

This is a repository copy of *Climate and fragmentation affect forest structure at the southern border of Amazonia*.

White Rose Research Online URL for this paper: http://eprints.whiterose.ac.uk/128852/

Version: Accepted Version

Article:

Reis, SM, Marimon, BS, Marimon Junior, BH et al. (9 more authors) (2018) Climate and fragmentation affect forest structure at the southern border of Amazonia. Plant Ecology and Diversity, 11 (1). pp. 13-25. ISSN 1755-0874

https://doi.org/10.1080/17550874.2018.1455230

© 2018 Botanical Society of Scotland and Taylor & Francis. This is an Accepted Manuscript of an article published by Taylor & Francis in Plant Ecology & Diversity on 25 Apr 2018, available online: http://www.tandfonline.com/10.1080/17550874.2018.1455230. Uploaded in accordance with the publisher's self-archiving policy.

Reuse

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.



eprints@whiterose.ac.uk https://eprints.whiterose.ac.uk/

Climate and fragmentation affect forest structure in the southern border of Amazonia

Simone Matias Reis^{a,b}, Beatriz Schwantes Marimon^{b,c,*}, Ben Hur Marimon Junior^{b,c}, Paulo S. Morandi^{a,b}, Edmar Almeida de Oliveira^b, Fernando Elias^c, Eder Carvalho das Neves^c, Bianca de Oliveira^c, Denis Silva Nogueira^c, Ricardo Keichi Umetsu^c, Ted R. Feldpausch^{c,d} and Oliver L. Phillips^a ^aSchool of Geography, University of Leeds, Leeds, United Kingdom; ^bPrograma de Pós-Graduação da Rede de Biodiversidade e Biotecnologia da Amazônia Legal (BIONORTE), Universidade do Estado de Mato Grosso-Universidade Federal do Amazonas, Nova Xavantina, Mato Grosso, Brazil; ^cUniversidade do Estado de Mato Grosso, Programa de Pós-graduação em Ecologia e Conservação, Nova Xavantina, MT, Brazil; ^dUniversity of Exeter, Geography, College of Life and Environmental Sciences, Exeter, United Kingdom *Corresponding author. Email: biamarimon@unemat.br

25 Abstract

Background: The remaining forests in the extensive contact zone between southern
Amazonia (seasonal rain forest) and the Cerrado (savanna) biomes are at risk due to
intense land-use and climate change.

Aims: To explore the vulnerability of these transitional forests to changes in land use and climate, we evaluated the effects of fragmentation and climatic variables on forest structure.

32 **Methods**: We measured the diameter and height of 14,185 trees with diameter ≥ 10 cm 33 at 24 forest plots distributed over an area of 25,000 km². For each plot, we obtained data 34 on contemporary fragmentation and climatic variables.

Results: Forest structure variables (height, diameter, height:diameter allometry,
biomass) varied significantly both within and among plots. The height, H:D and
biomass of trees were positively correlated with annual precipitation and fragment area.

38 **Conclusions**: The association between forest structure and precipitation indicates that 39 these forests plots are likely to be vulnerable to dry season intensification anticipated for 40 the southern edge of the Amazon. Additionally, the reduction in the fragment area may 41 contribute to reductions in forest biomass and tree height, and consequently ecosystem 42 carbon stocks. Given the likely susceptibility of these forests, urgent conservation action 43 is needed to prevent further habitat degradation.

44

45 Keywords: allometry; Amazon arc of deforestation; biomass; climate change; habitat
46 fragmentation; precipitation; stem diameter; tree height; transition zone

47

48 Introduction

49 Across the Earth's biomes, environmental conditions are expected to be 50 more variable close to the edges than in the core area of each biome, posing potentially 51 ecological and evolutionary challenges to biota towards their biogeographical edges 52 (Safriel et al. 1994; Kark and van Rensburg 2006; Kark et al. 2008). This may be 53 particularly the case in regions subject to rapid environmental change, of which perhaps 54 the most extreme example are the forests of the southern edge of the Amazon rain forest 55 biome, an area affected by high deforestation rates and subject to significant recent and 56 forecast climate change. Thus, here the advance of the agricultural frontier has already 57 resulted in converting most forest to pasture and cropland, increasingly fragmenting the 58 landscape over the last few decades (Alencar et al. 2004, 2015; Nogueira et al. 2008). 59 The remaining forests are subject to recent climate change, including lengthening of the 60 dry season and increasing incidence of strong droughts (Marengo et al. 2011; Gloor et 61 al. 2015; Feldpausch et al. 2016), trends which are expected to intensify further (e.g. 62 Boisier et al. 2015). The land surface temperature has been rising steadily recently, 63 especially in the south and east of the Amazon region (Jiménez-Muñoz et al. 2013), and 64 the effects of these climatic changes may be exacerbated by changes in land use (Aragão 2012; Silvério et al. 2015). Finally, research elsewhere in Amazonia clearly has 65 66 indicated that the structure of the tropical forest vegetation is affected by both climate change (e.g. Phillips et al. 2010; Feldpausch et al. 2016) and fragmentation of habitats 67 68 (e.g. Laurance et al. 1997, 2000; Laurance 2004).

69

70

71

72

Yet few studies have evaluated structural variation among the forests in the southern border region of the Amazon forest biome and its covariation with climate and landscape factors. Exceptions include one analysis of the effects of the interaction between droughts and wildfires on tree mortality at one experimental site (Brando et al. 73 2014), and a landscape study which showed that habitat fragmentation, combined with 74 droughts, increased the susceptibility of the forests to fire (Alencar et al. 2015). We are 75 not aware of a single study that has evaluated the effects of habitat fragmentation and 76 different climate variables across the region's forests using direct, on-the-ground 77 measurement of vegetation structural variables such as tree diameter, height, and 78 biomass.

79 Habitat fragmentation, by decreasing fragment size and increasing forest 80 edges and numbers of fragments, may modify the forest structure in the remaining 81 fragments (Fahring 2003; Haddad et al. 2015). For example, fragment edges are subject 82 to a greater incidence of insolation and increased velocity of winds, resulting in higher 83 temperatures and a drier microclimate than the forest interior (D'Angelo et al. 2004; 84 Laurance 2004; Haddad et al. 2015), which increases tree mortality rates, principally for 85 larger trees (Laurance et al. 2000; Laurance 2004). The death of bigger trees reduces 86 total biomass, height, mean diameter and basal area, especially in the smaller fragments 87 and the areas closest to the forest edge, although with some mortality effects also 88 propagating a few hundred meters into the forest (Laurance 2004; Haddad et al. 2015; 89 Rocha-Santos et al. 2016). Recently, it has even been suggested, based on interpretation 90 of pantropical satellite imagery, that in tropical forests the negative effects on standing 91 biomass and forest structure penetrate as much as 1.5 km into forests (Chaplin-Kramer 92 et al. 2015).

93

In addition to landscape-scale factors, regional climate is related to variation in the forest structure (e.g. Banin et al. 2015). For example, where precipitation and 94 95 temperature are higher, forests generally have taller trees that accumulate more biomass 96 (Koch et al. 2004; Way and Oren 2010; Feldpausch et al. 2011; Pan et al. 2013; Chave

97 et al. 2014). However, in the very warmest forests the forest structural responses are 98 unclear. There is some evidence that here plants may photosynthesise less and expend 99 more energy on respiration, so potentially accumulating less biomass (Lloyd and 100 Farquhar 2008; Lewis et al. 2013). However, the temperature sensitivity of key 101 respiration processes appears to decline in warmer environments (Atkin et al. 2015, 102 Heskel et al. 2016), rather than increasing exponentially as simple Q_{10} formulations in 103 earlier global vegetation models suggested (Cox et al. 2000), suggesting that the overall 104 sensitivity of biomass stocks to high temperatures might be weaker than many models 105 indicated.

Extreme drought events may alter the forest structure. Drought causes mortality, principally in the bigger trees, which are more susceptible to damage in their vascular system (Phillips et al. 2010; Rowland et al. 2015; Bennett et al. 2015; Feldpausch et al. 2016). During drought events, tropical trees may also grow less (e.g. Worbes 1999; Doughty et al. 2015), and if droughts are prolonged or repeated forests eventually accumulate less biomass (Feldpausch et al. 2016; Rowland et al. 2015).

112 In the context of regional land-use and climatic changes in southern Amazonia, 113 and the projected high regional climate sensitivity to global warming (IPCC 2015), it is 114 therefore extremely important to understand how the forest structure is affect by abiotic 115 factors. It may for example help to improve the conservation measures to protect the 116 remaining forest fragments. In this study, we evaluated whether, and to what extent, 117 climatic factors and fragmentation determine variation in the forest structure of the 118 southern Amazon border. We assembled data from permanent plots established across 119 the region close to the natural border of Amazonia with the neighboring Cerrado 120 (savanna) biome, to test hypotheses related to the variation in the forest structure and 121 the factors that determine this variation. We addressed two questions. First, does habitat 122 fragmentation affect the forest structure? We expected that forest cover loss and forest 123 plots present in smaller fragments and/or nearer the edge would have trees with lower 124 height and smaller diameter stems, or with smaller height:diameter (H:D) allometric 125 relationships and reduced biomass, since work elsewhere has shown mortality rates are 126 greater in smaller, more edge-affected fragments, especially for bigger trees (e.g. 127 Laurance et al. 1997, 1998, 2000; Laurance 2004; Chaplin-Kramer et al. 2015). Second, 128 how does the forest structure vary in relation to the climate? We expected that the 129 height and the diameter of stems, the H:D ratio, and biomass were all greater in forest 130 plots that have greater precipitation, and consequently less deficit water, since the 131 greater water availability favours the height growth of the trees, accumulating more 132 biomass (e.g. Feldpausch et al. 2011; Pan et al. 2013; Chave et al. 2014).

