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1 Root biomass, root : shoot ratio and belowground carbon stocks in the open savannahs of
2 Roraima, Brazilian Amazonia

3

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15

16 Abstract

17

18 Biomass of roots, the root : shoot ratio (ratio of below to aboveground biomass) and carbon
 19 stocks below ground (to 100 cm depth) are estimated in different open savannah
 20 environments in the extreme north of the Brazilian Amazon. Sampling was conducted in
 21 permanent plots established in two open savannah areas in the state of Roraima. We
 22 identified four phytopedounits in the 27 plots sampled in two areas: four in dry grasslands on
 23 Argisol/Ultisol soils (DG-Arg), eight in dry grasslands on Latosol/Oxisol soils (DG-Lts), five
 24 in a mosaic of grasslands with savannah-parkland on Latosol/Oxisol soils (GP-Lts) and ten in
 25 seasonally flooded (wet) grasslands on Hydromorphic/Entisol soils (WG-Hyd). Fine roots (<
 26 2 mm diameter) dominated the 0-100 cm vertical profile in the four phytopedounits (>
 27 92.5%). Biomass of the roots in WG-Hyd ($29.52 \pm 7.15 \text{ Mg ha}^{-1}$) was significantly higher as
 28 compared to the other phytopedounits studied, although the carbon stocks did not differ
 29 among the phytopedounits ($6.20\text{-}7.21 \text{ MgC ha}^{-1}$). The largest concentration of roots was
 30 found in the upper three 10-cm sections of the soil profile, ranging from 56.3 to 82.9% in the
 31 four environments. The root : shoot ratio based only on living biomass of roots with diameter
 32 $\geq 2 \text{ mm}$ (standard IPCC methodology) ranged from zero for seasonally flooded grasslands to
 33 0.07-0.20 for unflooded grasslands on clay soils. The results indicate that the root : shoot
 34 ratio (expansion factor) for belowground biomass in open savannah ecosystems in the
 35 northern Amazon are low and differ from the default values used in Brazil's reference report
 36 to the Climate Convention.

37

38 Introduction

39

40 Savannah is a common type of vegetation in the tropics, including the Neotropics
 41 (Solbrig *et al.* 1996; Furley 1999). Their terrestrial coverage has been estimated to be 16-19
 42 $\times 10^6 \text{ km}^2$, depending on the ecogeographical definitions used (Scholes and Hall 1996; Asner
 43 *et al.* 2004). Most studies on the vertical and horizontal structure of these ecosystems are
 44 aimed at identifying structural patterns associated with biological diversity and aboveground
 45 biomass or carbon stocks. Root biomass and carbon are often not reported on a small scale
 46 because they demand so much time and effort to sample. However, even with the small
 47 number of studies, some reviews suggest that roots in tropical savannah and grassland
 48 ecosystems represent a major compartment for carbon accumulation (Jackson *et al.* 1996;
 49 Mokany *et al.* 2006). Estimates of these stocks are important for national inventories of
 50 greenhouse gases under the UN Framework Convention on Climate Change (UN-FCCC).

51 Generally, distribution, production and accumulation of roots are related to water
 52 availability in the soil (climatic seasonality), which is a variable with strong temporal
 53 fluctuations in the more superficial soil layers in savannahs and grasslands (San José *et al.*
 54 1982; Baruk 1994; Delitti *et al.* 2001). Other factors have been investigated in order to
 55 understand which processes regulate subterranean biomass, such as physionomic structure
 56 (Sarmiento and Vera 1979; Castro and Kauffman 1998), nutrient availability (Kellman and
 57 Sanmugadas 1985; February and Higgins 2010), human alterations (Fiala and Herrera 1988),
 58 fire (Menaut and Cesar 1979; Castro-Neves 2007) and grazing (Pandey and Singh 1992;
 59 Milchunas and Lauenroth 1993; McNaughton *et al.* 1998). These studies make it possible to
 60 assess parameters such as root : shoot ratio (the "expansion factor" used for inferring
 61 belowground biomass from aboveground biomass measurements) and rates of growth of root
 62 biomass under different successional paths, which are crucial to the understanding of
 63 belowground carbon allocation. However, no study has been conducted in Amazonian
 64 savannahs.

65 In savannahs, the thicker roots represent an important reservoir of carbon at greater
66 depths, especially in physiognomies that are more densely populated by trees (Abdala *et al.*
67 1998; Schenck and Jackson 2002). However, these open-vegetation environments are
68 characterised by having large grassy expanses and a low-density tree component. Most roots
69 in these ecosystems are located in the most superficial layers of the soil and are characterised
70 by having small diameter (< 2 mm) (Knoop and Walker 1985). It is estimated that the
71 different forms of savannahs (open and wooded) contain ~ 20% of all of the fine-root
72 biomass on Earth to a depth of 30 cm (Jackson *et al.* 1997). This category of roots has a high
73 rate of replenishment in tropical grasslands and savannahs, making it a critical component in
74 sequestering atmospheric carbon by allowing constant accumulation of organic matter in the
75 soil (Stanton 1988; Gill and Jackson 2000; Chen *et al.* 2004). However, the
76 Intergovernmental Panel on Climate Change (IPCC 2006) suggests that this category of roots
77 (< 2 mm in diameter) should not be included as part of the "belowground biomass" because it
78 is difficult to distinguish it empirically from soil organic matter.

79 Savannahs, grasslands and other natural non-forest ecosystems occupy ~200,000 km²
80 (about 5%) of the Brazilian Amazonia (Santos *et al.* 2007). Although the area of these
81 ecosystems is substantial, all existing studies on the biomass of roots in natural vegetation in
82 the Amazonian biome in Brazil are from forest ecosystems due to the much larger area of
83 forests (Klinge 1973; Thompson *et al.* 1992; Luizão *et al.* 1992; Nepstad *et al.* 1994; Cattanio
84 *et al.* 2004). Estimates of the temporal dynamics and spatial distribution of roots in different
85 phytophysionomies of Amazon savannahs are nonexistent, representing an important lacuna
86 in our knowledge about this potential below-ground carbon reservoir in the Amazon. These
87 data are important for understanding the role of these environments in mitigating global
88 warming (IPCC 2007).

89 The first Brazilian inventory of greenhouse gases, which was submitted to the Climate
90 Convention (UN-FCCC) in 2004, did not consider the roots, explaining that "the
91 consideration of carbon below ground (roots) is complex and was not included in this
92 inventory" (Brazil, MCT 2004, p. 146). Of course, considering the carbon stock in roots to be
93 zero, and consequently considering the emission from this source to be zero after clearing the
94 vegetation, represents a substantial underestimation, especially in savannahs, where the great
95 majority of the biomass is below ground. The second Brazilian inventory used the default
96 root : shoot ratios values presented by IPCC (2003, p. 3.109, Table 3.4.3) to estimate total
97 biomass for all grassland and savannah environments listed in Brazil, MCT (2010, pp. 236-
98 237). This was done both for the *cerrados* of central Brazil and for Amazon savannahs. The
99 root : shoot ratio is an expansion factor used to estimate below-ground biomass from above-
100 ground biomass (IPCC 2006, p. 6.7). However, use of the IPCC default values for
101 calculating below-ground biomass in open-vegetation systems under different environmental
102 conditions can cause undesired distortion in the final values for total biomass.

103 Within this context, our goal was to estimate the biomass of roots in different open
104 savannah environments in Roraima in view of the combination of two effects
105 (phytophysionomic structure and soil class, forming a "phytopedounit"). The phytopedounits
106 presented here are similar to the "landscape system units" defined by Sombroek *et al.* (2000).
107 A classification including both vegetation and soil effects is important in order to prevent
108 phytophysionomies with the same structure on different soil classes from being analysed as
109 the same ecological unit. This strategy includes spatial variations that occur along different
110 edaphic gradients that may affect the modelling of root biomass (Espeleta and Clark 2007).

111 The current study included the following questions: (i) Does the allocation of total
112 carbon to roots differ among open savannahs on different phytopedounits? (ii) Does the
113 vertical distribution of root biomass differ among these environments?, and (iii) Does the
114 ratio between the biomass of roots (total and ≥ 2 mm) and the aerial biomass (root : shoot)

115 differ among the phytopedounits investigated? Our results represent an opportunity to
 116 reformulate the estimates of below-ground biomass and carbon stocks in Amazonian
 117 savannas, providing appropriate regional values for open vegetation systems with low
 118 densities of trees and shrubs.

119

120 **Material and Methods**

121

122 *Savannahs of Roraima*

123

124 Savannahs of Roraima are part of the Rio Branco-Rupununi complex located in the
 125 triple frontier between Brazil, Venezuela and Guyana (Beard 1953; Eden 1970). Altogether,
 126 these continuous savannahs cover $68 \times 10^3 \text{ km}^2$, the Brazilian part being approximately $43 \times$
 127 10^3 km^2 (~ 63%) (Barbosa *et al.* 2007; Barbosa and Campos 2011). In general, these
 128 savannahs are located on poor soils with high frequency of fire and strong climatic
 129 seasonality that directly influences the fluctuation of the water table and the
 130 phytophysionomic structure (Miranda *et al.* 2002; Barbosa and Fearnside 2005a). The climate
 131 type of this whole region is Awi according to the Köppen classification, with average rainfall
 132 of approximately $1650 \text{ mm year}^{-1}$; the peak of the dry period is between December and March
 133 and the rainy period between May and August (Barbosa 1997). These savannahs have a wide
 134 variety of phytophysionomies ranging from grasslands that are totally devoid of trees to
 135 densely populated types on different soil classes (Brazil Projeto RADAMBRASIL 1975;
 136 Barbosa and Fearnside 2005b). The Venezuelan *llanos* have structure and species
 137 composition that are similar to those of the savannahs of Roraima (San José and Fariñas
 138 1983; Medina and Silva 1990) and neither of these should be confused with the savannahs
 139 (*cerrados*) of central Brazil (Eiten 1978).

140

141 *Study areas*

142

143 The study was carried out in two savannah areas that have sample grids for a
 144 Research Program on Biodiversity (PPBio): (i) Água Boa Experimental Station (AB) and (ii)
 145 Cauamé or “Monte Cristo” Campus (MC) (Fig. 1). The grids are composed of walking trails
 146 in the North-South (N-S) and East-West (E-W) directions that cross the area at intervals of
 147 500 m. All sampling was performed based on permanent plots (10 m \times 250 m) that are
 148 systematically distributed at points equidistant from the intersections of E-W and N-S trails.
 149 Each plot is an independent sampling unit that follows the contour line established beginning
 150 from the initial picket. This configuration was adopted to minimise the effects of topographic
 151 variability in each plot (Magnusson *et al.* 2005). All plots are individually classified by soil
 152 class and vegetation physiognomy. The general descriptions of the sample sites are given
 153 below:

154

155 *** Figure 1 ***

156

157 *Água Boa Experimental Station (AB)*

158

159 This experimental station of the Brazilian Enterprise for Agriculture and Ranching
 160 Research (EMBRAPA-Roraima) is located approximately 35 km south of the city of Boa
 161 Vista, on the BR-174 Highway (02° 51' 49" N, 02° 53' 6" N and 60° 44' 14" W, 42° 60' 27"
 162 W). The grid area is 616 ha and relief is typically flat with an average altitude of 77.7 ± 1.3
 163 m. Seventeen of the 22 terrestrial plots in this grid were sampled. The soil classes determined

164 by Brazil's National Soil Survey and Conservation Service (SNLCS) indicate that most of the
165 area has low fertility and high aluminium toxicity (Brazil SNLCS 1996).

