

Diversity, floristic composition, and structure of the woody vegetation of the Cerrado in the Cerrado–Amazon transition zone in Mato Grosso, Brazil

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Abstract We compared the diversity and species composition and the structure of the vegetation of three distinct Cerrado phytophysiognomies (Cerradão, Dense Cerrado, and Typical Cerrado) in the Cerrado-Amazon transition, Mato Grosso (Brazil). Species richness (observed and estimated) in the Cerradão and Dense Cerrado was higher than that recorded in the Typical Cerrado. Species diversity, based on a Rényi profile, was highest in the Dense Cerrado, in comparison with the other phytophysiognomies. We recorded a higher number of exclusive species in the Cerradão and a greater similarity (Morisita and Sørensen indices) between this vegetation type and the Dense Cerrado. While individuals were tallest in the Cerradão and Dense Cerrado and lowest in the Typical Cerrado, there was no difference among phytophysiognomies in mean diameter. A gradient in decreasing species richness and diversity (hypothesis 1) and vegetation vertical structure (hypothesis 3) was expected for the Cerradão-Dense Cerrado-Typical Cerrado; however, neither hypothesis was supported by the results. The Cerradão and Dense Cerrado were most similar in species composition not confirming

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hypothesis 2, which predicted that the two savanna vegetation types (Dense Cerrado and Typical Cerrado) were more similar to one another than either is to the woodland (Cerradão). Overall, the similarities among the three study communities depended on the type of parameter analyzed. While the species richness and the vertical and structure of the vegetation of the Cerradão and Dense Cerrado are closely similar, the Cerradão and Typical Cerrado are more similar in their species diversity. With regard to the floristic composition, Dense Cerrado occupies an intermediate position between Cerradão and Typical Cerrado.

Keywords Conservation · Floristic similarity · Indicator species · Mean height

Introduction

The South American Cerrado biome is the most diverse and species rich of all the world's savannas (Mendonça et al. 2008), due to the marked heterogeneity of habitats and soils (Ribeiro and Walter 2008; Reatto et al. 2008), the occurrence of bushfires (Hoffmann and Franco 2003; Hoffmann et al. 2012), and landscape relief (Oliveira Filho et al. 1989; Marimon et al. 1998). In addition to these complex environmental factors, the contact zones among Cerrado and four other Brazilian biomes (Amazon, Atlantic Rain forest, Pantanal, and Caatinga) (Ribeiro and Walter 2008) can contribute to the tree species diversity.

In the Cerrado–Amazon contact zones of ecological tension, the interchange of species contributes to increased richness and influences the species composition (Castro and Martins 1999; Méio et al. 2003). The most extensive zone of contact is located in southern and southeastern Amazon (Marimon et al. 2006; Miranda et al. 2006;

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Ribeiro and Walter 2008). Here, the vegetation of both biomes intermingle, forming a complex matrix (Askew et al. 1970; Marimon et al. 2006) with a singular distribution of tree species along the hyperdynamic vegetation in the transitional Cerrado–Amazon landscapes (Marimon et al. 2014). As this region also coincides with the "Arc of Deforestation" in the Brazilian agricultural frontier, it is of considerable importance for the conservation of the country's biodiversity (Nogueira et al. 2008; Domingues and Bermann 2012).

Scientific studies of the Cerrado-Amazon transition zone in eastern Mato Grosso began with the classic, descriptive works of the Xavantina-Cachimbo Expedition (Askew et al. 1970; Eiten 1972; Ratter et al. 1973). The local vegetation is composed predominantly of semi-deciduous forests, known as transitional forest (Marimon et al. 2006; Balch et al. 2008; Hoffmann et al. 2012), the composition of which is influenced primarily by the Amazon rainforest, and only to a lesser extent by the Cerrado (Méio et al. 2003). A number of other vegetation types and subtypes (sensu Ribeiro and Walter 2008) can be found in the region, with a predominance of Cerrado species (Askew et al. 1970; Marimon Junior and Haridasan 2005; Maracahipes et al. 2011). More recent studies suggest that the savanna formations of the transition zone are characterized by higher diversity than similar areas on the central Brazilian plateau (Felfili et al. 2002; Kunz et al. 2009; Gomes et al. 2011; Lenza et al. 2011; Maracahipes et al. 2011).

The Cerradão is the only woodland habitat sensu lato found in the Cerrado (sensu Ribeiro and Walter 2008) and is characterized both by generalist tree species, as well as those more typical of either the savanna or forest formations of the Cerrado (Oliveira Filho and Ratter 1995; Marimon Junior and Haridasan 2005; Ribeiro and Walter 2008; Solórzano et al. 2012). The savanna formations of the Cerrado include Dense Cerrado and Typical Cerrado, which can be differentiated in terms of the density of the vegetation, and canopy cover and height, but tend to be very similar in terms of species composition (Ribeiro and Walter 2008). Tree cover in the Dense Cerrado is between 50 and 70 %, with a canopy height of 5-8 m (Ribeiro and Walter 2008). The Typical Cerrado is the most widespread vegetation subtype in the Cerrado sensu stricto (Ribeiro and Walter 2008) and is characterized by tree cover of 20-50 %, with a canopy of 3-6 m in height (Ribeiro and Walter 2008).

