{Monitoring length}^{1/3}$
51 263 $- 1$; Recruitment rate, $\text{Area}^{1/5}$; Stem turnover, $\text{Area}^{1/3} + \text{Monitoring length}^{1/4} - 1$.

264 In order to investigate in more detail the relationships between the response
 265 variables (AGB, AGWP, etc) and hydrological variables, we used loess (locally-
 266 weighted) regressions. We used partial-dependence plots to visualize the shape of the
 267 relationships between response and predictor variables. To visualize interactions, climate
 268 and soil texture were divided in three classes based on the standard deviation around the
 269 mean of each of these variables.

270 To describe the climate and water table effects, we used the following data
 271 subdivisions of WTD and MCWD, made to provide an idea of the variation in forest
 272 structure and dynamics among the extremes of these gradients. We recognize that in
 273 nature the forest response is not abrupt or categorized, and the continuous responses are
 274 shown in the regression models. Shallower and deeper water tables were defined using a
 275 5-m depth threshold. We chose this division because groundwater ≤ 5 m depth is where
 276 most roots are potentially in direct contact with the groundwater or the capillary fringe
 277 (Fan & Miguez-Macho, 2010; Fan et al., 2017). We also ran boosted regression trees for
 278 the relationship between WTD and all response variables (Fig. S1) to check if this value
 279 was supported by the data. Wet (MCWD > -160 mm) and dry (MCWD < -160 mm)
 280 forests were divided based on the MCWD average in our data set (see the histograms in
 281 Fig. S2). To test whether there was a significant statistical difference in forest structure
 282 and dynamics between the shallow and deep water table subgroups, or dry and wet
 283 climates, we used unpaired *Welch two-sample t-tests* for unequal sized samples.

284 All analyses were conducted in R version 3.6.1 software. We used the *BiomasaFP*
 285 R package (Lopez-Gonzalez et al. 2015) to calculate AGB, AGWP and AGB mortality.
 286 Multicollinearity was tested using the package *performance* (Lüdecke et al., 2021);
 287 LOESS regressions were calculated with package *ggplot2* (Wickham, 2011); multiple
 288 linear regressions with package *car* (Fox et al., 2018); the interaction plots with the
 289 package *interactions* (Bauer & Curran, 2005); and boosted regression trees with the
 290 packages *rpart* (Milborrow, 2019) and *gmb* (Greenwell et al., 2019).

291 292 **Results**

293 294 *How does the structure and dynamics of Amazonian forest vary with the water table* 295 *depth and climatic water deficit?*

296 Based on the simple relationships between WTD and forest dynamics and
 297 biomass, shallower water tables (depth < 5 m) on average decreased the forest biomass
 298 productivity ($t = -5.62$; $df = 342$; $p < 0.01$) and biomass stocks ($t = -6.28$; $df = 342$; $p <$
 299 0.01) of Amazon forests (Figures 1a and 1b, respectively). Shallower water table forests
 300 had on average 18% lower biomass productivity ($4.5 \text{ Mg ha}^{-1} \text{ yr}^{-1}$) and 23% lower
 301 biomass stock (234.6 Mg ha^{-1}) than those on deeper water tables ($5.5 \text{ Mg ha}^{-1} \text{ yr}^{-1}$ and
 302 306.9 Mg ha^{-1} , respectively). Also, based on the simple relationships between MCWD
 303 and forest dynamics and biomass, climatically drier sites (MCWD < -160 mm) had 21%
 304 lower biomass productivity ($4.5 \text{ Mg ha}^{-1} \text{ yr}^{-1}$; $t = -7.67$; $df = 342$; $p < 0.01$) and 24%
 305 lower biomass stock (240.2 Mg ha^{-1} ; $t = -7.01$; $df = 342$; $p < 0.01$) than those in wetter
 306 climates ($5.7 \text{ Mg ha}^{-1} \text{ yr}^{-1}$, 314.3 Mg ha^{-1}) (Figures 2a and 2b, respectively). Thus, the

negative direct effects of climatic-water deficit (MCWD) were only slightly stronger than the negative effects of excess soil water associated with shallow water tables.

Stem mortality rate (2.6% yr⁻¹, Fig. 1c; $t = 3.40$; $df = 342$; $p < 0.01$) and stem turnover (2.4% yr⁻¹, Fig. 1d; $t = 3.62$; $df = 342$; $p < 0.01$) were higher in shallower water table forests than in those with deeper water tables (2.1% yr⁻¹ and 2.0 % yr⁻¹, respectively). Conversely, stem mortality rate (2.8% yr⁻¹; $t = 7.21$; $df = 342$; $p < 0.01$), recruitment rate (2.3% yr⁻¹; $t = 3.62$; $df = 342$; $p < 0.01$) and stem turnover (2.5% yr⁻¹; $t = 6.24$; $df = 342$; $p < 0.01$) were higher in drier than in wet climates (1.9% yr⁻¹, 1.8% yr⁻¹ and 1.9% yr⁻¹, respectively) (Figures 2d, 2e and 2f).

The greatest biomass stocks were found in the eastern and northeastern portions of the Amazon, which combine, on average, intermediate MCWD, deep water table and clayey soils (Figures 3c, 3e and 3a, respectively). Biomass productivity was higher in the western portion of the basin and on the Guiana shield, associated with wetter climates (Fig. 3f). Within the Guiana shield, higher productivity was associated with deep water tables (Fig. 3d). Beyond these trends already captured by regression analyses, the maps depict the large local variation (i.e., within sites) of biomass stock and productivity, largely due to intra-site (between plot) variation in topography and consequently in WTD.

How does water table depth interact with climatic-water deficit and soil texture to influence Amazonian forest biomass?

A significant interaction between WTD and MCWD was detected only for AGWP. The best model (Table S2) fit of the interaction divides MCWD data into three groups, based on the standard deviation around the mean, following a gradient from wetter (blue line) to drier climates (red line). Shallow water table forests had lower AGWP than deeper water table forests when under drier climates, with this difference decreasing in wet climates (Fig. 4). The very low biomass productivity of some plots (< 2 Mg ha⁻¹ yr⁻¹) is related to vegetation structure, as in these sites most trees are very thin and therefore have lower productivity. Additional analysis showed that excluding these plots does not change the Amazon-wide pattern of the interactive effects of water table depth and climate in productivity (Fig. S4).

Despite the average negative effect of shallow water table on forest productivity within dry climates, the more complex interactions between soil texture, MCWD and water table depth suggest a contribution of soil drainage to forest functioning (Fig. 5). These interactions show that forest productivity was lower in shallower water table conditions in dry climates when the soil is less clayey, as compared to deeper water table conditions in the same climate (red line, Fig. 5a). However, when the soil was more clayey dry-climate forests with shallower water table had greater productivity than their climatic equivalents on deeper water tables (red line, Fig. 5c). The data coverage of some combinations of climate, water table and soil texture were low (especially clayey soils under dry climates and shallow water table), what may limit the interpretation of this result. We also note some non-linear trends in wet climates and sandier soils, where AGWP in low where the water table is very shallow (< 2 m) but increases to reach a peak in the range of 2 to 8 m depth (Fig. 5a).

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3 351 The variation in AGB, mortality and turnover rates was related to the interaction
4 352 between MCWD and clay content, with less-clayey and climatically drier sites having
5 353 lower AGB, whereas mortality and turnover are higher in those sites (Fig. S3).
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7 354
8 355 *The effects of other factors*

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10 356 The well-known effects of soil fertility on forest dynamics were detected in the
11 357 multiple linear models. Above-ground woody productivity and biomass mortality rate
12 358 increased with soil fertility (Table S2). Soil fertility also affects mortality, recruitment
13 359 rates and stem turnover, which were higher on more fertile soils (Table S2). The effects
14 360 of maximum temperature in the multiple-regression models were detected only for
15 361 biomass stock, with sites with higher maximum temperature having lower biomass
16 362 stock (Table S2).
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19 364
20 364 **Discussion**

21 365 Our study demonstrates for the first time the large-scale effects of water table
22 366 depth on the structure and dynamics of the Amazon forests, based on a unique
23 367 combination of ground-plot data and water table depth modelling. Amazon forests with
24 368 shallower water tables had, on average, lower biomass productivity, lower biomass
25 369 stock, higher stem mortality and higher turnover. Amazon forests with drier climates
26 370 had, on average, lower biomass productivity, lower biomass stock, higher stem
27 371 mortality and higher turnover. This indicates that an excess of water, as well as a deficit,
28 372 has a detrimental effect on forest functioning.
29 373

30 373 Our results show that the landscape-scale patterns of Amazonian forest structure
31 374 and dynamics are affected by groundwater and its interaction with climatic conditions.
32 375 Therefore, WTD is an especially important environmental variable to be considered in
33 376 modelling the effects of climate change on vegetation (Fan et al., 2013; Fan & Miguez-
34 377 Macho, 2011; Roebroek et al., 2020; Taylor et al., 2013).
35 378

36 378
37 379 *Effects of water table depth and the long-term average climatic water deficit on the*
38 380 *structure and dynamics of Amazon forests*

39 381 We hypothesized that shallow water tables impose constraints on plant
40 382 development under generally wet climates of Amazonia, through excess soil water and
41 383 consequent oxygen limitation. Our results support this hypothesis since, on average,
42 384 sites with shallow water table tended to have lower biomass productivity (Fig. 1a).
43 385 However, there is high variability in AGWP, with some sites having high biomass
44 386 productivity despite the shallow water table. Therefore, it is important to explore the
45 387 mechanisms which may lead to the two extremes of low and high biomass productivity
46 388 in shallow water tables. To help understand the lower productivity, we must review the
47 389 response of soils and plants to waterlogging, the condition prevailing to various degrees
48 390 – seasonal to permanent - in many of the shallow water table sites. When soils are
49 391 waterlogged, most of the soil spaces are occupied with water, and the metabolism of
50 392 roots and microorganisms quickly consumes the available oxygen and produces carbon
51 393 dioxide. As oxygen is depleted, roots and aerobic microorganisms lose most of their
52 394 capacity to produce energy through aerobic respiration (Gibbs & Greenway, 2003). In

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3 395 this case, the major pathway to energy production is alcoholic fermentation, which has a
4 396 much lower yield (2 mols ATP per glyucose molecule) than respiration (36 ATP), and
5 397 thus severely limits plant growth (Setter & Belford, 1990; Kreuzwieser & Rennenberg,
6 398 2014). Low oxygen levels also reduce root permeability (North et al., 2004; Vandeleur
7 399 et al., 2005), generating a cascade of responses that reduce stomatal conductance and
8 400 thus limit photosynthesis (Lopez & Kursar, 1999, 2003; Parent et al., 2008; Pezeshki,
9 401 2001). Low photosynthetic activity and consequent low growth is well documented in
10 402 periodically flooded forests (Parolin, 2000; Waldhoff et al., 1998), although this a more
11 403 extreme condition than the soil waterlogging examined here. Given the various
12 404 deleterious effects of excess water on plant metabolism and physiology, most tree
13 405 growth occurs during the windows when water table levels decrease and anoxia is
14 406 relieved, mostly in the dry season. Such growth windows have been described in
15 407 flooded areas, where the largest diameter growth occurs in the non-flooded period
16 408 (Schöngart et al., 2002; 2004). Therefore, the period with environmental conditions
17 409 suitable for growth is shorter in shallow water table, and therefore, on average, biomass
18 410 productivity is lower in these locations than in deep water table (but, see next section,
19 411 these patterns change when combined with climate).

20 412 For vegetation dynamics, we found higher mortality and stem turnover in
21 413 shallow water table sites, as we had hypothesized. Poorly drained sites have higher
22 414 mortality rates due to weak plant anchorage caused by the groundwater layer that
23 415 prevents deep root growth, and this is also generally associated with loose soil texture
24 416 (Gale & Barfod, 1999; Toledo et al., 2011). This low adherence to the soil increases the
25 417 tree's susceptibility to uprooting (Madelaine *et al.*, 2007). Forests with waterlogged soils
26 418 have higher proportions of uprooting as the tree mode of death, whereas forests on well-
27 419 drained soils have higher proportions of trees dying standing (Gale & Hall, 2001). The
28 420 effects of excess water on forest structure and dynamics are well described in the
29 421 literature for floodplain forests (Simone et al., 2003; Godoy et al., 1999; Parolin et al.,
30 422 2004; Piedade et al., 2013; Schöngart et al., 2004), but little is known about the effects
31 423 of shallow water tables on *terra-firme* forests. In local studies, paired comparisons of
32 424 shallow and deep water tables within the same wet macroclimate have shown similar
33 425 patterns of lower biomass productivity and basal area (Castilho et al., 2006; Castilho et
34 426 al., 2010; Ferry et al., 2010), higher tree mortality (Ferry et al., 2010; Toledo et al.,
35 427 2011) and recruitment rates (Ferry *et al.*, 2010) in seasonally waterlogged shallower
36 428 water table forests than on deeper water table hilltops, as we now find here to occur at
37 429 an Amazon-wide scale. In a global analysis, based on remote sensing data, water table
38 430 depth was associated with forest productivity, stimulating or hindering vegetation
39 431 growth depending on climate (Roebroek *et al.*, 2020), and our large-scale on-the-ground
40 432 assessment of this effect supports those results for the Amazonian forests, but here with
41 433 above-ground wood productivity data.

42 434
43 435 *Interactions among water table depth, climatic water deficit and soil properties*
44 436 *influence Amazon-forest structure and function*

45 437 Our results also agree with a well-described average effect of increasing climate
46 438 seasonality lowering productivity and biomass stock, and increasing stem turnover

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3 439 (Álvarez-Dávila *et al.*, 2017; Malhi *et al.*, 2004, 2006; Saatchi *et al.*, 2007; Vilanova *et*
4 440 *al.*, 2018). The effects of soil fertility were in line with those described in the literature,
5 441 in which forest dynamics and especially above-ground woody productivity were greater
6 442 on more fertile soils (Baker *et al.*, 2003; Malhi *et al.*, 2004; Quesada *et al.*, 2012; Banin
7 443 *et al.*, 2014; Esquivel-Muelbert *et al.*, 2020). However, neither soil properties, nor
8 444 climatic or groundwater conditions alone fully explain the distribution of biomass and
9 445 vegetation growth in our study or worldwide (Baraloto *et al.*, 2011; Quesada *et al.*,
10 446 2012; Fan, 2015).

11 447 We hypothesized that an interaction of these factors would provide a better
12 448 description of the vegetation patterns, with drier regions with shallow water table
13 449 having higher biomass productivity, while in wetter climates shallow water tables
14 450 would result in excess water and lower productivity, however, this is not what we
15 451 found. The combination of shallow water table and dry climate resulted in lower
16 452 biomass productivity. This outcome may result from an aspect of the water availability
17 453 that was not accounted in this study - the temporal fluctuation of the water table. The
18 454 available WTD product gives what is expected to be the average water table depth of
19 455 each pixel, but there may be varying degrees of temporal fluctuation modulated by
20 456 climate, topographic and geomorphological conditions (Costa *et al.* 2022). In the drier
21 457 climates, the seasonal fluctuation of the water table tends to be higher (Miguez-Macho
22 458 & Fan, 2012; Costa *et al.* 2022), so plants may be exposed to stresses of both water
23 459 deficit in the dry season and water excess in the wet season, giving rise to the worst
24 460 scenario for growth. In the wet season the rise in the water table may lead to anoxic
25 461 stress. In the dry season, when the water table level drops, the shallow plant root
26 462 systems characteristic of these environments may not access the groundwater and go
27 463 through water deficit stress, also limiting the biomass accumulation.

28 464 Also contrary to our general hypothesis, the limitation of biomass productivity
29 465 given by the combination of wet climate and shallow water table occurred only where
30 466 the water table is very shallow (< 2 m deep), which is where most fine roots tend to be
31 467 (Jackson *et al.*, 1996). This seems to restrict the pure anoxic limitation of productivity to
32 468 a smaller range of very wet conditions than previously hypothesized here. Still under
33 469 wet climates, we see high biomass production in the intermediate shallow water table
34 470 (2-5 m, Fig. 5a) that may be a consequence of an interaction of the tree functional traits
35 471 typically selected under wet environments (lower wood density, higher xylem vessel
36 472 diameter and higher specific leaf area, review in Costa *et al.* 2022, aligned with faster
37 473 resource acquisition and growth) and the potentially moist, instead of anoxic conditions,
38 474 during a large period of the year. The number of plots within each combination of
39 475 climate, water table and soil conditions is relatively low here, and there is a clear need
40 476 for more work to improve the evaluation of these potential non-linearities in the
41 477 response of forest productivity to the determinants of water availability.

42 478 A full accounting of the factors affecting soil moisture also requires consideration
43 479 of soil properties, especially soil texture (Richter & Babbar, 1991; Quesada *et al.*,
44 480 2012). In general, the ecological effects of the soil water regime will depend on the
45 481 degree of soil saturation in the wet months, the degree and frequency of water deficit
46 482 periods, the water-holding capacity of the soil, and the root distribution in the soil

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3 483 (Franco & Dezzio, 1994). By having higher aggregation particles, clayey soils have
4 484 better water-holding capacity (Richter & Babbar, 1991), therefore, clay soils should
5 485 increase the time interval between precipitation inputs and groundwater recharge, while
6 486 predominantly sandy soils should have faster groundwater level responses to
7 487 precipitation. Our results suggest a contribution of clayey texture in increasing
8 488 productivity in dry climates with shallow water table (Fig. 5 C). However, here too the
9 489 dataset lacks complete coverage of the relevant environmental combinations, limiting
10 490 our conclusions.

11 491 *Limitations of this study*

12 492
13 493 While this and other work points to a key role for water table depth and
14 494 consequent soil hydrology in shaping the structure and composition of tropical forests
15 495 (e.g. Damasco et al., 2013; Jirka et al., 2007; Moulatlet et al., 2014; Schiatti et al.,
16 496 2013; Sousa et al., 2020; and see a review in Costa et al. 2022), precise measurement of
17 497 water table depth and its fluctuation is still limited due to the challenge of installation of
18 498 equipment and periodic monitoring in the field. The alternative for large-scale analytical
19 499 studies like these is to use water table depth models, such as the Fan et al. (2013) model
20 500 used here. These, however, come with limitations as they condense the full micro-
21 501 spatial variation of hydrology in a relatively coarse spatial resolution (here ~ 270 m). A
22 502 further difficulty is that vegetation-monitoring plots may not be designed to detect
23 503 variation in hydrological environments, such that varying hydrological conditions may
24 504 occur within the same plot (see Magnusson et al., 2005 for a design that minimizes this
25 505 problem). These imprecisions probably limit our capacity to detect the local effects of
26 506 water table depth on forest functioning, so that effects in nature may eventually prove to
27 507 be even stronger than shown here.