166 Most of the grid is seasonally flooded grasslands with various species of Poaceae and
167 Cyperaceae (Araújo and Barbosa 2007). In this area the soils are typically hydromorphic and
168 of sandy texture due to an association of Gleysols with quartzo-arenitic Neosols (Entsols). A
169 smaller part of the grid has two types of savannah on clay soils that are not exposed to
170 periodic flooding (dry grasslands) and are characterised by the high density of the tree-bush
171 component: (i) low density (< 5% canopy cover), represented by grassland savannahs mixed
172 with scrubby savannah (shrublands) and (ii) medium density (5-20%), characterised by
173 shrublands mixed with grassland and savannah-parkland. In this sector of the grid the soil is
174 well drained and problems of flooding are not present.

175

176 *Cauamé Campus (MC)*

177

178 The Cauamé Campus, known as "Monte Cristo" (MC), belongs to the Federal
179 University of Roraima (UFRR) and is situated approximately 15 km north of the city of Boa
180 Vista on the BR-174 Highway segment that leads to the border with Venezuela (02° 38' 07" N
181 02° 40' 11" N and 60° 49' 25" W, 60° 52' 28" W). The grid has an area of 498 ha with an
182 average altitude of 77.3 ± 4.9 m. The relief is flat to gently rolling and is derived from the
183 Apoteri Geological Formation. This area is the most densely wooded type on clay soils. Ten
184 of the 12 terrestrial plots in this grid were sampled. The soil classes were determined by
185 Benedetti *et al.* (2011), indicating that this grid has soils with better drainage as compared to
186 the Água Boa grid.

187

188 *Sampling design and procedures*

189

190 The sampling period for collections and field assessments was between 03.06.2009
191 and 27.02.2010. Samples were paired in all plots between the rainy and the dry season. We
192 adopted this criterion in order to avoid distortions that would either under- or over-estimate
193 biomass depending on the collection period. This was necessary because there is strong
194 seasonal variation in root production in grassland and savannah areas (Neill 1992).

195 Our first goal was to quantify total aboveground biomass through direct methods (for
196 herbaceous vegetation) and indirect methods (for trees and bushes). We sampled roots using
197 two methods: (a) direct (destructive) to understand the vertical distribution of small-diameter
198 roots, which are generally associated with grasses and herbs, and (b) indirect (regression) to
199 calculate the total biomass of the root crown in the tree-bush component. Although the term
200 "root crown" is usually used to refer to roots located immediately below the surface of the
201 soil under the main stems of the plants (Snowdon *et al.* 2000), we use this term to specify
202 coarse roots at the transition point between stem and soil, including all roots ≥ 10 mm in
203 diameter up to 1 m depth. The term "root crown" has been used by Abdala *et al.* (1998) to
204 refer to roots in this diameter and depth range that are located directly beneath the aerial
205 portion of the tree, thereby distinguishing these roots from roots of the same diameter located
206 in the open spaces between the trees. However, in the case of open savannahs in Roraima,
207 where trees are widely spaced and root diameter distributions are dominated by small and
208 medium-diameter roots, the biomass of roots ≥ 10 mm in diameter is negligible in the open
209 spaces, and a separate category for these roots would have minimal effect on the overall total.

210 Total aerial biomass was estimated from the sum of its two components: (i)
211 herbaceous and (ii) tree-bush. Herbaceous biomass was defined as "grasses" (Poaceae,
212 Cyperaceae, seedlings, small dicots and litter) and woody individuals with diameter at the
213 base (Db) < 2 cm, measured at 2 cm above the ground. We sampled this group by

214 establishing four subsampling points in each permanent plot. The first subsample was
 215 established just to the right (R) of the 50-m picket, perpendicularly at a distance of 5 m from
 216 the reference line for the central trail in the permanent plot. This procedure was performed
 217 alternately using the picket at 100 m (L-left), 150 m (R) and 200 m (L).

218 After marking the four points we used a 1-m² metal frame to delimit the area for
 219 destructive sampling. All individuals in this group within the metal frame were cut close to
 220 the ground using metal blades. They were then weighed to obtain the wet weight
 221 corresponding to the subsample point. A composite sample of herbaceous biomass (80-150 g)
 222 was brought to the laboratory for determination of its dry weight after drying in an oven at
 223 70-75°C until constant weight. The total herbaceous biomass in each plot was estimated by
 224 discounting the water content from the total fresh weight of each subsample and then
 225 calculating a simple mean of the four subsamples.

226 To estimate the total carbon corresponding to the herbaceous biomass we used the
 227 carbon content (%C), in the form of a weighted average of the different components of this
 228 group as described by Barbosa (2001). The weighted average of %C was calculated
 229 separately for each experiment station: 34.4% (MC) and 36.2% (AB).

230 Live tree-bush biomass was defined as the group of woody individuals composed of
 231 two vertical strata (tree and bush or shrub) as set in Miranda *et al.* (2002) and Barbosa *et al.*
 232 (2005). The area used for sampling the arboreal stratum was 10 m × 250 m, while the shrubs
 233 were sampled in a sub-plot (2 m × 250 m). The central trail of the plot was always used as the
 234 baseline for the sampling. All individuals in the tree-bush group were identified
 235 taxonomically and inventoried by measuring biometric parameters: Db = diameter of the base
 236 of the stem measured at 2 cm above the ground; D₃₀ = diameter of the stem measured at 30
 237 cm height; Dc = diameter of the canopy calculated as the average of the largest and smallest
 238 individual crown diameter; Ht = total height, defined as the distance from the insertion of the
 239 stem in the ground to the top of the canopy. These parameters were used to indirectly
 240 estimate the biomass of each tree-bush individual based on the regression model developed
 241 by Barbosa and Fearnside (2005b) for savannahs in Roraima. Tree-bush biomass of each plot
 242 was derived from the sum of all individual biomasses.

243 The carbon corresponding to the tree-bush biomass of species inventoried in the two
 244 grids was estimated from data derived from Barbosa (2001) for biomass of savannah species
 245 in Roraima, according to the weighting given in on-line supplementary material A.

246
 247 *Total biomass of roots*

248
 249 *Direct method (destructive)*

250
 251 The sampling of root biomass using the direct (destructive) method came from the
 252 same four subsampling points established for herbaceous biomass estimates. The goal of this
 253 method was to obtain mean data for each plot at five depths in the soil column: 0-10 cm, 10-
 254 20 cm, 20-30 cm, 30-40 cm and 40-50 cm. This method checks the inventory and the vertical
 255 distribution pattern of small-diameter roots present along the altitudinal gradient in the plots.
 256 In general, these roots are associated with grasses, herbs and the lateral roots of the small-
 257 diameter individuals of the tree-bush component.

258 Each subsample was obtained at the exact geometric centre of the metal frame used to
 259 delimit the area for destructive sampling of the herbaceous biomass. We used a soil collector
 260 measuring 0.8 m in length by 0.1 m in diameter adapted for collecting soil at depths up to 0.5
 261 m. Each sample was placed directly in a plastic bag identified individually by depth, and was
 262 then weighed to obtain the net weight in the field. The samples were then forwarded to the
 263 laboratory for separation of the roots and for determination of air-dried weight. These weights

264 allowed calculation of soil density (dry weight of the soil divided by its saturated volume) for
 265 each 10-cm section of the soil column.

266 We separated live roots manually, packing them in plastic bags identified by the plot,
 267 the subsample point and the depth in the profile. After this initial screening and separation,
 268 the residual soil was subjected to the floatation process as suggested by McKell *et al.* (1961).
 269 This method consists of adding water to the residual soil so that the lighter plant material that
 270 was not visible in the first separation would float and could be collected and added to the
 271 roots separated in the previous stage. After this process the roots were placed in a drying
 272 oven at 70-75°C until they attained constant weight.

273 Throughout the process all of the collected material was separated by diameter
 274 category (d) using the classes suggested by Abdala *et al.* (1998): $d < 2$ mm (very fine + fine
 275 roots) $2 \leq d < 10$ mm (medium) and $d \geq 10$ mm (coarse). After sorting, washing and drying,
 276 all categories were weighed individually to obtain the mean biomass of roots by diameter
 277 category, depth section, plot and phytopedunit.

278

279 *Indirect method (root crown)*

280

281 This method was to estimate the biomass of the “root crown” (as defined above) of
 282 the individual trees or bushes with $D_{30} \geq 2$ cm. In open savannah ecosystems of Roraima the
 283 tree-bush component is present at low density, and our destructive method therefore cannot
 284 provide values for the biomass of the root crown of trees and shrubs. We therefore used an
 285 indirect method, applying the linear regression of Abdala *et al.* (1998) to estimate this
 286 category. We assumed that the values derived from the regression correspond to the “root
 287 crown” (roots ≥ 10 mm in diameter) connected to the trees and bushes with $D_{30} \geq 2$ cm up to
 288 1-m depth.

289

290 *Laboratory analyses*

291

292 All of the biomass of roots collected by the direct method was separated by plot and
 293 vertical section of depth, and then ground using a knife mill. The samples were then sent to
 294 the Soil and Plant Thematic Laboratory at the National Institute for Research in the Amazon
 295 (INPA), Manaus, Amazonas, Brazil, for determination of carbon content (%C). The %C was
 296 determined using a CHN Auto-Analyzer (Vario MAX, Elementar, Germany). This equipment
 297 performs the analysis by combustion at high temperatures, followed by reduction (Nelson and
 298 Sommers 1996). In the case of %C for the root crowns of trees and shrubs we used the same
 299 values for aboveground tree-bush biomass.

300

301 *Data analysis*

302

303 The differences between all values obtained for the herbaceous and tree-bush
 304 components and for the total biomass in each environment were verified using the Kruskal-
 305 Wallis non-parametric test ($H_{0.05}$) (Zar 1999). In cases where the null hypothesis (equal
 306 means) was rejected, the Student-Newman-Keuls (SNK) test was applied for multiple
 307 comparisons ($p < 0.05$).

308 All values for root biomass (total biomass and biomass by diameter category) were
 309 transformed into units of weight per unit area, for each 10-cm section of the soil profile.
 310 Using the results for the biomass of roots for each section of the 0-50 cm soil profile, we
 311 derived an estimate for the 50-100 cm section. This estimate was to combine the information
 312 from this method with the same profile (0-100 cm) adopted for calculating the root crown.
 313 For both, each result obtained from destructive subsamples (0-50 cm) was applied using an

314 exponential model ($Y = a \times b \times e^{-X}$), with a unique value for the 50-100 cm section for each
 315 subsample (see Jackson *et al.* 1996).

316 Different sections up to 1 m depth were summed to determine the biomass in the
 317 vertical soil column. We also used the Kruskal-Wallis test ($H_{0.05}$) to assess biomass
 318 differences (by diameter category and total) in the soil column and the vertical distribution
 319 patterns of biomass in all environments. The total biomass per unit area for roots to 1 m depth
 320 was added to the values estimated by regression for root crowns in the tree-bush stratum.