Local studies comparing Cerrado vegetation types and subtypes have provided important insights into the formation of distinct habitats and the conservation and management of the biome's biodiversity (Oliveira Filho et al. 1989; Marimon et al. 1998; Marimon Junior and Haridasan 2005; Gomes et al. 2011). In their comparison of Cerradão and Typical Cerrado, Marimon Junior and Haridasan (2005) found marked differences in species composition and vegetation structure. In Typical and Rocky Cerrado, Gomes et al. (2011) found marked differences in the most important species in terms of community structure. In their study of Cerradão, Dense Cerrado, and Typical Cerrado in São Paulo (Brazil), Pinheiro and Durigan (2012) recorded structural differences in basal area classes, canopy cover, and the diameter and height of the largest trees. These findings emphasize the importance of understanding both the species composition and structure of the vegetation for the interpretation of the features of the different phytophysiognomies.

In the present study, our main objective was to characterize and compare the diversity, species composition, and structure of the woody vegetation of three Cerrado sites, representing the Cerradão (forest facies), Dense Cerrado, and Typical Cerrado, in the Cerrado-Amazon transition zone in Mato Grosso (Brazil). We tested three hypotheses related to the distributions of tree species in the transitional Cerrado-Amazon transitional landscapes: (1) because of the co-existence of savanna and forest species in the Cerradão, there is a decreasing gradient in tree species richness and diversity from this vegetation to Dense and Typical Cerrado; (2) the species composition and horizontal structure of the vegetation is more similar between Typical and Dense Cerrado than between either of these and the Cerradão woodland; (3) there is a decreasing gradient in height, diameter, density, and total basal area of the trees from the Cerradão and the Typical Cerrado.

Materials and methods

Study area

The study focused on three Cerrado sites with distinct vegetation types, which represent a gradient in density, height, and the shrubby component of the canopy in the Cerrado-Amazon transition areas. One site was representative of the Cerradão (CO: 12°49'26.8"S, 051°46'06.0"W), one of the Dense Cerrado (DC-12°49'07.6"S, 051°46'12.3"W), and one of the Typical Cerrado (TC: 12°50'02.5"S, 051°45′55.9″W) sensu Ribeiro and Walter (2008), located in the Cerrado-Amazon transition zone in the municipality of Ribeirão Cascalheira, in the State of Mato Grosso, Brazil (Fig. 1). The local climate is characterized by a well-defined rainy season, from May to September, with rainfall peaking from November to February, and dry conditions, including soil water deficits during the rest of the year (Silva et al. 2008).

Fig. 1 Location of the study area (*filled square*) in the municipality of Ribeirão Cascalheira, Mato Grosso (Brazil), where we sampled three different types of woody Cerrado vegetation in the Cerrado–Amazon transition zone



Data collection

We established a 100 m \times 100 m plot at each of the three sites divided into 25 subplots of 20 m \times 20 m. All live woody plants with diameter at 30 cm above the ground (D₃₀) of at least 5 cm were identified and marked with aluminum tags. We measured the total height of each individual with a laser rangefinder and stem diameter with a measuring tape. For individuals with multiple stems, all stems were measured for the subsequent calculation of the quadratic diameter (Moro and Martins 2011). We conducted the data collection in three field campaigns with 5 days each, the first one in July 2013 (Dense Cerrado) and the other two in September 2013 (Cerradão and Typical Cerrado).

Whenever possible, we identified the species in the field, but when necessary, we collected specimens for comparison with those available in the Herbarium NX, of the collection James Alexander Ratter (State University of Mato Grosso at Nova Xavantina) and in appropriate references. The system suggested by the Angiosperm Phylogeny Group (APG III 2009) was used to classify the families, with the taxonomy revised and updated based on the List of Brazilian Flora (2014).

Data analysis

We compared the species richness recorded at the three sites using the rarefaction technique with the standardization of sampling effort by the number of individuals recorded (Gotelli and Colwell 2001; Magurran 2011), using the S_{obs} *Mao Tau* estimator with 1000 randomizations. The species richness at the three sites was also estimated using Jackknife 1, which was the estimator that

correlated most closely with observed richness (Walther and Moore 2005). We used diversity profiles based on the Rényi exponential series (Tóthmérész 1995) to compare diversity at the three study sites, in which alpha corresponds to Shannon's index and alpha 2 to Simpson's index (Melo 2008).

We evaluated the similarity in species composition among the three sites using the quantitative Morisita-Horn (Cm_H) coefficient (Magurran 2011) and the binary (qualitative) coefficient of Sørensen (Ss) (Brower and Zar 1984). A two-way indicator species analysis (ISA) (TWINSPAN: Hill et al. 1975) was used to classify the vegetation, with eigenvalues above 0.3 being considered to represent strong ecological divisions (Hill 1979; Felfili et al. 2007, 2011).