28 508 Also, while we could account for the major trends, there was large variation in
29 509 biomass-productivity, and some shallow water table plots had high biomass productivity
30 510 ($> 5 \text{ Mg ha}^{-1} \text{ yr}^{-1}$). Such unexpected variation suggests we have still not accounted for
31 511 all the key variables and processes, with additional variation related to species
32 512 composition and functional traits being obvious candidates. Species composition and
33 513 dominant functional traits differ across the hydrological environments within the same
34 514 climate (Schiatti et al., 2013; Cosme et al., 2017), but it is not known whether they are
35 515 filtered similarly across soil hydrology under different macroclimates, or soil vs.
36 516 macroclimate interactions that could potentially change the responses of shallow water
37 517 table forests under different climates. This is an important subject to address in future
38 518 studies because it could suggest ways to mitigate carbon losses.

39 519 *Final considerations*

40 520
41 521 The Amazon hydrological cycle is already changing due to climate change and
42 522 these are projected to intensify in the future (Gloor et al., 2015). To predict ecological
43 523 impacts and mitigate their effects on the Amazon forests, it is essential to assess the
44 524 functioning and ecology of forests at the ecosystem level. Improved understanding of
45 525 the effects of local hydrology on forest functioning is also key to plan the conservation
46 526 and management on the scales at which landscapes are normally exploited. Our results

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3 527 indicate the need to protect some critical environments with shallow water table forests
4 528 as buffers against the negative effects of climate change. They also provide indications
5 529 of critical missing factors when modelling the biomass dynamics of Amazonia.

7 530 By analyzing long-term forest monitoring records from across the 6 million km²
8 531 expanse of lowland Amazonia, we find a significant, large-scale control of forest
9 532 structure and dynamics by water table depth. Both water excess and water deficit hinder
10 533 vegetation development. Above-ground productivity is suppressed, tree mortality
11 534 increased and thus biomass stocks are reduced in shallow water table forests. These key
12 535 effects of water table depth have typically been neglected in large-scale studies (e.g.
13 536 Malhi et al., 2015, 2006; Saatchi et al., 2007), but must be considered in global
14 537 environmental modelling to better understand the relative contribution of the key drivers
15 538 of Amazon forest structure and dynamics and the ecosystem functions they provide.
16 539

20 540 **Data Availability Statement**

22 541 Data for the analyses are available as a ForestPlots.net data package at
23 542 http://doi.org/10.5521/forestplots.net/2022_2

References

- A., R., Freeze & Cherry, J.A. (1979) *Groundwater*, Englewood Cliffs, N.J: Prentice-Hall.
- Abatzoglou, J.T., Dobrowski, S.Z., Parks, S.A. & Hegewisch, K.C. (2018) TerraClimate, a high-resolution global dataset of monthly climate and climatic water balance from 1958-2015. *Scientific Data*, **5**.
- Álvarez-Dávila, E., Cayuela, L., González-Caro, S., Aldana, A.M., Stevenson, P.R., Phillips, O., Cogollo, Á., Peñuela, M.C., Von Hildebrand, P., Jiménez, E., Melo, O., Londoño-Vega, A.C., Mendoza, I., Velásquez, O., Fernández, F., Serna, M., Velázquez-Rua, C., Benítez, D. & Rey-Benayas, J.M. (2017) Forest biomass density across large climate gradients in northern South America is related to water availability but not with temperature. *PLoS ONE*, **12**, 1–16.
- Aragão, L.E.O.C., Malhi, Y., Roman-Cuesta, R.M., Saatchi, S., Anderson, L.O. & Shimabukuro, Y.E. (2007) Spatial patterns and fire response of recent Amazonian droughts. *Geophysical Research Letters*, **34**, 1–5.
- Baker, T.R., Burslem, D.F.R.P. & Swaine, M.D. (2003) Associations between tree growth, soil fertility and water availability at local and regional scales in Ghanaian tropical rain forest. *Journal of Tropical Ecology*, **19**, 109–125.
- Banin, L., Lewis, S.L., Lopez-Gonzalez, G., Baker, T.R., Quesada, C.A., Chao, K.J., Burslem, D.F.R.P., Nilus, R., Abu Salim, K., Keeling, H.C., Tan, S., Davies, S.J., Monteagudo Mendoza, A., Vásquez, R., Lloyd, J., Neill, D.A., Pitman, N. & Phillips, O.L. (2014) Tropical forest wood production: A cross-continental comparison. *Journal of Ecology*, **102**, 1025–1037.
- Baraloto, C., Rabaud, S., Molto, Q., Blanc, L., Fortunel, C., Hérault, B., Dávila, N., Mesones, I., Rios, M., Valderrama, E. & Fine, P.V.A. (2011) Disentangling stand and environmental correlates of aboveground biomass in Amazonian forests. *Global Change Biology*, **17**, 2677–2688.
- Bauer, D.J. & Curran, P.J. (2005) Probing Interactions in Fixed and Multilevel Regression: Inferential and Graphical Techniques. *Multivariate Behavioral Research*, **40**, 373–400.
- Brienen, R.J.W., Phillips, O.L., Feldpausch, T.R., Gloor, E., Baker, T.R., Lloyd, J., Lopez-Gonzalez, G., Monteagudo-Mendoza, A., Malhi, Y., Lewis, S.L., Vásquez Martínez, R., Alexiades, M., Álvarez Dávila, E., Alvarez-Loayza, P., Andrade, A., Aragão, L.E.O.C., Araujo-Murakami, A., Arets, E.J.M.M., Arroyo, L., Aymard C., G.A., Bánki, O.S., Baraloto, C., Barroso, J., Bonal, D., Boot, R.G.A., Camargo, J.L.C., Castilho, C. V., Chama, V., Chao, K.J., Chave, J., Comiskey, J.A., Cornejo Valverde, F., da Costa, L., de Oliveira, E.A., Di Fiore, A., Erwin, T.L., Fauset, S., Forsthofer, M., Galbraith, D.R., Grahame, E.S., Groot, N., Hérault, B., Higuchi, N., Honorio Coronado, E.N., Keeling, H., Killeen, T.J., Laurance, W.F., Laurance, S., Licona, J., Magnussen, W.E., Marimon, B.S., Marimon-Junior, B.H., Mendoza, C., Neill, D.A., Nogueira, E.M., Núñez, P., Pallqui Camacho, N.C., Parada, A., Pardo-Molina, G., Peacock, J., Peña-Claros, M., Pickavance, G.C., Pitman, N.C.A., Poorter, L., Prieto, A., Quesada, C.A., Ramírez, F., Ramírez-Angulo, H., Restrepo, Z., Roopsind, A., Rudas, A., Salomão, R.P., Schwarz, M., Silva, N., Silva-Espejo, J.E., Silveira, M., Stropp, J., Talbot, J., ter Steege, H., Teran-Aguilar, J., Terborgh, J., Thomas-Caesar, R., Toledo, M., Torello-Raventos, M., Umetsu, R.K., van der Heijden, G.M.F., van der Hout, P., Guimarães Vieira, I.C., Vieira, S.A., Vilanova, E., Vos, V.A. & Zagt, R.J. (2015) Long-term decline of the Amazon carbon sink. *Nature*, **519**, 344–348.
- Canadell, J., Jackson, R.B., Ehleringer, J.B., Mooney, H.A., Sala, O.E. & Schulze, E.-D.

- (1996) Maximum rooting depth of vegetation types at the global scale. *Oecologia*, **108**, 583–595.
- Castilho, C. V., Magnusson, W.E., de Araújo, R.N.O. & Luizão, F.J. (2010) Short-Term Temporal Changes in Tree Live Biomass in a Central Amazonian Forest, Brazil. *Biotropica*, **42**, 95–103.
- Castilho, C. V., Magnusson, W.E., de Araújo, R.N.O., Luizão, R.C.C., Luizão, F.J., Lima, A.P. & Higuchi, N. (2006) Variation in aboveground tree live biomass in a central Amazonian Forest: Effects of soil and topography. *Forest Ecology and Management*, **234**, 85–96.
- Chave, J., Coomes, D., Jansen, S., Lewis, S.L., Swenson, N.G. & Zanne, A.E. (2009) Towards a worldwide wood economics spectrum. *Ecology Letters*, **12**, 351–366.
- Chave, J., Réjou-Méchain, M., Búrquez, A., Chidumayo, E., Colgan, M.S., Delitti, W.B.C., Duque, A., Eid, T., Fearnside, P.M., Goodman, R.C., Henry, M., Martínez-Yrizar, A., Mugasha, W.A., Muller-Landau, H.C., Mencuccini, M., Nelson, B.W., Ngomanda, A., Nogueira, E.M., Ortiz-Malavassi, E., Pélissier, R., Ploton, P., Ryan, C.M., Saldarriaga, J.G. & Vieilledent, G. (2014) Improved allometric models to estimate the aboveground biomass of tropical trees. *Global Change Biology*, **20**, 3177–3190.
- Chitra-Tarak, R., Xu, C., Aguilar, S., Anderson-Teixeira, K.J., Chambers, J., Detto, M., Faybishenko, B., Fisher, R.A., Knox, R.G., Koven, C.D., Kueppers, L.M., Kunert, N., Kupers, S.J., McDowell, N.G., Newman, B.D., Paton, S.R., Pérez, R., Ruiz, L., Sack, L., Warren, J.M., Wolfe, B.T., Wright, C., Wright, S.J., Zailaa, J. & McMahon, S.M. (2021) Hydraulically-vulnerable trees survive on deep-water access during droughts in a tropical forest. *New Phytologist*, **231**, 1798–1813.
- Cosme, L.H.M., Schiatti, J., Costa, F.R.C. & Oliveira, R.S. (2017) The importance of hydraulic architecture to the distribution patterns of trees in a central Amazonian forest. *New Phytologist*, **215**.
- Costa, A. da, Albuquerque, J.A., Costa, A. da, Pértile, P. & Silva, F.R. da (2013) Water retention and availability in soils of the State of Santa Catarina-Brazil: effect of textural classes, soil classes and lithology. *Revista Brasileira de Ciência do Solo*, **37**, 1535–1548.
- Costa, F.R.C., Schiatti, J., Stark, S.C. & Smith, M.N. (2022) The other side of tropical forest drought: do shallow water table regions of Amazonia act as large-scale hydrological refugia from drought? *New Phytologist*.
- Damasco, G., Vicentini, A., Castilho, C. V., Pimentel, T.P. & Nascimento, H.E.M. (2013) Disentangling the role of edaphic variability, flooding regime and topography of Amazonian white-sand vegetation. *Journal of Vegetation Science*, **24**, 384–394.
- Ellison, D., Morris, C.E., Locatelli, B., Sheil, D., Cohen, J., Murdiyarso, D., Gutierrez, V., Noordwijk, M. van, Creed, I.F., Pokorny, J., Gaveau, D., Spracklen, D. V., Tobella, A.B., Ilstedt, U., Teuling, A.J., Gebrehiwot, S.G., Sands, D.C., Muys, B., Verbist, B., Springgay, E., Sugandi, Y. & Sullivan, C.A. (2017) Trees, forests and water: Cool insights for a hot world. *Global Environmental Change*, **43**, 51–61.
- Esquivel-Muelbert, A., Phillips, O.L., Brien, R.J.W., Fauset, S., Sullivan, M.J.P., Baker, T.R., Chao, K.-J., Feldpausch, T.R., Gloor, E., Higuchi, N., Houwing-Duistermaat, J., Lloyd, J., Liu, H., Malhi, Y., Marimon, B., Marimon Junior, B.H., Monteagudo-Mendoza, A., Poorter, L., Silveira, M., Torre, E.V., Dávila, E.A., del Aguila Pasquel, J., Almeida, E., Loayza, P.A., Andrade, A., Aragão, L.E.O.C., Araujo-Murakami, A., Arets, E., Arroyo, L., Aymard C., G.A., Baisie, M., Baraloto, C., Camargo, P.B., Barroso, J., Blanc, L., Bonal, D., Bongers, F., Boot,

- R., Brown, F., Burban, B., Camargo, J.L., Castro, W., Moscoso, V.C., Chave, J., Comiskey, J., Valverde, F.C., da Costa, A.L., Cardozo, N.D., Di Fiore, A., Dourdain, A., Erwin, T., Llampazo, G.F., Vieira, I.C.G., Herrera, R., Honorio Coronado, E., Huamantupa-Chuquimaco, I., Jimenez-Rojas, E., Killeen, T., Laurance, S., Laurance, W., Levesley, A., Lewis, S.L., Ladvoat, K.L.L.M., Lopez-Gonzalez, G., Lovejoy, T., Meir, P., Mendoza, C., Morandi, P., Neill, D., Nogueira Lima, A.J., Vargas, P.N., de Oliveira, E.A., Camacho, N.P., Pardo, G., Peacock, J., Peña-Claros, M., Peñuela-Mora, M.C., Pickavance, G., Pipoly, J., Pitman, N., Prieto, A., Pugh, T.A.M., Quesada, C., Ramirez-Angulo, H., de Almeida Reis, S.M., Rejou-Machain, M., Correa, Z.R., Bayona, L.R., Rudas, A., Salomão, R., Serrano, J., Espejo, J.S., Silva, N., Singh, J., Stahl, C., Stropp, J., Swamy, V., Talbot, J., ter Steege, H., Terborgh, J., Thomas, R., Toledo, M., Torres-Lezama, A., Gamarra, L.V., van der Heijden, G., van der Meer, P., van der Hout, P., Martinez, R.V., Vieira, S.A., Cayo, J.V., Vos, V., Zagt, R., Zuidema, P. & Galbraith, D. (2020) Tree mode of death and mortality risk factors across Amazon forests. *Nature Communications*, **11**, 5515.
- Esquivel-Muelbert, A., Baker, T.R., Dexter, K.G., Lewis, S.L., Brienen, R.J.W., Feldpausch, T.R., Lloyd, J., Monteagudo-Mendoza, A., Arroyo, L., Álvarez-Dávila, E., Higuchi, N., Marimon, B.S., Marimon-Junior, B.H., Silveira, M., Vilanova, E., Gloor, E., Malhi, Y., Chave, J., Barlow, J., Bonal, D., Davila Cardozo, N., Erwin, T., Fauset, S., Hérault, B., Laurance, S., Poorter, L., Qie, L., Stahl, C., Sullivan, M.J.P., ter Steege, H., Vos, V.A., Zuidema, P.A., Almeida, E., Almeida de Oliveira, E., Andrade, A., Vieira, S.A., Aragão, L., Araujo-Murakami, A., Arets, E., Aymard C, G.A., Baraloto, C., Camargo, P.B., Barroso, J.G., Bongers, F., Boot, R., Camargo, J.L., Castro, W., Chama Moscoso, V., Comiskey, J., Cornejo Valverde, F., Lola da Costa, A.C., del Aguila Pasquel, J., Di Fiore, A., Fernanda Duque, L., Elias, F., Engel, J., Flores Llampazo, G., Galbraith, D., Herrera Fernández, R., Honorio Coronado, E., Hubau, W., Jimenez-Rojas, E., Lima, A.J.N., Umetsu, R.K., Laurance, W., Lopez-Gonzalez, G., Lovejoy, T., Aurelio Melo Cruz, O., Morandi, P.S., Neill, D., Núñez Vargas, P., Pallqui Camacho, N.C., Parada Gutierrez, A., Pardo, G., Peacock, J., Peña-Claros, M., Peñuela-Mora, M.C., Petronelli, P., Pickavance, G.C., Pitman, N., Prieto, A., Quesada, C., Ramírez-Angulo, H., Réjou-Méchain, M., Restrepo Correa, Z., Roopsind, A., Rudas, A., Salomão, R., Silva, N., Silva Espejo, J., Singh, J., Stropp, J., Terborgh, J., Thomas, R., Toledo, M., Torres-Lezama, A., Valenzuela Gamarra, L., van de Meer, P.J., van der Heijden, G., van der Hout, P., Vasquez Martinez, R., Vela, C., Vieira, I.C.G. & Phillips, O.L. (2019) Compositional response of Amazon forests to climate change. *Global Change Biology*, **25**, 39–56.
- Esteban, E.J.L., Castilho, C. V., Melgaço, K.L. & Costa, F.R.C. (2020) The other side of droughts: wet extremes and topography as buffers of negative drought effects in an Amazonian forest. *New Phytologist*, nph.17005.
- Fan, Y. (2015) Groundwater in the Earth's critical zone: Relevance to large-scale patterns and processes. *Water Resources Research*, **51**, 3052–3069.
- Fan, Y., Li, H. & Miguez-Macho, G. (2013) Global patterns of groundwater table depth. *Science*, **339**, 940–943.
- Fan, Y. & Miguez-Macho, G. (2011) A simple hydrologic framework for simulating wetlands in climate and earth system models. *Climate Dynamics*, **37**, 253–278.
- Fan, Y. & Miguez-Macho, G. (2010) Potential groundwater contribution to Amazon evapotranspiration. *Hydrology and Earth System Sciences*, **14**, 2039–2056.
- Fan, Y., Miguez-Macho, G., Jobbágy, E., Jackson, R.B. & Otero-Casal, C. (2017)

Hydrologic regulation of plant rooting depth. *Proceedings of the National Academy of Sciences*, **114**, 10572–10577.

