

The scale of effect depends on operational definition of forest cover-evidence from terrestrial mammals of the Brazilian savanna

Christophe Amiot, Cyntia Cavalcante Santos, Damien Arvor, Beatriz Bellón, Hervé Fritz, Clément Harmange, Jeffrey D D Holland, Isabel Melo, Jean-Paul Metzger, Pierre-Cyril Renaud, et al.

▶ To cite this version:

Christophe Amiot, Cyntia Cavalcante Santos, Damien Arvor, Beatriz Bellón, Hervé Fritz, et al.. The scale of effect depends on operational definition of forest cover-evidence from terrestrial mammals of the Brazilian savanna. Landscape Ecology, 2021, 36 (4), pp.973-987. 10.1007/s10980-021-01196-9. halshs-03131932

HAL Id: halshs-03131932 https://shs.hal.science/halshs-03131932

Submitted on 21 Oct 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

1 The scale of effect depends on forest definition – evidence from terrestrial

2 mammals of the Brazilian savanna

4	<u>Amiot</u> Christophe ^{ag} ,	Santos Cavalcante	Cyntia ^{a e j} , <u>Arvor</u>	<u>:</u> Damien ^b , <u>B</u>	<u>Bellón</u> Beatriz ^{a g k} ,	<u>Fritz</u>
	<u> </u>		J ,			

- 5 Hervé^{c g i}, <u>Harmange</u> Clément ^{a g}, <u>Holland</u> Jeffrey D. ^d, <u>Melo</u> Isabel ^e, <u>Metzger</u> Jean Paul ^f, <u>Renaud</u>
- 6 Pierre-Cyril^{ag}, <u>Roque</u> Fabio de Oliveira^{eh}, <u>Souza</u> Franco Leandro^e, <u>Pays</u> Olivier^{ag}*
- 7
- 8 ^a LETG-Angers, UMR 6554 CNRS, Université d'Angers, 49045 Angers, France.
- 9 ^b LETG-Rennes, UMR 6554 CNRS, Université de Rennes 2, 35043 Rennes, France
- 10 ° CNRS, Université Lyon, Université Lyon 1, Laboratoire de Biométrie et Biologie Evolutive UMR
- 11 5558, F-69622 Villeurbanne, France.
- ¹² ^d Department of Entomology, Purdue University, West Lafayette, Indiana, 47907 USA.
- 13 ^e Bioscience Institute, Federal University of Mato Grosso do Sul, Cidade Universitária, 79060-300
- 14 Campo Grande, MS, Brazil
- ¹⁵ ^fUniversidade de São Paulo, Departamento de Ecologia, São Paulo, SP, Brazil
- 16 ^g REHABS International Research Laboratory, CNRS-Université Lyon 1-Nelson Mandela
- 17 University, George Campus, Madiba drive 6531 George, South Africa.
- 18 ^h Centre for Tropical Environmental and Sustainability Science (TESS) and College of Science and
- 19 Engineering, James Cook University, Cairns, Qld, Australia
- 20 ⁱ Sustainability Research Unit, Nelson Mandela University, Port Elizabeth, South Africa
- 21 ^jWetlands International Brazil, Campo Grande, Mato Grosso do Sul, Brazil
- ^k Department of Environmental Science, Rhodes University, Makhanda 6140, South Africa
 23
- 24 * Corresponding author: Olivier Pays, LETG-Angers, UMR 6554 CNRS, Université d'Angers,
- 25 Campus Belle Beille, 2 Bd Lavoisier, 49045 Angers, France.
- e-mail address: olivier.pays@univ-angers.fr (O. Pays), phone: +33 (0) 241 735 261.
- 27
- 28 Author ORCID:
- 29 Amiot Christophe 0000-0002-4788-0928
- 30 Santos Cavalante Cyntia 0000-0002-8207-9263
- 31 Arvor Damien 0000-0002-3017-9625

- 32 Bellón Beatriz 0000-0002-9620-6200
- 33 Fritz Hervé 0000-0002-7106-3661
- 34 Harmange Clément 0000-0001-5207-021X
- 35 Holland Jeffrey D. 0000-0003-4889-6363
- 36 Metzger Jean Paul 0000-0002-0087-5240
- 37 Renaud Pierre-Cyril 0000-0003-1776-4923
- 38 Roque Fabio de Oliveira 0000-0001-5635-0622
- 39 Souza Franco Leandro 0000-0002-7041-4036
- 40 Pays Olivier 0000-0001-8268-1804
- 41