We used an ISA to identify the species considered to be indicators of a given vegetation type, together with a Monte Carlo test to verify the significance of the classification (Dufrêne and Legendre 1997; McCune and Mefford 1999). We described the horizontal structure of the vegetation using the absolute density of individuals and the Importance Value Index (IVI), as proposed by Curtis and McIntosh (1950) and Mueller-Dombois and Ellenberg (1974).

We used an analysis of variance (ANOVA) to compare trunk diameter, tree height, and mean density per subplot among sites. When the assumptions of normality and homogeneity of variance were not satisfied, we used an ANOVA with Welch's F test for heterogeneous variables (Zar 2010). All analyses were complemented with Tukey's post hoc test.

Structural parameters were analyzed in FITOPAC 2.1.2 (Shepherd 2009), diversity and similarity were evaluated in PC-ORD version 6.07 (McCune and Mefford 2011), TWINSPAN and ISA were run in PAST (*PAleontological*

STatistics) version 2.15 (Hammer et al. 2001), while R 3.0.2 (R Core Team 2013) was used for all other analyses. In all cases, the normality of the data was assessed using the Shapiro-Wilk test, and the homogeneity of variance was confirmed by Levene's test (Zar 2010), and a 5 % significance level was considered.

Results

We sampled a total of 5946 individuals at the three sites, belonging to 115 species, 80 genera, and 40 families. We recorded 2186 individuals, 91 species, 67 genera, and 35 families in the Cerradão (CO) and 2070 individuals of 91 species, 68 genera, and 37 families in the Dense Cerrado (DC). In the Typical Cerrado (TC), we sampled 1690 individuals belonging to 76 species, 56 genera, and 32 families (Table 1).

The species richness estimated by rarefaction (Fig. 2) was lowest in the Typical Cerrado (76 species), but remarkably similar between the Cerradão (86.7 ± 8.9 species) and the Dense Cerrado (86.8 ± 8.7 species). Observed species richness (76) in the Typical Cerrado was the most similar (89 %) to the estimated number (85.0 ± 12.9). In the case of the Cerradão, observed richness was 91 species, 84 % of the estimated value (108.0 ± 11.4 species), while in the Dense Cerrado, observed richness was also 91 species, 81 % of the estimated number (112.0 ± 12.7 species).

The Dense Cerrado had the highest diversity in comparison with the Cerradão and Typical Cerrado (Fig. 3). The other two vegetation types were closely similar to one another.

Of the total of 115 species recorded in the present study, 54 (47 %) were common to all three sites, and 26 (23 %) were exclusive to two of the sites. The largest number of exclusive species was recorded in the CO plot, followed by TC and DC (Fig. 4; Table 1). The highest similarity values were found between the Cerradão and Dense Cerrado, and the lowest similarity values were found between Cerradão and Typical Cerrado, considering both the quantitative Morisita index and the qualitative Sørensen index (Fig. 4). The TWINSPAN classification initially grouped all the 25 Cerradão subplots with four of the Dense Cerrado plots (eigenvalue = 0.33), while all the remaining subplots were grouped together (Fig. 5). A total of 24 indicator species were recorded in the Cerradão, 16 in the Dense Cerrado, and 20 in the Typical Cerrado. Despite these large numbers of indicator species at each site (Monte Carlo, p < 0.001), the large numbers of species common to two or more sites contributed to high indices of similarity (Table 1; Fig. 4).

The most structurally important species (highest IVI values: see Table 1) in the Cerradão were *Myrcia*

splendens (Sw.) DC., Pterodon pubescens (Benth.) Benth., Xylopia sericea A.St.-Hil., Roupala montana Aubl., Syagrus comosa (Mart.) Mart., Emmotum nitens (Benth.) Miers, Mezilaurus crassiramea (Meisn.) Taub. ex Mez, Pouteria ramiflora (Mart.) Radlk., Qualea grandiflora Mart., and Tachigali vulgaris L.G.Silva & H.C.Lima (53.6 % of the total). In the Dense Cerrado, the most important species were E. nitens, Mouriri elliptica Mart., S. comosa, M. splendens, R. montana, P. ramiflora, X. sericea, Myrcia multiflora (Lam.) DC., Bowdichia virgilioides Kunth, and M. crassiramea (50.9 % of the total), while in the Typical Cerrado, they were S. comosa, Davilla elliptica A.St.-Hil., Qualea parviflora Mart., M. crassiramea, Byrsonima verbascifolia A.St.-Hil., Myrcia rimosa Cambess., Eugenia dysenterica DC., Byrsonima coccolobifolia Kunth, and Curatella americana L. (49.4 % of the total). Among the ten species with the highest IVI in each phytophysiognomy, seven were common among the Cerradão and the Dense Cerrado, and only two were common between the three phytophysiognomies (Table 1). These results suggest a higher similarity, with regard to horizontal structure, between the Cerradão and Dense Cerrado and lesser resemblance between these two phytophysiognomies and the Typical Cerrado.