- Fauset, S., Johnson, M.O., Gloor, M., Baker, T.R., Monteagudo M., A., Brien, R.J.W., Feldpausch, T.R., Lopez-Gonzalez, G., Malhi, Y., Ter Steege, H., Pitman, N.C.A., Baraloto, C., Engel, J., Pétronelli, P., Andrade, A., Camargo, J.L.C., Laurance, S.G.W., Laurance, W.F., Chave, J., Allie, E., Vargas, P.N., Terborgh, J.W., Ruokolainen, K., Silveira, M., Aymard C., G.A., Arroyo, L., Bonal, D., Ramirez-Angulo, H., Araujo-Murakami, A., Neill, D., Hérault, B., Dourdain, A., Torres-Lezama, A., Marimon, B.S., Salomão, R.P., Comiskey, J.A., Réjou-Méchain, M., Toledo, M., Licona, J.C., Alarcón, A., Prieto, A., Rudas, A., Van Der Meer, P.J., Killeen, T.J., Marimon Junior, B.H., Poorter, L., Boot, R.G.A., Stergios, B., Torre, E.V., Costa, F.R.C., Levis, C., Schiatti, J., Souza, P., Groot, N., Arets, E., Moscoso, V.C., Castro, W., Coronado, E.N.H., Peña-Claros, M., Stahl, C., Barroso, J., Talbot, J., Vieira, I.C.G., Van Der Heijden, G., Thomas, R., Vos, V.A., Almeida, E.C., Davila, E.Á., Aragão, L.E.O.C., Erwin, T.L., Morandi, P.S., De Oliveira, E.A., Valadão, M.B.X., Zagt, R.J., Van Der Hout, P., Loayza, P.A., Pipoly, J.J., Wang, O., Alexiades, M., Cerón, C.E., Huamantupa-Chuquimaco, I., Di Fiore, A., Peacock, J., Camacho, N.C.P., Umetsu, R.K., De Camargo, P.B., Burnham, R.J., Herrera, R., Quesada, C.A., Stropp, J., Vieira, S.A., Steininger, M., Rodríguez, C.R., Restrepo, Z., Muelbert, A.E., Lewis, S.L., Pickavance, G.C. & Phillips, O.L. (2015) Hyperdominance in Amazonian forest carbon cycling. *Nature Communications*, **6**, 1–9.
- Fearnside, P.M. (2008) Amazon Forest maintenance as a source of environmental services. *Anais da Academia Brasileira de Ciências*, **80**, 101–114.
- Ferry, B., François, M., Bontemps, J.D., Blanc, L. & Freycon, V. (2010) Higher treefall rates on slopes and waterlogged soils result in lower stand biomass and productivity in a tropical rain forest. *Journal of Ecology*, **98**, 106–116.
- ForestPlots.net, Blundo, C., Carilla, J., Grau, R., Malizia, A., Malizia, L., Osinaga-Acosta, O., Bird, M., Bradford, M., Catchpole, D., Ford, A., Graham, A., Hilbert, D., Kemp, J., Laurance, S., Laurance, W., Ishida, F.Y., Marshall, A., Waite, C., Woell, H., Bastin, J.-F., Bauters, M., Beekman, H., Boeckx, P., Bogaert, J., De Canniere, C., de Haulleville, T., Doucet, J.-L., Hardy, O., Hubau, W., Kearsley, E., Verbeeck, H., Vleminckx, J., Brewer, S.W., Alarcón, A., Araujo-Murakami, A., Arets, E., Arroyo, L., Chavez, E., Fredericksen, T., Villaroel, R.G., Sibauty, G.G., Killeen, T., Licona, J.C., Lléigue, J., Mendoza, C., Murakami, S., Gutierrez, A.P., Pardo, G., Peña-Claros, M., Poorter, L., Toledo, M., Cayo, J.V., Viscarra, L.J., Vos, V., Ahumada, J., Almeida, E., Almeida, J., de Oliveira, E.A., da Cruz, W.A., de Oliveira, A.A., Carvalho, F.A., Obermuller, F.A., Andrade, A., Carvalho, F.A., Vieira, S.A., Aquino, A.C., Aragão, L., Araújo, A.C., Assis, M.A., Gomes, J.A.M.A., Baccaro, F., de Camargo, P.B., Barni, P., Barroso, J., Bernacci, L.C., Bordin, K., de Medeiros, M.B., Broggio, I., Camargo, J.L., Cardoso, D., Carniello, M.A., Rochelle, A.L.C., Castilho, C., Castro, A.A.J.F., Castro, W., Ribeiro, S.C., Costa, F., de Oliveira, R.C., Coutinho, I., Cunha, J., da Costa, L., da Costa Ferreira, L., da Costa Silva, R., da Graça Zacarias Simbine, M., de Andrade Kamimura, V., de Lima, H.C., de Oliveira Melo, L., de Queiroz, L., de Sousa Lima, J.R., do Espírito Santo, M., Domingues, T., dos Santos Prestes, N.C., Carneiro, S.E.S., Elias, F., Eliseu, G., Emilio, T., Farrapo, C.L., Fernandes, L., Ferreira, G., Ferreira, J., Ferreira, L., Ferreira, S., Simon, M.F., Freitas, M.A., García, Q.S., Manzatto, A.G., Graça, P., Guilherme, F., Hase, E., Higuchi, N., Iguatemy, M., Barbosa, R.I., Jaramillo, M., Joly, C., Klipel, J., do Amaral, I.L., Levis, C., Lima, A.S., Dan,

1
 2
 3 M.L., Lopes, A., Madeiros, H., Magnusson, W.E., dos Santos, R.M., Marimon, B.,
 4 Junior, B.H.M., Grillo, R.M.M., Martinelli, L., Reis, S.M., Medeiros, S., Meira-
 5 Junior, M., Metzker, T., Morandi, P., do Nascimento, N.M., Moura, M., Müller,
 6 S.C., Nagy, L., Nascimento, H., Nascimento, M., Lima, A.N., de Araújo, R.O.,
 7 Silva, J.O., Pansonato, M., Sabino, G.P., de Abreu, K.M.P., Rodrigues, P.J.F.P.,
 8 Piedade, M., Rodrigues, D., Rodrigues Pinto, J.R., Quesada, C., Ramos, E.,
 9 Ramos, R., Rodrigues, P., de Sousa, T.R., Salomão, R., Santana, F., Scaranello,
 10 M., Bergamin, R.S., Schietti, J., Schöngart, J., Schwartz, G., Silva, N., Silveira, M.,
 11 Seixas, C.S., Simbine, M., Souza, A.C., Souza, P., Souza, R., Sposito, T., Junior,
 12 E.S., do Vale, J.D., Vieira, I.C.G., Villela, D., Vital, M., Xaud, H., Zanini, K.,
 13 Zartman, C.E., Ideris, N.K.H., Metali, F. binti H., Salim, K.A., Saparudin, M.S.,
 14 Serudin, R.M., Sukri, R.S., Begne, S., Chuyong, G., Djuikouo, M.N., Gonmadje,
 15 C., Simo-Droissart, M., Sonké, B., Taedoung, H., Zemagho, L., Thomas, S., Baya,
 16 F., Saiz, G., Espejo, J.S., Chen, D., Hamilton, A., Li, Y., Luo, T., Niu, S., Xu, H.,
 17 Zhou, Z., Álvarez-Dávila, E., Escobar, J.C.A., Arellano-Peña, H., Duarte, J.C.,
 18 Calderón, J., Bravo, L.M.C., Cuadrado, B., Cuadros, H., Duque, A., Duque, L.F.,
 19 Espinosa, S.M., Franke-Ante, R., García, H., Gómez, A., González-M., R.,
 20 Idárraga-Piedrahíta, Á., Jimenez, E., Jurado, R., Oviedo, W.L., López-Camacho,
 21 R., Cruz, O.A.M., Polo, I.M., Paky, E., Pérez, K., Pijachi, A., Pizano, C., Prieto,
 22 A., Ramos, L., Correa, Z.R., Richardson, J., Rodríguez, E., Rodríguez M., G.M.,
 23 Rudas, A., Stevenson, P., Chudomelová, M., Dancak, M., Hédl, R., Lhota, S.,
 24 Svatek, M., Mukinzi, J., Ewango, C., Hart, T., Yakusu, E.K., Lisingo, J., Makana,
 25 J.-R., Mbayu, F., Toirambe, B., Mukendi, J.T., Kvist, L., Nebel, G., Báez, S.,
 26 Céron, C., Griffith, D.M., Andino, J.E.G., Neill, D., Palacios, W., Peñuela-Mora,
 27 M.C., Rivas-Torres, G., Villa, G., Demissie, S., Gole, T., Gonfa, T., Ruokolainen,
 28 K., Baisie, M., Bénédet, F., Betian, W., Bezard, V., Bonal, D., Chave, J., Droissart,
 29 V., Gurllet-Fleury, S., Hladik, A., Labrière, N., Naisso, P., Réjou-Méchain, M.,
 30 Sist, P., Blanc, L., Burban, B., Derroire, G., Dourdain, A., Stahl, C., Bengone,
 31 N.N., Chezeaux, E., Ondo, F.E., Medjibe, V., Mihindou, V., White, L., Culmsee,
 32 H., Rangel, C.D., Horna, V., Wittmann, F., Adu-Bredu, S., Affum-Baffoe, K., Foli,
 33 E., Balinga, M., Roopsind, A., Singh, J., Thomas, R., Zagt, R., Murthy, I.K.,
 34 Kartawinata, K., Mirmanto, E., Priyadi, H., Samsuedin, I., Sunderland, T., Yassir,
 35 I., Rovero, F., Vinceti, B., Hérault, B., Aiba, S.-I., Kitayama, K., Daniels, A.,
 36 Tuagben, D., Woods, J.T., Fitriadi, M., Karolus, A., Khoon, K.L., Majalap, N.,
 37 Maycock, C., Nilus, R., Tan, S., Siteo, A., Coronado G., I., Ojo, L., de Assis, R.,
 38 Poulsen, A.D., Sheil, D., Pezo, K.A., Verde, H.B., Moscoso, V.C., Oroche, J.C.C.,
 39 Valverde, F.C., Medina, M.C., Cardozo, N.D., de Rutte Corzo, J., del Aguila
 40 Pasquel, J., Llampazo, G.F., Freitas, L., Cabrera, D.G., Villacorta, R.G., Cabrera,
 41 K.G., Soria, D.G., Saboya, L.G., Rios, J.M.G., Pizango, G.H., Coronado, E.H.,
 42 Huamantupa-Chuquimaco, I., Huasco, W.H., Aedo, Y.T.H., Peña, J.L.M.,
 43 Mendoza, A.M., Rodriguez, V.M., Vargas, P.N., Ramos, S.C.P., Camacho, N.P.,
 44 Cruz, A.P., Arevalo, F.R., Huaymacari, J.R., Rodriguez, C.R., Paredes, M.A.R.,
 45 Bayona, L.R., del Pilar Rojas Gonzales, R., Peña, M.E.R., Revilla, N.S., Shareva,
 46 Y.C.S., Trujillo, R.T., Gamarra, L.V., Martinez, R.V., Arenas, J.V., Amani, C., Ifo,
 47 S.A., Bocko, Y., Boundja, P., Ekoungoulou, R., Hockemba, M., Nzala, D.,
 48 Fofanah, A., Taylor, D., Bañares-de Dios, G., Cayuela, L., la Cerda, Í.G., Macía,
 49 M., Stropp, J., Playfair, M., Wortel, V., Gardner, T., Muscarella, R., Priyadi, H.,
 50 Rutishauser, E., Chao, K.-J., Munishi, P., Bánki, O., Bongers, F., Boot, R.,
 51 Fredriksson, G., Reitsma, J., ter Steege, H., van Andel, T., van de Meer, P., van der
 52 Hout, P., van Nieuwstadt, M., van Ulf, B., Veenendaal, E., Vernimmen, R.,
 53
 54
 55
 56
 57
 58
 59
 60

- Zuidema, P., Zwerts, J., Akite, P., Bitariho, R., Chapman, C., Gerald, E., Leal, M., Mucunguzi, P., Abernethy, K., Alexiades, M., Baker, T.R., Banda, K., Banin, L., Barlow, J., Bennett, A., Berenguer, E., Berry, N., Bird, N.M., Blackburn, G.A., Brearley, F., Brienen, R., Burslem, D., Carvalho, L., Cho, P., Coelho, F., Collins, M., Coomes, D., Cuni-Sanchez, A., Dargie, G., Dexter, K., Disney, M., Draper, F., Duan, M., Esquivel-Muelbert, A., Ewers, R., Fadrique, B., Fauset, S., Feldpausch, T.R., França, F., Galbraith, D., Gilpin, M., Gloor, E., Grace, J., Hamer, K., Harris, D., Jeffery, K., Jucker, T., Kalamandeen, M., Klitgaard, B., Levesley, A., Lewis, S.L., Lindsell, J., Lopez-Gonzalez, G., Lovett, J., Malhi, Y., Marthens, T., McIntosh, E., Melgaço, K., Milliken, W., Mitchard, E., Moonlight, P., Moore, S., Morel, A., Peacock, J., Peh, K.S.-H., Pendry, C., Pennington, R.T., de Oliveira Pereira, L., Peres, C., Phillips, O.L., Pickavance, G., Pugh, T., Qie, L., Riutta, T., Roucoux, K., Ryan, C., Sarkinen, T., Valeria, C.S., Spracklen, D., Stas, S., Sullivan, M., Swaine, M., Talbot, J., Taplin, J., van der Heijden, G., Vedovato, L., Willcock, S., Williams, M., Alves, L., Loayza, P.A., Arellano, G., Asa, C., Ashton, P., Asner, G., Brncic, T., Brown, F., Burnham, R., Clark, C., Comiskey, J., Damasco, G., Davies, S., Di Fiore, T., Erwin, T., Farfan-Rios, W., Hall, J., Kenfack, D., Lovejoy, T., Martin, R., Montiel, O.M., Pipoly, J., Pitman, N., Poulsen, J., Primack, R., Silman, M., Steininger, M., Swamy, V., Terborgh, J., Thomas, D., Umunay, P., Uriarte, M., Torre, E.V., Wang, O., Young, K., Aymard C., G.A., Hernández, L., Fernández, R.H., Ramírez-Angulo, H., Salcedo, P., Sanoja, E., Serrano, J., Torres-Lezama, A., Le, T.C., Le, T.T. & Tran, H.D. (2021) Taking the pulse of Earth's tropical forests using networks of highly distributed plots. *Biological Conservation*, 108849.
- Fox, J., Weisberg, S., Adler, D., Bates, D., Baud-bovy, G., Ellison, S., Firth, D., Friendly, M., Gorjanc, G., Graves, S., Heiberger, R., Laboissiere, R., Monette, G., Murdoch, D., Ogle, D., Ripley, B. & Venables, W. (2018) Package 'car': Companion to Applied Regression. *CRAN Repository*.
- Franco, W. & Dezzio, N. (1994) Soils and soil water regime in the terra firme-caatinga forest complex near San Carlos de Río Negro, state of Amazonas, Venezuela. *Interciencia-Caracas-*, **19**, 305–305.
- Gale, N. & Barfod, A.S. (1999) Canopy tree mode of death in a western Ecuadorian rain forest. *Journal of Tropical Ecology*, **15**, 415–436.
- Gale, N. & Hall, P. (2001) Factors determining the modes of tree death in three Bornean rain forests. *Journal of Vegetation Science*, **12**, 337–348.
- Gibbs, J. & Greenway, H. (2003) Review: Mechanisms of anoxia tolerance in plants. I. Growth, survival and anaerobic catabolism. *Functional Plant Biology*, **30**, 1.
- Gloor, M., Barichivich, J., Ziv, G., Brienen, R., Schöngart, J., Peylin, P., Ladvoat Cintra, B.B., Feldpausch, T., Phillips, O. & Baker, J. (2015) Recent Amazon climate as background for possible ongoing and future changes of Amazon humid forests. *Global Biogeochemical Cycles*, **29**, 1384–1399.
- Gloor, M., Brienen, R.J.W., Galbraith, D., Feldpausch, T.R., Schöngart, J., Guyot, J.-L., Espinoza, J.C., Lloyd, J. & Phillips, O.L. (2013) Intensification of the Amazon hydrological cycle over the last two decades. *Geophysical Research Letters*, **40**, 1729–1733.
- Godoy, J.R., Petts, G. & Salo, J. (1999) Riparian flooded forests of the Orinoco and Amazon basins: A comparative review. *Biodiversity and Conservation*, **8**.
- Greenwell, B., Boehmke, B. & Cunningham, J. (2019) Package "gbm" - Generalized Boosted Regression Models. *CRAN Repository*.
- Grogan, J. & Galvão, J. (2006) Physiographic and floristic gradients across topography

- in transitional seasonally dry evergreen forests of southeast Pará, Brazil. *Acta Amazonica*, **36**, 483–496.
- Hengl, T., De Jesus, J.M., Heuvelink, G.B.M., Gonzalez, M.R., Kilibarda, M., Blagotić, A., Shangguan, W., Wright, M.N., Geng, X., Bauer-Marschallinger, B., Guevara, M.A., Vargas, R., MacMillan, R.A., Batjes, N.H., Leenaars, J.G.B., Ribeiro, E., Wheeler, I., Mantel, S. & Kempen, B. (2017) *SoilGrids250m: Global gridded soil information based on machine learning*.
- Hillel, D. (1998) Environmental Soil Physics: Fundamentals, Applications, and Environmental Considerations. *Environmental Soil Physics*.
- Hodnett, M.G., Vendrame, I., De O. Marques Filho, A., Oyama, M.D. & Tomasella, J. (1997) Soil water storage and groundwater behaviour in a catenary sequence beneath forest in central Amazonia: I. Comparisons between plateau, slope and valley floor. *Hydrology and Earth System Sciences*, **1**, 265–277.
- Ivanov, V.Y., Hutrya, L.R., Wofsy, S.C., Munger, J.W., Saleska, S.R., De Oliveira, R.C. & De Camargo, P.B. (2012) Root niche separation can explain avoidance of seasonal drought stress and vulnerability of overstory trees to extended drought in a mature Amazonian forest. *Water Resources Research*, **48**, 1–21.
- Jackson, R.B., Canadell, J., Ehleringer, J.R., Mooney, H.A., Sala, O.E. & Schulze, E.D. (1996) A global analysis of root distributions for terrestrial biomes. *Oecologia*, **108**, 389–411.
- Jirka, S., McDonald, A.J., Johnson, M.S., Feldpausch, T.R., Couto, G., Riha, S.J., Andrew, J., Mark, S., Ted, R., Eduardo, G. & Susan, J. (2007) Relationships between soil hydrology and forest structure and composition in the southern Brazilian Amazon. *Journal of Vegetation Science*, **18**, 183–194.
- Kreuzwieser, J. & Rennenberg, H. (2014) Molecular and physiological responses of trees to waterlogging stress. *Plant, Cell & Environment*, **37**, n/a-n/a.
- Law, B., Falge, E., Gu, L., Baldocchi, D., Bakwin, P., Berbigier, P., Davis, K., Dolman, A., Falk, M., Fuentes, J., Goldstein, A., Granier, A., Grelle, A., Hollinger, D., Janssens, I., Jarvis, P., Jensen, N., Katul, G., Mahli, Y., Matteucci, G., Meyers, T., Monson, R., Munger, W., Oechel, W., Olson, R., Pilegaard, K., Paw U, K., Thorgeirsson, H., Valentini, R., Verma, S., Vesala, T., Wilson, K. & Wofsy, S. (2002) Environmental controls over carbon dioxide and water vapor exchange of terrestrial vegetation. *Agricultural and Forest Meteorology*, **113**, 97–120.
- Lewis, S.L., Lopez-Gonzalez, G., Sonké, B., Affum-Baffoe, K., Baker, T.R., Ojo, L.O., Phillips, O.L., Reitsma, J.M., White, L., Comiskey, J.A., Djukouo K, M.N., Ewango, C.E.N., Feldpausch, T.R., Hamilton, A.C., Gloor, M., Hart, T., Hladik, A., Lloyd, J., Lovett, J.C., Makana, J.R., Malhi, Y., Mbago, F.M., Ndangalasi, H.J., Peacock, J., Peh, K.S.H., Sheil, D., Sunderland, T., Swaine, M.D., Taplin, J., Taylor, D., Thomas, S.C., Votere, R. & Wöll, H. (2009) Increasing carbon storage in intact African tropical forests. *Nature*, **457**, 1003–1006.
- Lewis, S.L., Phillips, O.L., Sheil, D., Vinceti, B., Timothy, R., Brown, S., Graham, A.W., Higuchi, N., Hilbert, D.W., William, F., Lejoly, J., Malhi, Y., Monteagudo, A., Vargas, P.N.N., N, N.S.M., Terborgh, J.W., Martínez, R.V., Baker, T.R., Hilbert, W. & Laurance, W.F. (2004) Tropical forest tree mortality, recruitment when and comparison rates: calculation, interpretation census intervals vary. *Journal of Ecology*, **92**, 929–944.
- Llopert, M., Reboita, M., Coppola, E., Giorgi, F., da Rocha, R. & de Souza, D. (2018) Land Use Change over the Amazon Forest and Its Impact on the Local Climate. *Water*, **10**, 149.