133

134 Materials and methods

135 Study area

We studied 24 forest plots distributed in the so-called 'arc of deforestation' (Nogueira et al. 2008) over an area of ca. 25,000 km² (Figure 1 and Table 1). The regional climate is of the Aw (tropical with dry winters) and Am (tropical monsoon) types in the Köppen classification system (Alvares et al. 2013), and originally supported evergreen or semi-evergreen forest vegetation in all cases. Mean annual precipitation and temperature range from 1511 to 2353 mm and from 24.1 to 27.3 °C, respectively (Table 1).



143

Figure 1. Location of the forests sampled in the southern Amazon border, between eastern and northern Mato Grosso and southern Pará, Brazil, showing the approximate biome boundaries based in IBGE (2004). The classification of forest and no forest was based on the PRODES (Amazon Deforestation Monitoring Project) (INPE 2017). All plots sampled lie within mature, evergreen or semi-evergreen forest fragments.

149

Table 1. Characteristics of plots sampled in different tropical forest ecosystems at the southern Amazon border. FA, fragment area; DE, distance to the forest edge; Prec, total mean annual precipitation; Temp, mean annual temperature; TB, total above-ground biomass per hectare; PP, private properties; and CU, conservation unit. In this study, we used codes ('Plot code') to represent the different types of vegetation: FEP, floresta estacional perenifólia (seasonal evergreen forest), FTP, floresta estacional perenifólia em terra preta de índio (seasonal evergreen forest on anthropogenic black earth); FES,

157 floresta estacional semidecidual (seasonal semi-deciduous forest); FOA, floresta 158 ombrófila aberta (open rainforest); and FSI, floresta sazonalmente inundável 159 (seasonally flooded forest). Equivalent forest plot codes are given to indicate 160 equivalency to those codes used in the ForestPlots.net database (Lopez-Gonzalez et al. 161 2011) where the data have been deposited.

Plot	Forest plot	Geographical	Local	AF	DE	Prec	Temp	ТВ
Code	code	coordinate		(ha)	(m)	(mm)	(°C)	(Mg)
FEP-01	FLO-01	-12.8S and -51.9W	PP	870	1,030	1613	25.5	111.1
FEP-02	FLO-02	-12.8S and -51.9W	PP	2,035	1,000	1621	25.6	144.7
FEP-03	TAN-02	-13.1S and -52.4W	PP	8,432	990	1625	24.9	143.5
FEP-04	TAN-03	-12.8S and -52.3W	PP	16,901	520	1679	25.1	127.4
FEP-05	TAN-04	-12.9S and -52.4W	PP	16,901	329	1662	25	138.3
FEP-06	FRP-01	-11.5S and -51.5W	PP	45,459	3,600	1634	26.9	135.1
FEP-07	POA-01	-11.0S and -52.2W	PP	9,789	1,180	1772	26.1	140.1
FES-01	VCR-02	-14.8S and -52.2W	PP	4,968	1,350	1511	25.2	196.8
FES-02	GAU-02	-13.4S and -53.3W	PP	3,499	160	1701	24.1	91.7
FES-03	SAT-01	-9.8S and -50.5W	PP	17,624	90	1821	26.7	121.8
FES-04	SAA-01	-9.8S and -50.4W	PP	13,039	860	1815	26.8	187.7
FES-05	SAA-02	-9.6S and -50.4W	PP	15,680	2,980	1778	26.6	166.3
FOA-01	SIP-01	-11.4S and -55.3W	PP	12,066	900	1848	25.1	79.2
FOA-02	ALF-01	-9.6S and -55.9W	CU	17,628	5,440	2350	25.5	98.8
FOA-03	ALF-02	-9.6S and -55.9W	CU	17,628	5,410	2353	25.6	160.5
FSI-01	PEA-01	-12.1S and -50.8W	CU	21	1	1631	27.3	133.7
FSI-02	PEA-02	-12.3S and -50.7W	CU	378	1	1637	27.2	154.7
FSI-03	PEA-03	-12.4S and -50.9W	CU	164	1	1621	27.1	131.4
FSI-04	PEA-04	-12.4S and -50.7W	CU	605	1	1637	27.1	210.4
FSI-05	PEA-07	-12.5S and -50.9W	CU	5	1	1621	27.1	226.8
FSI-06	PEA-08	-12.5S and -50.7W	CU	8	1	1632	27	222.5
FTP-01	GAU-04	-13.1S and -53.3W	PP	234	150	1795	24.7	145.8
FTP-02	GAU-05	-13.0S and -52.9W	PP	29,560	2,720	1757	24.9	250.2
FTP-03	GAU-06	-13.3S and -53.4W	PP	85	80	1729	24.7	176.9

162

163 Forest fragments

The largest and best preserved regional fragments of mature forests were selected for the study, using Google Earth imagery in order to capture regional variation in floristics and physiognomy, and with at least three plots for each forest type. All forest fragments are surrounded by extensive cattle-ranching or soybean fields. The fragments surveyed varied in size from 5 to 45,459 ha (Table 1). 169

170 Forest structure

171 In each fragment we established an inventory plot of 1 ha, which was 172 subdivided into 25 contiguous subplots of 20 m x 20 m. The forest plots were 173 established between 2008 and 2016 within the private properties and in conservation 174 units; locations varied between 1 and 5440 m from the nearest edge of the fragment. Six 175 plots were seasonally flooded (Table 1) and occasionally affected by fire; the others 176 have no recent record of fire and were either on anthropogenic black earth (terra preta 177 de índio), open rain forests, seasonal evergreen forests, or seasonal semi-deciduous 178 forests (Table 1). For this study, we used the latest available censuses between 2013 and 179 2016.

180 We identified and tagged all the woody individuals with a diameter at breast 181 height (1.3 m) of \geq 10 cm, for a total of 14,185 (range = 338-1599; standard deviation = 182 31) trees and at least 410 (range = 9-135; standard deviation = 256) taxa identified to 183 species level. We identified species in the field or by comparison of collections with 184 herbarium (NX, UFMT, UB and IAN) material of known identity, and with the help of 185 specialists. After identification, the material was incorporated into Herbarium NX, Nova 186 Xavantina, Mato Grosso (Coleção Zoobotânica James Alexander Ratter). We 187 determined the classification of families based on APG III (Angiosperm Phylogeny 188 Group 2009) and reviewed and updated the nomenclature of the taxa using the Lista de 189 Espécies da Flora do Brasil (http://floradobrasil.jbrj.gov.br/2015).

We measured the diameter of each tree following standard protocols of theRAINFOR network (http://www.rainfor.org/). We measured the total height using a

Leica DISTO laser measurement device. Data were deposited in the ForestPlots.netforest monitoring database (Lopez-Gonzalez et al. 2011).

194

195 Habitat fragmentation

196 To evaluate the effect of habitat fragmentation on forest structure, we 197 measured distance from each plot to the forest edge, the size of each fragment and the 198 forest cover in surrounding landscapes. Whenever possible we measured the distance to 199 the nearest edge in the field. When this was not possible, we estimated this value using 200 Google Earth, which provided a spatial resolution of approximately 20 to 30 m 201 depending on available imagery, and based on our own detailed knowledge, having 202 explored the local context of each plot on foot. In our definition of forest habitat edge, 203 we included all other vegetation and land-use such as plantations, pastures, and roads at 204 least 25 m wide, as well as natural grasslands in the six floodplain forests.

We calculated the area of the fragment where each plot was located using Google Earth and ZONUM software (http://zonums.com/online/kmlArea/). These edge and fragment data were collected at the closest possible date to the field sampling and in no case were they collected more than 2 years after the last forest census.

We calculated the percentage of forest cover surrounding each plot using buffers of radius size of 1000 m (314 ha), following recommendations of Rocha-Santos et al. (2016). For this we used the land-based metrics in the Fragstats software, that computes descriptors of forest patch and landscape attributes (McGarigal and Cushman 2002).

214

215 Climate variables

To evaluate the climate effect on the forest structure, we obtained data on 19 216 217 bioclimatic variables (Table S1) from the WorldClim 1.4 database, with a horizontal 218 resolution of ca. 1 km (Hijmans et al. 2005). We also used data from the Tropical 219 Rainfall Monitoring Mission (TRMM) (NASA 2012) to derive the mean of the annual 220 maximum climatological water deficit (MCWD) (Aragão et al. 2007) between January 221 1999 and December 2011, including the droughts of 2005, 2007 and 2010 (Figure S2). 222 To estimate this, we first calculated MCWD for each year, and then took the mean of all 223 years. MCWD was defined as the most negative value of climatological water deficit 224 (precipitation lower than evapotranspiration) among all the months in each year.

225

226 Data analysis

In each plot, we calculated the minimum, maximum, median, and 95 percentile of tree diameter (D), height (H) and their allometric (H:D) relationship. We also calculated the weighted Lorey's height values, based on basal area per subplot, using the equation

231 $\sum AB_i * H_i / \sum AB_i$,

where AB_i is the basal area of an individual and H_i is its height (e.g. Saatchi et al. 2011). To evaluate the H:D relationship, independently of disturbance, such as the damage caused by recently-opened clearings, we excluded from the analyses all trees with broken stems or those with more than 50% of the crown broken off.