Mean tree height was significantly higher (Welch's $F_{(2,3935)} = 626.70, P < 0.01$ in the Cerradão (5.5 ± 2.4 m, SD) in comparison with the Dense (4.5 \pm 2.3 m) and Typical Cerrado $(3.0 \pm 1.98 \text{ m})$. However, mean diameter varied little ($F_{(2,3884)} = 0.66$, P = 0.52) among the Cerradão $(9.5 \pm 4.7 \text{ cm})$, Dense Cerrado $(9.6 \pm 5.0 \text{ cm})$, or Typical Cerrado (9.5 \pm 4.5 cm). Despite these similarities in the diameter of individuals, the mean densities recorded in the subplots at the three sites varied significantly ($F_{(2,72)} = 28.09$; P < 0.01), with the lowest density being recorded in the Typical Cerrado (67.6 \pm 9.4 inds ha⁻¹), with much higher values being recorded in the Cerradão (87 ± 8.85 inds ha⁻¹) and Dense Cerrado (83 \pm 11.0 inds ha⁻¹). The lower density recorded in the Typical Cerrado also accounts for the reduced basal area recorded in this vegetation type (14.5 $\text{m}^2 \text{ha}^{-1}$) in comparison with the Cerradão $(19.3 \text{ m}^2 \text{ ha}^{-1})$ and Dense Cerrado (19.1 $\text{m}^2 \text{ha}^{-1}$).

Discussion

The high species richness recorded in the present study (91 in Cerradão and Dense Cerrado; 76 species in Typical Cerrado) has also been found in other cerrado communities in eastern Mato Grosso. For example, in Cerradão, Kunz et al. (2009) registered 95 species, Franczak et al. (2011) found between 84 and 87 species, and Solórzano et al. (2012) sampled 124 species. In Cerrado *sensu stricto*, Nogueira et al. (2001) observed 88 species, Felfili et al.

Table 1	Density and	importance	of the v	voody pl	ant species	; (D ₃₀ ≥	2 5 cm)	found	at three	sites	in the	Cerrado-Amazon	transition	zone	in
Ribeirão	Cascalheira,	Mato Grosse	o (Brazil)											

Species	Family		Absolute density			Importance Value Index		
		CO	DC	TC	СО	DC	TC	
1. Myrcia splendens (Sw.) DC*	Myrtaceae	383	162	9	32.61	16.22	2.41	
2. Pterodon pubescens (Benth.) Benth*	Fabaceae	236	9	_	25.25	1.59	_	
3. Xylopia sericea A.StHil*	Annonaceae	136	100	22	21.18	11.83	4.14	
4. Roupala montana Aubl*	Proteaceae	170	114	32	16.64	13.59	6.67	
5. Syagrus comosa (Mart.) Mart***	Arecaceae	177	187	315	16.25	18.07	42.40	
6. Emmotum nitens (Benth.) Miers**	Icacinaceae	84	206	31	11.89	34.43	6.80	
7. Mezilaurus crassiramea (Meisn.) Taub. ex Mez	Lauraceae	46	34	45	11.21	7.06	10.90	
8. Pouteria ramiflora (Mart.) Radlk**	Sapotaceae	45	68	17	8.93	13.36	5.02	
9. Qualea grandiflora Mart*	Vochysiaceae	57	25	2	8.90	3.73	0.60	
10. Tachigali vulgaris L.G.Silva & H.C.Lima*	Fabaceae	63	23	18	8.04	3.39	5.31	
11. Antonia ovata Pohl*	Loganiaceae	70	3	1	7.45	0.60	0.26	
12. Aspidosperma multiflorum A.DC*	Apocynaceae	48	3	_	6.55	0.72	_	
13. Vatairea macrocarpa (Benth.) Ducke*	Fabaceae	37	4	4	5.55	0.92	1.30	
14. Euplassa inaequalis (Pohl) Engl*	Proteaceae	24	13	_	5.34	3.46	_	
15. Bowdichia virgilioides Kunth***	Fabaceae	16	37	41	5.15	7.97	10.50	
16. Mouriri elliptica Mart**	Melastomataceae	30	157	18	5.01	22.06	4.28	
17. Miconia albicans (Sw.) Triana*	Melastomataceae	44	10	6	4.99	1.65	1.23	
18. Eriotheca gracilines (K.Schum.) A.Robyns*	Malvaceae	24	4	8	4.54	0.53	4.72	
19. Salvertia convallariodora A.StHil.	Vochysiaceae	18	14	15	4.36	2.45	5.52	
20. Oualea parviflora Mart***	Vochvsiaceae	16	58	90	3.85	6.72	14.80	
21. Aspidosperma tomentosum Mart**	Apocynaceae	25	35	4	3.71	6.21	1.01	
22. Davilla elliptica A.StHil***	Dilleniaceae	25	32	213	3.61	4.08	22.70	
23. Oualea multiflora Mart*	Vochysiaceae	26	14	_	3.58	2.91	_	
24 Eugenia dysenterica DC	Myrtaceae	23	28	42	3.51	4.39	9.08	
25 Machaerium acutifolium Vogel*	Fabaceae	22	_	1	3.43	_	0.26	
26 Heisteria ovata Benth*	Olacaceae	19	7	1	3.13	1 40	0.25	
27. Schoenfig brasiliensis A DC*	Schoenfiaceae	26	2	_	3.04	0.54	_	
28. Hymenaea stigonocarna Mart ex Havne**	Fabaceae	12	- 43	8	2.95	6.19	1 84	
29. Xylonia aromatica (Lam.) Mart	Annonaceae	18	50	35	2.93	5 70	6 14	
30 Diospyros hispida A DC	Fhenaceae	15	28	12	2.95	3.62	2.17	
31 Curatella americana L ***	Dilleniaceae	9	14	48	2.59	3.62	8 47	
32 Buchenavia tomentosa Fichler	Combretaceae	11	11	10	2.33	2 51	2.18	
33 Handroanthus ochraceus (Cham) Mattos**	Bignoniaceae	9	26	10	2.35	5 22	2.10	
34 Peltogyne confertiflora (Mart. ex. Hayne) Benth**	Fabaceae	10	14	_	2.36	2.63	_	
35. Ouratea hexasperma (A St -Hil) Baill**	Ochnaceae	12	32	4	2.26	4.09	0.91	
36 Guanira graciliflora (Mart. ex. Schmidt) Lundell*	Nyctaginaceae	9	_	_	2.13	_	_	
37 Hirtella glandulosa Spreng	Chrysobalanaceae	9	6	3	2.12	1 39	0.70	
38 Licania humilis Cham & Schltdl**	Chrysobalanaceae	10	24	6	2.05	4 51	1 73	
39 Rourea induta Planch**	Connaraceae	11	34	18	1.93	4.66	3 23	
40 Byrsoning coccolabifalia Kunth***	Malnighiaceae	9	37	56	1.95	4.00	8 49	
41 Vochysia rufa Mart	Vochysiaceae	9	8	10	1.00	1.26	2 34	
42 Aspidosperma macrocarpon Mart*	Apocynaceae	8	2	1	1.72	0.51	0.39	
43. Dalbergia miscolobium Benth	Fahaceae	5	2	_	1.32	0.57	_	
44 Miconia stenostachva DC*	Melastomataceae	7	-	1	1.20	_	0.28	
45 Vochysia haenkeana Mart*	Vochysiaceae	, 6	_	_	1.27	_	_	
46. Brosimum gaudichaudii Trécul*	Moraceae	5	_	_	1.05	_	_	
0		-						