- 1
2
3 Lopez-Gonzalez, G., Lewis, S.L., Burkitt, M. & Phillips, O.L. (2011) ForestPlots.net: A
4 web application and research tool to manage and analyse tropical forest plot data.
5 *Journal of Vegetation Science*, **22**, 610–613.
- 6 Lopez-Gonzalez, G., Sullivan, M.J.P. & Baker, T.R. (2015) BiomasaFP: Tools for
7 analysing data downloaded from ForestPlots.net. *R package version*.
- 8 Lopez, O.R. & Kursar, T.A. (2003) Does flood tolerance explain tree species
9 distribution in tropical seasonally flooded habitats? *Oecologia*, **136**, 193–204.
- 10 Lopez, O.R. & Kursar, T.A. (1999) Flood tolerance of four tropical tree species. *Tree*
11 *Physiology*, **19**, 925–932.
- 12 Lüdecke, D., Ben-Shachar, M., Patil, I., Waggoner, P. & Makowski, D. (2021)
13 Performance: An R Package for Assessment, Comparison and Testing of Statistical
14 Models. *Journal of Open Source Software*, **6**.
- 15 Madelaine, C., Péliissier, R., Vincent, G., Molino, J.F., Sabatier, D., Prévost, M.F. & De
16 Namur, C. (2007) Mortality and recruitment in a lowland tropical rain forest of
17 French Guiana: Effects of soil type and species guild. *Journal of Tropical Ecology*,
18 **23**, 277–287.
- 19 Magnusson, W., Braga-Neto, R., Pezzini, F., Baccaro, F., Bergallo, H., Penha, J.,
20 Rodrigues, D., Verdade, L.M., Lima, A., Albernaz, A.L., Hero, J.M., Ben Lawson,
21 B., Castilho, C., Drucker, D., Franklin, E., Mendonça, F., Costa, F.R.C., Galdino,
22 G., Castley, G., Zuanon, J., Vale, J., Santos, J.L.C., Luizão, R., Cintra, R., Barbosa,
23 R.I., Lisboa, A., Koblitz, R.V., Cunha, C.N. & Ponte, A.R.M. (2013)
24 *Biodiversidade e monitoramento ambiental integrado - Biodiversity and integrated*
25 *environmental monitoring*, 1 ed. Áttema Editorial :: Assessoria e Design, Manaus.
- 26 Magnusson, W.E., Lima, A.P., Luizão, R., Luizão, F., Costa, F.R.C., Castilho, C.V. de
27 & Kinupp, V.F. (2005) RAPELD: a modification of the Gentry method for
28 biodiversity surveys in long-term ecological research sites. *Biota Neotropica*, **5**,
29 21–26.
- 30 Malhi, Y., Aragao, L.E.O.C., Galbraith, D., Huntingford, C., Fisher, R., Zelazowski, P.,
31 Sitch, S., McSweeney, C. & Meir, P. (2009) Exploring the likelihood and
32 mechanism of a climate-change-induced dieback of the Amazon rainforest.
33 *Proceedings of the National Academy of Sciences*, **106**, 20610–20615.
- 34 Malhi, Y., Baker, T.R., Phillips, O.L., Almeida, S., Alvarez, E., Arroyo, L., Chave, J.,
35 Czimczik, C.I., Di Fiore, A., Higuchi, N., Killeen, T.J., Laurance, S.G., Laurance,
36 W.F., Lewis, S.L., Montoya, L.M.M., Monteagudo, A., Neill, D.A., Vargas, P.N.,
37 Patino, S., Pitman, N.C.A., Quesada, C.A., Salomao, R., Silva, J.N.M., Lezama,
38 A.T., Martínez, R.V., Terborgh, J., Vinceti, B. & Lloyd, J. (2004) The above-
39 ground coarse wood productivity of 104 Neotropical forest plots. *Global Change*
40 *Biology*, **10**, 563–591.
- 41 Malhi, Y., Doughty, C.E., Goldsmith, G.R., Metcalfe, D.B., Girardin, C.A.J., Marthews,
42 T.R., del Aguila-Pasquel, J., Aragão, L.E.O.C., Araujo-Murakami, A., Brando, P.,
43 da Costa, A.C.L., Silva-Espejo, J.E., Farfán Amézquita, F., Galbraith, D.R.,
44 Quesada, C.A., Rocha, W., Salinas-Revilla, N., Silvério, D., Meir, P. & Phillips,
45 O.L. (2015) The linkages between photosynthesis, productivity, growth and
46 biomass in lowland Amazonian forests. *Global Change Biology*, **21**, 2283–2295.
- 47 Malhi, Y., Wood, D., Baker, T.R., Wright, J., Phillips, O.L., Cochrane, T., Meir, P.,
48 Chave, J., Almeida, S., Arroyo, L., Higuchi, N., Killeen, T.J., Laurance, S.G.,
49 Laurance, W.F., Lewis, S.L., Monteagudo, A., Neill, D.A., Vargas, P.N., Pitman,
50 N.C.A., Quesada, C.A., Salomão, R., Silva, J.N.M., Lezama, A.T., Terborgh, J.,
51 Martínez, R.V. & Vinceti, B. (2006) The regional variation of aboveground live
52 biomass in old-growth Amazonian forests. *Global Change Biology*, **12**, 1107–

- 1
2
3 1138.
- 4 Marengo, J.A. & Espinoza, J.C. (2016) Extreme seasonal droughts and floods in
5 Amazonia: Causes, trends and impacts. *International Journal of Climatology*, **36**,
6 1033–1050.
- 7 Miguez-Macho, G. & Fan, Y. (2012) The role of groundwater in the Amazon water
8 cycle: 2. Influence on seasonal soil moisture and evapotranspiration. *Journal of*
9 *Geophysical Research: Atmospheres*, **117**.
- 10 Milborrow, S. (2019) Plotting rpart trees with the rpart.plot package.
11 [Http://Www.Milbo.Org/Rpart-Plot/Prp.Pdf](http://www.milbo.org/rpart-plot/prp.pdf).
- 12 Moeslund, J.E., Arge, L., Bøcher, P.K., Dalgaard, T., Odgaard, M. V., Nygaard, B. &
13 Svenning, J.-C. (2013) Topographically controlled soil moisture is the primary
14 driver of local vegetation patterns across a lowland region. *Ecosphere*, **4**, art91.
- 15 Moulatlet, G.M., Costa, F.R.C., Rennó, C.D., Emilio, T. & Schiatti, J. (2014) Local
16 hydrological conditions explain floristic composition in lowland amazonian
17 forests. *Biotropica*, **46**, 395–403.
- 18 Moulatlet, G.M., Zuquim, G., Figueiredo, F.O.G., Lehtonen, S., Emilio, T.,
19 Ruokolainen, K. & Tuomisto, H. (2017) Using digital soil maps to infer edaphic
20 affinities of plant species in Amazonia: Problems and prospects. *Ecology and*
21 *Evolution*, **7**, 8463–8477.
- 22 Nobre, A.D., Cuartas, L.A., Hodnett, M., Rennó, C.D., Rodrigues, G., Silveira, A.,
23 Waterloo, M. & Saleska, S. (2011) Height Above the Nearest Drainage – a
24 hydrologically relevant new terrain model. *Journal of Hydrology*, **404**, 13–29.
- 25 North, G.B., Martre, P. & Nobel, P.S. (2004) Aquaporins account for variations in
26 hydraulic conductance for metabolically active root regions of *Agave deserti* in
27 wet, dry, and rewetted soil. *Plant, Cell and Environment*, **27**.
- 28 Pan, Y., Birdsey, R.A., Fang, J., Houghton, R., Kauppi, P.E., Kurz, W.A., Phillips,
29 O.L., Shvidenko, A., Lewis, S.L., Canadell, J.G., Ciais, P., Jackson, R.B., Pacala,
30 S.W., McGuire, A.D., Piao, S., Rautiainen, A., Sitch, S. & Hayes, D. (2011) A
31 Large and Persistent Carbon Sink in the World’s Forests. *Science*, **333**, 988–993.
- 32 Parahyba, R. da B.V., Araújo, M.D.S.B. de, Almeida, B.G. de, Rolim Neto, F.C.,
33 Sampaio, E.V.S.B. & Caldas, A.M. (2019) Water retention capacity in Arenosols
34 and Ferralsols in a semiarid area in the state of Bahia, Brazil. *Anais da Academia*
35 *Brasileira de Ciências*, **91**.
- 36 Parent, C., Capelli, N., Berger, A., Crèvecoeur, M. & Dat, J. (2008) An overview of
37 plant responses to soil waterlogging. *Plant Stress*, **2**, 20–27.
- 38 Parolin, P. (2012) Diversity of adaptations to flooding in trees of amazonian
39 floodplains. *Pesquisas Botanica*, **63**.
- 40 Parolin, P. (2000) Phenology and CO₂-assimilation of trees in Central Amazonian
41 floodplains. *Journal of Tropical Ecology*, **16**, 465–473.
- 42 Parolin, P., Simone, O., Haase, K., Waldhoff, D., Rottenberger, S., Kuhn, U.,
43 Kesselmeier, J., Kleiss, B., Schmidt, W., Piedade, M.T.F. & Junk, W.J. (2004)
44 Central Amazonian Floodplain Forests: Tree Adaptations in a Pulsing System. *The*
45 *Botanical Review*, **70**, 357–380.
- 46 Pezeshki, S.R. (2001) Wetland plant responses to soil flooding. *Environmental and*
47 *Experimental Botany*, **46**, 299–312.
- 48 Phillips, O.L., Aragao, L.E.O.C., Lewis, S.L., Fisher, J.B., Lloyd, J., Lopez-Gonzalez,
49 G., Malhi, Y., Monteagudo, A., Peacock, J., Quesada, C.A., van der Heijden, G.,
50 Almeida, S., Amaral, I., Arroyo, L., Aymard, G., Baker, T.R., Banki, O., Blanc, L.,
51 Bonal, D., Brando, P., Chave, J., de Oliveira, A.C.A., Cardozo, N.D., Czimczik,
52 C.I., Feldpausch, T.R., Freitas, M.A., Gloor, E., Higuchi, N., Jimenez, E., Lloyd,
53
54
55
56
57
58
59
60

- G., Meir, P., Mendoza, C., Morel, A., Neill, D.A., Nepstad, D., Patino, S., Penuela, M.C., Prieto, A., Ramirez, F., Schwarz, M., Silva, J., Silveira, M., Thomas, A.S., Steege, H. Ter, Stropp, J., Vasquez, R., Zelazowski, P., Davila, E.A., Andelman, S., Andrade, A., Chao, K.-J., Erwin, T., Di Fiore, A., C., E.H., Keeling, H., Killeen, T.J., Laurance, W.F., Cruz, A.P., Pitman, N.C.A., Vargas, P.N., Ramirez-Angulo, H., Rudas, A., Salamao, R., Silva, N., Terborgh, J. & Torres-Lezama, A. (2009) Drought Sensitivity of the Amazon Rainforest. *Science*, **323**, 1344–1347.
- Phillips, O.L., Baker, T.R., Brienen, R. & Feldpausch, T.R. (2010) Field manual for plot establishment and remeasurement. (<http://www.geog.leeds.ac.uk/projects/rainfor/>).
- Phillips, O.L. & Brienen, R.J.W. (2017) Carbon uptake by mature Amazon forests has mitigated Amazon nations' carbon emissions. *Carbon Balance and Management*, **12**, 1.
- Phillips, O.L., Hall, P., Gentry, A.H., Sawyer, S.A. & Vasquez, R. (1994) Dynamics and species richness of tropical rain forests. *Proceedings of the National Academy of Sciences*, **91**, 2805–2809.
- Piedade, M.T.F., Wittmann, F., Parolin, P. & Junk, W.J. (2013) *Impactos ecológicos da inundação e seca na vegetação das áreas alagáveis amazônicas. Eventos climáticos extremos na Amazônia: causas e conseqüências*, pp. 409–461. São Paulo, Brasil.
- Pokhrel, Y.N., Fan, Y. & Miguez-Macho, G. (2014) Potential hydrologic changes in the Amazon by the end of the 21st century and the groundwater buffer. *Environmental Research Letters*, **9**, 084004.
- Quesada, C.A., Lloyd, J., Schwarz, M., Patiño, S., Baker, T.R., Czimczik, C., Fyllas, N.M., Martinelli, L., Nardoto, G.B., Schmerler, J., Santos, A.J.B., Hodnett, M.G., Herrera, R., Luizão, F.J., Arneith, A., Lloyd, G., Dezzeo, N., Hilke, I., Kuhlmann, I., Raessler, M., Brand, W.A., Geilmann, H., Filho, J.O.M., Carvalho, F.P., Filho, R.N.A., Chaves, J.E., Cruz, O.F., Pimentel, T.P. & Paiva, R. (2010) Variations in chemical and physical properties of Amazon forest soils in relation to their genesis. *Biogeosciences*, **7**, 1515–1541.
- Quesada, C.A., Phillips, O.L., Schwarz, M., Czimczik, C.I., Baker, T.R., Patiño, S., Fyllas, N.M., Hodnett, M.G., Herrera, R., Almeida, S., Alvarez Dávila, E., Arneith, A., Arroyo, L., Chao, K.J., Dezzeo, N., Erwin, T., Di Fiore, A., Higuchi, N., Honorio Coronado, E., Jimenez, E.M., Killeen, T., Lezama, A.T., Lloyd, G., López-González, G., Luizão, F.J., Malhi, Y., Monteagudo, A., Neill, D.A., Núñez Vargas, P., Paiva, R., Peacock, J., Peñuela, M.C., Peña Cruz, A., Pitman, N., Priante Filho, N., Prieto, A., Ramírez, H., Rudas, A., Salomão, R., Santos, A.J.B., Schmerler, J., Silva, N., Silveira, M., Vásquez, R., Vieira, I., Terborgh, J. & Lloyd, J. (2012) Basin-wide variations in Amazon forest structure and function are mediated by both soils and climate. *Biogeosciences*, **9**, 2203–2246.
- Rennó, C.D., Nobre, A.D., Cuartas, L.A., Soares, J.V., Hodnett, M.G., Tomasella, J. & Waterloo, M.J. (2008) HAND, a new terrain descriptor using SRTM-DEM: Mapping terra-firme rainforest environments in Amazonia. *Remote Sensing of Environment*, **112**, 3469–3481.
- Richter, D.D. & Babbar, L.I. (1991) *Soil Diversity in the Tropics. Environment International*, pp. 315–389.
- Rinke, P., Delaney, K., Garcia-Gonzalez, P. & Godby, R.W. (2004) Image states in metal clusters. *Forest Ecology and Management*, **310**, 994–1004.
- Rocha, H.R., Goulden, M.L., Miller, S.D., Menton, M.C., Pinto, L.D.V.O., De Freitas, H.C. & E Silva Figueira, A.M. (2004) Seasonality of water and heat fluxes over a tropical forest in eastern Amazonia. *Ecological Applications*, **14**, S22–S32.