We also calculated the mean, median, and total biomass of trees per plot. We estimated the biomass (B) based on the Pantropical model revised by Chave et al. (2014), which is derived from the equation in Chave et al. (2005), that is,

239 $B = 0.0673 \text{ x} (\rho D^2 H)^{0.976},$

240 where D is the diameter in cm, H is the total height of the tree in m, and ρ is 241 the density of the wood. We obtained wood density values from the ForestPlots 242 database (https://www.ForestPlots.net/). We chose this equation to calculate the 243 biomass because it is the most robust approach, given that it takes into consideration the 244 diameter and height of each tree.

245 We developed a correlation matrix of the Kendall's tau values of the 246 environmental and forest structure variables mentioned above (Table S3). Multiple 247 variables share similar source data, leading to high correlation amongst them, so we 248 excluded those with greatest correlations (r > 0.7) to avoid repetition of largely 249 redundant forest structure and climate variables (Tables S3 and S4). For all variables, 250 the maximum values and the 95 percentiles were highly correlated; we included only 251 the 95 percentile in order to avoid the influence of outliers. Finally, we excluded 252 predictor variables that correlated poorly (r < 0.1) with the vegetation descriptors 253 (Tables S3 and S4).

To verify possible differences among all forest plots in the structural variables (95 percentiles of the D, H and H:D, and mean B), we applied the Kruskal-Wallis analysis of variance with the Dunnett post hoc test and a Bonferroni correction (Zar 2010).

We evaluated the influence of habitat fragmentation and climatic variables on forest structure using simple correlation and Generalised Linear Models (GLM). We also included in the models the forest type for each forest plot. Simple correlation showed that, six seasonally flooded plots and two plots on anthropogenic black earth were unduly influential, with extreme structure and covarying extreme climatic and fragmentation conditions. To avoid these outliers driving the regional results weexcluded them from the GLM and correlation analyses described above.

265 To build the GLM, we first standardised the data and removed the 266 collinearities on the basis of Variance Inflation Factors (VIFs) of less than 10 (Quinn 267 and Keough 2002). We conducted model selection using the Akaike's Information 268 Criterion (AIC), with a model considered to be the best if it had the lowest AIC value 269 (Barton 2016). To access the spatial autocorrelation in the residuals for each model we 270 used Moran's I. Here, no spatial dependence was detected among plots, indicating that 271 the data were not spatially structured (Figure S5). Thus, we considered the plots as 272 independent samples in our subsequent analyses.

We conducted the analyses using SAM 4.0 program (Rangel et al. 2010) and R 2.15.1 (R Core Team 2012). The applied R packages were vegan (Oksanen et al. 2016), spdep (Bivand et al. 2013), spacemakeR (Dray 2013), MuMIn (Barton 2016) and packfor (Dray et al. 2016). We adopted a 5% significance level for all analyses and used 999 randomisations for the permutation methods.

278

279 **Results**

280 Forest structure

In general, the three open rainforest plots (FOA-01-03), a forest plot on anthropogenic black earth (FTP-01), were significantly taller than the six seasonally flooded forest plots (FSI-01-06), three seasonal semi-deciduous forest (FES-01-02-05) (Figure 2 and Table S6) and like the other 11 forest plots (FEP-01-07; FES-03-04 and FTP-02-03). The H:D ratio varied in a similar fashion to tree height, with the lowest ratios (i.e., the lowest heights for a given diameter) being recorded in two of the seasonally flooded forest plots (FSI-05 and FSI-06). Tree diameter and biomass did not
vary systematically among the plots, except for FSI-03, which had lower diameter and
biomass than the most of others plots (Figure 2).



Figure 2. Variation in the vertical structure of forests at the southern Amazon border. Box-plots show subplot-level values in each location, statistical comparisons are made for among-forest analyses based on the non-parametric Kruskal-Wallis test (H). The complementary pair-wise analysis of forest structure is provided in Table S7. \blacksquare = FTP (seasonal evergreen forest on anthropogenic black earth), \blacksquare = FOA (open rainforest), \blacksquare = FEP (seasonal evergreen forest), \blacksquare = FES (seasonal semi-deciduous forest), \square = FSI (seasonally flooded forest).

298



The structural variables were associated with the precipitation and with fragment area and distance from the edge (Figure 3 and Table 2). Tree height, allometry (H:D) and biomass all correlated positively with precipitation and fragment area (Figure 303 3). Tree height also correlated with the MCWD (Figure 3). Tree diameter did not correlate with any of the variables. Additionally, the precipitation and MCWD 305 correlated positively with the fragment area (P < 0.05; Kendal's $\tau = 0.44$ and 0.60, 306 respectively).

307

Table 2. The relationship between environmental variables and forest structure, using generalised linear models, of the southern Amazonia forests, Brazil. DE, distance to the edge; PrecWM, precipitation of wettest month; H:D, allometric H:D ratio; FES, seasonal semi-deciduous forest-plots; FOA, open rainforest-plots. Significant effects (P

F	actors	Estimate	Standard	t	Р
Height 95 percen	tile				
Iı	ntercept	-0.276	0.109	-2.531	0.003
F	ES	-0.008	0.161	-0.050	0.961
F	OA	1.392	0.328	4.249	0.001
Р	recWM	0.431	0.140	3.082	0.010
Diameter 95 perc	entile				
Iı	ntercept	-0.356	0.290	-1.228	0.243
F	ES	0.039	0.445	0.089	0.931
F	OA	1.715	0.530	3.237	0.007
H:D 95 percentil	e				
Iı	ntercept	< 0.001	0.174	< 0.001	1.000
Ľ	ЭE	-0.785	0.302	-2.597	0.023
Р	recWM	1.260	0.302	4.167	0.001
Mean biomass					
Iı	ntercept	-0.540	0.166	-3.249	0.007
F	ES	0.244	0.257	0.949	0.361
F	OA	2.291	0.303	7.555	<0.001

312 ≤ 0.05) are shown in bold type.



Figure 3. Significant ($P \le 0.05$) relationships between forest structure and climatic and fragmentation variables of the southern Amazon border forest plots. H95 = height 95 percentile, A95 = allometric ratio (H:D) 95 percentile, MB = mean biomass (Mg), FA = fragment area (ha), PrecWM = precipitation of wettest month (mm), MCWD = maximum climatological water deficit (mm).

314

Based on the best GLM models for each forest structure variable, forest type and precipitation were most related to tree height (Table 2). Forest type was also a strongly related to tree diameter and biomass. Annual mean precipitation and distance from the edge were important factors for mean plot H:D (Table 2). The percentage of forest cover around each plot was not selected in the best models and was not correlated with any forest structure variables. All plots presented more than 50% forest cover in surrounding landscapes.

Precipitation and MCWD were not selected in the same model, indicating that each had similar (but inverse) effects on forest structure. Thus, all structural parameters affected positively by precipitation (Table 2) are affected negatively by moisture stress (MCWD) (Table S7).

332

333 Discussion

Our results show that the forests of the southern border zone of Amazonia vary remarkably in their structure, principally in terms of their tree height and tree height:diameter ratio. Most of the structural variation in these forests was statistically related to fragment area and precipitation, supporting our overall expectations and largely consistent with our hypotheses. Here we briefly first discuss this overall variability and its potential ultimate drivers, before proceeding to discuss the results in more detail.

341

342 Structural variability of the forests of the southern Amazon border zone

343 Our general expectation was that climatic variation in the region would be a 344 fundamental determinant of the variability in forest structure here, principally because 345 drought events and seasonality may be more intense at the southern border in relation to 346 the core area of the Amazonas basin with evergreen non-seasonal rain forests (Lewis et 347 al. 2011). In particular, water deficit may kill large trees (McIntyre et al. 2015), taller 348 trees tend to be most affected by these conditions (Rowland et al. 2015). As these trees 349 die and break-up or fall, large clearings are opened, favouring the establishment of 350 species of different ecological groups (Lawton and Putz 1988). The frequent formation 351 of clearings in these hyperdynamic transitional forests, as documented by Marimon et 352 al. (2014), may thus also contribute to the structural variability found here. Finally, the 353 forests of the southern border of the Amazon are located within a mosaic of vegetation 354 types with many species typical of the adjacent biomes (Ratter et al. 1973), which may 355 have a direct influence on the structural diversity of these forests.

356

357 Seasonally flooded forest plots

358 The lowest height and H:D allometric ratio in the seasonally flooded forest 359 plots may be explained by their smaller fragment size and proximity to edges. These 360 factors as well as higher temperatures and lower precipitation (Table 1) may intensify 361 the fire effects. Fires in the wider grassland matrix can penetrate into forest fragments 362 and increase tree mortality, as observed in a recent study in these forest plots 363 (Maracahipes et al. 2014). It therefore appears likely that the combined effects of 364 reduced fragment area and precipitation and higher temperatures, together with fire and 365 its potential interactions with droughts (Brando et al. 2014), contribute to forest 366 structure here.

367

368 Response of the forest structure to the fragmentation and climate variables

369 Temperature appears to be an important factor determining the height of the 370 trees worldwide, including potentially in tropical forests (Koch et al. 2004; Way and 371 Oren 2010; Feldpausch et al. 2011; Lines et al. 2012; Pan et al. 2013), but here the 372 absence of a clear statistical relationship between structure and temperature (P > 0.05, 373 Kendal's $\tau = 0.31$) suggests it is not critical at the southern Amazon transition zone. 374 Rather, in our study the greater forest heights, H:D ratio and biomass that were 375 observed with increasing precipitation suggest water supply is the dominant climate 376 control on forest structure, and is consistent with some work elsewhere in the tropics 377 (e.g. Alvarez et al. 2017), given especially that tropical plants tend to grow faster and 378 taller as water is more available (Vlam et al. 2014; Givnish et al. 2014). In addition to 379 apparent effects of annual rainfall, we also found that climatological water deficit was 380 associated with reduced investment by the trees in height growth, consistent with the 381 hypothesis that tree height is constrained by the availability of water (Ryan et al. 2004; 382 Givnish et al. 2014). A significant positive correlation was also found between 383 precipitation and tree height along a precipitation gradient in Australia, which Givnish 384 et al. (2014) related to the increase in leaf area and rates of photosynthesis with 385 increasing precipitation.