321 Carbon allocation in aerial biomass and roots was calculated from the multiplication
 322 of each of these groups by the corresponding carbon fraction. For calculating the root : shoot
 323 ratio for each phytopedounit we used the values of live aboveground and belowground
 324 biomass and carbon (direct and indirect methods). We also carried out a separate analysis for
 325 roots with diameter ≥ 2 mm (direct + indirect methods). The purpose of this second analysis
 326 was to provide values for Amazon open savannahs that could be used in the national
 327 inventory, as recommended by the IPCC (2006, p. 8).

328

329 Results

330

331 *Aboveground biomass and carbon*

332

333 Four phytopedounits were observed in the 27 plots sampled in the two experimental
 334 grids in open savannahs in Roraima. The main tree-bush species present in dry grasslands
 335 both on Argisols (DG-Arg) and on Latosol (DG-Lts), as well as in the more densely wooded
 336 landscapes (GP-Lts), were *Curatella americana* L. (Dilleniaceae), *Byrsonima crassifolia* (L.)
 337 Kunth (Malpighiaceae) and *B. coccolobifolia* Kunth (Malpighiaceae). In WG-Hyd woody
 338 individuals were not found with $D_{30} \geq 2$ cm.

339 Herbaceous and tree-bush biomasses differ significantly among environments (Table
 340 1). The total herbaceous biomass of WG-Hyd (9.01 ± 2.86 Mg ha⁻¹) was the largest value
 341 among all of the phytopedounits, although it only differed from the DG-Arg environment. No
 342 significant difference was detected between the largest total biomass (WG-Hyd) and the other
 343 phytopedounits due to a greater presence of the tree-bush component in the dry grasslands
 344 and in the mosaic of grasslands with parkland savannahs.

345

346 *** Table 1 ***

347

348 The WG-Hyd phytopedounit had the largest carbon stock in the herbaceous
 349 component (3.26 ± 1.04 MgC ha⁻¹), but GP-Lts (3.47 ± 0.89 MgC ha⁻¹) had the largest total
 350 aboveground carbon stock (Fig. 2). In this environment the tree-bush component represents a
 351 greater proportion of the biomass and has greater carbon content per unit of weight.

352

353 *** Figure 2 ***

354

355 *Belowground total biomass and carbon*

356

357 The total biomass of roots determined for the 0-100 cm profile (direct + indirect
 358 method) of WG-Hyd (29.52 ± 7.15 Mg ha⁻¹) was greater as compared to the other
 359 phytopedounits studied (Table 2). The phytopedounits with tree-bush biomass all have the
 360 smallest values for root biomass. Fine roots (< 2 mm) in diameter dominated the 0-100 cm
 361 vertical profile for the four open savannah phytopedounits evaluated in Roraima. The
 362 concentration of this category reached 100% in WG-Hyd and was between 92.5 and 97.9% in
 363 the other environments. The medium-diameter roots (2-10 mm) were found in three

364 landscapes that contained tree-bush biomass, with no significant differences being detected
 365 between these phytopedounits for this diameter category. Coarse roots (≥ 10 mm) were
 366 determined only by the indirect method, with no concentration of this diameter class being
 367 detected by the direct (destructive) method.

368

369 *** Table 2 ***

370

371 The carbon content (%C) in the root biomass measured by the direct method varied
 372 from 24.8% in WG-Hyd, where herbaceous biomass predominated on sandy soil, to 31.7% in
 373 GP-Lts, which was the environment with the greatest presence of tree-bush biomass on clay
 374 soil (Table 3).

375

376 *** Table 3 ***

377

378 *Vertical distribution*

379

380 The vertical distribution of root biomass for the four phytopedounits evaluated in
 381 open savannahs of Roraima, measured by the direct method, followed a pattern of
 382 exponential decrease, with the greatest values in the 0-10 cm section, and smaller values in
 383 the subsequent sections (Fig. 3). The largest concentration of roots (fine + medium) was
 384 found in the first three sections of the vertical profile of the soil (0-30 cm), ranging from 56.0
 385 to 64.6% in the four environments. Taking into consideration only these three sections of
 386 depth, the biomass of the roots of WG-Hyd was significantly different from the other
 387 environments that had a tree-bush component.

388

389 *** Figure 3 ***

390

391 *Root : Shoot ratio (biomass and carbon)*

392

393 WG-Hyd was the environment with the highest absolute root : shoot ratio, taking into
 394 consideration the total aboveground and belowground live biomass (Table 4). The values of
 395 the root : shoot ratios calculated on the basis of carbon were smaller than those calculated on
 396 the basis of biomass in all phytopedounits. Using only the carbon values for roots ≥ 2 mm in
 397 diameter, the root : shoot ratio have the highest values in GP-Lts and DG-Arg (both with high
 398 tree-bush biomass).

399

400 *** Table 4 ***

401

402 **Discussion**

403

404 *Aboveground biomass and carbon*

405

406 All of the environments evaluated had tree-bush biomass values within the expected
 407 range for open savannahs in Roraima (0.05 - 3.64 Mg ha⁻¹) (Barbosa and Fearnside 2005b). On
 408 the other hand, our values for total herbaceous biomass are closer to the values found by
 409 Castro and Kauffman (1998) (6.0 - 7.5 Mg ha⁻¹) and Castro-Neves (2007) (6.2 - 10.4 Mg ha⁻¹)
 410 for *cerrado* areas near Brasília (in central Brazil) than to those found by Barbosa and
 411 Fearnside (2005b) for open savannahs in Roraima (2.55 - 4.18 Mg ha⁻¹). Both studies in
 412 *cerrado* carried out their sampling at the peak of the dry period because they were interested
 413 in estimating the emission of greenhouse gases by burning. Our research took measures over

414 the wet and dry periods to avoid biasing the mean for each environment. In the short term this
 415 method entails higher values for herbaceous biomass due to more favourable edaphic
 416 conditions in the wet season and even in the early months of the dry season, as in the case of
 417 WG-Hyd. This environment is characterised by the dominance of a grassy stratum that makes
 418 use of the soil moisture in order to expand its above-ground biomass. In the long term,
 419 savannah ecosystems where water availability is not limiting or that are protected from fire
 420 tend to increase their belowground biomass (Sarmiento 1984; San José *et al.* 1998).

421 Our results imply that, while the herbaceous and tree-bush components are
 422 heterogeneous amongst themselves, the results for aboveground total biomass and carbon of
 423 each phytopedonunit may be considered homogeneous, representing the same set of open
 424 savannah environments in Roraima. The phytophysionomic groups with low or average
 425 density of trees and bushes have total biomass and carbon that are similar to exclusively
 426 grassy environments under periodic flooding, regardless of the soil type.

427

428 *Below-ground biomass and carbon*

429

430 Our values for total biomass of live roots (direct + indirect methods) for WG-Hyd are
 431 higher than the 11.4-18.9 Mg ha⁻¹ presented by Sarmiento and Vera (1979) for savannah
 432 gradients between grasslands and woodlands in the Venezuelan *llanos* up to 2 m depth.
 433 However, despite differences in the sampling depth, more than 90% of the roots found in the
 434 study by Sarmiento and Vera are located in the top 60 cm, which is a value very close to 76-
 435 83% of our study to 50 cm. On the other hand, the biomass of roots derived from studies in
 436 the *cerrado* of central Brazil is larger. Abdala *et al.* (1998) estimated a total value of 41.1 Mg
 437 ha⁻¹ (live + dead) in a 6.2-m profile for a *cerrado* “*sensu stricto*” on dark red latosol near
 438 Brasília, of which ~ 23.3 Mg ha⁻¹ were in the first 50 cm (excluding the root crowns).
 439 Similarly, Castro and Kauffman (1998) found values ranging from 16.3 to 52.9 Mg ha⁻¹ (2 m)
 440 for live roots in different savannah types ranging from grassland (*campo limpo* or “clean
 441 field”) to woodland (*cerrado denso* or “dense *cerrado*”), also located close to Brasília, with ~
 442 80% concentrated in the first 50 cm of depth.

443 Differences between our results for total belowground live biomass in Roraima and
 444 those for the *cerrado* of central Brazil are clearly due to sampling being performed at sites
 445 with different phytophysionomies, depths and burning schemes. Despite this contrast, it is
 446 possible to infer that, regardless of the depth or savannah type, the total biomass of live roots
 447 in areas of open savannah in Roraima with a low or medium presence of the tree-bush
 448 component are closer to those in the Venezuelan *llanos* than to those of the *cerrado* of central
 449 Brazil. This should be expected since both the Venezuelan *llanos* and the open savannahs of
 450 Roraima have similar species composition, physionomic structure, soil type and rainfall
 451 regime (San José and Fariñas 1983; Medina and Silva 1990; Miranda *et al.* 2002).

452 Another important inference is that the WG-Hyd phytopedonunit, which is grassy and
 453 seasonally flooded, can have a large absolute increment in the biomass of live roots even in
 454 hydromorphic soils. This observation was also made by Menaut and Cesar (1979) when they
 455 investigated 7 types of savannah in Lamto (Ivory Coast), also indicating that the biomass in
 456 wooded environments is almost always constant regardless of the density of trees. This
 457 contrasts with the general conclusions of Castro and Kauffman (1998) in the *cerrado* of
 458 central Brazil, indicating that dominance of aboveground woody biomass is reflected in
 459 increased belowground biomass. In our study, total carbon allocated to roots did not differ
 460 between the phytopedonunits evaluated in open savannahs of Roraima, supporting the idea of
 461 uniformity among the open environments studied with low or no tree density.

462 The concentration of fine roots in the first layers of the soil in tropical savannah and
 463 grasslands is a pattern detected globally. Oliveira *et al.* (2005) observed that up to 1 m depth

464 fine roots represented ~ 90% of the total determined for two types of *cerrado* (*campo limpo*
 465 “grassland” and *campo sujo* “scrubby savannah”) in central Brazil. In a general review,
 466 Jackson *et al.* (1996) calculated 57% (9.90 Mg ha⁻¹) as the average proportion of fine roots in
 467 the upper 30 cm of soil in tropical savannahs and grasslands. Our study indicates that in open
 468 savannahs of Roraima these figures are higher in absolute terms and can reach values almost
 469 double the general average found by Jackson and collaborators for fine roots to 30 cm depth
 470 (11.5-19.1 Mg ha⁻¹ or 55-65% for the 0-100 cm profile).

471 The most significant example is the WG-Hyd savannah type, which is seasonally
 472 flooded and has 100% fine roots (< 2 mm) throughout the sampled soil column. The plants in
 473 this type of environment are fully adapted to soils with sandy texture, periodic flooding and
 474 aluminium toxicity, but this savannah type has the largest biomass of roots even under these
 475 unfavourable edaphic conditions. In part, this expansion is explained by the prolonged
 476 maintenance of moisture in the soil in these phytopedounits even during the dry season. WG-
 477 Hyd has the largest concentration of roots between 0 and 10 cm depth (26.4%) and the lowest
 478 between 50 and 100 cm depth (17.1%), suggesting that the exploitation of nutrients in this
 479 soil is very superficial. Environments with greater presence of grasses are more efficient in
 480 absorbing water and nutrients in the upper soil layers because of the high concentrations of
 481 fine roots (Knoop and Walker 1985). In addition, sandy soils can also have a positive effect
 482 on root biomass increase as compared to soils with more clayey soil texture (Silver *et al.*
 483 2000). Roots with smaller diameter have higher surface area relative to their size or weight
 484 and are more effective in capturing water and nutrients (Newman 1966; Vitousek and
 485 Sanford 1986). Nutrient-poor tropical environments therefore tend to have larger quantities of
 486 fine roots in the upper layers of the soil, with high rates of replacement (turnover rate), and
 487 better capacity to absorb nutrients (Jordan and Escalante 1980; Priess *et al.* 1999).