- 1
2
3 Roebroek, C.T.J., Melsen, L.A., Hoek van Dijke, A.J., Fan, Y. & Teuling, A.J. (2020)
4 Global distribution of hydrologic controls on forest growth. *Hydrology and Earth*
5 *System Sciences*, **24**, 4625–4639.
- 6 Saatchi, S.S., Houghton, R.A., dos Santos Alvalá, R.C., Soares, J. V. & Yu, Y. (2007)
7 Distribution of aboveground live biomass in the Amazon basin. *Global Change*
8 *Biology*, **13**, 816–837.
- 9 Schietti, J., Emilio, T., Rennó, C.D., Drucker, D.P., Costa, F.R.C., Nogueira, A.,
10 Baccaro, F.B., Figueiredo, F., Castilho, C. V., Kinupp, V., Guillaumet, J.L.,
11 Garcia, A.R.M., Lima, A.P. & Magnusson, W.E. (2013) Vertical distance from
12 drainage drives floristic composition changes in an Amazonian rainforest. *Plant*
13 *Ecology and Diversity*, **7**, 241–253.
- 14 Schöngart, J., Junk, W.J., Piedade, M.T.F., Ayres, J.M., Hüttermann, A. & Worbes, M.
15 (2004) Teleconnection between tree growth in the Amazonian floodplains and the
16 El Niño-Southern Oscillation effect. *Global Change Biology*, **10**, 683–692.
- 17 Schöngart, J., Piedade, M.T.F., Ludwigshausen, S., Horna, V. & Worbes, M. (2002)
18 Phenology and stem-growth periodicity of tree species in Amazonian floodplain
19 forests. *Journal of Tropical Ecology*, **18**, 581–597.
- 20 Setter, T. & Belford, B. (1990) Waterlogging : how it reduces plant growth and how
21 plants can overcome its effects. *Journal of the Department of Agriculture, Western*
22 *Australia, Series*, **4**.
- 23 Sheil, D., Burslem, D.F.R.P. & Alder, D. (1995) The Interpretation and
24 Misinterpretation of Mortality Rate Measures. *British Ecological Society*, **83**, 331–
25 333.
- 26 Shuttleworth, W.J. (1988) Evaporation from Amazonian rainforest. *Proceedings - Royal*
27 *Society of London, Series B*, **233**, 321–346.
- 28 Simone, O., Junk, W.J. & Schmidt, W. (2003) Central Amazon Floodplain Forests:
29 Root Adaptations to Prolonged Flooding. *Russian Journal of Plant Physiology*, **50**,
30 848–855.
- 31 Sousa, T.R., Schietti, J., Coelho de Souza, F., Esquivel-Muelbert, A., Ribeiro, I.O.,
32 Emílio, T., Pequeno, P.A.C.L., Phillips, O. & Costa, F.R.C. (2020) Palms and trees
33 resist extreme drought in Amazon forests with shallow water tables. *Journal of*
34 *Ecology*, **108**, 2070–2082.
- 35 Stahl, C., Burban, B., Bompy, F., Jolin, Z.B., Sermage, J. & Bonal, D. (2010) Seasonal
36 variation in atmospheric relative humidity contributes to explaining seasonal
37 variation in trunk circumference of tropical rain-forest trees in French Guiana.
38 *Journal of Tropical Ecology*, **26**, 393–405.
- 39 ter Steege, H., Pitman, N.C.A., Sabatier, D., Baraloto, C., Salomão, R.P., Guevara, J.E.,
40 Phillips, O.L., Castilho, C. V., Magnusson, W.E., Molino, J.F., Monteagudo, A.,
41 Vargas, P.N., Montero, J.C., Feldpausch, T.R., Coronado, E.N.H., Killeen, T.J.,
42 Mostacedo, B., Vasquez, R., Assis, R.L., Terborgh, J., Wittmann, F., Andrade, A.,
43 Laurance, W.F., Laurance, S.G.W., Marimon, B.S., Marimon, B.H., Vieira, I.C.G.,
44 Amaral, I.L., Brienen, R., Castellanos, H., López, D.C., Duivenvoorden, J.F.,
45 Mogollón, H.F., Matos, F.D.D.A., Dávila, N., García-Villacorta, R., Diaz, P.R.S.,
46 Costa, F., Emilio, T., Levis, C., Schietti, J., Souza, P., Alonso, A., Dallmeier, F.,
47 Montoya, A.J.D., Piedade, M.T.F., Araujo-Murakami, A., Arroyo, L., Gribel, R.,
48 Fine, P.V.A., Peres, C.A., Toledo, M., Aymard C., G.A., Baker, T.R., Cerón, C.,
49 Engel, J., Henkel, T.W., Maas, P., Petronelli, P., Stropp, J., Zartman, C.E., Daly,
50 D., Neill, D., Silveira, M., Paredes, M.R., Chave, J., Lima Filho, D.D.A.,
51 Jørgensen, P.M., Fuentes, A., Schöngart, J., Valverde, F.C., Di Fiore, A., Jimenez,
52 E.M., Mora, M.C.P., Phillips, J.F., Rivas, G., Van Andel, T.R., Von Hildebrand,

P., Hoffman, B., Zent, E.L., Malhi, Y., Prieto, A., Rudas, A., Ruschell, A.R., Silva, N., Vos, V., Zent, S., Oliveira, A.A., Schutz, A.C., Gonzales, T., Nascimento, M.T., Ramirez-Angulo, H., Sierra, R., Tirado, M., Medina, M.N.U., Van Der Heijden, G., Vela, C.I.A., Torre, E.V., Vriesendorp, C., Wang, O., Young, K.R., Baider, C., Balslev, H., Ferreira, C., Mesones, I., Torres-Lezama, A., Giraldo, L.E.U., Zagt, R., Alexiades, M.N., Hernandez, L., Huamantupa-Chuquimaco, I., Milliken, W., Cuenca, W.P., Pauletto, D., Sandoval, E.V., Gamarra, L.V., Dexter, K.G., Feeley, K., Lopez-Gonzalez, G. & Silman, M.R. (2013) Hyperdominance in the Amazonian tree flora. *Science*, **342**.

Sullivan, M.J.P., Lewis, S.L., Affum-Baffoe, K., Castilho, C., Costa, F., Sanchez, A.C., Ewango, C.E.N., Hubau, W., Marimon, B., Monteagudo-Mendoza, A., Qie, L., Sonké, B., Martinez, R.V., Baker, T.R., Brienen, R.J.W., Feldpausch, T.R., Galbraith, D., Gloor, M., Malhi, Y., Aiba, S.I., Alexiades, M.N., Almeida, E.C., de Oliveira, E.A., Dávila, E.Á., Loayza, P.A., Andrade, A., Vieira, S.A., Aragão, L.E.O.C., Araujo-Murakami, A., Arets, E.J.M.M., Arroyo, L., Ashton, P., Aymard C, G., Baccaro, F.B., Banin, L.F., Baraloto, C., Camargo, P.B., Barlow, J., Barroso, J., Bastin, J.F., Batterman, S.A., Beeckman, H., Begne, S.K., Bennett, A.C., Berenguer, E., Berry, N., Blanc, L., Boeckx, P., Bogaert, J., Bonal, D., Bongers, F., Bradford, M., Brearley, F.Q., Brncic, T., Brown, F., Burban, B., Camargo, J.L., Castro, W., Céron, C., Ribeiro, S.C., Moscoso, V.C., Chave, J., Chezeaux, E., Clark, C.J., de Souza, F.C., Collins, M., Comiskey, J.A., Valverde, F.C., Medina, M.C., da Costa, L., Dančák, M., Dargie, G.C., Davies, S., Cardozo, N.D., de Haulleville, T., de Medeiros, M.B., Del Aguila Pasquel, J., Derroire, G., Di Fiore, A., Doucet, J.L., Dourdain, A., Droissart, V., Duque, L.F., Ekoungoulou, R., Elias, F., Erwin, T., Esquivel-Muelbert, A., Fauset, S., Ferreira, J., Llampazo, G.F., Foli, E., Ford, A., Gilpin, M., Hall, J.S., Hamer, K.C., Hamilton, A.C., Harris, D.J., Hart, T.B., Hédli, R., Herault, B., Herrera, R., Higuchi, N., Hladik, A., Coronado, E.H., Huamantupa-Chuquimaco, I., Huasco, W.H., Jeffery, K.J., Jimenez-Rojas, E., Kalamandeen, M., Djuikouo, M.N.K., Kearsley, E., Umetsu, R.K., Kho, L.K., Killeen, T., Kitayama, K., Klitgaard, B., Koch, A., Labrière, N., Laurance, W., Laurance, S., Leal, M.E., Levesley, A., Lima, A.J.N., Lisingo, J., Lopes, A.P., Lopez-Gonzalez, G., Lovejoy, T., Lovett, J.C., Lowe, R., Magnusson, W.E., Malumbres-Olarte, J., Manzatto, A.G., Marimon, B.H., Marshall, A.R., Marthews, T., de Almeida Reis, S.M., Maycock, C., Melgaço, K., Mendoza, C., Metali, F., Mihindou, V., Milliken, W., Mitchard, E.T.A., Morandi, P.S., Mossman, H.L., Nagy, L., Nascimento, H., Neill, D., Nilus, R., Vargas, P.N., Palacios, W., Camacho, N.P., Peacock, J., Pendry, C., Peñuela Mora, M.C., Pickavance, G.C., Pipoly, J., Pitman, N., Playfair, M., Poorter, L., Poulsen, J.R., Poulsen, A.D., Preziosi, R., Prieto, A., Primack, R.B., Ramirez-Angulo, H., Reitsma, J., Réjou-Méchain, M., Correa, Z.R., de Sousa, T.R., Bayona, L.R., Roopsind, A., Rudas, A., Rutishauser, E., Abu Salim, K., Salomão, R.P., Schiatti, J., Sheil, D., Silva, R.C., Espejo, J.S., Valeria, C.S., Silveira, M., Simo-Droissart, M., Simon, M.F., Singh, J., Soto Shareva, Y.C., Stahl, C., Stropp, J., Sukri, R., Sunderland, T., Svátek, M., Swaine, M.D., Swamy, V., Taedoumg, H., Talbot, J., Taplin, J., Taylor, D., Ter Steege, H., Terborgh, J., Thomas, R., Thomas, S.C., Torres-Lezama, A., Umunay, P., Gamarra, L.V., van der Heijden, G., van der Hout, P., van der Meer, P., van Nieuwstadt, M., Verbeek, H., Vernimmen, R., Vicentini, A., Vieira, I.C.G., Torre, E.V., Vleminckx, J., Vos, V., Wang, O., White, L.J.T., Willcock, S., Woods, J.T., Wortel, V., Young, K., Zagt, R., Zemagho, L., Zuidema, P.A., Zwerts, J.A. & Phillips, O.L. (2020) Long-term

- thermal sensitivity of Earth's tropical forests. *Science (New York, N.Y.)*, **368**, 869–874.
- Talbot, J., Lewis, S.L., Lopez-gonzalez, G., Brienen, R.J.W., Monteagudo, A., Baker, T.R., Feldpausch, T.R., Malhi, Y., Vanderwel, M., Araujo, A., Arroyo, L.P., Chao, K., Erwin, T., Heijden, G. Van Der, Keeling, H., Killeen, T., Neill, D., Núñez, P., Alexander, G., Gutierrez, P., Pitman, N., Alberto, C., Silveira, M. & Phillips, O.L. (2014) Forest Ecology and Management Methods to estimate aboveground wood productivity from long-term forest inventory plots. *Forest Ecology and Management*, **320**, 30–38.
- Tanco, R. & Kruse, E. (2001) Prediction of seasonal water table fluctuations in La Pampa and Buenos Aires, Argentina. *Hydrogeology Journal*, **9**, 339–347.
- Taylor, R.G., Scanlon, B., Döll, P., Rodell, M., van Beek, R., Wada, Y., Longuevergne, L., Leblanc, M., Famiglietti, J.S., Edmunds, M., Konikow, L., Green, T.R., Chen, J., Taniguchi, M., Bierkens, M.F.P., MacDonald, A., Fan, Y., Maxwell, R.M., Yechieli, Y., Gurdak, J.J., Allen, D.M., Shamsudduha, M., Hiscock, K., Yeh, P.J.-F., Holman, I. & Treidel, H. (2013) Ground water and climate change. *Nature Climate Change*, **3**, 322–329.
- Toledo, J.J., Magnusson, W.E., Castilho, C. V. & Nascimento, H.E.M. (2011a) How much variation in tree mortality is predicted by soil and topography in Central Amazonia? *Forest Ecology and Management*, **262**, 331–338.
- Toledo, M., Poorter, L., Peña-Claros, M., Alarcón, A., Balcázar, J., Leño, C., Licona, J.C., Llanque, O., Vroomans, V., Zuidema, P. & Bongers, F. (2011b) Climate is a stronger driver of tree and forest growth rates than soil and disturbance. *Journal of Ecology*, **99**, 254–264.
- Tomasella, J., Hodnett, M.G., Cuartas, L.A., Nobre, A.D., Waterloo, M.J. & Oliveira, S.M. (2008) The water balance of an Amazonian micro-catchment: the effect of interannual variability of rainfall on hydrological behaviour. *Hydrological Processes*, **22**, 2133–2147.
- Tomlinson, P.B. (1990) *The Structural Biology of Palms*.
- Vandeleur, R., Niemietz, C., Tilbrook, J. & Tyerman, S.D. (2005) Roles of aquaporins in root responses to irrigation. *Plant and Soil*, **274**.
- Vilanova, E., Ramírez-Angulo, H., Torres-Lezama, A., Aymard, G., Gámez, L., Durán, C., Hernández, L., Herrera, R., van der Heijden, G., Phillips, O.L. & Ettl, G.J. (2018) Environmental drivers of forest structure and stem turnover across Venezuelan tropical forests. *PLOS ONE*, **13**, e0198489.
- Waldhoff, D., Junk, W.J. & Furch, B. (1998) Responses of three central Amazonian tree species to drought and flooding under controlled conditions. *International Journal of Ecology and Environmental Sciences*, **24**, 237–252.
- Webb, W., Szarek, S., Lauenroth, W., Kinerson, R. & Smith, M. (1978) Primary Productivity and Water Use in Native Forest, Grassland, and Desert Ecosystems. *Ecology*, **59**, 1239–1247.
- Whittaker, R.H. (1975) *Communities and ecosystems - Second edition*.
- Wickham, H. (2011) ggplot2. *Wiley Interdisciplinary Reviews: Computational Statistics*, **3**.
- Zanne, A.E., Lopez-Gonzalez, G., Coomes, D. a., Ilic, J., Jansen, S., Lewis, S.L.S.L., Miller, R.B.B., Swenson, N.G.G., Wiemann, M.C.C. & Chave, J. (2009) *Data from: Towards a worldwide wood economics spectrum*.
- Zhao, M. & Running, S.W. (2010) Drought-induced reduction in global terrestrial net primary production from 2000 through 2009. *Science*, **329**, 940–943.
- Zipper, S.C., Soylu, M.E., Booth, E.G. & Loheide, S.P. (2015) Untangling the effects of

1
2
3 shallow groundwater and soil texture as drivers of subfield-scale yield variability.
4 *Water Resources Research*, **51**, 6338–6358.

5 Zuquim, G., Stropp, J., Moulatlet, G.M., Van doninck, J., Quesada, C.A., Figueiredo,
6 F.O.G., Costa, F.R.C., Ruokolainen, K. & Tuomisto, H. (2019) Making the most of
7 scarce data: Mapping soil gradients in data-poor areas using species occurrence
8 records. *Methods in Ecology and Evolution*, **10**, 788–801.
9
10
11
12
13
14
15
16
17
18
19
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21
22
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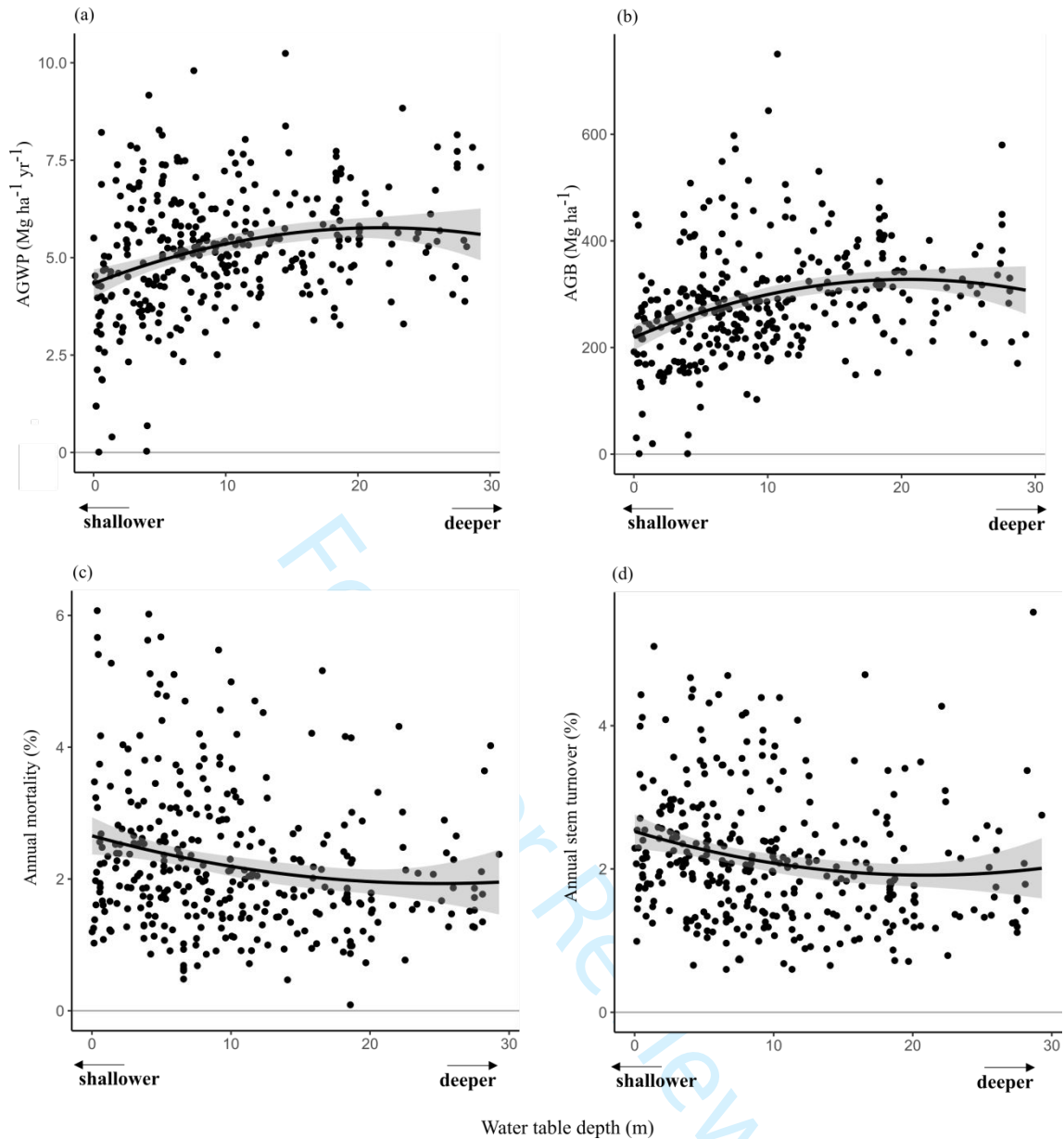


Figure 1. Impact of water table depth on (A) biomass productivity; (B) biomass stock; (C) mortality rate; and (D) stem turnover in Amazonian forests. LOESS regression was used to adjust the relationships between the response variables and WTD. The shaded region shows the confidence interval of the regression.