The negative correlation between the cumulative water deficit and tree height may be related to the mortality of the largest individuals during extreme drought events (Phillips et al. 2010). Such droughts have been directly observed in the study region in 2005, 2007, and 2010 (e.g. Brando et al. 2014), and these have indeed tended to kill larger trees (Phillips et al. 2010; Feldpausch et al. 2016), as is often the case with droughts in other tropical forests (Bennett et al. 2015). In Amazonia, recent strong droughts appear also to be a major cause of the recent basin-wide increase in tree

393 mortality rates (Phillips et al. 2009; Brienen et al. 2015). In the near future, more 394 frequent extreme droughts, especially if combined with warming of the Amazon region 395 and thermal peaks in El Niño events such as in 2015-16 (Jiménez-Muñoz et al. 2016), 396 may therefore have profound implications for the forest structure of the southern 397 Amazon border, located as they are in a region that is already naturally close to their 398 distributional and hydraulic limits. In this scenario, large trees are more susceptible to 399 damage to the xylem, which can ultimately result in the death of the plant (e.g. McIntyre 400 et al. 2015) and eventually lead to forests of lower stature (McDowell et al. 2008; 401 Rowland et al. 2015). Trees being smaller in drier areas with greater water deficiency is 402 directly be related to conservative modifications in the hydraulic structure of the plants 403 under hydrological stress to avoid embolism (e.g. Lines et al. 2012, Claeys and Inzé 404 2013). Thus, as have recently argued in both tropical and temperate zone contexts (e.g. 405 Stegen et al. 2011; Banin et al. 2012; McIntyre et al. 2015) it is likely that trees in 406 forests subject either to more extreme climatic events, or to more disturbance (including 407 seasonally flooded habitats), or both, will in general tend to be shorter at a given diameter in order to avoid risks of hydraulic and/or mechanical failure, whereas trees in 408 409 forests with high rainfall, such as our FOA-01 and FOA-02, will have greater heights 410 and hence greater biomass.

Besides the correlation with the climatic variables, both height and the biomass of trees were positively correlated with fragment area. This result may be related to the incidence of wind in smaller fragments which have a higher proportion of forest edge (D'Angelo et al. 2004; Laurance 2004; Haddad et al. 2015). These disturbances are known to be able to generate high mortality, especially of the tallest trees (Laurance et al. 2000; Laurance 2004), and consequently in our dataset such edge417 generated disturbances may have affected the height and biomass of trees. Elsewhere, 418 local climatic changes as a result of fragmentation can reduce the density and diversity 419 of species (Mantyka-Pringle et al. 2012). Such effects can also increase the 420 susceptibility of fragmented forest structure and their biota to fire (Laurance and 421 Williamson 2001; Laurance 2004). In the southern Amazon region, these different 422 effects are all likely to be relevant, but clearly further analysis is needed, including long-423 term monitoring evaluation of the climatic and dynamic processes in these forests.

424

425 **Conclusions**

426 Our analysis across different locations, spanning a large part of the southern 427 Amazon zone, suggests climate sensitivity in forest structure here. Climate change, and 428 especially any reduction in annual or seasonal precipitation, is thus likely to have a 429 significant effect on the forest structure in the southern border of the Amazon. 430 Secondly, our results also suggest that the effects of reduction in the annual 431 precipitation may be further exacerbated in smaller fragments. This suggests that habitat 432 fragmentation may intensify the negative effects of climate change and burning in 433 forests in the southern Amazon border, resulting in a substantial risk of increases in tree 434 mortality. Given the likely susceptibility of the remaining southern Amazon border 435 forests to environmental change, strong conservation strategies are urgently needed to guarantee the persistence of these habitats, especially for the smaller fragments and 436 437 those close to agricultural frontiers.

438

439 Acknowledgements

440 We are grateful to the team of the Laboratório de Ecologia Vegetal - Plant 441 Ecology Laboratory at the UNEMAT campus in Nova Xavantina, especially to 442 Henrique Augusto Mews, Nayane Cristina Prestes, Ana Paula Gomes da Silva, Laís 443 Fernandes de Souza Neves, Mônica Forsthofer and Leonardo Maracahipes, for help in 444 collecting field data. We also thank the National Council for Scientific and 445 Technological Development (CNPq) for financial support of the projects PELD 446 "Cerrado-Amazon Forest Transition: ecological and socio-environmental bases for 447 Conservation" (stage II) - process nr. 403725/2012-7), PVE "special visiting researcher" 448 (401279/2014-6), PPBIO "Phytogeography of the Amazon-Cerrado Transition Zone" 449 (457602/2012-0) and FAPEMAT – Fundação de Amparo à Pesquisa do Estado de Mato 450 Grosso (164131/2013). We also thank CNPq for research productivity grants to B.S. 451 Marimon and B.H. Marimon Junior, a post-doctoral scholarship to D. Nogueira, and 452 international doctoral grants to S.M.A. Reis and P.S. Morandi. We are also grateful to 453 CAPES and FAPEMAT for scholarships to S.M.A. Reis, P.S. Morandi, F. Elias, E.A. 454 Oliveira and E.C. Neves.

455

456 Notes on contributors

457 Simone Matias Reis is a Ph.D. student focussing on the effects of biotic and abiotic458 factors on forest structure in the southern border of Amazonia.

459 Beatriz Schwantes Marimon is professor and an ecologist with experience in working 460 with structure and dynamics of different vegetation types of the southern Amazon 461 boundaries in Central Brazil. Her current work focuses on ecology and management of 462 forests in the transition zone between Cerrado and Amazon forest biomes, and currently 463 has great interest in understanding changes in tropical plants communities as a result of464 climate change.

Ben Hur Marimon-Junior is professor. He has experience in forest ecology, studying
biogeochemical cycles, carbon stocks, pyrogenic carbon in mineral nutrition of plants,
biodiversity and ecosystem functions. His current work focuses on understanding
changes in tropical plants communities as a result of climate change.

- 469 Paulo S. Morandi is a Ph.D. student focussing on structural patterns, diversity and
 470 distribution of species under the effect of environmental variables and vegetation
 471 succession in the Amazon-Cerrado transition.
- 472 Edmar Almeida de Oliveira is a Ph.D. student focussing on historical effect of soil
- 473 disturbance and pyrogenic carbon on the composition and structure of forests in
- 474 southern Amazonia.
- 475 Fernando Elias is a Ph.D. student focussing on the role of climate, land management

and landscape on the resilience of secondary forests in the Amazon.

- 477 Eder Carvalho das Neves is M.Sc. student focussing on the physiology of forests in the478 Amazon-Cerrado transition.
- 479 Bianca de Oliveira is a biologist and conducted her master's research on nutrient480 cycling in the Amazonia-Cerrado transition.

481 Denis Nogueira is a post-doctoral researcher focusing on forest dynamics of the482 Amazon-Cerrado transition in Mato Grosso.

- 483 Ricardo Keichi Umetsu is a lecturer. He has experience in environmental planning with
- 484 emphasis on water resources, hydrographic basins, hydrogeochemistry of streams, plant
- 485 ecology and landscape.

486 Ted R. Feldpausch is a lecturer in tropical ecology. His research focuses on the ecology487 and effects of global change on tropical forests.

488 Oliver L. Phillips is professor of tropical ecology. He leads the RAINFOR network of489 scientists in Amazonian forests.

490

491 **References**

- 492 Alencar A, Nepstad DC, McGrath D, Moutinho P, Pacheco P, Diaz MDCV, Soares
- 493 Filho B. 2004. Desmatamento na Amazônia: indo além da "emergência crônica".

494 Intituto de Pesquisa Ambiental da Amazônia, Belém.

- Alencar AA, Brando PM, Asner GP, Putz FE. 2015. Landscape fragmentation, severe
 drought, and the new Amazon forest fire regime. Ecological Applications 25: 1493–
 1505.
- 498 Alvares CA, Stape JL, Sentelhas PC, de Moraes G, Leonardo J, Sparovek G. 2013.
- Köppen's climate classification map for Brazil. Meteorologische Zeitschrift 22: 711–
 728.
- 501 Álvarez-Dávila E, Cayuela L, González-Caro S, Aldana AM, Stevenson PR, Phillips O,
- 502 Cogollo A, Peñuela MC, von Hildebrand P, Jiménez E, et al. 2017. Forest biomass
 503 density across large climate gradients in northern South America is related to water

availability but not with temperature. PLOS ONE 12:e0171072.

Angiosperm Phylogeny Group. 2009. An update of the angiosperm Phylogeny group
classification for the orders and families of flowering plants: APG III. Bot J Linn Soc
161: 105–121.