488 The larger-diameter roots (≥ 2 mm) are essential for the calculation of the total
 489 belowground biomass and carbon stock, even in environments with low tree-bush density, as
 490 in the case of open savannahs in Roraima. The direct method allowed us to sample medium-
 491 diameter roots (2-10 mm) under conditions of lateral rooting. Adding this medium-root
 492 biomass to the biomass determined by the indirect method for coarse roots (≥ 10 mm)
 493 indicates that 0-7.5% (0-1.65 Mg ha⁻¹) of the total belowground biomass in savannahs with
 494 low tree-bush density in the far northern part of Amazonia is live roots with ≥ 2 mm
 495 diameter. In the more-wooded environments of the *cerrado* of central Brazil, the biomass of
 496 this component can reach values > 20 Mg ha⁻¹ (Abdala *et al.* 1998; Castro-Neves 2007),
 497 depending on the tree-bush structure and density.

498 The smaller carbon content (%C) in roots found in the 0-50 cm soil column suggests a
 499 direct relationship with the large quantity of fine roots found in all of the savannah types
 500 studied. For example, Manlay *et al.* (2002) also found low values for carbon content (29.8-
 501 35.1% C) for fine roots under agricultural crops established in savannah areas in West Africa.
 502 Carbon content values lower than 40% are not common in the literature, but can be expected
 503 where the material analysed does not have lignified parenchyma. Fine roots are characterised
 504 as non-ligneous, almost all being without bark and with a short life cycle (McClaugherty *et*
 505 *al.* 1982). These smaller-diameter roots die steadily throughout the year and quickly
 506 disappear from the system (Yavitt and Wright 2001) providing an important source of
 507 organic matter and mineral nutrients for maintenance and functioning of ecosystems (Luizão
 508 *et al.* 1992).

509 Gill and Jackson (2000) presented a range of 0.64 - 0.88 for the turnover rate in open
 510 environments in tropical zones (grasslands, shrublands and wetlands). Taking the midpoint of
 511 this range (0.76) and applying the results derived for the total carbon stock of roots in the
 512 phytopedounits evaluated in Roraima (Table 3), we estimate an annual carbon cycling on the
 513 order of 4.7-5.5 MgC ha⁻¹ (0 - 100 cm). In temperate forest ecosystems it is estimated that ~

514 1/3 of this carbon is used in the production of new roots (Nadelhoffer and Raich 1992), but
 515 there are no estimates of this for tropical Amazonian savannahs and grasslands.

516

517 *Vertical distribution*

518

519 The distribution pattern of root biomass within the vertical soil profile observed in
 520 four phytopedounits was typically exponential with most of the roots concentrated in the
 521 surface layers. This pattern is the same as that observed in other studies of Neotropical
 522 savannahs and grasslands in northern South America (Sarmiento and Vera 1979; San José *et al.*
 523 *et al.* 1982), in Central America (Kellman and Sanmugadas 1985; Fiala and Herrera 1988) and
 524 in the Brazilian *cerrados* (Abdala *et al.* 1998; Castro and Kauffman 1998; Delitti *et al.* 2001;
 525 Rodin 2004; Oliveira *et al.* 2005; Castro-Neves 2007; Paiva and Faria 2007). This is also an
 526 overall global pattern observed in other ecosystems where the great majority of roots is
 527 concentrated in the top 30 cm of the soil (Jackson *et al.* 1996; Schenk and Jackson 2002a).

528 In our case the extinction or decay coefficient (β) cannot be calculated using the
 529 formula of Gale and Grigal (1987) because our data were divided into 10-cm sections only up
 530 to 50 cm depth, whereas at least 100 cm would need to be so divided for a calculation of this
 531 type (see Jackson *et al.* 1996). However, based on the decay pattern in the current data up to
 532 30 cm depth, we suggest that all environments investigated have a superficial root
 533 distribution, with WG-Hyd (grassy environment on sandy soil) being most prominent ($\beta_{30\text{ cm}}$
 534 = 0.95) as compared to tree-bush environments on clay soils ($\beta_{30\text{ cm}}$ = 0.96).

535 These values are lower than those given in the general review of Jackson *et al.* (1996)
 536 for the tropical savannah and grassland biome (0.972), but this could be a reflection of the
 537 small number of studies available at the time of the review (5 in Africa, 1 in India and 1 in
 538 Cuba). Recent investigations in the *cerrado* of central Brazil found values of 0.97 (*cerrado*
 539 *stricto sensu*) and 0.99 (*campo sujo* “shrublands”) (Rodin 2004), and ranging from 0.88 to
 540 0.92 for *cerrado stricto sensu* under different burning regimes (Castro-Neves 2007). This
 541 variation in values indicates that β is very variable and is highly dependent on the time of
 542 sampling (dry or wet season), soil type (clay or sand), drainage of the environment
 543 (hydromorphy) and phytophysionomy (grassy or different forms of wooded savannah).

544

545 *Root : Shoot ratio*

546

547 Use of the root : shoot ratio as an indicator of the relationship between the below-
 548 ground and aerial biomass (total live) is very important because it can serve as an estimator of
 549 below-ground carbon based on a simple biometric survey of above-ground biomass with
 550 lower costs (Schenk and Jackson 2002b). Realistic root : shoot ratios are necessary to
 551 improve the accuracy of estimates of root biomass and to estimate the effects of management
 552 and land-use change in national inventories of greenhouse gas emissions (Mokany *et al.*
 553 2006). In our study we calculate the root : shoot ratio based on biomass and carbon. This
 554 latter form provides ecological values closer to reality for calculation of stock, production and
 555 ecosystem productivity. This is because the carbon content (%C) of the different above-
 556 ground components is not the same as that applied to below-ground biomass. In ecosystems
 557 where the biomass of fine roots is overwhelmingly superior to the other categories, as in the
 558 case on the open savannahs of Roraima, the carbon content can be lower, causing the root :
 559 shoot ratio based on biomass to not represent the ecosystem faithfully.

560 The values of the root : shoot ratio based on total live biomass varied between 2.7 and
 561 3.8, reflecting discrepancies between the total values above and below ground for all
 562 phytopedounits. Higher ratios (3 to 5) were determined in the savannahs in Lamto (Ivory
 563 Coast), indicating greater total belowground biomass to a depth of 1 m as compared to aerial

564 biomass (Menaut and Cesar 1979). However, these values are extremely variable and
 565 dependent on the depth of sampling. In the *cerrado* of central Brazil, Castro and Kauffman
 566 (1998) found high values for savannahs with low tree density (5.6-7.7) and smaller values for
 567 more wooded phytophysionomies (2.6 - 2.9) to 2 m depth, even without including any
 568 estimate for the biomass of the root crowns. Thus, although the phytopedounits investigated
 569 in Roraima are limited by the low density of tree individuals, our values are closer to those of
 570 the wooded *cerrado* environments of Castro and Kauffman (1998) than to grassland
 571 environments. Our results indicate that the total biomass of roots (0-100 cm depth) is a
 572 component of great importance in the open savannah environments of Roraima, representing
 573 2.4-3.3 times the total carbon allocated to above-ground biomass.

574 The IPCC (2006, p. 4.72) suggests that fine roots (< 2 mm) are an integral part of the
 575 soil and, therefore, should be considered in the calculations of soil carbon. To have a valid
 576 correction for this it is necessary to disaggregate the results and use only the categories of
 577 roots ≥ 2 mm in diameter. This is required to prevent double counting of inventory values
 578 derived for soil carbon stocks. Thus, using our results for roots with diameter ≥ 2 mm for
 579 open savannah phytopedounits studied in Roraima, the root : shoot ratio, based on biomass, is
 580 between 0 (seasonally flooded grasslands) and 0.07-0.20 (grasslands with low tree density),
 581 or between 0 and 0.08-0.24, based on carbon (see Table 4). These values are lower than those
 582 indicated as the default values by the IPCC (2003, p. 3.109, Table 3.4.3; 2006, p. 6.8, Table
 583 6.1) for sub-tropical/ tropical grassland (1.6), woodland/savannah (0.5) and shrubland (2.8).
 584 However, the IPCC strongly suggests that default values only be used when the country does
 585 not have regional values that better reflect the ecosystem (IPCC 2006, p. 6.8).

586 The second Brazilian inventory used the default values for all grassland and savannah
 587 environments as listed in Brazil, MCT (2010, pp. 236-237). This was done both for the
 588 *cerrados* of central Brazil (for which published estimates of below-ground biomass existed)
 589 and for Amazon savannahs (for which the present study provides the first estimates).
 590 Although few in number, it is possible to make inferences about the root : shoot ratio for
 591 Brazilian savannahs, including *cerrados* (Table 5). For example, the Brazilian estimates for
 592 root : shoot ratio (roots ≥ 2 mm) vary tremendously depending on the vegetation type, fire
 593 regime, seasonality of the water table, soil class and sampling depth. Environments in central
 594 Brazil with greater above-ground biomass and that are not influenced by the water table have
 595 root : shoot ratios from 7 to 27 times higher than those in Amazonian grasslands and
 596 savannahs with low arboreal biomass.

597
 598 *** Table 5 ***
 599

600 We therefore propose a reformulation of the calculations for the next Brazilian
 601 national inventory. We suggest a minimum standardization of 1 m depth for the estimates of
 602 below-ground biomass and carbon, in addition to region-specific values for root : shoot
 603 ratios, with different ratios, for Amazonian grasslands/savannahs and for central Brazilian
 604 *cerrados*. This calculation strategy would bring advantages in avoiding the use of empirical
 605 default values from the IPCC (2003; 2006), thereby providing more realistic values for total
 606 biomass and carbon for ecosystems with open vegetation in Brazil.

607 608 **Conclusions**

609
 610 The total biomass of roots of seasonally flooded grasslands is higher than the root
 611 biomass of grasslands with low tree-bush density, although the total below-ground carbon
 612 stock does not differ among phytopedounits.

613 The vertical distribution pattern of root biomass follows an exponential model, with
 614 the largest concentration of roots being in the more superficial layers of the soil. This pattern
 615 does not differ among phytopedounits.

616 The total biomass of roots (direct + indirect methods) in open savannah environments
 617 of Roraima represents a pool 2.4 - 3.3 times the total carbon stocked in above-ground
 618 biomass.

619 The expansion factor (root : shoot ratio) used by IPCC for below-ground biomass in
 620 roots ≥ 2 mm diameter, starting from live above-ground biomass, is zero for seasonally
 621 flooded grasslands of Roraima (in northern Amazonia). For unflooded grasslands with low
 622 densities of trees the values of this factor range from 0.07 to 0.20 on a biomass basis, or from
 623 0.08 to 0.24 on a carbon basis.