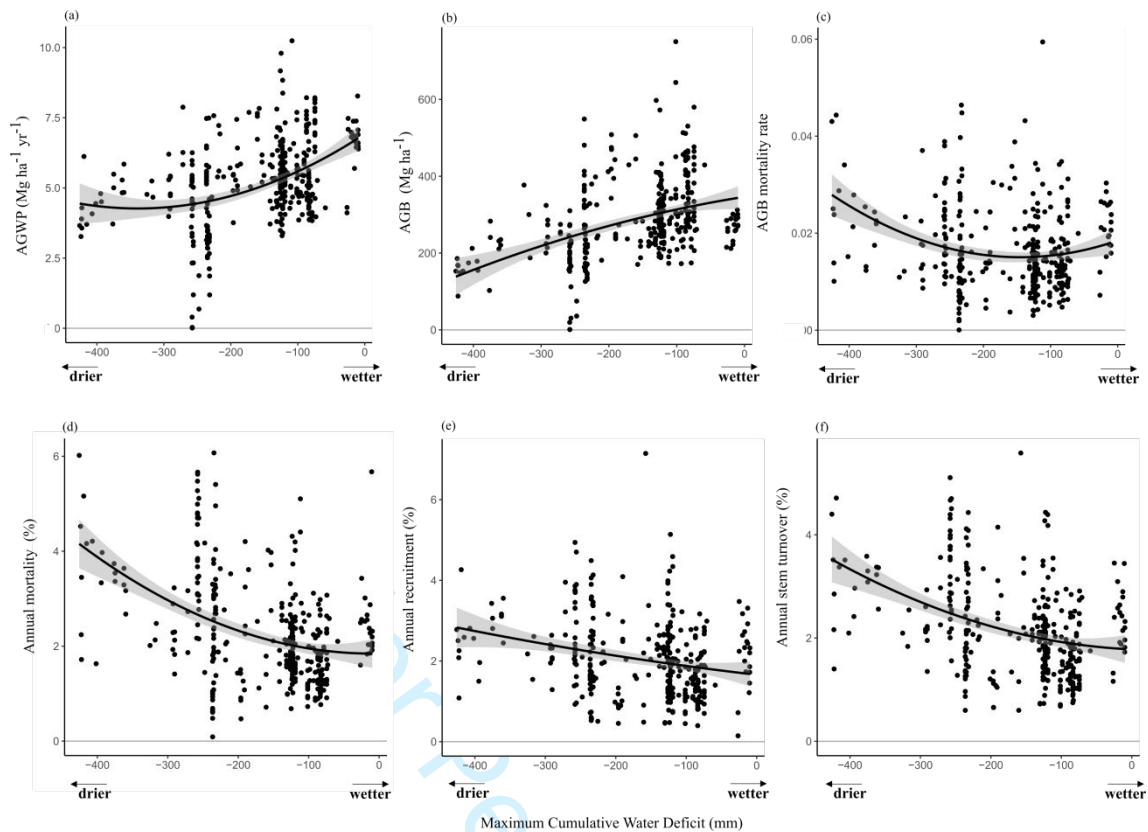
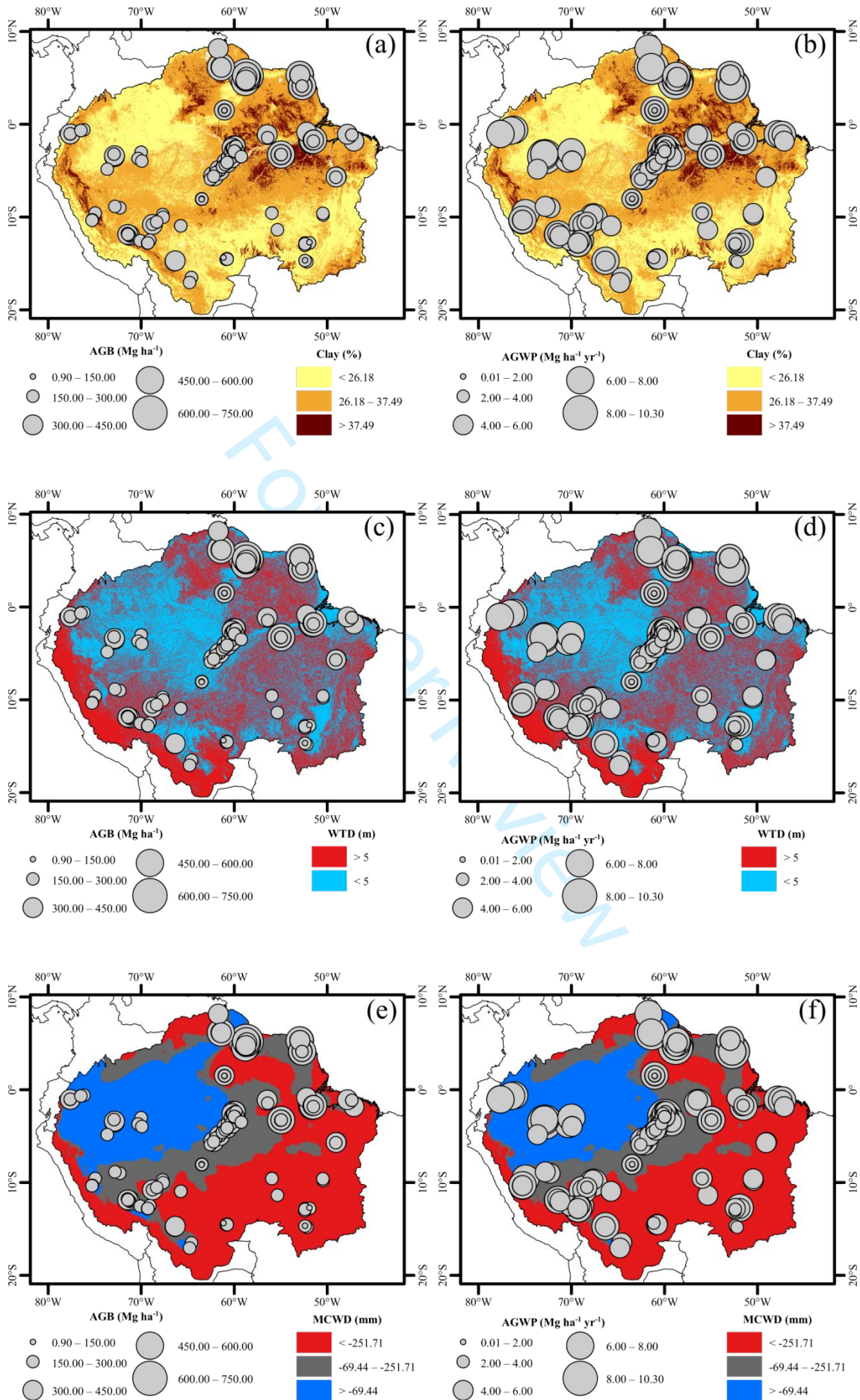


Figure 2. Impact of the maximum cumulative water deficit on (A) biomass productivity; (B) biomass stock; (C) biomass mortality rate; (D) mortality rate; (E) recruitment rate; and (F) stem turnover in Amazonian forests. LOESS regression was used to adjust the relationships between the response variables and MCWD. The shaded region shows the confidence interval of the regression.



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3 **Figure 3.** Spatial variation in forest biomass and productivity across Amazonia. Plots a
4 and b display these metrics against a background of clay content; c and d the WTD
5 background; and e and f the MCWD background. The clay content and MCWD classes
6 were defined based on the standard deviation around the mean of each of these
7 variables, shallow and deep water tables follow the definitions of Fan & Miguez-Macho
8 (2010). These classes are the same those used in Figures 4 and 5. Gray dots represent
9 plots with size proportional to the biomass stock or productivity.
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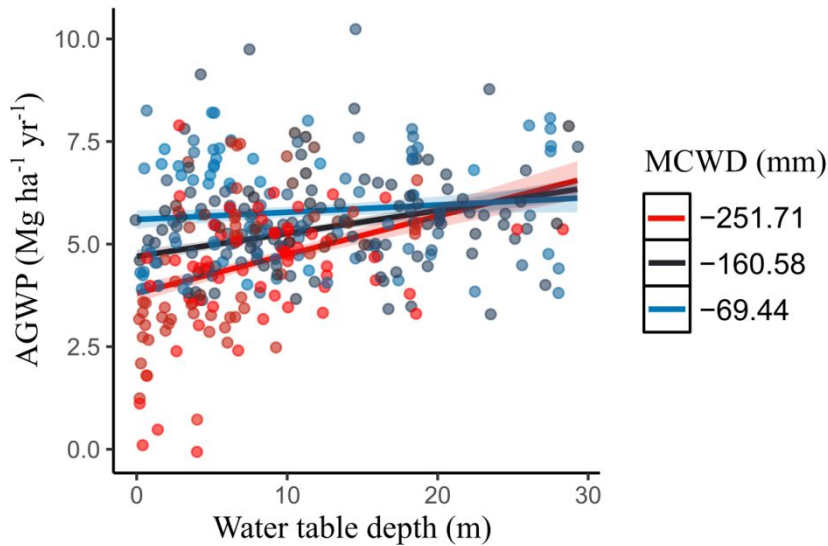


Figure 4. Partial-dependence plot of the interaction between MCWD and water table depth on biomass productivity. In order to visualize interactions, climate was divided in three classes based on the standard deviation around the mean. Red colour is for plots with MCWD values less than one standard deviation below the mean; black is for plots with MCWD values within one standard deviation of the mean; and blue is for plots with MCWD values greater than one standard deviation above the mean. Shaded regions represent confidence intervals.

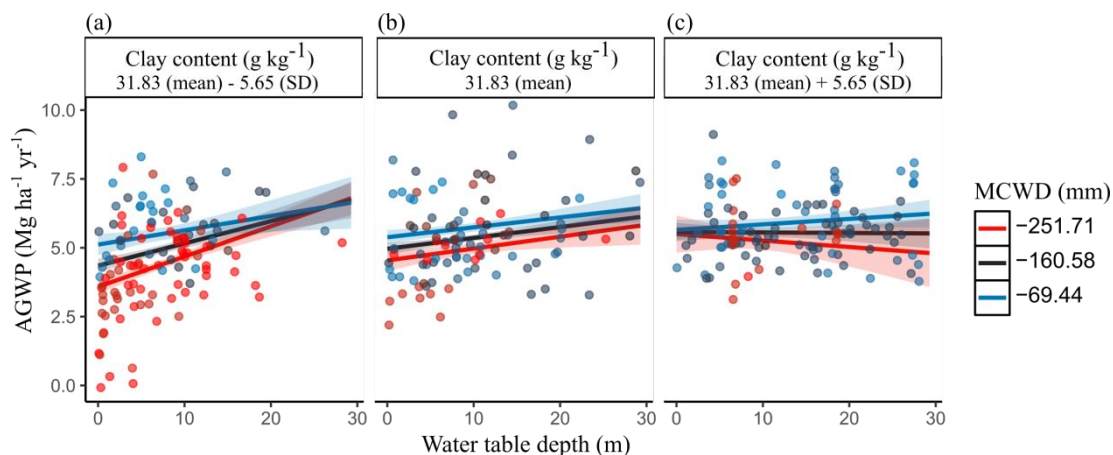


Figure 5. Partial-dependence plots derived from multiple-regression models investigating the effects of interactions among clay content, water table depth and MCWD on biomass productivity in Amazonian forests. (a) Partial plots of the interaction in less clayey soil; (b) Partial effect of the interaction in moderately clayey soil; and (c) Partial effect of the interaction in more clayey soil. In order to visualize interactions, climate and soil texture were divided in three classes based on the standard deviation around the mean. Red colour is for plots with MCWD values less than one standard deviation below the mean; black is for plots with MCWD values within one standard deviation of the mean; and blue is for plots with MCWD values greater than one standard deviation above the mean. Shaded regions represent confidence intervals.

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Supporting Information

Table S1 – List of permanent Amazon forest monitoring plots used in this study.