Table 1 continued

Species	Family		Absolute density			Importance Value Index		
		СО	DC	TC	СО	DC	TC	
47. Byrsonima pachyphylla A.Juss***	Malpighiaceae	5	42	36	1.02	5.3	5.93	
48. Tabebuia aurea (Silva Manso) Benth. & Hook.f. ex S.Moore	Bignoniaceae	4	10	9	1.02	2.75	1.98	
49. Byrsonima basiloba A.Juss.	Malpighiaceae	4	8	4	1.02	1.53	1.02	
50. Connarus suberosus Planch***	Connaraceae	5	3	57	1.01	0.59	8.24	
51. Guapira noxia (Netto) Lundell*	Nyctaginaceae	4	_	_	0.96	_	_	
52. Leptolobium dasycarpum Vogel	Fabaceae	4	1	_	0.90	0.20	_	
53. Chaetocarpus echinocarpus (Baill.) Ducke	Peraceae	4	4	_	0.89	0.55	_	
54. Aspidosperma nobile Müll.Arg.	Apocynaceae	4	2	_	0.88	0.28	_	
55. Annona coriacea Mart*	Annonaceae	5	_	_	0.88	_	_	
56. Luetzelburgia praecox (Harms) Harms*	Fabaceae	4	_	_	0.84	_	_	
57. Virola sebifera Aubl**	Myristicaceae	4	16	_	0.83	2.10	_	
58. Hancornia speciosa Gomes	Apocynaceae	4	3	_	0.82	0.61	_	
59. <i>Myrcia</i> sp.	Myrtaceae	4	11	11	0.81	1.85	2.43	
60. Simarouba versicolor A.StHil.	Simaroubaceae	3	1	_	0.72	0.33	_	
61. Himatanthus sucuuba (Spruce ex Müll.Arg.) Woodson	Apocynaceae	3	1	_	0.72	0.20	_	
62. Cybianthus detergens Mart.	Primulaceae	3	3	1	0.65	0.59	0.36	
63. Maprounea guianensis Aubl.	Euphorbiaceae	3	3	1	0.65	0.60	0.29	
64. Salacia crassifolia (Mart. ex Schult.) G.Don***	Celastraceae	3	10	34	0.63	1.75	6.59	
65. Myrcia bella Cambess	Mvrtaceae	3	1	4	0.62	0.21	0.89	
66 Myrcia rimosa Cambess***	Myrtaceae	3	42	85	0.60	4.75	10.30	
67. Heteropterys byrsonimifolia A Juss	Malpighiaceae	3	_	3	0.60	_	0.74	
68. Svagrus flexuosa (Mart.) Becc**	Arecaceae	3	11	1	0.48	1.95	0.23	
69 Dimorphandra mollis Benth	Fabaceae	3	1	8	0.48	0.23	2.12	
70 Couepia grandiflora (Mart & Zucc) Benth	Chrysobalanaceae	2	_	_	0.44	_	_	
71 Lafoensia pacari A St -Hil**	Lythraceae	2	40	2	0.43	5 35	0.37	
72 Fugenia gemmiflora O Berg***	Myrtaceae	2	5	39	0.41	1.01	7 87	
73. Aspidosperma subincanum Mart	Apocynaceae	2	_	_	0.36		-	
74 Ferdinandusa elliptica (Pohl) Pohl	Rubiaceae	2	1	_	0.34	0.21	_	
75. Miconia sp. 2	Melastomataceae	1	1	_	0.34	0.21	_	
76 Byrsonima varbascifolia (L.) DC***	Malpighiaceae	1	16	64	0.28	3 73	10.6	
77. Miconia sp. 1	Malastomataceae	1	1	04	0.20	0.20	10.0	
77. Miconiu sp. 1 78. Cybistar antisyphilitica (Mort.) Mort	Pignoniaceae	1	1	_	0.27	0.20	-	
70. Enterolohium gummifarum (Mart.) LE Machr	Eabacasa	1	1	_	0.20	-	-	
20 Diplontarys publicatala (A Juss) WP Anderson & C C Davis	Malpighiagaaa	1	1	_	0.24	0.21	-	
81 Annong grassiflorg Mort**	Annonaccae	1	-	-	0.22	-	-	
22 Eugenia gunata O Dana***	Murrtagaga	1	20	10	0.21	1.22	4.50	