624 The standardization of the minimum sampling depth and the use of region-specific
 625 values for root : shoot ratios to calculate below-ground biomass in grasslands and savannahs
 626 is advantageous because it provides more realistic values of total biomass and carbon.

627

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629

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638

639 **References**

640 Abdala GC, Caldas LS, Haridasan M, Eiten G (1998) Above and belowground organic matter
 641 and root:shoot ratio in a cerrado in Central Brazil. *Brazilian Journal of Ecology* **2**, 11-
 642 23.

643 Araújo ACO, Barbosa RI (2007) Riqueza e diversidade do estrato arbóreo-arbustivo de duas
 644 áreas de savanas em Roraima, Amazônia Brasileira. *Mens Agitat* **2(1)**, 11-18.

645

646 Asner GP, Elmore AJ, Olander LP, Martin RE, Harris AT (2004) Grazing systems,
 647 ecosystem responses, and global change. *Annual Review of Environment and*
 648 *Resources* **29**, 261–299.

649

650 Barbosa RI (1997) 'Distribuição das chuvas em Roraima.' In: Barbosa RI, Ferreira EJJ,
 651 Castellon EG (Eds.), *Homem, Ambiente e Ecologia no Estado de Roraima*. INPA:
 652 Manaus, Amazonas, Brazil, pp. 325-335.

653

654 Barbosa RI (2001) 'Savanas da Amazônia: emissão de gases do efeito estufa e material
 655 particulado pela queima e decomposição da biomassa acima do solo, sem a troca do
 656 uso da terra, em Roraima, Brasil.' Ph.D. Thesis in Ecology, Instituto Nacional de
 657 Pesquisas da Amazônia / Universidade do Amazonas: Manaus, Amazonas, Brazil.

658

- 659 Barbosa RI, Campos C (2011) Detection and geographical distribution of clearing areas in the
660 savannas ('lavrado') of Roraima using Google Earth web tool. *Journal of Geography*
661 *and Regional Planning* **4(3)**, 122-136.
662
- 663 Barbosa RI, Fearnside PM (2005a) Fire frequency and area burned in the Roraima savannas
664 of Brazilian Amazonia. *Forest Ecology and Management* **204**, 371-384.
665
- 666 Barbosa RI, Fearnside PM (2005b) Above-ground biomass and the fate of carbon after
667 burning in the savannas of Roraima, Brazilian Amazonia. *Forest Ecology and*
668 *Management* **216**, 295-316.
669
- 670 Barbosa RI, Nascimento SP, Amorim PAF, Silva RF (2005) Notas sobre a composição
671 arbóreo-arbustiva de uma fisionomia das savanas de Roraima, Amazônia Brasileira.
672 *Acta Botanica Brasilica* **19**, 323-329.
673
- 674 Barbosa RI, Campos C, Pinto F, Fearnside PM (2007) The "Lavrados" of Roraima:
675 Biodiversity and Conservation of Brazil's Amazonian Savannas. *Functional*
676 *Ecosystems and Communities* **1(1)**, 29-41.
677
- 678 Baruch Z (1994) Responses to drought and flooding in tropical forage grasses I. Biomass
679 allocation, leaf growth and mineral nutrients. *Plant and Soil* **164**, 87-96.
680
- 681 Beard JS (1953) The savanna vegetation of northern tropical America. *Ecological*
682 *Monographs* **23**, 149-215.
683
- 684 Benedetti UG, Vale Jr JF, Schaefer CEGR, Melo VF, Uchôa SCP (2011) Gênese, química e
685 mineralogia de solos derivados de sedimentos Pliopleistocênicos e de rochas
686 vulcânicas básicas em Roraima, norte amazônico. *Revista Brasileira de Ciência do*
687 *Solo* **35**, 299-312.
688
- 689 Brazil, MCT (2004) 'Brazil's Initial National Communication to the United Nations
690 Framework Convention on Climate Change.' MCT (Ministério de Ciência e
691 Tecnologia): Brasília, Brazil.
692
- 693 Brazil, MCT (2010) 'Segunda Comunicação Nacional do Brasil à Convenção-Quadro das
694 Nações Unidas sobre Mudança do Clima.' MCT (Ministério de Ciência e
695 Tecnologia): Brasília, Brazil. (volume 1).
696 (http://www.mct.gov.br/upd_blob/0213/213909.pdf).
697
- 698 Brazil, Projeto RADAMBRASIL (1975) 'Levantamento dos Recursos Naturais (Vol. 8).'
699 Ministério das Minas e Energia: Rio de Janeiro, Brazil.
700
- 701 Brazil, SNLCS (1996) 'Levantamento Semidetalhado dos Solos e Aptidão Agrícola das terras
702 do Campo Experimental Água Boa do CPAF-RR, Estado de Roraima.' Empresa
703 Brasileira de Pesquisa Agropecuária (EMBRAPA), Serviço Nacional de
704 Levantamento e Conservação do Solo (SNLCS): Rio de Janeiro, Brazil.
705
- 706 Castro EA, Kauffman JB (1998) Ecosystem structure in the Brazilian Cerrado: a vegetation
707 gradient of aboveground biomass, root mass and consumption by fire. *Journal of*
708 *Tropical Ecology* **14**, 263-283.

- 709
710 Castro-Neves BM (2007) 'Efeito de queimadas em áreas de cerrado Strictu Sensu e na
711 biomassa de raízes finas.' Thesis in ecology, Universidade de Brasília: Brasília,
712 Brazil.
- 713
714 Cattanio JH, Anderson AB, Rombold JS, Nepstad DC (2004) Phenology, litterfall, growth,
715 and root biomass in a tidal floodplain forest in the Amazon estuary. *Revista Brasileira*
716 *de Botânica* **27**, 703-712.
- 717
718 Chen X, Eamus D, Hutley LB (2004) Seasonal patterns of fine-root productivity and turnover
719 in a tropical savanna of northern Australia. *Journal of Tropical Ecology* **20**, 221–224.
720
- 721 Delitti WBC, Pausas JG, Burger DM (2001) Belowground biomass seasonal variation in two
722 Neotropical savannahs (Brazilian Cerrados) with different fire histories. *Annals of*
723 *Forest Science* **58**, 713–721.
- 724
725 Eden M (1970) Savanna vegetation in the northern Rupununi, Guyana. *The Journal of*
726 *Tropical Geography* **30**, 17-28.
- 727
728 Eiten G (1978) Delimitation of the Cerrado concept. *Plant Ecology* **36**, 169-178.
729
- 730 Espeleta JF, Clark DA (2007) Multi-scale variation in fine-root biomass in a tropical rain
731 forest: a seven-year study. *Ecological Monographs* **77**, 377-404.
732
- 733 February EC, Higgins SI (2010) The distribution of tree and grass roots in savannas in
734 relation to soil nitrogen and water. *South African Journal of Botany* **76**, 517-523.
735
- 736 Fiala K, Herrera R (1988) Living and dead belowground biomass and its distribution in some
737 savanna communities in Cuba. *Folia Geobotanica* **23**, 225-237.
738
- 739 Furley P (1999) The nature and diversity of neotropical savanna vegetation with particular
740 reference to the Brazilian cerrados. *Global Ecology and Biogeography* **8**, 223-241.
741
- 742 Gale MR, Grigal DF (1987) Vertical distributions of northern tree species in relation to
743 successional status. *Canadian Journal of Forest Research* **17**, 829–834.
744
- 745 Gill RA, Jackson RB (2000) Global patterns of root turnover for terrestrial ecosystems. *New*
746 *Phytologist* **147**, 13-31.
747
- 748 IPCC (2003). '2003 IPCC Good Practice Guidance for Land Use, Land-Use Change and
749 Forestry', Penman J, Gytarsky M, Hiraishi T, Krug T, Kruger D, Pipatti R, Buendia L,
750 Miwa K, Ngara T, Tanabe K, Wagner F (Eds.), Institute for Global Environmental
751 Strategies (IGES) for the Intergovernmental Panel on Climate Change (IPCC):
752 Kanagawa, Japan.
753
- 754 IPCC (2006) '2006 IPCC Guidelines for National Greenhouse Gas Inventories. Prepared by
755 the National Greenhouse Gas Inventories Programme', Eggleston HS, Buendia L,
756 Miwa K, Ngara T, Tanabe K (Eds.), Intergovernmental Panel on Climate Change
757 (IPCC) and Institute for Global Environmental Strategies (IGES): Kanagawa, Japan.
758

- 759 IPCC (2007) 'Climate Change 2007: Synthesis Report. Intergovernmental Panel on Climate
760 Change (IPCC Plenary XXVII; Valencia, Spain, 12-17 November 2007)'. Working
761 Group contributions to the Fourth Assessment Report - IPCC: Geneva, Switzerland.
762
- 763 Jackson RB, Canadell J, Ehleringer JR, Mooney HA, Sala OE, Schulze ED (1996) A global
764 analysis of root distributions for terrestrial biomes. *Oecologia* **108**, 389-411.
765
- 766 Jackson RB, Mooney HA, Schulze ED (1997) A global budget for fine root biomass, surface
767 area, and nutrient contents. *Ecology* **94**, 7362-7366.
768
- 769 Jordan CF, Escalante G (1980) Root productivity in an Amazonian rain forest. *Ecology* **61**,
770 14-18.
771
- 772 Kellman M, Sanmugadas K (1985) Nutrient retention by savanna ecosystems I. retention in
773 the absence of fire. *Journal of Ecology* **73**, 935-951.
774
- 775 Klinge H (1973) Root mass estimation in lowland tropical rain forests of central Amazonia
776 Brazil. *Tropical Ecology* **14**, 29-38.
777
- 778 Knoop WT, Walker BH (1985) Interactions of woody and herbaceous vegetation in a
779 southern African savanna. *Journal of Ecology* **73**, 235-253.
780
- 781 Luizão F, Luizão R, Chauvel A (1992) Premiers résultats sur la dynamique des biomasses
782 racinaires et microbiennes dans un latosol d'Amazonie centrale (Brésil) sous forêt et
783 sous pâturage. *Cahiers Orstom, serie Pédologie* **27**, 69-79.
784
- 785 Magnusson WE, Lima AP, Luizão R, Luizão F, Costa FRC, Castilho CV, Kinupp VF (2005)
786 RAPELD: A modification of the Gentry method for biodiversity surveys in long-term
787 ecological research sites. *Biota Neotropica* **5(2)**,
788 <http://www.biotaneotropica.org.br/v5n2/pt/abstract?point-of-view+bn01005022005>.
789
- 790 Manlay RJ, Chotte JL, Masse D, Laurent JY, Feller C (2002) Carbon, nitrogen and
791 phosphorus allocation in agro-ecosystems of a West African savanna. III. Plant and
792 soil components under continuous cultivation. *Agriculture, Ecosystems and*
793 *Environment* **88**, 249-269.
794
- 795 McLaugherty CA, Aber JD, Melillo JM (1982) The role of fine roots in the organic matter
796 and nitrogen budgets of two forested ecosystems. *Ecology* **63**, 1481-1490.
797
- 798 McKell CM, Wilson AM, Jones MB (1961) A flotation method for easy separation of roots
799 from soil samples. *Agronomy Journal* **253**, 56-57.
800
- 801 McNaughton SJ, Banyikwa FF, McNaughton MM (1998) Root biomass and productivity in a
802 grazing ecosystem: the Serengeti. *Ecology* **79**, 587-592.
803
- 804 Medina E, Silva JF (1990) Savannas of northern South America: a steady state regulated by
805 water-fire interactions on a background of low nutrient availability. *Journal of*
806 *Biogeography* **17**, 403-413.
807