- 509 2007. Spatial patterns and fire response of recent Amazonian droughts. Geophysical 510 Research Letters 34.
- 511 Aragão LE. 2012. Environmental science: The rainforest's water pump. Nature 489: 512 217–218.
- Atkin OK, Bloomfield KJ, Reich PB, Tjoelker MG, Asner GP, Bonal D, Bönisch G, 514 Bradford MG, Cernusak LA, Cosio EG, et al. 2015. Global variability in leaf 515 respiration in relation to climate, plant functional types and leaf traits. New 516 Phytologist 206:614-636.
- 517 Banin L, Feldpausch TR, Phillips OL, Baker TR, Lloyd J, Affum-Baffoe K, Arets
- 518 EJMM, Berry NJ, Bradford M, Brienen RJW, et al. 2012. What controls tropical
- 519 forest architecture? Testing environmental, structural and floristic drivers. Global 520 Ecology and Biogeography 21: 1179–1190.
- 521 Banin LF, Phillips OL, Lewis SL. 2015. Tropical Forests. Routledge Handbook of 522 Forest Ecology 5: 56–75.
- 523 Barton K. 2016. MuMIn: Multi-model inference. R package; [cited 2017 Fev 20].
- 524 Available from: https://cran.r-project.org/web/packages/MuMIn/MuMIn.pdf.
- 525 Bennett AC, McDowell NG, Allen CD, Anderson-Teixeira KJ. 2015. Larger trees suffer
- 526 most during drought in forests worldwide. Nature plants 1: 15139.
- Bivand RS, Hauke J, Kossowski T. 2013. Computing the Jacobian in Gaussian spatial 527 528 autoregressive models: An illustrated comparison of available methods.
- 529 Geographical Analysis 45: 150–179.

513

530	Boisier JP, Ciais P, Ducharne A, Guimberteau M. 2015. Projected strengthening of
531	Amazonian dry season by constrained climate model simulations. Nature Climate
532	Change 5: 656–660.

- 533 Brando PM, Balch JK, Nepstad DC, Morton DC, Putz FE, Coe MT, Silvério D, Macedo
- 534 MN, Davidson EA, Nóbrega CC, et al. 2014. Abrupt increases in Amazonian tree
- mortality due to drought-fire interactions. Proceedings of the National Academy of
 Sciences 111: 6347–6352.
- 537 Brienen RJW, Phillips OL, Feldpausch TR, Gloor E, Baker TR, Lloyd J, Lopez-538 Gonzalez G, Monteagudo-Mendoza A, Malhi Y, Lewis SL, et al. 2015. Long-term
- 539 decline of the Amazon carbon sink. Nature 519: 344–348.
- 540 Chaplin-Kramer R, Ramler I, Sharp R, Haddad NM, Gerber JS, West PC, Mandle L,
- 541 Engstrom P, Baccini A, Sim S, et al. 2015. Degradation in carbon stocks near
 542 tropical forest edges. Nature communications 6.
- 543 Chave J, Andalo C, Brown S, Cairns MA, Chambers JQ, Eamus D, Folster H, Fromard
- 544 F, Higuchi N, Kira T, et al. 2005. Tree allometry and improved estimation of carbon
- 545 stocks and balance in tropical forests. Oecologia 145: 87–99.
- 546 Chave J, Réjou-Méchain M, Búrquez A, Chidumayo E, Colgan MS, Delitti WB, Duque

547 A, Eid T, Fearnside PM, Goodman RC, et al. 2014. Improved allometric models to

- 548 estimate the aboveground biomass of tropical trees. Global change biology 20: 3177–
- 549 3190.
- 550 Claeys H, Inzé D. 2013. The agony of choice: how plants balance growth and survival
- under water-limiting conditions. Plant Physiology 162: 1768-1779.

- 552 Cox PM, Betts RA, Jones CD, Spall SA, Totterdell IJ. 2000. Acceleration of global
 553 warming due to carbon-cycle feedbacks in a coupled climate model. Nature 408:
 554 184-187.
- 555 D'Angelo SA, Andrade ACS, Laurance SG, Laurance WF, Mesquita RCG. 2004.
- Inferred causes of tree mortality in fragmented and intact Amazonian forests. Journal
 of Tropical Ecology 20: 243–246.
- 558 Doughty CE, Metcalfe DB, Girardin CAJ, Amézquita FF, Cabrera DG, Huasco WH,
- 559 Silva-Espejo JE, Araujo-Murakami A, da Costa MC, Rocha W, et al. 2015. Drought
- 560 impact on forest carbon dynamics and fluxes in Amazonia. Nature 519: 78–82.
- 561 Dray S. 2013. spacemakeR: Spatial modelling. R package version 0.0-5/r113. https://R-
- 562 Forge.R-project.org/projects/sedar/
- 563 Dray S, Legendre P, Blanchet G. 2016. packfor: Forward Selection with permutation
 564 (Canoco p. 46). R package version 0.0-8/r136. https://R-Forge.R-
- 565 project.org/projects/sedar/
- 566 Feldpausch TR, Banin L, Phillips OL, Baker TR, Lewis SL, Quesada CA, Affum-
- 567 Baffoe K, Arets EJMM, Berry NJ, Bird M, et al. 2011. Height-diameter allometry of
- tropical forest trees. Biogeosciences 8: 1081–1106.
- Fahrig L. 2003. Effects of habitat fragmentation on biodiversity. Annual Review of
 Ecology, Evolution, and Systematics 34: 487–515.
- 571 Feldpausch TR, Phillips OL, Brienen RJW, Gloor E, Lloyd J, Lopez-Gonzalez G,
- 572 Monteagudo-Mendoza A, Malhi Y, Alarcón A, Álvarez Dávila E, et al. 2016.
- 573 Amazon forest response to repeated droughts. Global Biogeochemical Cycles 30: 1–
- 574 19.

- Givnish TJ, Wong SC, Stuart-Williams H, Holloway-Phillips M, Farquhar GD. 2014.
 Determinants of maximum tree height in Eucalyptus species along a rainfall gradient
 in Victoria, Australia. Ecology 95: 2991–3007.
- 578 Gloor M, Barichivich J, Ziv G, Brienen R, Schöngart J, Peylin P, Cintra BBL,
- 579 Feldpausch T, Phillips O, Baker J. 2015. Recent Amazon climate as background for
- possible ongoing and future changes of Amazon humid forests. GlobalBiogeochemical Cycles 29: 1384–1399.
- 582 Haddad NM, Brudvig LA, Clobert J, Davies KF, Gonzalez A, Holt RD, Lovejoy TE,
- 583 Sexton JO, Austin MP, Collins CD, et al. 2015. Habitat fragmentation and its lasting
- impact on Earth's ecosystems. Science Advances 1: e1500052.
- 585 Heskel MA, O'Sullivan OS, Reich PB, Tjoelker MG, Weerasinghe LK, Penillard A,
- Egerton JJ, Creek D, Bloomfield KJ, Xiang J, et al. 2016. Convergence in the
 temperature response of leaf respiration across biomes and plant functional types.
 Proceedings of the National Academy of Sciences 113: 3832–3837.
- Hijmans RJ, Cameron SE, Parra JL, Jones PG, Jarvis A. 2005. Very high resolution
 interpolated climate surfaces for global land areas. International journal of
 climatology 25: 1965–1978.
- 592 IBGE Instituto Brasileiro de Geografia e Estatística. 2004. Área territorial brasileira;
 593 [cited 2017 Jun 25]. Available from: http://www.ibge.gov.br/
- 594 INPE - Instituto Nacional de Pesquisas Espaciais. 2017. Coordenação geral de 595 observação da terra. Projeto PRODES: monitoramento da Floresta Amazônica 596 Brasileira Satélite; [cited] 2017 Jun 29]. Available from: por 597 http://www.obt.inpe.br/prodes/index.php

- 598 IPCC Intergovernmental Panel on Climate Change. 2015. Climate change 2014:
 599 mitigation of climate change. Cambridge: Cambridge University Press.
- 600 Jiménez-Muñoz JC, Mattar C, Barichivich J, Santamaría-Artigas A, Takahashi K, Malhi
- 601 Y, Sobrino JA, van der Schrier G. 2016. Record-breaking warming and extreme
- drought in the Amazon rainforest during the course of El Niño 2015-2016. Scientific
- 603 Reports 6: 33130.
- Jiménez-Muñoz JC, Sobrino JA, Mattar C, Malhi Y. 2013. Spatial and temporal patterns
- 605 of the recent warming of the Amazon forest. Journal of Geophysical Research:
 606 Atmospheres 118: 5204–5215.
- Kark S, van Rensburg BJ. 2006. Ecotones: marginal or central areas of transition? Israel
 Journal of Ecology & Evolution 52: 29–53.
- 609 Kark S, Hadany L, Safriel UN, Noy-Meir I, Eldredge N, Tabarroni C, Randi E. 2008.
- How does genetic diversity change towards the range periphery? Evolutionary
 Ecology Research 10: 391–414.
- Koch GW, Sillett SC, Jennings GM, Davis SD. 2004. The limits to tree height. Nature
 428: 851–854.
- 614 Laurance WF, Ferreira LV, Rankin-de Merona JM, Laurance SG. 1998. Rain forest
- 615 fragmentation and the dynamics of Amazonian tree communities. Ecology 79: 2032–
 616 2040.
- 617 Laurance WF, Laurance SG, Ferreira LV, Merona JMR, Gascon C. 1997. Biomass
- 618 Collapse in Amazonian Forest Fragments. Science 278: 1117–1118.
- 619 Laurance WF, Delamônica P, Laurance SG, Vasconcelos HL, Lovejoy TE. 2000.
- 620 Conservation: rainforest fragmentation kills big trees. Nature 404: 836–836.