82. Alahamag disaalar Daann	Furthershieses	1	0	10	0.21	0.21	5.59	
85. Alchornea alscolor Poepp.	Dubiosos	1	1	-	0.21	0.21	-	
84. <i>Tocoyena Jormosa</i> (Cham. & Schudi.) K.Schum.	Fabaaaaa	1	1	3	0.21	0.20	0.69	
85. Copaijera ci. marin Hayne	Pabaceae	1	-	-	0.21	-	-	
80. Friaericia cinnamomea (DC.) L.G.Lonmann	Bignoniaceae	1	-	-	0.21	-	-	
87. Protium heptaphyllum (Aubl.) Marchand	Burseraceae	1	1	-	0.21	0.20	-	
88. Myrcia multifiora (Lam.) DC**	Myrtaceae	1	20	19	0.20	8.05	3.50	
89. <i>Pseudobombax longiflorum</i> (Mart. & Zucc.) A.Robyns	Malvaceae	1	5	-	0.20	0.53	-	
90. Byrsonima intermedia A.Juss.	Malpighiaceae	1	2	2	0.20	0.42	0.55	
91. Erythroxylum sp.	Erythroxylaceae	1	_	-	0.20	-	-	
92. Strychnos pseudoquina A.StHil.	Loganiaceae	-	5	7	-	1.19	3.39	
93. Eugenia punicifolia (Kunth) DC***	Myrtaceae	-	7	12	-	1.15	2.75	

Diversity, floristic composition, and structure of the woody vegetation of the Cerrado in the...

Table	1	continued
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Species	Family	Absolute density			Importance Value Index		
		СО	DC	TC	СО	DC	TC
94. Erythroxylum tortuosum Mart**	Erythroxylaceae	_	4	_	_	0.81	_
95. Stryphnodendron coriaceum Benth.	Fabaceae	_	3	2	_	0.61	0.47
96. Andira cujabensis Benth***	Fabaceae	_	2	6	_	0.53	1.73
97. Stryphnodendron rotundifolium Mart***	Fabaceae	_	2	16	_	0.40	2.73
98. Plathymenia reticulata Benth.	Fabaceae	_	2	4	_	0.39	0.97
99. Mouriri pusa Gardner	Melastomataceae	_	1	-	_	0.29	-
100. Tapirira guianensis Aubl	Anacardiaceae	_	1	_	_	0.28	_
101. Agonandra brasiliensis Miers ex Benth. & Hook.f	Opiliaceae	_	1	_	_	0.26	_
102. Caryocar brasiliense Cambess	Caryocaraceae	_	1	5	_	0.25	0.83
103. Cordiera elliptica (Cham.) Kuntze***	Rubiaceae	_	1	9	_	0.21	2.07
104. Erythroxylum engleri O.E.Schulz	Erythroxylaceae	_	1	5	_	0.20	1.01
105. Himatanthus obovatus (Müll.Arg.) Woodson	Apocynaceae	_	1	3	_	0.20	0.68
106. Eremanthus cf. goyazensis (Gardner) Sch.Bip.	Asteraceae	_	1	2	_	0.20	0.50
107. Zeyheria montana Mart.	Bignoniaceae	_	1	2	_	0.20	0.45
108. Copaifera langsdorffii Desf.	Fabaceae	_	1	-	_	0.20	-
109. Andira vermifuga (Mart.) Benth***	Fabaceae	_	_	15	_	_	3.88
110. Myrcia variabilis DC***	Myrtaceae	_	_	11	_	_	1.99
111. Chomelia ribesioides Benth. ex A.Gray***	Rubiaceae	_	_	4	_	_	0.98
112. Tachigali aurea Tul.	Fabaceae	_	_	2	_	_	0.47
113. Kielmeyera coriacea Mart. & Zucc	Calophyllaceae	_	-	2	_	-	0.46
114. Protium unifoliolatum Engl.	Burseraceae	_	_	2	_	_	0.34
115. Alibertia edulis (Rich.) A.Rich	Rubiaceae	_	_	1	_	-	0.23
Total		2186	2070	1690	300	300	300