- 808 Menaut JC, Cesar J (1979) Structure and primary productivity of Lamto Savannas, Ivory
809 Coast. *Ecology* **60**, 1197-1210.
- 810
- 811 Milchunas DG, Lauenroth WK (1993) Quantitative effects of grazing on vegetation and soils
812 over a global range of environments. *Ecological Monographs* **63**, 327-366.
- 813
- 814 Miranda IS, Absy ML, Rebêlo GH (2002) Community structure of woody plants of Roraima
815 savannahs, Brazil. *Plant Ecology* **164**, 109-123.
- 816
- 817 Mokany K, Raison RJ, Prokushkin AS (2006) Critical analysis of root:shoot ratios in
818 terrestrial biomes. *Global Change Biology* **12**, 84-96.
- 819
- 820 Nadelhoffer KJ, Raich JW (1992) Fine root production estimates and belowground carbon
821 allocation in forest ecosystems. *Ecology* **73**, 1139-1147.
- 822
- 823 Neill C (1992) Comparison of soil coring and ingrowth methods for measuring belowground
824 production. *Ecology* **73**, 1918-1921.
- 825
- 826 Nelson DW, Sommers LE (1996) 'Total carbon and total nitrogen.' In: Sparks DL (ed.),
827 Methods of Soil Analysis: Part 3 – Chemical Methods. SSSA Book Series No. 5,
828 SSSA & ASA: Madison, Wisconsin.
- 829
- 830 Nepstad D, Carvalho CR, Davidson EA, Jipp PH, Lefebvre PA, Negreiros GH, Silva ED,
831 Stone TA, Trumbore SA, Vieira S (1994) The role of deep roots in the hydrological
832 and carbon cycles of Amazonian forests and pastures. *Nature* **372**, 666-669.
- 833
- 834 Newman EI (1966) A method of estimating the total length of root in a sample. *Journal of*
835 *Applied Ecology* **3**, 139-145.
- 836
- 837 Oliveira RS, Bezerra L, Davidson ED, Pinto F, Klink CA, Nepstad D, Moreira A (2005)
838 Deep root function in soil water dynamics in cerrado savannas of central Brazil.
839 *Functional Ecology* **19**, 574-581.
- 840
- 841 Paiva OA, Faria GE (2007) Estoque de carbono do solo sob cerrado sensu stricto no Distrito
842 Federal, Brasil. *Revista Trópica (Ciências Agrárias e Biológicas)* **1(1)**, 59-65.
- 843
- 844 Pandey CB, Singh JS (1992) Rainfall and grazing effects on net primary productivity in a
845 tropical savanna, India. *Ecology* **73**, 2007-2021.
- 846
- 847 Priess J, Then C, Fölster H (1999) Litter and fine-root production in three types of tropical
848 premontane rain forest in SE Venezuela. *Plant Ecology* **143**, 171-187.
- 849
- 850 Rodin P (2004) 'Distribuição da biomassa subterrânea e dinâmica de raízes finas em
851 ecossistemas nativos e em uma pastagem plantada no Cerrado do Brasil Central.'
852 MSc Thesis, Universidade de Brasília, Instituto de Ciências Biológicas: Brasília,
853 Brazil.
- 854
- 855 San José JJ, Berrade F, Ramirez J (1982) Seasonal changes of growth mortality and
856 disappearance of belowground root biomass in the *Trachypogon* savanna grass. *Acta*
857 *Oecologia (Oecologia Plantarum)* **3**, 347-352.

- 858
859 San José JJ, Fariñas MR (1983) Change in tree density and species composition in a protected
860 Trachypogon savanna, Venezuela. *Ecology* **64**, 447-453.
861
- 862 San José JJ, Montes RA, Fariñas MR (1998) Carbon stocks and fluxes in a temporal scaling
863 from a savanna to a semi-deciduous forest. *Forest Ecology and Management* **105**,
864 251-262.
865
- 866 Santos CPF, Valles GF, Sestini MF, Hoffman P, Dousseau SL, Homem de Mello AJ (2007)
867 'Mapeamento dos Remanescentes e Ocupação Antrópica no Bioma Amazônia.' In:
868 Anais do XIII Simpósio Brasileiro de Sensoriamento Remoto (Florianópolis, Brasil,
869 21-26 abril 2007), Instituto Nacional de Pesquisas Espaciais (INPE), São José dos
870 Campos, Brazil, pp. 6941-6948.
871 <http://mar.te.dpi.inpe.br/rep/dpi.inpe.br/sbsr@80/2006/11.18.01.25?mirror=dpi.inpe.br/banon/2003/12.10.19.30.54&metadatarpository=dpi.inpe.br/sbsr@80/2006/11.18.01.25.31>
872
873
874
- 875 Sarmiento G (1984) 'The Ecology of Neotropical Savannas.' (Harvard University Press:
876 Cambridge, Massachusetts).
877
- 878 Sarmiento G, Vera M (1979) Composición, estructura, biomasa y producción primaria de
879 diferentes sabanas en los Llanos occidentales de Venezuela. *Boletín de la Sociedad*
880 *Venezolana de Ciencia Natural* **34(136)**, 5-41.
881
- 882 Scholes RJ, Hall DO (1996) 'The carbon budget of tropical savannas, woodlands and
883 grasslands.' In: Breyer AI, Hall DO, Melillo JM, Agren GI (Eds.), *Global*
884 *Change: Effects on Coniferous Forests and Grasslands*. SCOPE (v. 56). Wiley &
885 Sons: Chichester, UK. pp. 69-100.
886
- 887 Schenk HJ, Jackson RB (2002a) The global biogeography of roots. *Ecological Monographs*
888 **72**, 311-328.
889
- 890 Schenk HJ, Jackson RB (2002b) Rooting depths, lateral root spreads and below-
891 ground/above-ground allometries of plants in water-limited ecosystems. *Journal of*
892 *Ecology* **90**, 480-494.
893
- 894 Silver WL, Neff J, McGroddy M, Veldkamp E, Keller M, Cosme R (2000) Effects of soil
895 texture on belowground carbon and nutrient storage in a lowland Amazonian forest
896 ecosystem. *Ecosystems* **3**, 193-209.
897
- 898 Snowdon P, Eamus D, Gibbons P, Khanna P, Keith H, Raison J, Kirschbaum M (2002)
899 'Synthesis of allometrics, review of root biomass and design of future woody biomass
900 sampling strategies.' Technical Report no. 17 (National Carbon Accounting System):
901 Australia.
902
- 903 Solbrig OT, Medina E, Silva JF (1996) 'Biodiversity and tropical savanna properties: a global
904 view.' In: Mooney HA, Cushman JH, Medina E, Sala OE, Schulze ED (Eds.),
905 *Functional Roles of Biodiversity: A Global Perspective*. SCOPE (ICSU / UNEP).
906 Wiley & Sons: Chichester, UK. pp. 185-211.
907

- 908 Sombroek WG, Fearnside PM, Cravo M (2000) 'Geographic assessment of carbon stored in
909 Amazonian terrestrial ecosystems and their soils in particular.' In: Lal R, Kimble JM,
910 Stewart BA (Eds.), *Global Climate Change and Tropical Ecosystems*. Advances in
911 Soil Science, CRC Press: Boca Raton, Florida. pp. 375-389.
912
- 913 Stanton NL (1988) The underground in grasslands. *Annual Review of Ecology and*
914 *Systematics* **19**, 573-89.
915
- 916 Thompson J, Proctor J, Viana V, Milliken W, Ratter JA, Scott DA (1992) Ecological studies
917 on a lowland evergreen rain forest on Maraca Island, Roraima, Brazil. I. Physical
918 environment, forest structure and leaf chemistry. *Journal of Ecology* **80**, 689-703.
919
- 920 Vitousek PM, Sanford RL (1986) Nutrient cycling in moist tropical forest. *Annual Review of*
921 *Ecology and Systematics* **17**, 137-167.
922
- 923 Yavitt JB, Wright SJ (2001) Drought and irrigation effects on fine root dynamics in a
924 Tropical Moist Forest, Panama. *Biotropica* **33**, 421-434.
925
- 926 Zar JH (1999) 'Biostatistical Analysis (4th Ed.)' (Prentice-Hall: Englewood Cliffs, New
927 Jersey).

928 FIGURE LEGENDS

929

930 Fig. 1 – Location of the two sample areas (AB = Água Boa; MC = Monte Cristo) established
931 in savannahs (*lavrado*) in Roraima, Brazil.

932

933 Fig. 2 – Distribution of aboveground biomass carbon stock in two components (herbaceous
934 and tree-bush) for the four phytotopedounits sampled in open savannahs of Roraima, Brazil.

935

936 Fig. 3 - Vertical distribution of root biomass (Mg ha^{-1}) by depth interval as estimated by the
937 direct method (50-100 cm, calculated by exponential regression) in four open savannah
938 phytotopedounits evaluated in Roraima. Values with the same letter in each depth interval
939 have no significant difference between means, as determined the by SNK test; $p < 0.05$.

940

941

1 **Table 1. Aerial biomass distribution by group in different phytopedounits in two grids**
 2 **of open savannahs in Roraima, Brazil (mean \pm SD Mg ha⁻¹).** Values in parentheses
 3 represent the plot's live component (Mg ha⁻¹) with the litter (dead biomass) already
 4 discounted. Different lower-case letters indicate distinct significance among values in each
 5 column (SNK test; $p < 0.05$).

Phytopedounit (A)	Number of Plots (n)		Herbaceous	Tree-bush	Total
	AB	MC			
DG-Arg	0	4	5.25 \pm 0.36 a (4.59)	1.06 \pm 0.68 bc	6.31 \pm 0.88 a (5.65)
DG-Lts	5	3	6.74 \pm 1.91 ab (5.89)	0.60 \pm 1.08 b	7.34 \pm 1.96 a (6.50)
GP-Lts	2	3	6.10 \pm 2.78 ab (5.34)	2.76 \pm 1.59 c	8.87 \pm 2.43 a (8.10)
WG-Hyd	10	0	9.01 \pm 2.86 b (7.65)	0.00 a	9.01 \pm 2.86 a (7.65)

7
 8 (A) Dry grasslands on Argisols = DG-Arg; Dry grasslands on Latosols = DG-Lts;
 9 Mosaic of grasslands with savannah-parkland on Latosols = GP-Lts; Wet grasslands on
 10 Hydromorphic soils = WG-Hyd (AB = Água Boa and MC = Cauamé/Monte Cristo).

11

1 **Table 2. Distribution of root biomass (mean \pm SD) by diameter category (0-100 cm).**

2 Different lower-case letters in each column indicate a distinct difference among values

3 (SNK test; $p < 0.05$).