- 621 Laurance WF, Williamson GB. 2001. Positive feedbacks among forest fragmentation,
- drought, and climate change in the Amazon. Conservation Biology 15: 1529–1535.
- 623 Laurance WF. 2004. Forest-climate interactions in fragmented tropical landscapes.
- 624 Philosophical Transactions of the Royal Society of London B: Biological Sciences
- 625 359: 345–352.
- Lawton RO, Putz FE. 1988. Natural disturbance and gap-phase regeneration in a
 wind-exposed tropical cloud forest. Ecology 69: 764–777.
- Lewis SL, Brando PM, Phillips OL, van der Heijden GM, Nepstad D. 2011. The 2010
 amazon drought. Science 331: 554-554
- 630 Lewis SL, Sonké B, Sunderland T, Begne SK, Lopez-Gonzalez G, van der Heijden,
- 631 GM, Phillips OL, Affum-Baffoe K, Baker TR, Banin L, et al. 2013. Above-ground
- biomass and structure of 260 African tropical forests. Philosophical Transactions of
 the Royal Society B 368: 20120295.
- Lines ER, Zavala MA, Purves DW, Coomes DA. 2012. Predictable changes in
 aboveground allometry of trees along gradients of temperature, aridity and
 competition. Global Ecology and Biogeography 21: 1017–1028.
- 637 Lista de Espécies da Flora do Brasil. 2016. Jardim Botânico do Rio de Janeiro; [cited
 638 2016 Jul 17]. Available from: http://floradobrasil.jbrj.gov.br/2016
- 639 Lloyd J, Farquhar GD. 2008. Effects of rising temperatures and [CO₂] on the
 640 physiology of tropical forest trees. Philosophical Transactions of the Royal Society
- 641 B: Biological Sciences 363: 1811–1817.
- 642 Lopez-Gonzalez G, Lewis SL, Burkitt M, Phillips OL. 2011. ForestPlots.net: a web
- 643 application and research tool to manage and analyse tropical forest plot data. Journal
- of Vegetation Science 22: 610–613.

Mantyka-pringle CS, Martin TG, Rhodes JR. 2012. Interactions between climate and
habitat loss effects on biodiversity: a systematic review and meta-analysis. Global
Change Biology 18: 1239–1252.

648 Maracahipes L, Marimon BS, Lenza E, Marimon-Junior BH, de Oliveira EA, Mews

- 649 HA, Gomes L, Feldpausch TR. 2014. Post-fire dynamics of woody vegetation in
- seasonally flooded forests (impucas) in the Cerrado-Amazonian Forest transition
 zone. Flora-Morphology, Distribution, Functional Ecology of Plants 209: 260–270.
- Marengo JA, Tomasella J, Alves LM, Soares WR, Rodrigues DA. 2011. The drought
- 800 of 2010 in the context of historical droughts in the Amazon region. GeophysicalResearch Letters 38.
- 655 Marimon BS, Marimon-Junior BH, Feldpausch TR, Oliveira-Santos C, Mews HA,
- 656 Lopez-Gonzalez G, Lloyd J, Franczak D, de Oliveira EA, Maracahipes L, et al. 2014.

Disequilibrium and hyperdynamic tree turnover at the forest–cerrado transition zone
in southern Amazonia. Plant Ecology & Diversity 7: 281–292.

659 McDowell N, Pockman T, Allen CD, Breshears DD, Cobb N, Kolb T, Plaut J, Sperry J,

West A, Williams DG. 2008. Mechanisms of plant survival and mortality during
drought: why do some plants survive while others succumb to drought? New
Phytologist 178: 719–739.

- McGarigal K, Cushman S. 2002. Comparative evaluation of experimental approaches to
 the study of habitat fragmentation effects. Ecological Applications 12: 335–345.
- McIntyre PJ, Thorne JH, Dolanc CR, Flint AL, Flint LE, Kelly M, Ackerly DD. 2015.
- 666 Twentieth-century shifts in forest structure in California: Denser forests, smaller
- trees, and increased dominance of oaks. Proceedings of the National Academy of
- 668 Sciences 112: 1458–1463.

- 671 http://disc.sci.gsfc.nasa.gov/precipitation/documentation/TRMM_README/TRMM
- 672 _3B43_readme.shtml, edited, NASA Distrib. Active Arch. Cent., Goddard Space
- 673 Flight Cent. Earth Sci., Greenbelt, Md.
- Nogueira EM, Fearnside PM, Nelson BW, Barbosa RI, Keizer EWH. 2008. Estimates
 of forest biomass in the Brazilian Amazon: new allometric equations and adjustments
 to biomass from wood-volume inventories. Forest Ecology and Management 256:
 1853–1857.
- 678 Oksanen J, Blanchet FG, Friendly M, Kindt R, Legendre P, McGlinn D, Minchin PR,
- O'Hara RB, Simpson GL, Solymos P, et al. 2016. vegan: Community Ecology
 Package. R package version 2.4-0. https://CRAN.R-project.org/package=vegan
- 681 Pan Y, Birdsey RA, Phillips OL, Jackson RB. 2013. The structure, distribution, and
- biomass of the world's forests. Annual Review of Ecology, Evolution, and
 Systematics 44: 593–622.
- 684 Phillips OL, Aragão LEOC, Lewis SL, Fisher JB, Lloyd J, López-González G, Malhi Y,
- 685 Monteagudo A, Peacock J, Quesada CA, et al. 2009. Drought Sensitivity of the
- 686 Amazon Rainforest. Science 323:1344–1347.
- 687 Phillips OL, Van Der Heijden G, Lewis SL, López-González G, Aragão LE, Lloyd J,
- Malhi Y, Monteagudo A, Almeida S, Dávila EA, et al. 2010. Drought-mortality
- relationships for tropical forests. New Phytologist 187: 631–646.
- 690 Quinn GP, Keough PR. 2002. Experimental design and data analysis for biologists.
- 691 Cambridge: Cambridge University Press.

- Rangel TF, Diniz-Filho JAF, Bini LM. 2010. SAM: a comprehensive application for
 spatial analysis in macroecology. Ecography 33: 46–50.
- 694 Ratter JA, Richards PW, Argent G, Gifford DR. 1973. Observations on the vegetation
- of the northeastern Mato Grosso. I. The woody vegetation types of the Xavantina-
- 696 Cachimbo Expedition area. Philosophical Transactions of the Royal Society of
- 697 London B 266: 449–492.
- 698 Rocha-Santos L, Pessoa MS, Cassano CR, Talora DC, Orihuela RL, Mariano-Neto E,
- 699 Morante-Filho JC, Faria D, Cazetta E. 2016. The shrinkage of a forest: landscape-
- scale deforestation leading to overall changes in local forest structure. Biological
- 701 Conservation 196: 1-9.
- 702 Rowland L, da Costa ACL, Galbraith DR, Oliveira RS, Binks OJ, Oliveira AAR, Pullen
- AM, Doughty CE, Metcalfe DB, Vasconcelos SS, et al. 2015. Death from drought in
 tropical forests is triggered by hydraulics not carbon starvation. Nature 528: 119–
 122.
- Ryan MG, Binkley D, Fownes JH, Giardina CP, Senock RS. 2004. An experimental test
 of the causes of forest growth decline with stand age. Ecological Monographs 74:
 393–414.
- 709 Saatchi SS, Harris NL, Brown S, Lefsky M, Mitchard ET, Salas W, Zuttaa BR,
- 710 Buermannb W, Lewis SL, Hagenf S, et al. 2011. Benchmark map of forest carbon
- 711 stocks in tropical regions across three continents. Proceedings of the National
- 712 Academy of Sciences 108: 9899–9904.
- 713 Safriel UN, Volis S, Kark S. 1994. Core and peripheral populations and global climate
- change. Israel Journal of Plant Sciences 42: 331–345.

Silvério DV, Brando PM, Macedo MN, Beck PS, Bustamante M, Coe MT. 2015.
Agricultural expansion dominates climate changes in southeastern Amazonia: the
overlooked non-GHG forcing. Environmental Research Letters 10: 104015.

718 Stegen JC, Swenson NG, Enquist BJ, White EP, Phillips OL, Jørgensen PM, Weiser

- 719 MD, Monteagudo Mendoza A, Núñez Vargas P. 2011. Variation in above-ground
- forest biomass across broad climatic gradients. Global Ecology and Biogeography20: 744–754.
- Vlam M, Baker PJ, Bunyavejchewin S, Zuidema PA. 2014. Temperature and rainfall
 strongly drive temporal growth variation in Asian tropical forest trees. Oecologia
 174: 1449–1461.
- Way DA, Oren R. 2010. Differential responses to changes in growth temperature
 between trees from different functional groups and biomes: a review and synthesis of
 data. Tree physiology 30: 669–688.

728 Worbes M. 1999. Annual growth rings, rainfall-dependent growth and long-term growth

- patterns of tropical trees from the Caparo Forest Reserve in Venezuela. Journal of
 Ecology 87: 391–403.
- 731 Zar JH. 2010. Biostatistical analysis. New Jersey: Prentice Hall.