CO Cerradão, DC dense Cerrado, TC typical Cerrado. The species with the highest densities and IVI values at each site are shown in bold type. The species are listed in decreasing order of the IVI values recorded at the Cerradão site

* Indicator species of the Cerradão (n = 24; 20.9 %)

** Indicator species of the Dense Cerrado (n = 16; 13.9 %)

*** Indicator species of the Typical Cerrado (n = 20; 17.4 %)

(2002) found 80 species, and Marimon Junior and Haridasan (2005) recorded 77 species, and Gomes et al. (2011) registered 79 species. Lenza et al. (2011) also found evidence of marked differences in the species composition of these transitional cerrados in comparison with those located towards the center of the biome. Felfili et al. (2002) concluded that the unique features of these Cerrados were related to the influence of Amazonian ecosystems. Together, these findings emphasize the floristic importance of the Cerrados of this region for the conservation of its woody flora.

The relatively low observed and estimated species richness recorded for the Typical Cerrado in comparison with the other two vegetation types appears to be typical of the Cerrado in general, given that Pinheiro and Durigan (2012) obtained similar results in a study of the Cerrado in São Paulo. In addition, data on the Cerradão from the same region as the present study (Kunz et al. 2009; Franczak et al. 2011; Solórzano et al. 2012) indicate similarly high levels of species richness in comparison with areas of Typical Cerrado (Felfili et al. 2002; Marimon Junior and Haridasan 2005; Gomes et al. 2011). This indicates the existence of a gradient of species richness between the more open and woodland habitats associated with density and basal area.

The relatively higher species diversity recorded in the Dense Cerrado in comparison with the other two vegetation types may reflect the more equitable distribution of individuals among species in these types and the lower species richness for Typical Cerrado. The higher diversity of the Dense Cerrado compared to the others may also be related to the intermediate nature of this vegetation type, between the Cerradão and Typical Cerrado. This may be because neither the species characteristic of forest formations (which are abundant in the Cerradão) nor those that predominate in more open savannas (abundant in the Typical



Fig. 2 Rarefaction curve (S_{obs} - *Mao Tau*) based on the number of individuals recorded in the woody Cerrado vegetation of the Cerrado-Amazon transition zone in Ribeirão Cascalheira, Mato Grosso (Brazil). *CO* Cerradão, *DC* dense Cerrado, and *TC* typical Cerrado. The 95 % confidence interval is shaded in gray



Fig. 3 Diversity profiles prepared using the Rényi series, comparing three types of woody Cerrado vegetation in the Cerrado–Amazon transition zone in Ribeirão Cascalheira, Mato Grosso (Brazil). *CO* Cerradão (*continuous line*), *DC* dense Cerrado (*dotted line*), *TC* typical Cerrado (*dashed line*) and (H' = 1 alpha corresponds to Shannon's index and D = 2 alpha 2 to Simpson's index)

Cerrado) were dominant in the Dense Cerrado. Previous comparative studies of Cerrado habitats, such as those of Pinheiro and Durigan (2012), Marimon Junior and Haridasan (2005), and Costa and Araújo (2001), have applied only a single diversity index, rather than a variety of indices that weight common and rare species differently. Based on this condition, we would recommend the application of diversity profiles for the better understanding of structural differences in comparative studies, as also suggested by Melo (2008).

These results show a decrease in plant richness from Cerradão and Dense to Typical Cerrado, but not for diversity. The fact that tree species diversity in Dense Cerrado was higher than the others demonstrates, in sum, that Dense Cerrado is an intermediate situation of species composition and possibly a temporal stage between Cerradão and Typical Cerrado. This state can reveal evidence of shifts between forest and savanna (forest advances into Cerrado) over the last millennium as previously recorded by Ab'Saber (1977, 2002) and recently investigated by the TROBIT Project (Saez, Lloyd, Marimon Junior et al., unpublished data) in the same area of this study.

The high similarity of the species found at the three sites, especially in qualitative terms, and the large proportion of species in common (approximately 50 %) could be a consequence of the geographic proximity of the sites, which are separated by distances of only 0.5–3 km, and reinforces the lack of any specific identity for these communities. A number of authors have found that the woody vegetation of adjacent cerrado to forest communities may be highly similar, irrespective of the similarities in substrate conditions (Silva et al. 2006; Gomes et al. 2011; Mews et al. 2014). In the present study, this is probably due to the physicochemical similarities of the soil among sites and the interchange of species—especially the more generalist forms—among communities.