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|--------|--------|--------------|-----------|------------|
| Bolivia | BEE-01 | -16.53 | -64.58 | 180 | 1 | 4 |
| Bolivia | BEE-05 | -16.53 | -64.58 | 180 | 1 | 4 |
| Bolivia | CHO-01 | -14.39 | -61.15 | 170 | 1 | 2 |
| Bolivia | EBB-05 | -14.76 | -66.34 | 210 | 1 | 2 |
| Bolivia | EBB-09 | -14.73 | -66.32 | 220 | 1 | 2 |
| Bolivia | HCC-21 | -14.53 | -60.74 | 729 | 1 | 7 |
| Bolivia | HCC-22 | -14.53 | -60.73 | 747 | 1 | 7 |
| Bolivia | HCC-23 | -14.56 | -60.75 | 809 | 1 | 5 |
| Bolivia | HCC-24 | -14.57 | -60.75 | 735 | 1 | 5 |
| Bolivia | LFB-01 | -14.58 | -60.83 | 245 | 1 | 8 |
| Bolivia | LFB-02 | -14.58 | -60.83 | 227 | 1 | 9 |
| Bolivia | RET-05 | -10.97 | -65.72 | 160 | 1 | 10 |
| Bolivia | RET-06 | -10.97 | -65.72 | 160 | 1 | 10 |
| Bolivia | RET-08 | -10.97 | -65.72 | 160 | 1 | 10 |
| Bolivia | RET-09 | -10.97 | -65.72 | 160 | 1 | 10 |
| Bolivia | SCT-01 | -17.09 | -64.77 | 248 | 1 | 7 |
| Brazil | ALF-01 | -9.6 | -55.94 | 269 | 1 | 5 |
| Brazil | ALF-02 | -9.58 | -55.92 | 277 | 1 | 4 |
| Brazil | BDF-01 | -2.34 | -60.1 | 75 | 2 | 6 |
| Brazil | BDF-03 | -2.42 | -59.85 | 75 | 1 | 6 |
| Brazil | BDF-04 | -2.43 | -59.85 | 75 | 1 | 6 |
| Brazil | BDF-05 | -2.43 | -59.85 | 75 | 1 | 6 |
| Brazil | BDF-06 | -2.41 | -59.86 | 75 | 3 | 6 |
| Brazil | BDF-07 | -2.4 | -59.9 | 75 | 1 | 6 |
| Brazil | BDF-08 | -2.4 | -59.9 | 75 | 1 | 6 |
| Brazil | BDF-09 | -2.4 | -59.85 | 75 | 1 | 5 |
| Brazil | BDF-10 | -2.39 | -59.86 | 75 | 2 | 5 |
| Brazil | BDF-11 | -2.38 | -59.85 | 75 | 3 | 5 |
| Brazil | BDF-12 | -2.39 | -59.85 | 75 | 2 | 5 |
| Brazil | BDF-13 | -2.4 | -59.91 | 75 | 9 | 6 |
| Brazil | BDF-14 | -2.36 | -59.97 | 75 | 1 | 7 |
| Brazil | BNT-01 | -2.64 | -60.16 | 73 | 1 | 22 |
| Brazil | BNT-02 | -2.64 | -60.15 | 73 | 1 | 22 |
| Brazil | BNT-04 | -2.63 | -60.15 | 73 | 1 | 21 |
| Brazil | BNT-05 | -2.63 | -60.17 | 73 | 1 | 5 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|--------|--------|--------------|-----------|------------|
| Brazil | BNT-06 | -2.63 | -60.17 | 73 | 1 | 5 |
| Brazil | BNT-07 | -2.63 | -60.17 | 73 | 1 | 5 |
| Brazil | CAX-06 | -1.72 | -51.46 | 20 | 1 | 6 |
| Brazil | CAX-08 | -1.85 | -51.47 | 10 | 1 | 8 |
| Brazil | CNA-10 | -8.07 | -63.49 | 89 | 1 | 2 |
| Brazil | CNA-11 | -8.07 | -63.48 | 89 | 1 | 2 |
| Brazil | CNA-12 | -8.07 | -63.47 | 86 | 0.992 | 2 |
| Brazil | CNA-13 | -8.07 | -63.46 | 89 | 0.966 | 2 |
| Brazil | CNA-14 | -8.07 | -63.45 | 75 | 0.981 | 2 |
| Brazil | CNA-20 | -8.08 | -63.49 | 87 | 0.99 | 2 |
| Brazil | CNA-21 | -8.08 | -63.48 | 87 | 0.999 | 2 |
| Brazil | CNA-22 | -8.08 | -63.47 | 87 | 0.999 | 2 |
| Brazil | CNA-23 | -8.08 | -63.46 | 86 | 0.989 | 2 |
| Brazil | CNA-24 | -8.08 | -63.45 | 73 | 0.989 | 2 |
| Brazil | CNA-30 | -8.09 | -63.49 | 86 | 0.979 | 2 |
| Brazil | CNA-31 | -8.09 | -63.48 | 83 | 0.951 | 2 |
| Brazil | CNA-32 | -8.09 | -63.47 | 87 | 0.997 | 2 |
| Brazil | CNA-33 | -8.09 | -63.46 | 78 | 0.951 | 2 |
| Brazil | CNA-34 | -8.09 | -63.45 | 72 | 0.967 | 2 |
| Brazil | CNA-40 | -8.1 | -63.49 | 73 | 0.969 | 2 |
| Brazil | CNA-41 | -8.1 | -63.48 | 83 | 0.977 | 2 |
| Brazil | CNA-42 | -8.1 | -63.47 | 78 | 0.98 | 2 |
| Brazil | CNA-43 | -8.1 | -63.46 | 78 | 0.998 | 2 |
| Brazil | CNA-44 | -8.1 | -63.45 | 73 | 1 | 2 |
| Brazil | CNA-50 | -8.11 | -63.49 | 84 | 0.981 | 2 |
| Brazil | CNA-51 | -8.11 | -63.48 | 82 | 0.995 | 2 |
| Brazil | CNA-52 | -8.11 | -63.47 | 74 | 0.992 | 2 |
| Brazil | CNA-53 | -8.11 | -63.46 | 67 | 0.985 | 2 |
| Brazil | CNA-54 | -8.11 | -63.45 | 66 | 0.976 | 2 |
| Brazil | CNA-60 | -8.12 | -63.49 | 86 | 0.993 | 2 |
| Brazil | CNA-61 | -8.12 | -63.48 | 83 | 0.971 | 2 |
| Brazil | CNA-62 | -8.12 | -63.47 | 80 | 0.996 | 2 |
| Brazil | CNA-64 | -8.12 | -63.45 | 77 | 0.998 | 2 |
| Brazil | CPP-01 | -1.84 | -47.1 | 73 | 1 | 2 |
| Brazil | DOI-01 | -10.57 | -68.32 | 203 | 1 | 10 |
| Brazil | DOI-02 | -10.55 | -68.31 | 203 | 1 | 8 |
| Brazil | DUK-10 | -2.93 | -59.97 | 85 | 1 | 3 |
| Brazil | DUK-11 | -2.93 | -59.96 | 108 | 1 | 3 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|-------|--------|--------------|-----------|------------|
| Brazil | DUK-12 | -2.92 | -59.95 | 109 | 1 | 3 |
| Brazil | DUK-13 | -2.92 | -59.94 | 104 | 1 | 3 |
| Brazil | DUK-14 | -2.92 | -59.93 | 89 | 1 | 3 |
| Brazil | DUK-15 | -2.92 | -59.92 | 70 | 1 | 3 |
| Brazil | DUK-16 | -2.92 | -59.91 | 59 | 1 | 3 |
| Brazil | DUK-17 | -2.92 | -59.9 | 54 | 1 | 3 |
| Brazil | DUK-21 | -2.94 | -59.97 | 61 | 1 | 3 |
| Brazil | DUK-22 | -2.94 | -59.96 | 98 | 1.1 | 4 |
| Brazil | DUK-23 | -2.93 | -59.95 | 75 | 1 | 6 |
| Brazil | DUK-24 | -2.93 | -59.94 | 81 | 1.1 | 3 |
| Brazil | DUK-25 | -2.93 | -59.93 | 65 | 1 | 3 |
| Brazil | DUK-26 | -2.93 | -59.92 | 96 | 1 | 3 |
| Brazil | DUK-27 | -2.93 | -59.91 | 59 | 1 | 3 |
| Brazil | DUK-28 | -2.93 | -59.9 | 45 | 1 | 3 |
| Brazil | DUK-31 | -2.94 | -59.96 | 81 | 1.1 | 3 |
| Brazil | DUK-32 | -2.94 | -59.95 | 61 | 1 | 6 |
| Brazil | DUK-33 | -2.94 | -59.94 | 53 | 1 | 6 |
| Brazil | DUK-34 | -2.94 | -59.94 | 55 | 1 | 6 |
| Brazil | DUK-35 | -2.94 | -59.93 | 88 | 1 | 6 |
| Brazil | DUK-36 | -2.94 | -59.92 | 74 | 1 | 3 |
| Brazil | DUK-37 | -2.94 | -59.91 | 40 | 1 | 3 |
| Brazil | DUK-38 | -2.94 | -59.9 | 51 | 1 | 3 |
| Brazil | DUK-41 | -2.95 | -59.96 | 46 | 1 | 6 |
| Brazil | DUK-42 | -2.95 | -59.95 | 55 | 1 | 3 |
| Brazil | DUK-43 | -2.95 | -59.94 | 100 | 1 | 6 |
| Brazil | DUK-44 | -2.95 | -59.93 | 105 | 1 | 6 |
| Brazil | DUK-45 | -2.95 | -59.93 | 98 | 1 | 6 |
| Brazil | DUK-46 | -2.95 | -59.92 | 92 | 1 | 3 |
| Brazil | DUK-47 | -2.95 | -59.91 | 66 | 1 | 3 |
| Brazil | DUK-48 | -2.95 | -59.91 | 66 | 1 | 3 |
| Brazil | DUK-51 | -2.96 | -59.96 | 56 | 1 | 3 |
| Brazil | DUK-52 | -2.96 | -59.95 | 93 | 1 | 3 |
| Brazil | DUK-53 | -2.96 | -59.94 | 73 | 1 | 6 |
| Brazil | DUK-56 | -2.96 | -59.91 | 60 | 1 | 6 |
| Brazil | DUK-57 | -2.95 | -59.91 | 90 | 1 | 3 |
| Brazil | DUK-58 | -2.95 | -59.9 | 39 | 1 | 3 |
| Brazil | DUK-62 | -2.97 | -59.95 | 53 | 1 | 5 |
| Brazil | DUK-63 | -2.97 | -59.94 | 60 | 1 | 6 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|--------|--------|--------------|-----------|------------|
| Brazil | DUK-64 | -2.97 | -59.93 | 100 | 1 | 3 |
| Brazil | DUK-65 | -2.97 | -59.92 | 75 | 1 | 3 |
| Brazil | DUK-66 | -2.96 | -59.91 | 77 | 1 | 3 |
| Brazil | DUK-67 | -2.96 | -59.9 | 53 | 1 | 3 |
| Brazil | DUK-68 | -2.96 | -59.9 | 60 | 1 | 3 |
| Brazil | DUK-71 | -2.98 | -59.96 | 58 | 1 | 5 |
| Brazil | DUK-72 | -2.98 | -59.95 | 84 | 1 | 5 |
| Brazil | DUK-73 | -2.98 | -59.94 | 101 | 1 | 3 |
| Brazil | DUK-74 | -2.98 | -59.93 | 92 | 1 | 6 |
| Brazil | DUK-75 | -2.97 | -59.92 | 100 | 1 | 3 |
| Brazil | DUK-76 | -2.97 | -59.91 | 100 | 1 | 3 |
| Brazil | DUK-77 | -2.97 | -59.9 | 95 | 1 | 3 |
| Brazil | DUK-78 | -2.97 | -59.89 | 63 | 1 | 3 |
| Brazil | DUK-81 | -2.99 | -59.96 | 52 | 1 | 6 |
| Brazil | DUK-82 | -2.99 | -59.95 | 49 | 1 | 5 |
| Brazil | DUK-83 | -2.99 | -59.94 | 63 | 1 | 3 |
| Brazil | DUK-84 | -2.98 | -59.93 | 56 | 1 | 3 |
| Brazil | DUK-85 | -2.98 | -59.92 | 102 | 1 | 3 |
| Brazil | DUK-87 | -2.98 | -59.9 | 47 | 1 | 3 |
| Brazil | DUK-88 | -2.98 | -59.89 | 95 | 1 | 3 |
| Brazil | DUK-91 | -3 | -59.95 | 60 | 1 | 3 |
| Brazil | DUK-92 | -3 | -59.94 | 87 | 1 | 3 |
| Brazil | DUK-93 | -3 | -59.94 | 87 | 1 | 3 |
| Brazil | DUK-94 | -2.99 | -59.93 | 56 | 1 | 3 |
| Brazil | DUK-95 | -2.99 | -59.92 | 85 | 1 | 3 |
| Brazil | DUK-96 | -2.99 | -59.91 | 57 | 1 | 3 |
| Brazil | DUK-98 | -2.99 | -59.89 | 97 | 1 | 3 |
| Brazil | FEC-01 | -10.07 | -67.62 | 204 | 1 | 5 |
| Brazil | FLO-01 | -12.81 | -51.85 | 377 | 1 | 5 |
| Brazil | FLO-02 | -12.75 | -51.88 | 366 | 1 | 4 |
| Brazil | GMT-01 | -1.11 | -47.8 | 50 | 1 | 3 |
| Brazil | IPM-20 | -3.69 | -60.33 | 50 | 1 | 3 |
| Brazil | IPM-21 | -4.62 | -61.24 | NA | 1 | 3 |
| Brazil | IPM-22 | -3.68 | -60.32 | 44 | 1 | 3 |
| Brazil | IPM-25 | -3.67 | -60.31 | 44 | 1 | 3 |
| Brazil | IPM-26 | -4.98 | -61.57 | 61 | 1 | 3 |
| Brazil | IPM-27 | -4.99 | -61.56 | 50 | 1 | 3 |
| Brazil | IPM-28 | -3.67 | -60.3 | 44 | 1 | 3 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|--------|--------|--------------|-----------|------------|
| Brazil | IPM-37 | -4.16 | -60.72 | 37 | 1 | 3 |
| Brazil | IPM-39 | -4.14 | -60.73 | 40 | 1 | 3 |
| Brazil | IPM-40 | -4.15 | -60.73 | 30 | 1 | 3 |
| Brazil | IPM-42 | -4.38 | -60.95 | 45 | 1 | 3 |
| Brazil | IPM-43 | -4.38 | -60.94 | 48 | 1 | 3 |
| Brazil | IPM-46 | -4.4 | -60.92 | 46 | 1 | 3 |
| Brazil | IPM-55 | -4.6 | -61.26 | 51 | 1 | 3 |
| Brazil | IPM-62 | -4.99 | -61.56 | 57 | 1 | 3 |
| Brazil | IPM-63 | -5 | -61.55 | 58 | 1 | 3 |
| Brazil | IPM-64 | -5 | -61.54 | 59 | 1 | 3 |
| Brazil | IPM-79 | -5.25 | -61.96 | 62 | 1 | 3 |
| Brazil | IPM-80 | -5.25 | -61.96 | 63 | 1 | 3 |
| Brazil | IPM-86 | -5.63 | -62.19 | 69 | 1 | 3 |
| Brazil | IPM-87 | -5.63 | -62.19 | 65 | 1 | 3 |
| Brazil | IPM-88 | -5.64 | -62.18 | 70 | 1 | 3 |
| Brazil | IPM-98 | -5.95 | -62.51 | 69 | 1 | 3 |
| Brazil | IPM-99 | -5.94 | -62.52 | 70 | 1 | 3 |
| Brazil | JAC-01 | -2.61 | -60.21 | 73 | 5 | 8 |
| Brazil | JAC-02 | -2.62 | -60.2 | 73 | 5 | 8 |
| Brazil | JRI-01 | -0.89 | -52.19 | 150 | 1 | 6 |
| Brazil | MRB-01 | -5.73 | -49.05 | 90 | 2 | 4 |
| Brazil | MRB-02 | -5.72 | -49.03 | 90 | 2 | 4 |
| Brazil | MRB-03 | -5.7 | -49 | 90 | 2 | 4 |
| Brazil | MTH-01 | -8.88 | -72.79 | 246 | 1 | 6 |
| Brazil | NOC-01 | -3.52 | -59.27 | 14 | 0.25 | 4 |
| Brazil | NOC-02 | -3.52 | -59.27 | 14 | 0.25 | 4 |
| Brazil | NOC-03 | -3.52 | -59.27 | 17 | 0.25 | 4 |
| Brazil | NXV-02 | -14.7 | -52.35 | 250 | 0.5 | 3 |
| Brazil | NXV-06 | -14.72 | -52.36 | 346 | 0.47 | 6 |
| Brazil | POR-01 | -10.82 | -68.77 | 268 | 1 | 9 |
| Brazil | POR-02 | -10.8 | -68.77 | 268 | 1 | 8 |
| Brazil | PPB-01 | -1.18 | -47.32 | 50 | 1 | 2 |
| Brazil | PPB-02 | -1.18 | -47.32 | 50 | 1 | 2 |
| Brazil | PPB-03 | -1.18 | -47.32 | 50 | 1 | 2 |
| Brazil | PTB-01 | -1.17 | -56.41 | 180 | 1 | 2 |
| Brazil | PTB-02 | -1.48 | -56.39 | 40 | 1 | 2 |
| Brazil | RBR-01 | -5.93 | -62.52 | NA | 1 | 3 |
| Brazil | RFH-01 | -9.75 | -67.67 | 176 | 1 | 6 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|--------|--------|--------------|-----------|------------|
| Brazil | RST-01 | -9.04 | -72.27 | 279 | 1 | 6 |
| Brazil | SAA-01 | -9.79 | -50.43 | 177 | 1 | 4 |
| Brazil | SAA-02 | -9.64 | -50.45 | 207 | 1 | 3 |
| Brazil | SIP-01 | -11.41 | -55.32 | 385 | 0.76 | 3 |
| Brazil | STO-03 | -3.3 | -54.96 | 132 | 0.25 | 3 |
| Brazil | STO-05 | -3.34 | -54.98 | 170 | 0.25 | 3 |
| Brazil | STO-06 | -3.34 | -54.96 | 178 | 0.25 | 3 |
| Brazil | STO-07 | -3.32 | -54.96 | 163 | 0.25 | 3 |
| Brazil | TAN-02 | -13.08 | -52.38 | 382 | 1 | 5 |
| Brazil | TAN-03 | -12.83 | -52.35 | 356 | 1 | 5 |
| Brazil | TAN-04 | -12.92 | -52.37 | 389 | 1 | 5 |
| Brazil | TAP-50 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-51 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-52 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-53 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-54 | -3.31 | -54.95 | 100 | 0.25 | 3 |
| Brazil | TAP-55 | -3.31 | -54.95 | 100 | 0.25 | 3 |
| Brazil | TAP-56 | -3.31 | -54.95 | 100 | 0.25 | 3 |
| Brazil | TAP-57 | -3.31 | -54.95 | 100 | 0.25 | 3 |
| Brazil | TAP-58 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-59 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-60 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TAP-61 | -3.31 | -54.94 | 100 | 0.25 | 3 |
| Brazil | TEC-01 | -1.71 | -51.46 | 15 | 1 | 11 |
| Brazil | TEC-02 | -1.74 | -51.49 | 15 | 1 | 11 |
| Brazil | TEC-03 | -1.73 | -51.51 | 15 | 1 | 11 |
| Brazil | TEC-04 | -1.75 | -51.52 | 15 | 1 | 11 |
| Brazil | TEC-05 | -1.78 | -51.59 | 15 | 1 | 11 |
| Brazil | TEC-06 | -1.73 | -51.43 | 15 | 1 | 11 |
| Brazil | TEM-01 | -2.97 | -59.9 | 120 | 1 | 10 |
| Brazil | TEM-02 | -2.93 | -59.95 | 120 | 1 | 10 |
| Brazil | TEM-03 | -2.41 | -59.9 | 100 | 1 | 9 |
| Brazil | TEM-04 | -2.43 | -59.79 | 100 | 1 | 9 |
| Brazil | TEM-05 | -2.62 | -60.21 | 100 | 1 | 9 |
| Brazil | TEM-06 | -2.6 | -60.11 | 100 | 1 | 10 |
| Brazil | TIC-01 | -3.54 | -59.22 | 13 | 0.25 | 4 |
| Brazil | TIC-03 | -3.53 | -59.22 | 12 | 0.25 | 4 |
| Brazil | VCR-02 | -14.83 | -52.17 | 297 | 0.6 | 6 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|----------|-----------|-------|--------|--------------|-----------|------------|
| Brazil | VIR-11 | 1.49 | -61.01 | 94 | 1 | 3 |
| Brazil | VIR-12 | 1.49 | -61.02 | 53 | 1 | 3 |
| Brazil | VIR-13 | 1.49 | -61.03 | 50 | 1 | 3 |
| Brazil | VIR-14 | 1.49 | -61.03 | 52 | 1 | 3 |
| Brazil | VIR-15 | 1.49 | -61.04 | 48 | 1 | 3 |
| Brazil | VIR-22 | 1.48 | -61.02 | 130 | 1 | 3 |
| Brazil | VIR-23 | 1.48 | -61.03 | 58 | 1 | 3 |
| Brazil | VIR-24 | 1.48 | -61.03 | 49 | 1 | 3 |
| Brazil | VIR-25 | 1.48 | -61.04 | 49 | 1 | 3 |
| Brazil | VIR-31 | 1.47 | -61.01 | 55 | 0.5 | 3 |
| Brazil | VIR-32 | 1.47 | -61.02 | 56 | 1 | 3 |
| Brazil | VIR-33 | 1.47 | -61.03 | 58 | 1 | 3 |
| Brazil | VIR-34 | 1.47 | -61.03 | 56 | 1 | 3 |
| Brazil | VIR-35 | 1.47 | -61.04 | 49 | 0.5 | 3 |
| Brazil | VIR-41 | 1.46 | -61.01 | 52 | 0.9 | 3 |
| Brazil | VIR-42 | 1.46 | -61.02 | 55 | 1 | 3 |
| Brazil | VIR-43 | 1.46 | -61.03 | 57 | 1 | 3 |
| Brazil | VIR-45 | 1.46 | -61.04 | 50 | 0.5 | 3 |
| Brazil | VIR-51 | 1.45 | -61.01 | 50 | 1 | 3 |
| Brazil | VIR-52 | 1.45 | -61.02 | 50 | 1 | 3 |
| Brazil | VIR-53 | 1.45 | -61.03 | 49 | 1 | 3 |
| Brazil | VIR-54 | 1.45 | -61.03 | 48 | 1 | 3 |
| Brazil | VIR-55 | 1.45 | -61.04 | 48 | 1 | 3 |
| Brazil | VIR-61 | 1.44 | -61.01 | 52 | 1 | 3 |
| Brazil | VIR-62 | 1.44 | -61.02 | 49 | 1 | 3 |
| Brazil | VIR-63 | 1.44 | -61.03 | 49 | 1 | 3 |
| Brazil | VIR-65 | 1.44 | -61.04 | 50 | 1 | 3 |
| Colombia | AGP-01 | -3.72 | -70.31 | 120 | 1 | 5 |
| Colombia | AGP-02 | -3.72 | -70.3 | 120 | 1 | 5 |
| Colombia | LOR-01 | -3.06 | -69.99 | 94 | 1 | 3 |
| Colombia | LOR-02 | -3.06 | -69.99 | 93 | 0.52 | 4 |
| Colombia | LOR-03 | -3.06 | -69.99 | 93 | 0.48 | 3 |
| Colombia | ZAR-03 | -3.99 | -69.9 | 130 | 1 | 4 |
| Colombia | ZAR-04 | -3.99 | -69.91 | 146 | 1 | 3 |
| Ecuador | BOG-01 | -0.7 | -76.48 | 257 | 1 | 6 |
| Ecuador | BOG-02 | -0.7 | -76.47 | 284 | 1 | 6 |
| Ecuador | JAS-02 | -1.07 | -77.62 | 452 | 1 | 9 |
| Ecuador | JAS-03 | -1.08 | -77.61 | 384 | 1 | 9 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------------|-----------|-------|--------|--------------|-----------|------------|
| Ecuador | JAS-04 | -1.07 | -77.61 | 430 | 0.96 | 6 |
| Ecuador | TIP-02 | -0.63 | -76.14 | 243 | 0.8 | 6 |
| French Guiana | NOU-01 | 4.09 | -52.67 | 110 | 1 | 5 |
| French Guiana | NOU-02 | 4.09 | -52.67 | 110 | 1 | 5 |
| French Guiana | NOU-03 | 4.09 | -52.68 | 110 | 1 | 5 |
| French Guiana | NOU-04 | 4.09 | -52.68 | 110 | 1 | 5 |
| French Guiana | NOU-05 | 4.09 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-06 | 4.09 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-07 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-08 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-09 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-10 | 4.09 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-11 | 4.08 | -52.68 | 110 | 1 | 5 |
| French Guiana | NOU-12 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-13 | 4.08 | -52.68 | 127 | 1 | 4 |
| French Guiana | NOU-14 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-15 | 4.08 | -52.68 | 118 | 1 | 4 |
| French Guiana | NOU-16 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-17 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-18 | 4.08 | -52.68 | 123 | 1 | 4 |
| French Guiana | NOU-19 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-21 | 4.08 | -52.68 | 110 | 1 | 4 |
| French Guiana | NOU-22 | 4.08 | -52.67 | 110 | 1 | 4 |
| French Guiana | PAB-01 | 5.27 | -52.92 | 40 | 6.25 | 28 |
| French Guiana | PAR-20 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-21 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-22 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-23 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-24 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-25 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-26 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-27 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-28 | 5.28 | -52.92 | 20 | 0.49 | 7 |
| French Guiana | PAR-29 | 5.28 | -52.92 | 35 | 0.49 | 7 |
| Guyana | FMH-01 | 5.17 | -58.69 | 98 | 1 | 8 |
| Guyana | FMH-02 | 5.17 | -58.69 | 122 | 1 | 4 |
| Guyana | FMH-03 | 5.18 | -58.7 | 115 | 1 | 4 |
| Guyana | IWO-03 | 4.53 | -58.78 | 100 | 1 | 3 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|---------|-----------|--------|--------|--------------|-----------|------------|
| Guyana | IWO-09 | 4.61 | -58.73 | 116 | 1 | 3 |
| Guyana | IWO-12 | 4.73 | -58.72 | 61 | 1 | 3 |
| Guyana | IWO-21 | 4.63 | -58.74 | 97 | 1 | 5 |
| Guyana | IWO-22 | 4.62 | -58.72 | 139 | 1 | 5 |
| Guyana | PIB-05 | 5.02 | -58.62 | 93 | 1 | 4 |
| Guyana | PIB-06 | 5.01 | -58.62 | 81 | 1 | 4 |
| Guyana | PIB-12 | 5.03 | -58.6 | 94 | 1 | 4 |
| Peru | ALM-01 | -11.8 | -71.47 | 400 | 2 | 5 |
| Peru | BAR-01 | -11.9 | -71.42 | 345 | 0.813 | 5 |
| Peru | CUZ-01 | -12.54 | -69.06 | 190 | 1 | 9 |
| Peru | CUZ-02 | -12.54 | -69.06 | 190 | 1 | 9 |
| Peru | CUZ-03 | -12.53 | -69.05 | 190 | 1 | 9 |
| Peru | CUZ-04 | -12.54 | -69.05 | 190 | 1 | 9 |
| Peru | IND-01 | -3.52 | -72.85 | 111 | 1 | 2 |
| Peru | JEN-11 | -4.88 | -73.63 | 151 | 1 | 8 |
| Peru | LAS-02 | -12.57 | -70.09 | 235 | 1 | 4 |
| Peru | MNU-03 | -11.9 | -71.4 | 312 | 2 | 5 |
| Peru | MNU-04 | -11.9 | -71.4 | 358 | 1 | 6 |
| Peru | MNU-08 | -12 | -71.24 | 295 | 2 | 6 |
| Peru | MNU-09 | -12.04 | -71.21 | 332 | 2 | 5 |
| Peru | PAK-01 | -11.94 | -71.28 | 345 | 1 | 4 |
| Peru | PNY-04 | -10.34 | -75.25 | 414 | 1 | 5 |
| Peru | PNY-05 | -10.35 | -75.25 | 448 | 1.002 | 5 |
| Peru | PNY-07 | -10.35 | -75.26 | 414 | 1 | 5 |
| Peru | PNY-22 | -10.38 | -75.26 | 463 | 1.026 | 4 |
| Peru | RCS-05 | -9.62 | -74.93 | 251 | 1 | 3 |
| Peru | SUC-01 | -3.25 | -72.91 | 107 | 1 | 10 |
| Peru | SUC-02 | -3.25 | -72.9 | 98 | 1 | 10 |
| Peru | SUC-04 | -3.25 | -72.89 | 107 | 1 | 8 |
| Peru | SUC-05 | -3.26 | -72.89 | 118 | 1 | 8 |
| Peru | TAM-01 | -12.84 | -69.29 | 205 | 1 | 10 |
| Peru | TAM-02 | -12.83 | -69.29 | 210 | 1 | 13 |
| Peru | TAM-04 | -12.84 | -69.28 | 210 | 0.42 | 9 |
| Peru | TAM-05 | -12.83 | -69.27 | 220 | 1 | 12 |
| Peru | TAM-06 | -12.84 | -69.3 | 200 | 0.96 | 12 |
| Peru | TAM-07 | -12.83 | -69.26 | 225 | 1 | 10 |
| Peru | TAM-08 | -12.83 | -69.27 | 220 | 1 | 6 |
| Peru | TAM-09 | -12.83 | -69.28 | 199 | 1 | 5 |