Supplementary material

Variable abbreviation	Environmental predictors	Variable abbreviation	Vegetation descriptors
FA	Fragment area (ha)	MIH	Minimum height (m)
DE	Distance to the forest edge (m)	MAH	Maximum height (m)
FC	Forest cover (%)	MH	Median height (m)
MCWD	Maximum climatological water deficit (mm)	H95	Height 95 percentile (m)
Temp	Mean annual temperature (°C)	LH	Weighted Lorey's height
TempMDR	Mean diurnal range (°C)	MD	Median diameter (cm)
Isoter	Isothermality (°C)	MAD	Maximum diameter (cm)
TempSaz	Temperature seasonality (standard deviation *100) (°C)	D95	Diameter 95 percentile (cm)
TempWM	Max temperature of warmest month (°C)	MIA	Minimum allometric ratio (H:D)
TempCM	Min temperature of coldest month (°C)	MAA	Maximum allometric ratio (H:D)
TempAR	Temperature annual range (°C) TempWM - TempCM	MA	Median allometric ratio (H:D)
TempWeQ	Mean temperature of wettest quarter (°C)	A95	Allometric ratio (H:D) 95 percentile
TempDQ	Mean temperature of driest quarter (°C)	MB	Mean biomass (Mg ha)
TempWaQ	Mean temperature of warmest quarter (°C)	MEB	Median biomass (Mg ha)
TempCQ	Mean temperature of coldest quarter (°C)	TB	Total biomass (Mg ha)
Prec	Total annual precipitation (mm)		
PrecWM	Precipitation of wettest month (mm)	-	-
PrecDM	Precipitation of driest month (mm)	-	-
PrecSaz	Precipitation seasonality (Coefficient of Variation) (mm)	-	-
PrecWeQ	Precipitation of wettest quarter (mm)	-	-
PrecDQ	Precipitation of driest quarter (mm)	-	-
PrecWaQ	Precipitation of warmest quarter (mm)	-	-
PrecCQ	Precipitation of coldest quarter (mm)	-	-

Table S1. Environmental predictors and vegetation descriptors used in the analyses.



Figure S2. Mean of the maximum climatological water deficit (MCWD) (mm) in the Amazon basin between 1999 and 2011, in the context of the rest of Amazonia. Circles show the forest plots localization.

Table S3. Kendall tau correlations of the all 37 environmental and forest structure variables obtained to the forests of the southern Amazon border. FA = fragment area (ha), DE = distance to the edge (m), MCWD= maximum climatological water deficit (mm), Temp = mean annual temperature (°C), TempMDR = Mean diurnal range (°C), Isoter = Isothermality (°C), TempSaz = Temperature seasonality (standard deviation *100) (°C), TempWM = Max temperature of warmest month (°C), TempCM = Min temperature of coldest month (°C), TempDQ = Mean temperature annual range (°C) TempWM – TempCM, TempWeQ = Mean temperature of wettest quarter (°C), TempDQ = Mean temperature of driest quarter (°C), TempWaQ = Mean temperature of warmest quarter (°C), TempCQ = Mean temperature of coldest quarter (°C), TempWaQ = Mean temperature of warmest quarter (°C), TempCQ = Mean temperature of coldest quarter (°C), TempWaQ = Mean temperature of warmest quarter (°C), TempCQ = Mean temperature of coldest quarter (°C), Prec = Total annual precipitation (mm), PrecWM = Precipitation of wettest month (mm), PrecDM = Precipitation of driest quarter (mm), PrecDQ = Precipitation of driest quarter (mm), PrecDQ = Precipitation of driest quarter (mm), PrecWaQ = Precipitation of warmest quarter (mm), PrecCQ = Precipitation of coldest quarter (mm), MIH = Minimum height (m), MAH = Maximum height (m), MH = Median height (m), H95 = Height 95 percentile (m), LH = Weighted Lorey's height, MD = Median diameter (cm), MAD = Maximum diameter (cm), D95 = Diameter 95 percentile (cm), MIA = Minimum allometric ratio (H:D), MAA = Maximum allometric ratio (H:D), MA = Median allometric ratio (H:D), A95 = Allometric ratio (H:D) 95 percentile, MB = Mean biomass (Mg ha), MEB = Median biomass (Mg ha), TB = Total biomass (Mg ha). Significant correlations ($p \le 0.05$) are shown in bold type.

	FA	DE	MCWD	Temp	TempMDR	Isoter	TempSaz	TempWM	TempCM	TempAR	TempWeQ	TempDQ	TempWaQ	TempCQ	Prec	PrecWM	PrecDM
FA		0.55	0.61	-0.18	0.28	-0.17	-0.19	0.17	-0.16	0.29	-0.26	0.02	-0.20	-0.01	0.37	0.48	0.00
DE			0.51	-0.21	0.30	-0.25	-0.13	0.12	-0.20	0.32	-0.33	-0.09	-0.23	-0.10	0.23	0.44	0.00
MCWD				-0.13	0.24	-0.12	-0.35	0.34	-0.05	0.24	-0.27	0.13	-0.16	0.08	0.62	0.56	0.05
Temp					-0.82	0.51	-0.42	0.32	0.92	-0.80	0.85	0.75	0.96	0.78	-0.16	-0.37	0.17
TempMDR						-0.64	0.29	-0.17	-0.79	0.96	-0.89	-0.59	-0.85	-0.62	0.29	0.50	-0.12
Isoter							-0.41	0.20	0.56	-0.69	0.66	0.62	0.54	0.63	-0.18	-0.46	0.15
TempSaz								-0.72	-0.51	0.29	-0.27	-0.68	-0.39	-0.67	-0.30	-0.09	-0.24
TempWM									0.39	-0.15	0.18	0.57	0.30	0.55	0.43	0.22	0.34
TempCM Temp AP										-0.80	0.78	0.82	0.89	0.85	-0.10	-0.35	0.22
TempAK											-0.00	-0.59	-0.82	-0.05	0.50	0.54	-0.13
TempDO												0.00	0.00	0.03	0.09	-0.51	0.09
TempWaO													0.71	0.74	-0.19	-0.17	0.14
TempCO														0.74	0.05	-0.21	0.30
Prec															0.00	0.56	0.25
PrecWM																0.00	-0.12
PrecDM																	
PrecSaz																	
PrecWeQ																	
PrecDQ																	
PrecWaQ																	
PrecCQ																	
MAH																	
MIH																	
MH																	
П95 I Н																	
MAD																	
MD																	
D95																	
MAA																	
MIA																	
MA																	
A95																	
MB																	
MEB																	
TB																	

a	
Conti	nuation

	PrecSaz	PrecWeQ	PrecDQ	PrecWaQ	PrecCQ	MAH	MIH	MH	H95	LH	MAD	MD	D95	MAA	MIA	MA	A95	MB	MEB	TB
FA	-0.32	0.36	0.16	-0.07	0.31	0.35	0.23	0.55	0.51	0.44	0.30	0.20	0.05	0.32	0.42	0.47	0.51	0.31	0.43	0.20
DE	-0.30	0.29	0.23	-0.29	0.33	0.49	0.34	0.50	0.49	0.38	0.38	0.18	0.14	0.30	0.33	0.49	0.50	0.31	0.35	0.29
MCWD	-0.64	0.50	0.38	-0.04	0.48	0.39	0.42	0.57	0.46	0.51	0.46	0.18	0.12	0.33	0.23	0.56	0.56	0.32	0.44	0.28
Temp	0.19	-0.49	-0.05	0.15	-0.19	-0.35	-0.21	-0.31	-0.36	-0.35	-0.23	0.04	-0.33	-0.11	-0.46	-0.27	-0.29	-0.45	-0.10	-0.22
TempMDR	-0.36	0.60	0.16	-0.17	0.30	0.48	0.22	0.43	0.49	0.46	0.29	-0.04	0.42	0.17	0.54	0.42	0.42	0.55	0.19	0.37
Isoter	0.23	-0.56	-0.07	0.37	-0.27	-0.43	-0.08	-0.34	-0.46	-0.36	-0.16	0.03	-0.39	-0.21	-0.46	-0.30	-0.33	-0.45	-0.12	-0.33
TempSaz	0.28	0.03	-0.16	-0.02	-0.18	-0.02	-0.25	-0.19	-0.04	-0.11	-0.17	-0.08	0.16	-0.19	0.14	-0.25	-0.20	0.13	-0.20	-0.13
TempWM	-0.33	0.10	0.28	0.03	0.27	0.06	0.18	0.17	0.14	0.14	0.11	0.05	-0.05	0.19	-0.06	0.20	0.16	0.00	0.25	0.22
TempCM	0.15	-0.43	0.00	0.11	-0.13	-0.34	-0.14	-0.27	-0.34	-0.32	-0.19	0.04	-0.36	-0.06	-0.43	-0.23	-0.26	-0.44	-0.07	-0.21
TempAR	-0.35	0.63	0.13	-0.19	0.31	0.50	0.24	0.42	0.53	0.48	0.31	-0.03	0.45	0.18	0.53	0.41	0.43	0.59	0.22	0.39
TempweQ	0.36	-0.62	-0.16	0.29	-0.34	-0.49	-0.29	-0.46	-0.48	-0.47	-0.34	-0.04	-0.40	-0.23	-0.49	-0.42	-0.43	-0.53	-0.23	-0.36
TempDQ	-0.05	-0.30	0.10	0.12	0.00	-0.22	-0.01	-0.08	-0.19	-0.14	-0.04	0.09	-0.29	0.01	-0.35	-0.04	-0.08	-0.34	0.06	-0.12
TempwaQ	0.25	-0.51	-0.08	0.19	-0.22	-0.40	-0.21	-0.30	-0.38	-0.38	-0.25	0.04	-0.35	-0.15	-0.49	-0.31	-0.33	-0.40	-0.11	-0.20
Broo	-0.02	-0.54	0.07	0.13	-0.03	-0.23	-0.02	-0.10	-0.23	-0.10	-0.07	0.00	-0.55	-0.01	-0.37	-0.03	-0.09	-0.39	0.01	-0.15
PrecW/M	-0.05	0.59	0.44	0.11	0.55	0.50	0.24	0.44	0.50	0.50	0.20	0.11	0.19	0.30	0.31	0.59	0.40	0.59	0.30	0.59
PrecDM	-0.45	0.01	0.20	-0.00	0.40	0.52	0.50	0.51	0.70	0.75	0.35	0.03	0.29	0.44	0.02	0.52	0.55	0.55	0.35	0.57
PrecSaz	-0.29	-0.12	0.04	0.07	0.44	0.00	-0.00	-0.03	0.00	-0.01	0.02	-0.11	-0.12	-0.08	-0.19	0.15	-0.12 0.47	-0.09	-0.28	0.02
PrecWeO		-0.50	-0.02	0.12	-0.78	-0.30	0.28	-0.52	-0.45	0.40	-0.57	-0.11	0.12	0.20	0.17	0.51	0.47	0.27	0.20	-0.56
PrecDO			0.22	-0.09	0.41	0.42	0.28	0.34	0.03	0.01	0.20	0.04	0.29	0.37	0.42	0.32	0.32	0.30	0.29	0.31
PrecWaO				0.11	-0.13	-0.23	-0.32	-0.27	-0.06	-0.07	-0.10	_0.15	-0.05	-0.15	-0.06	-0.25	-0.20	-0.021	-0.12	-0.04
PrecCO					-0.15	0.20	0.32	0.23	-0.00 0 /0	-0.07	0.30	-0.07	0.00	0.10	0.18	0.25	0.20	0.02	0.12	0.04
МАН						0.40	0.35	0.42	0.42	0.41	0.50	-0.03	0.15	0.23	0.10	0.33	0.33	0.54	0.14	0.29
MIH							0.25	0.44	0.00	0.00	0.55	0.03	0.18	0.24	-0.06	0.30	0.40	0.26	0.14	0.18
MH								0.77	0.52	0.42	0.32	0.00	0.15	0.54	0.00	0.47	0.31	0.20	0.27	0.10
H95									0.70	0.30	0.31	0.10	0.15	0.30	0.35	0.00	0.07	0.27	0.47	0.55
II) I H										0.70	0.42	-0.01	0.37	0.27	0.28	0.40	0.45	0.00	0.27	0.55
MAD											0.54	0.01	0.71	0.19	0.00	0.31	0.35	0.37	0.27	0.00
MD												0.01	0.27	0.06	0.00	-0.04	0.06	0.19	0.10	-0.22
D95													0.27	-0.01	0.15	0.03	0.00	0.70	0.28	0.16
MAA														0.01	0.18	0.55	0.61	0.06	0.28	0.28
MIA															0.10	0.30	0.31	0.32	0.20	0.31
MA																0.00	0.87	0.12	0.30	0.32
A95																	0.07	0.24	0.39	0.37
MB																		0.21	0.35	0.39
MEB																				0.07
TB																				