Phytopedounit	Fine Roots	Medium Roots	Coarse Roots	Total
	(< 2 mm)	(2-10mm)	(\geq 10 mm)	
	Mg ha ⁻¹	Mg ha ⁻¹	Mg ha ⁻¹	Mg ha ⁻¹
DG-Arg	20.27 \pm 1.39 a	0.26 \pm 0.15 b	0.87 \pm 0.72 bc	21.40 \pm 2.47 a
DG-Lts	22.14 \pm 1.47 a	0.14 \pm 0.10 b	0.33 \pm 0.33 b	22.62 \pm 2.21 a
GP-Lts	20.49 \pm 1.69 a	0.39 \pm 0.15 b	1.26 \pm 0.22 c	22.14 \pm 4.90 a
WG-Hid	29.52 \pm 2.40 b	0.00 a	0.00 a	29.52 \pm 7.15 b

4

5

1 **Table 3. Carbon content (%) and total carbon stock (mean \pm SD) in roots of open**
 2 **savanna phytodounits in Roraima, Brazil (0-100 cm depth).** Mean %C was calculated
 3 by weighting between direct and indirect methods. There is no significant difference between
 4 values with the same letter on each line (SNK test; $p < 0.05$).

Phytodounit	Sub-Total (Direct Method)		Sub-Total (Indirect Method)		Total	
	MgC ha ⁻¹	%C	MgC ha ⁻¹	%C	MgC ha ⁻¹	%C
DG-Arg	6.28 \pm 0.29 a	31.10 \pm 2.90	0.41 \pm 0.34 bc	46.80	6.69 \pm 0.29 a	32.06
DG-Lts	6.04 \pm 1.15 a	27.03 \pm 2.63	0.21 \pm 0.14 b	46.73	6.25 \pm 1.12 a	27.69
GP-Lts	6.62 \pm 2.00 a	31.73 \pm 4.10	0.58 \pm 0.10 c	46.09	7.21 \pm 1.85 a	32.89
WG-Hid	7.10 \pm 1.65 a	24.80 \pm 6.16	0.00 a	-	7.10 \pm 1.65 a	24.80

6

1 **Table 4. Root : shoot ratio in different phytopedounits sampled in open savannas of**
2 **Roraima, Brazil for total (live) biomass and carbon, and separately for roots with ≥ 2**
3 **mm diameter .**

4

Phytopedounit	Root :Shoot			
	Total		≥ 2 mm	
	Biomass	Carbon	Biomass	Carbon
DG-Arg	3.79	3.33	0.20	0.24
DG-Lts	3.48	2.64	0.07	0.08
GP-Lts	2.73	2.41	0.20	0.24
WG-Hyd	3.86	2.56	0	0

5

6

1 **Table 5. Root : shoot ratio in the form used by the IPCC (ratio the biomass of roots**
 2 **≥ 2 mm in diameter to live aboveground biomass) from the current study and**
 3 **recalculated from published studies on other Brazilian savannas.**

4

Phytophysionomy	IBGE Legend ^A	Depth (m)	Root Mg ha ⁻¹ (≥ 2 mm)	Shoot Mg ha ⁻¹	R/S (≥ 2 mm)	Reference
<i>Cerrado Sensu Stricto</i>	Sa	6.2	21.40	34.58	0.62	B
<i>Campo limpo</i> (grassland)	Sg (Gr)	2.0	11.57	2.90	3.99	C
<i>Campo sujo</i> (shrublands)	Sg (Sh)	2.0	21.37	3.90	5.48	
<i>Cerrado Sensu Stricto</i> (open cerrado)	Sa	2.0	33.02	17.60	1.88	
<i>Cerrado Sensu Stricto</i> (dense cerrado)	Sd	2.0	37.56	18.40	2.04	
<i>Cerrado Sensu Stricto</i> (biennial precocious)	Sa	0.5-1.0	38.15	21.00	1.82	D
<i>Cerrado Sensu Stricto</i> (biennial modal)	Sa	0.5-1.0	43.61	29.00	1.50	
<i>Cerrado Sensu Stricto</i> (biennial late)	Sa	0.5-1.0	39.18	22.90	1.71	
<i>Cerrado Sensu Stricto</i> (quadrennial)	Sa	0.5-1.0	39.86	26.30	1.52	
Dry grassland (Argisol)	Sg (Gr)	1.0	1.14	5.65	0.20	E
Dry grassland (Latosol)	Sg (Gr)	1.0	0.47	6.50	0.07	
Mosaic grasslands/shrublands (Latosol)	Sg / Sp	1.0	1.65	8.10	0.20	
<i>Campo úmido</i> (Hydromorphic)	Sg (Hy)	1.0	0.00	7.65	0.00	

5

6 (A) Brazilian vegetation classification code determined by IBGE (1992).

7 (B) Abdala et al. (1998); includes live and dead roots.

8 (C) Castro and Kauffman (1998); does not include tap roots. These authors
 9 considered fine roots to be < 6 mm diameter. The biomass of roots < 2 mm was
 10 estimated as 29% of the total of roots in a 2 m profile according to Abdala et al. (1998).

11 (D) Castro-Neves (2007); uses Abdala et al. (1998) for calculation of coarse roots
 12 (0-100 cm) and a direct method for estimating fine roots up to 50 cm depth.

13 (E) This study.

Fig. 1

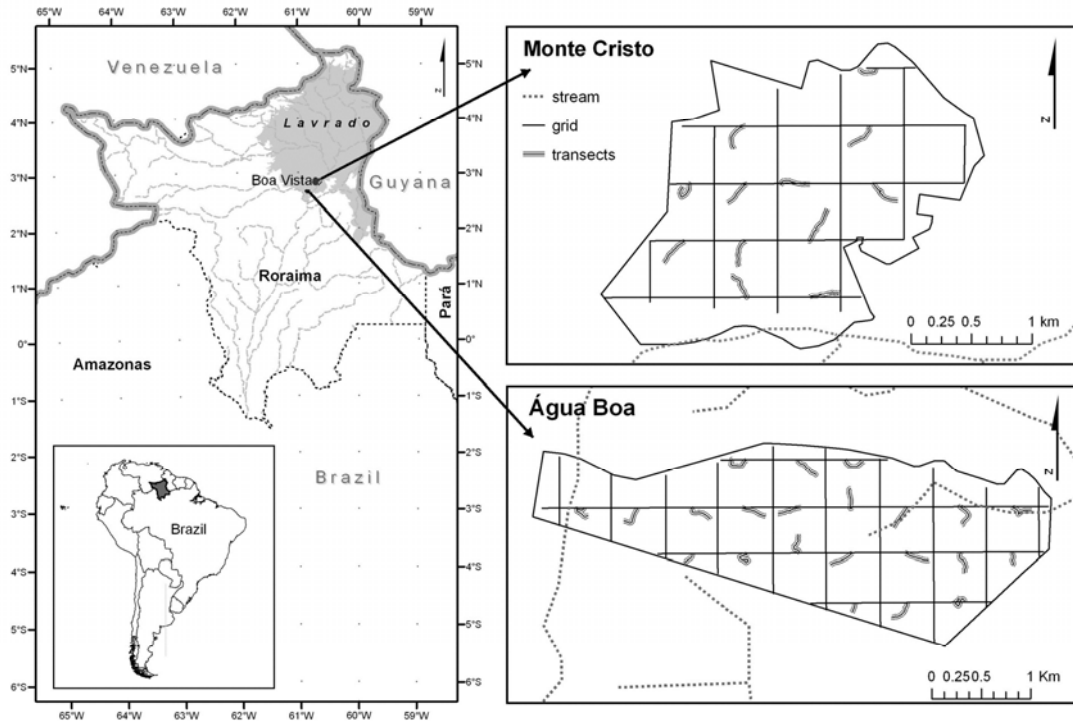


Fig. 2

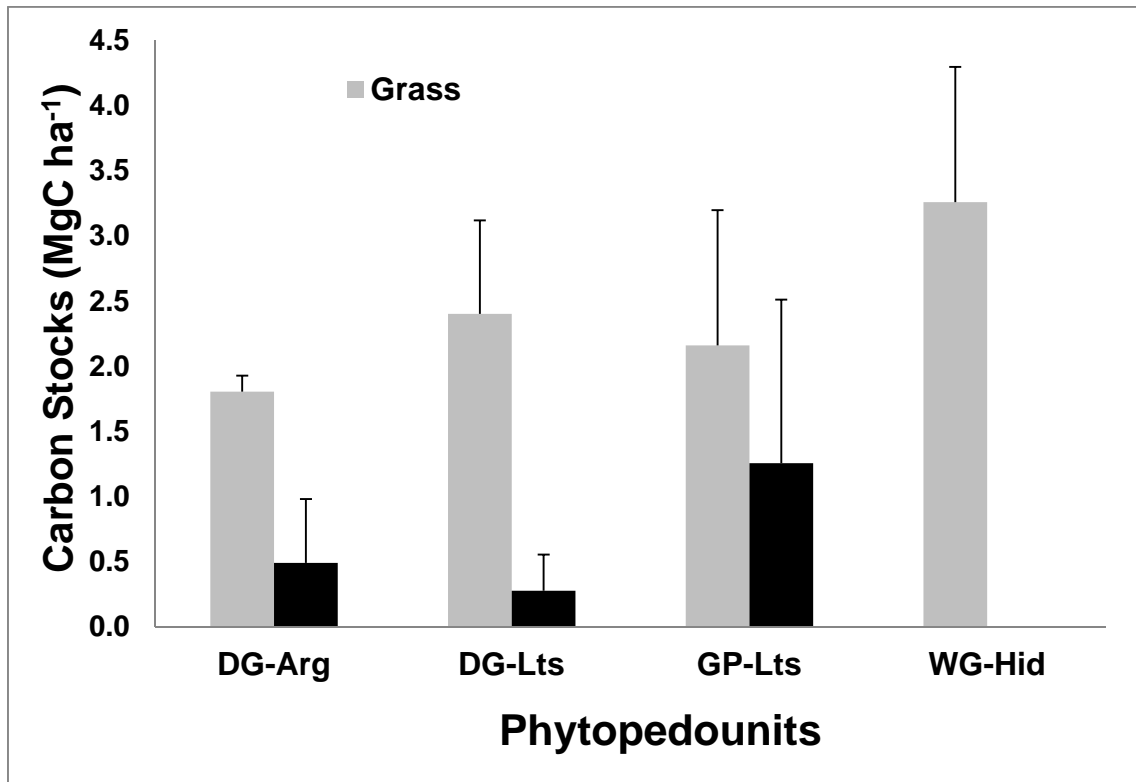
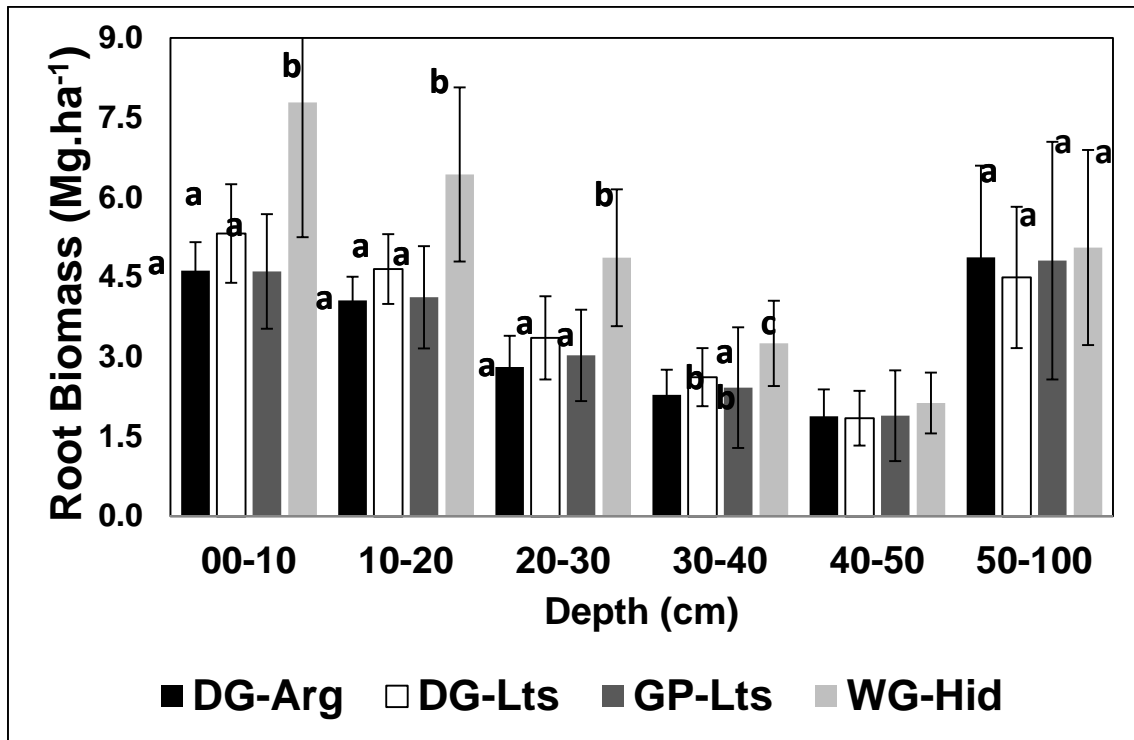