| Country | Plot Code | Lat. | Long. | Alt. (m asl) | Area (ha) | N censuses |
|-----------|-----------|-------|--------|--------------|-----------|------------|
| Peru | YAN-02 | -3.43 | -72.84 | 109 | 1 | 7 |
| Venezuela | ELD-01 | 6.11 | -61.41 | 220 | 0.25 | 26 |
| Venezuela | ELD-02 | 6.11 | -61.41 | 244 | 0.25 | 26 |
| Venezuela | ELD-04 | 6.09 | -61.35 | 366 | 0.25 | 26 |
| Venezuela | RIO-01 | 8.11 | -61.69 | 312 | 0.25 | 25 |

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Table S2 – Statistical summary of the multiple linear models tested to investigate the effects of environmental variables on vegetation structure and dynamics of Amazon forests. Standardized beta coefficients for the simple effects of variables (but not for interactions) are shown. Significant effects are shown in bold.

| Forest metric | Regression coefficients for predictors | P | R ² |
|-----------------------|--|------------------|----------------|
| AGWP | 0.45 MCWD | <0.001 | |
| | 0.18 water table depth | 0.008 | |
| | 0.39 clay content | <0.001 | |
| | -0.09 maximum temperature | 0.33 | |
| | 0.51 soil fertility | <0.001 | 0.36 |
| | -0.0004 water table depth*MCWD | <0.001 | |
| | 0.00007 water table depth*MCWD*clay | 0.01 | |
| | -0.00009 MCWD*clay | 0.55 | |
| AGB stock | 20.25 MCWD | <0.001 | |
| | 14.19 water table depth | 0.003 | |
| | 23.87 clay content | <0.001 | |
| | -18.06 maximum temperature | 0.007 | |
| | -4.20 soil fertility | 0.45 | 0.33 |
| | -0.01 water table depth*MCWD | 0.16 | |
| | 0.003 water table depth*MCWD*clay | 0.15 | |
| Annual mortality rate | -0.03 MCWD*clay | 0.02 | |
| | -0.27 MCWD | <0.001 | |
| | -0.08 water table depth | 0.11 | |
| | -0.17 clay content | 0.03 | |
| | 0.13 maximum temperature | 0.09 | |
| | 0.16 soil fertility | 0.01 | 0.29 |
| | 0.00005 water table depth*MCWD | 0.57 | |
| | 0.000008 water table depth*MCWD*clay | 0.67 | |
| 0.0005 MCWD*clay | <0.001 | | |

| Forest metric | Regression coefficients for predictors | P | R ² |
|-------------------------|--|------------------|----------------|
| | -0.14 MCWD | 0.03 | |
| | -0.05 water table depth | 0.38 | |
| | -0.13 clay content | 0.08 | |
| | 0.04 maximum temperature | 0.61 | |
| Annual recruitment rate | 0.13 soil fertility | 0.04 | 0.10 |
| | -0.0001 water table depth*MCWD | 0.13 | |
| | -0.000003 water table depth*MCWD*clay | 0.90 | |
| | 0.00002 MCWD*clay | 0.85 | |
| | -0.20 MCWD | <0.001 | |
| | -0.07 water table depth | 0.14 | |
| | -0.14 clay content | 0.03 | |
| | 0.09 maximum temperature | 0.14 | |
| Annual turnover rate | 0.16 soil fertility | 0.004 | 0.24 |
| | -0.00003 water table depth*MCWD | 0.66 | |
| | 0.000005 water table depth*MCWD*clay | 0.79 | |
| | 0.0003 MCWD*clay | 0.01 | |
| | -0.0008 MCWD | 0.14 | |
| | -0.0005 water table depth | 0.23 | |
| | -0.0007 clay content | 0.25 | |
| | 0.00007 maximum temperature | 0.92 | |
| AGB Mortality Rate | 0.002 soil fertility | 0.005 | 0.10 |
| | -0.0000008 water table depth*MCWD | 0.28 | |
| | 0.000003 water table depth*MCWD*clay | 0.08 | |
| | 0.000003 MCWD*clay | 0.004 | |

Table S3 – Multicollinearity test to check the associations between predictor variables in the linear models. Variance Inflation Factor (VIF) less than 5 indicates low multiple correlation of that predictor with others.

| Predictors | VIF | SE factor |
|---------------------|------|-----------|
| MCWD | 1.54 | 1.24 |
| Water table depth | 1.14 | 1.07 |
| Soil fertility | 1.64 | 1.28 |
| Clay content | 2.19 | 1.48 |
| Maximum temperature | 2.25 | 1.50 |

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Table S4 – Comparison of models with and without a random factor to assess spatial autocorrelation between plots. AIC0 is the model without random factor, AIC1 is the model with random factor and DIF is AIC0-AIC1.

| Forest metric | AIC0 | AIC1 | DIF |
|--------------------|---------|---------|--------|
| AGWP | 1268.53 | 1238.16 | 30.37 |
| AGB | 4037.11 | 3976.27 | 60.84 |
| Mortality rate | 1869.88 | 1607.16 | 262.72 |
| Recruitment rate | 863.77 | 800.41 | 63.66 |
| Turnover rate | 1387.79 | 1177.87 | 209.92 |
| AGB mortality rate | 4342.20 | 4393.22 | -51.02 |

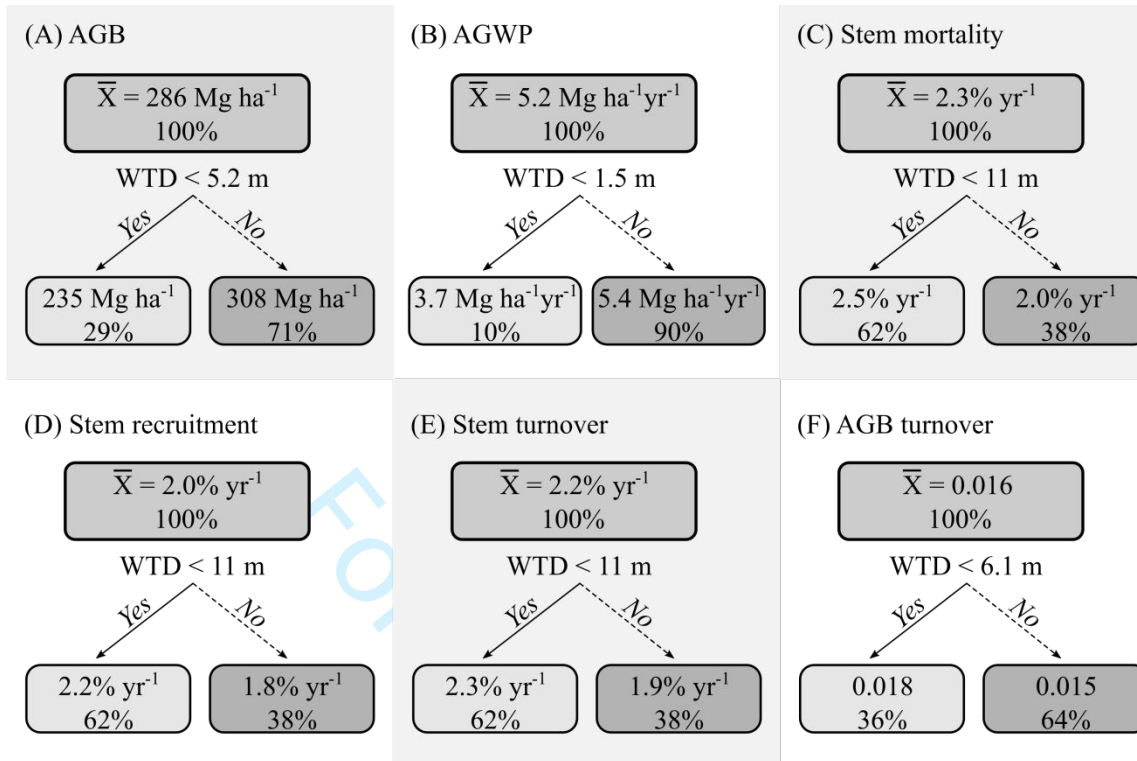


Figure S1. Regression trees showing the relationships and cutoff between water table depth and the response variables: (A) AGB stock, (B) AGWP, (C) Annual stem mortality, (D) Annual stem recruitment, (E) Annual stem turnover, and (F) AGB turnover.

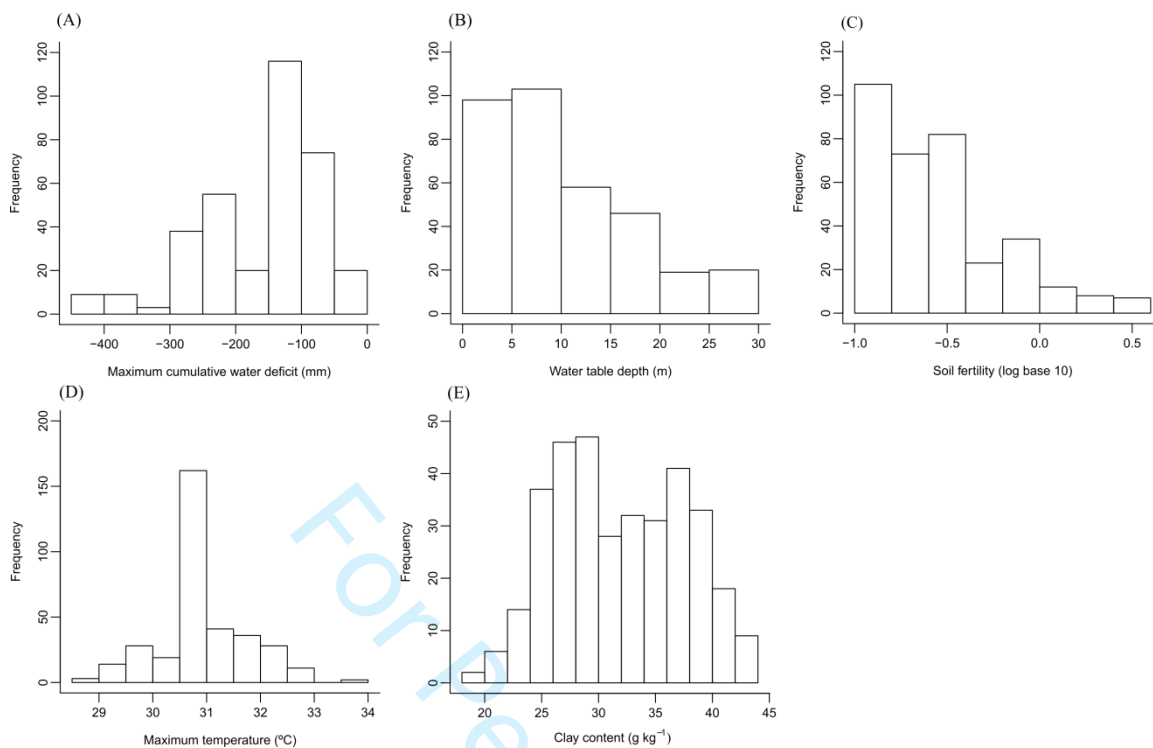
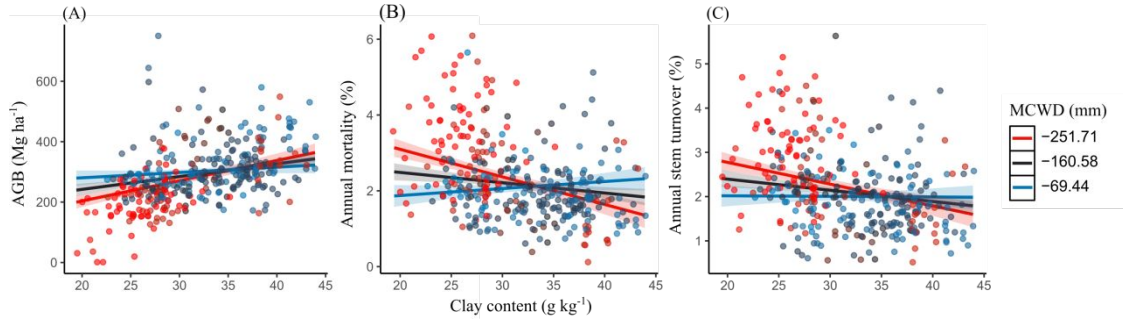
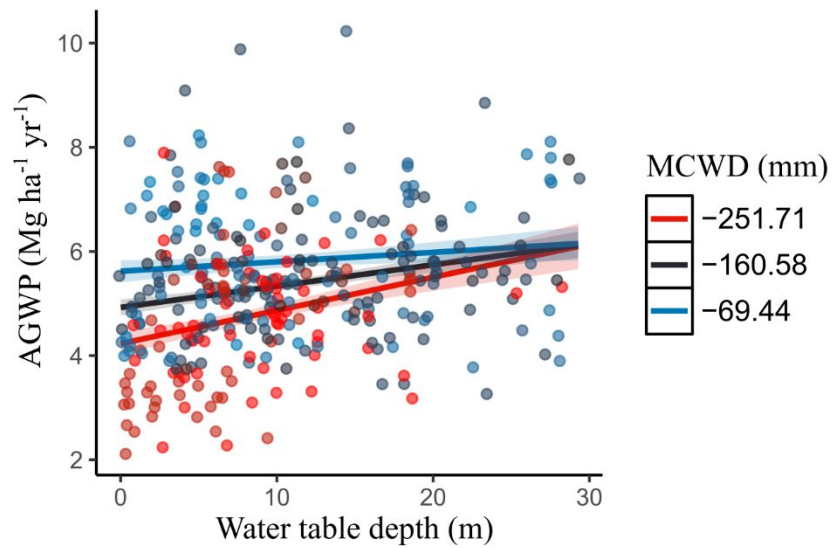


Figure S2. Histograms of the distribution of environmental variables in the plots used in this study are as follows: (A) Maximum cumulative water deficit; (B) Water table depth; (C) Soil fertility; (D) Maximum temperature; and (F) Clay content.



1 Figure S3. Multiple regression models investigating the effect of interactions among clay
 2 content and MCWD in Amazon forests. (A) biomass stock; (B) mortality rate; and (C)
 3 stem turnover. We used partial-dependence plots to visualize the shape of the
 4 relationships between response and predictor variables. In order to visualize interactions,
 5 clay content and climate deficit were divided in three classes based on the standard
 6 deviation around the mean of each of these variables. *Red* color is for plots with MCWD
 7 values less than one standard deviation below the mean; *black* is for plots with MCWD
 8 values within one standard deviation of the mean; and *blue* is for plots with MCWD
 9 values greater than one standard deviation above the mean. Shaded regions represent
 10 confidence intervals.



11 Figure S4. Partial-dependence plot of the interaction between MCWD and water table
12 depth on biomass productivity without plots with very low productivity (AGWP < 2 Mg
13 ha⁻¹ yr⁻¹). These were eight plots in white sand forests that are dominated by very thin
14 trees (<10 cm DBH). In order to visualize interactions, climate was divided in three
15 classes based on the standard deviation around the mean. Red colour is for plots with
16 MCWD values less than one standard deviation below the mean; black is for plots with
17 MCWD values within one standard deviation of the mean; and blue is for plots with
18 MCWD values greater than one standard deviation above the mean. Shaded regions
19 represent confidence intervals.

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3 Dear editor,
4

5 We are grateful for all review rounds, they were very important for the manuscript
6 improvement. The only request that was still pending was added to the manuscript.
7 Please see below.
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11 Sincerely,
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13 Thaiane Sousa, on behalf of all authors
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17 Detailed responses to the editor request:
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20 **EDITOR-IN-CHIEF'S COMMENTS TO AUTHORS**

21

22 Please provide a full Data Availability Statement, thank-you!
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26 **R: The Data Availability Statement was added to the manuscript. The DOI for the**
27 **data package was registered, this is currently not findable (ie. a draft), but will be**
28 **published once the paper embargo has been lifted and the paper is published.**
29

30 **Line 540:**
31

32 ***Data Availability Statement***
33

34 ***Data for the analyses are available as a ForestPlots.net data package at***
35 ***http://doi.org/10.5521/forestplots.net/2022_2***
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