Table S4. Pre-selected environmental and forest structure variables used in the analyses of the forest-plots of the southern Amazon border. FA = fragment area (ha), DE = distance to the edge (m), FC = forest cover (%), Temp = mean annual temperature (°C), PrecWM = precipitation of wettest month (mm), MCWD= maximum climatological water deficit (mm), MH= median height and H95 = 95 percentile, MD = median diameter and D95 = 95 percentile, MA = median allometric ratio (H:D) and A95 = 95 percentile, MB = mean biomass (Mg), and TB = total biomass.

Equart plata		Envir	onm	Vegetation descriptors						
Forest plots	FA	DE	FC	Temp	PrecWM	MCWD	H95	D95	A95	MB
FEP-01	870	1,030	99	25.5	291	-435.02	20.0	33.6	1.19	0.24
FEP-02	2,035	1,000	100	25.6	289	-435.02	20.1	36.6	1.34	0.25
FEP-03	8,432	990	98	24.9	285	-434.01	19.8	40	1.18	0.27
FEP-04	16,901	520	74	25.1	292	-428.93	20.0	37.8	1.26	0.25
FEP-05	16,901	329	100	25.0	291	-428.93	20.0	37.8	1.29	0.25
FEP-06	45,459	3,600	100	26.9	298	-411.82	22.0	41.4	1.19	0.32
FEP-07	9,789	1,180	100	26.1	309	-397.35	20.5	35.4	1.40	0.25
FES-01	4,968	1,350	78	25.2	274	-468.04	20.4	40.4	1.00	0.31
FES-02	3,499	160	69	24.1	283	-433.5	18.3	39.4	1.32	0.27
FES-03	17,624	90	58	26.7	293	-388.22	21.0	35.4	1.27	0.25
FES-04	13,039	860	88	26.8	289	-388.22	20.8	39.3	1.17	0.31
FES-05	15,680	2,980	100	26.6	278	-387.33	19.3	33.8	1.16	0.24
FOA-01	12,066	900	98	25.1	311	-420.38	25.3	44.8	1.37	0.39
FOA-02	17,628	5,440	100	25.5	390	-342.12	27.8	42.6	1.42	0.43
FOA-03	17,628	5,410	50	25.6	390	-342.12	28.1	42.3	1.37	0.41
FSI -01	21	1	-	27.3	273	-440.57	13.6	32.3	0.93	0.14
FSI -02	378	1	-	27.2	277	-454.52	15.0	35.2	0.92	0.19
FSI -03	164	1	-	27.1	273	-457.47	14.0	24.4	0.99	0.12
FSI -04	605	1	-	27.1	278	-454.52	15.7	28.1	1.02	0.15
FSI -06	5	1	-	27.1	274	-457.47	13.9	40.3	0.75	0.19
FSI -07	8	1	-	27.0	278	-444.82	15.6	45.0	0.77	0.3
FTP-01	234	150	38	24.7	308	-436.02	26.8	51.9	1.14	0.48
FTP-02	29,560	2,720	71	24.9	302	-429.99	22.0	34.7	1.16	0.29
FTP-03	85	80	30	24.7	294	-433.5	24.0	45.3	1.09	0.52



Figure S5. Spatial autocorrelation of the residuals of each model, based in Moran's I index for: A = height, B = diameter, C = allometric ratio (H:D), and D = biomass of the forests plots in the southern Amazon border.

Table S6. Comparison of the forest structure variables of the forests in the southern Amazon border, based on the Kruskal-Wallis nonparametric analysis of variance (H). MH= median height and H95 = 95 percentile, MD = median diameter and D95 = 95 percentile, MA = median allometric ratio (H:D) and A95 = 95 percentile, MB = mean biomass (Mg), and TB = total biomass. Values on different lines within the same column followed by different letters are significantly different based on Dunnett's post hoc test with the Bonferroni correction.

Forests	H95	D95	A95	MB		
FEP-01	19.3 afg	31.5 acd	1.17 aefghi	0.25 adef		
FEP-02	19.3 afg	33.1 abcd	1.27 afg	0.25 abdef		
FEP-03	19.0 fg	37.5 ab	1.10 deghi	0.27 abdef		
FEP-04	19.0 fg	33.9 abcd	1.20 afghi	0.25 abdef		
FEP-05	19.7 afg	35.1 abd	1.21 afgh	0.26 abdef		
FEP-06	20.4 afg	38.1 ab	1.12 defghi	0.33 abf		
FEP-07	19.9 afg	32.8 abcd	1.36 a	0.25 abdef		
FES-01	17.6 def	33.8 abcd	0.94 bcd	0.30 abdef		
FES-02	18.0 cdef	35.9 ab	1.26 afg	0.28 abef		
FES-03	20.1 afg	34.4 abcd	1.19 afghi	0.26 abdef		
FES-04	19.6 afg	38.2 ab	1.13 defghi	0.32 abf		
FES-05	18.2 ef	31.9 acd	1.12 defghi	0.25 adef		
FOA-01	24.0 a	38.8 ab	1.29 afg	0.39 ab		
FOA-02	25.7 a	39.4 ab	1.39 a	0.44 ab		
FOA-03	24.8 ag	38.3 ab	1.31 af	0.41 ab		
FSI-01	13.1 bc	30.5 acd	0.84 bc	0.18 cde		
FSI-02	14.2 bcde	31.6 acd	0.85 bc	0.20 cdef		
FSI-03	13.1 bcd	24.5 c	0.97 bcd	0.12 c		
FSI-04	14.3 bcde	27.0 cd	0.98 bcde	0.16 cd		
FSI-05	11.9 в	35.2 ab	0.66 b	0.23 acdef		
FSI-06	13.4 bcd	40.5 ab	0.61 b	0.32 abf		
FTP-01	23.5 ag	43.2 в	1.06 cdehi	0.47 b		
FTP-02	19.7 afg	33.1 abd	1.11 deghi	0.29 abef		
FTP-03	21.1 afg	42.8 ab	1.02 bcdei	0.52 abdef		

Table S7. Generalized linear models of the factors that influence forest structure of the vegetation in forest plots of the southern Amazon border. Temp = mean annual temperature, MCWD = maximum climatological water deficit, H:D = allometric H:D ratio, FES = seasonal semi-deciduous forest-plots, FOA = open rainforest-plots. Significant effects ($p \le 0.05$) are shown in bold type.

	Factors	Estimate	Standard	t	Р
Height 95 per	centile				
	Intercept	2.462	1.229	2.003	0.070
	FES	-0.206	0.177	-1.161	0.270
	FOA	1.848	0.262	7.060	0.000
	MCWD	0.007	0.003	2.340	0.039
H:D 95 percer	ntile				
	Intercept	8.630	2.679	3.221	0.007
	MCWD	0.021	0.007	3.230	0.007
	Temp	-0.497	0.230	-2.159	0.052