However, a small number of habitat specialist tree species may contribute to marked divergences in the species composition of adjacent communities. This may account for the reduced similarity observed between the Cerradão woodland and the Typical Cerrado savanna samples, in particular, and the larger number of exclusive Cerradão species. The Dense Cerrado was more similar to both Cerradão and Typical Cerrado, however, and subplots that grouped with both vegetation types in the TWINSPAN analysis, reinforcing the transitional and intermediate characteristics of this vegetation type, and the existence of a floristic gradient. This may also account for the reduced similarity between the Cerradão and the Typical Cerrado, observed in both the present and previous studies (Marimon Junior and Haridasan 2005; Pinheiro and Durigan 2012).

While all three phytophysiognomies presented an oligarchic pattern, that is, the predominance of an only a few species in the structure of the community, only two species—*S. comosa* (Mart.) Mart. and *M. crassiramea* (Meisn.) Taub. ex Mez—were among the ten most predominant at all three sites. Structural oligarchy is common in the plant communities of the Cerrado *sensu lato* (Ratter et al. 2003; Bridgewater et al. 2004; Felfili et al. 2008) and indicates that a small group of species is adapted to the environment and the competition with one another in such a way as to guarantee their dominance of the communities (Solórzano et al. 2012). In the present study, however, the reduced coincidence among more dense and forested **Fig. 4** Venn diagram for the distribution of the woody Cerrado plant species found in the Cerrado–Amazon transition zone in Ribeirão Cascalheira, Mato Grosso (Brazil). *CO* Cerradão, *DC* dense Cerrado, *TC* typical Cerrado, *N* number of species, *Cm_H* Morisita's similairty index, and *Ss* Sørensen's index





Fig. 5 The TWINSPAN classification of the 75 subplots sampled in three types of woody cerrado vegetation in the Cerrado–Amazon transition zone in Ribeirão Cascalheira, Mato Grosso (Brazil). *CO* Cerradão; *DC* dense Cerrado; and *TC* typical Cerrado. Other divisions are not shown here due to their low eigenvalues (<3)

phytophysiognomies (Cerradão and Dense Cerrado) and predominantly savannic community (Typical Cerrado) in the structurally dominant species and the high number of indicator species identified for each community reflect the structural differences among vegetation types. The distinct conditions found in each type, related to differences in tree density and height, appear to favor the predominance of different species in the three communities. In particular, *M. splendens* (Sw.) DC., *X. sericea* A.St.-Hil., *R. montana* Aubl., *E. nitens* (Benth.) Miers, and *P. ramiflora* (Mart.) Radlk. are the species most well adapted to the forested habitats (Cerradão and Dense Cerrado), while *D. elliptica* A.St.-Hil., *Q. parviflora* Mart., *B. verbascifolia* A.St.-Hil., *B. virgilioides* Kunth, *B. coccolobifolia* Kunth, *E.* *dysenterica* DC., and *C. americana* L. seem most adapted to the open savanna environments (Typical Cerrado).

Surprisingly, mean tree diameter did not vary significantly among sites, although the individuals in the Cerradão and Dense Cerrado were taller than those found in the Typical Cerrado, indicating that the plants in the denser vegetation types invest more in upward growth than the enlargement of their stems. In addition, despite the similarities in stem diameter, the highest basal area recorded in the Cerradão and Dense Cerrado was determined by the highest tree densities recorded in these vegetation types.

The first hypothesis-that there is a systematic Cerradão-Dense Cerrado-Typical Cerrado gradient of (decreasing) species richness and diversity-was not supported by the findings of this study, given that these parameters were very similar in the Cerradão and Dense Cerrado, and much higher than the values recorded for the Typical Cerrado. In addition, the highest species diversity was recorded in the Dense Cerrado, rather than the Cerradão. The second hypothesis was also denied, given that similarity index (Morisita-Horn and Sørensen) and common species between Dense and Typical Cerrado were lower than that between Dense Cerrado and Cerradão. Rather than that these results indicate that Dense Cerrado occupies an intermediate position between Cerradão and Typical Cerrado in terms of floristic composition. This is confirmed by TWINSPAN analysis which groups the Dense Cerrado plots with both Cerradão and Typical Cerrado plots. The third hypothesis, which predicted a gradient in structural parameters, the results once again led to rejection of the hypothesis, given the marked similarities of the Cerradão and Dense Cerrado, although tree height did vary systematically, decreasing from the Cerradão to the Dense Cerrado and then to the Typical Cerrado.

Overall, the similarities or differences among the three communities surveyed in the present study changed depending on the variables analyzed. While the Cerradão and Dense Cerrado were similar to one another in terms of species richness and the vertical and horizontal structure of the vegetation, the Cerradão and Typical Cerrado were similar in terms of species diversity. Finally, we observed a not predicted gradient in species composition, in which Cerradão and Typical Cerrado occupy the extremes and the Dense Cerrado a transitional position. From a conservation perspective, our results emphasize the importance of the protection of the three types of vegetation, not only for the preservation of beta diversity, but also the conservation of species that prefer either forested habitats or more open, savanna environments along the Cerrado-Amazon transition zone.

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