Fig. 3



1 **Supplementary Material A. Carbon concentration (%C) of the main tree and shrub**
2 **species by biomass category (leaves and diameter classes of wood pieces 'd') observed in**
3 **open savannahs in Roraima.**

Species	Category	Fraction	%C	Weighted %C
<i>Bowdichia virgilioides</i> Kunth (Fabaceae)	Leaves	0.1458	50.98	47.93
	d \geq 10cm	0.1406	47.33	
	5 \leq d<10cm	0.5826	47.62	
	d<5cm	0.1309	46.53	
<i>Byrsonima crassifolia</i> (L.) Kunth & <i>B. coccolobifolia</i> Kunth (Malpighiaceae)	Leaves	0.1998	51.66	47.86
	d \geq 10cm	0.0597	46.29	
	5 \leq d<10cm	0.5845	47.18	
	d<5cm	0.1560	46.15	
<i>B. verbascifolia</i> (L.) Rich. ex Juss. (Malpighiaceae)	Leaves	0.3723	50.01	48.52
	d \geq 10cm	-	-	
	5 \leq d<10cm	0.6058	47.72	
	d<5cm	0.0218	45.53	
<i>Curatella americana</i> L. (Dilleniaceae)	Leaves	0.1760	44.14	44.85
	d \geq 10cm	0.1235	44.99	
	5 \leq d<10cm	0.4718	45.24	
	d<5cm	0.2287	44.52	
<i>Hymatanthus articulatus</i> (Vahl) Woodson (Apocynaceae)	Leaves	0.2663	51.34	47.62
	d \geq 10cm	0.2191	45.33	
	5 \leq d<10cm	0.3883	47.01	
	d<5cm	0.1264	45.67	
Others (16 species)	Leaves	0.1549	50.18	46.28
	d \geq 10cm	0.1355	45.81	
	5 \leq d<10cm	0.5753	45.63	
	d<5cm	0.1343	45.07	

1 **Supplementary Material B. Density (number ha⁻¹) and basal area (cm² ha⁻¹) of the tree-bush component present in different**
2 **phytopedounits in two open savannah areas in Roraima (AB = Água Boa; MC = Cauamé/Monte Cristo).**

3
4

Phytopedounit	Plot (n)		Family	Species	Density (number ha ⁻¹)			Basal Area (cm ² ha ⁻¹)		
	AB	MC			D ₃₀ ≥5cm	2cm ≤D ₃₀ <5cm	Total	D ₃₀ ≥5cm	2cm ≤D ₃₀ <5cm	Total
DG-Arg	0	4	Apocynaceae	<i>Himatanthus articulatus</i>	1.04	15.63	16.7	43.85	437.47	481.3
			Dilleniaceae	<i>Curatella americana</i>	11.29	25.83	37.1	2955.44	441.02	3396.5
			Fabaceae	<i>Bowdichia virgilioides</i>	4.17	10.42	14.6	241.24	176.15	417.4
			Malpighiaceae	<i>Byrsonima coccolobifolia</i>	17.08	15.63	32.7	1369.77	278.21	1648.0
				<i>Byrsonima crassifolia</i>	27.63	31.25	58.9	2396.61	570.76	2967.4
			Proteaceae	<i>Roupala montana</i>	2.08	0.00	2.1	542.72	0.00	542.7
Total DG-Arg					63.3	98.8	162.0	7549.6	1903.6	9453.2
DG-Lts	5	3	Dilleniaceae	<i>Curatella americana</i>	5.10	2.50	7.6	1576.78	81.98	1658.8
			Malpighiaceae	<i>Byrsonima coccolobifolia</i>	8.76	5.00	13.8	564.44	85.14	649.6
				<i>Byrsonima crassifolia</i>	15.07	10.48	25.5	1420.49	154.15	1574.6
			Proteaceae	<i>Roupala montana</i>	0.50	0.00	0.5	30.09	0.00	30.1
Total DG-Lts					29.4	18.0	47.4	3591.8	321.3	3913.1
GP-Lts	2	3	Anonnaceae	<i>Xylopia aromatica</i>	0.80	0.00	0.8	69.33	0.00	69.3
			Apocynaceae	<i>Himatanthus articulatus</i>	3.20	4.00	7.2	605.50	42.10	647.6
			Dilleniaceae	<i>Curatella americana</i>	28.80	12.00	40.8	8119.37	324.90	8444.3
			Fabaceae	<i>Bowdichia virgilioides</i>	0.80	0.00	0.8	19.50	0.00	19.5
			Loganiaceae	<i>Antonia ovata</i>	0.00	8.00	8.0	0.00	125.30	125.3
			Malpighiaceae	<i>Byrsonima coccolobifolia</i>	4.80	0.00	4.8	380.80	0.00	380.8
				<i>Byrsonima crassifolia</i>	36.00	36.00	72.0	2729.77	792.71	3522.5
Proteaceae	<i>Roupala montana</i>	16.00	4.00	20.0	1972.58	64.18	2036.8			
Total GP-Lts					90.4	64.0	154.4	13896.8	1349.2	15246.0
WG-Hyd	10	0	-	-	0.0	0.0	0.0	0.0	0.0	0.0

1 **Supplementary Material C. Distribution of root biomass (mean \pm SD) by diameter category and method in the different vertical sections**

2 **of the soil (0-100 cm).** Different lower-case letters in each column indicate a distinct difference among values (SNK test; $p < 0.05$).

3

Phytopedounit	Depth (cm)	Fine roots (< 2mm)	Medium roots (2-10mm)	Subtotal (Direct Method)		Subtotal (Indirect Method ≥ 10 mm)	Total
		Mg.ha ⁻¹	Mg.ha ⁻¹	Mg.ha ⁻¹	%	Mg.ha ⁻¹	Mg.ha ⁻¹
DG-Arg	00-10	4.48 \pm 0.53	0.14 \pm 0.27	4.62 \pm 0.54	22.51		
	10-20	3.96 \pm 0.36	0.10 \pm 0.23	4.06 \pm 0.45	19.79		
	20-30	2.79 \pm 0.57	0.02 \pm 0.05	2.81 \pm 0.59	13.69		
	30-40	2.28 \pm 0.47	0.00	2.28 \pm 0.47	11.12	0.87 \pm 0.72 bc	21.40 \pm 2.47 a
	40-50	1.88 \pm 0.51	0.00	1.88 \pm 0.51	9.17		
	50-100	4.87 \pm 1.73	0.00	4.87 \pm 1.73	23.73		
Total DG-Arg		20.27 \pm 1.39 a	0.26 \pm 0.15 b	20.53 \pm 1.79 a	100.0		
DG-Lts	00-10	5.27 \pm 0.93	0.05 \pm 0.11	5.32 \pm 0.93	23.87		
	10-20	4.57 \pm 0.72	0.08 \pm 0.20	4.65 \pm 0.66	20.87		
	20-30	3.35 \pm 0.79	0.01 \pm 0.03	3.36 \pm 0.78	15.06		
	30-40	2.62 \pm 0.55	0.00	2.62 \pm 0.55	11.74	0.33 \pm 0.33 b	22.62 \pm 2.21 a
	40-50	1.85 \pm 0.51	0.00	1.85 \pm 0.51	8.28		
	50-100	4.49 \pm 1.33	0.00	4.49 \pm 1.33	20.17		
Total DG-Lts		22.14 \pm 1.47 a	0.14 \pm 0.10 b	22.28 \pm 2.41 a	100.0		
GP-Lts	00-10	4.50 \pm 1.03	0.11 \pm 0.19	4.61 \pm 1.08	22.07		
	10-20	4.00 \pm 0.96	0.12 \pm 0.20	4.12 \pm 0.96	19.74		
	20-30	2.96 \pm 0.84	0.07 \pm 0.12	3.03 \pm 0.86	14.50		
	30-40	2.35 \pm 1.02	0.07 \pm 0.18	2.42 \pm 1.14	11.59	1.26 \pm 0.22 c	22.14 \pm 4.9 a
	40-50	1.87 \pm 0.78	0.02 \pm 0.10	1.89 \pm 0.85	9.07		
	50-100	4.81 \pm 2.24	0.00	4.81 \pm 2.24	23.04		
Total GP-Lts		20.49 \pm 1.69 a	0.39 \pm 0.15 b	20.88 \pm 4.82 a	100.0		
WG-Hyd	00-10	7.79 \pm 2.53	0.00	7.79 \pm 2.53	26.37		
	10-20	6.43 \pm 1.64	0.00	6.43 \pm 1.64	21.78	0.00 a	29.52 \pm 7.15 b
	20-30	4.86 \pm 1.29	0.00	4.86 \pm 1.29	16.48		

30-40	3.25±0.80	0.00	3.25±0.80	11.02
40-50	2.13±0.57	0.00	2.13±0.57	7.22
50-100	5.06±1.84	0.00	5.06±1.84	17.13
Total WG-Hyd	29.52±2.40 b	0.00 a	29.52±2.40 b	100

4

5

Summary text

Root biomass, root : shoot ratio and below-ground carbon stocks in the open savannahs of northern Amazonia were estimated. The results indicate that the expansion factor for below-ground biomass in these ecosystems are low and differ from the default values used in Brazil's reference report to the Climate Convention.