42 Keywords

43 Scale of response; Tree cover; Spatial scale; Multi-scale model; Mammal; Herbivore; Brazil

44

46 Abstract

- 47 *Context* The underlying mechanisms determining the scale at which species interact with their
- 48 environment are still poorly known.
- 49 *Objective* We investigated the spatial extent at which landscape structure affects the occurrence of
- 50 four species of terrestrial mammal herbivores in the Brazilian savannas and tested whether those
- 51 scales could be explained by species ecological traits and habitat definition.
- 52 *Methods* Using maps of forest cover, camera trapping and occupancy modelling, we determined the
- 53 relations between three landscape metrics (percentage of forest cover, patch density and edge
- 54 density) and the occurrence of four species. To determine the optimal scale of effect for each
- 55 species, we computed landscape metrics at different spatial extents (from 0.5 to 10 km radius) from
- 56 camera trap locations and for three forest maps, considering different definitions of what is a
- 57 "forest" (minimum of tree cover of 25, 50 or 75% per pixel).
- 58 *Results* The occupancy models revealed scales of effect of 0.5 to 2 km, and those scales overlapped
- 59 highly among species. However, the strength of the effect depends highly on how forest is defined,
- 60 being stronger when forest was defined with greater tree cover, particularly for forest-dwelling
- 61 species.
- 62 Conclusions Besides biological traits, the way habitat is defined shapes our ability to detect scale of
- 63 effects. Thus, if we want to properly identify scales of effect for multiple species, it is necessary not
- 64 only to adopt a multi-scale approach, but also to use multiple definitions of habitat, considering
- 65 particularities of how each species interact with their environment.
- 66

67 Introduction

68

69 Identifying the spatial scale at which species respond to landscape structure, or the scale of effect 70 (Jackson and Fahrig 2012) has become an important scientific challenge when investigating 71 species-environment relationships (Fahrig et al. 2011; Redon et al. 2014; Miguet et al. 2016, 2017). 72 As species might respond to a specific landscape attribute at a particular scale, studies have pointed out the importance to consider multiple spatial scales when investigating the effects of habitat 73 74 change on species (Levin 1992). This is particularly relevant in the understanding of mechanisms 75 underlying the erosion of biodiversity, especially the mechanisms for which human activities shape 76 land use and land cover changes (Newbold et al. 2015; Püttker et al. 2020). Knowing the scale at 77 which species respond to landscapes should help decision-makers in designing management plan to 78 maintain and restore biodiversity and their habitats (Haines-Young 2009).

79 From an ecological standpoint, optimal spatial extents are mainly determined by the scale at 80 which ecological processes are expected to operate for the studied organisms. Huais (2018) argued 81 that ecologists do not usually know *a priori* which are those optimal spatial extents, mainly due to a 82 lack of a full understanding of the underlying biological processes (Jackson and Fahrig 2015). 83 However, several studies have clarified the link between spatial scale and the ecological responses 84 of species under investigation (Levin 1992; Saab 1999; Crawley and Harral 2001; Chase and 85 Leibold 2002: Leibold et al. 2004: Rahbek 2005: Gabriel et al. 2010: Delsol et al. 2018). For 86 instance, the effect of spatial extent has been particularly investigated in habitat selection (Fortin et 87 al. 2008) and animal movement (Fryxell et al. 2008). For large mammal herbivores, Mayor et al. 88 (2009) have reported spatial scales at which ecological and behavioural mechanisms underlying 89 habitat selection should be investigated. Analysing the activities of animals at fine scales (i.e., 1 -90 100 m) allows to investigate bite, feeding site and patch selection, while studies at local scales (100 91 m - 10 km) allow the understanding of habitat selection and home range, and finally, research at 92 broader, regional scales (>10 km) are relevant for migration and (meta)population dynamics 93 comprehension (Johnson 1980; Danell et al. 2006).

94 Without knowing *a priori* what is the appropriate scale of study, landscape ecologists 95 commonly assess landscape variables at multiple scales to select the scale that yields the best 96 species-landscape relationship (e.g. Brennan et al. 2002; Holland et al. 2004; Holland and Yang 97 2016; Melo et al. 2017; Huais 2018). Different methods or statistical analyses have been proposed 98 in the literature to test for the effect of landscape features on species attributes (e.g., richness, 99 abundance, composition) at different spatial scales and to extract their goodness of fit through an 100 appropriate criterion index (e.g., correlation and determination coefficients or AIC). Spatial scales are usually tested by varying spatial extent (e.g., the size of a buffer zone) over which the habitat 101

predictor variable (i.e., a landscape metric) is measured. The scale of effect is commonly derived by plotting the spatial scale (e.g., buffer size) against goodness of fit. This is sometimes challenging because there is a common lack of knowledge on 1) the suitable landscape metrics to investigate the ecological processes, and 2) the appropriate grain extent at which their relationships should be assessed.

107 Biological/ecological traits of species including for instance body size, dispersal distance, 108 home range and habitat specialisation may affect the scale of effect (Miguet et al 2016). Those traits 109 are particularly relevant when they are indicators of animal mobility, a key biological feature which 110 determines the relationship between species attributes and their surrounding space (Bowman et al. 111 2002). Although the empirical studies are not all in agreement, we might expect that the scale of 112 effect would be larger for larger-bodied species because they are more mobile with higher dispersal 113 distance and home range than smaller-bodied species (Jackson and Fahrig 2012; Miguet et al 2016). 114 Thus, when investigating the effect of landscape features on species occurrence, a general guideline 115 for empirical studies is to consider a radius of a landscape to be 4-9 times the median dispersal 116 distance or 0.3 - 0.5 times the maximum dispersal distance of a species. Thornton and Fletcher (2014) in their meta-analysis on birds reported a positive correlation between body size and the 117 spatial scale at which species responded to landscape variables. Finally, the scale of effect may 118 119 differ between specialist and generalist species of similar body-size although this literature is sparse 120 and not in agreement on whether habitat specialists would exhibit a larger or smaller scale of effect 121 (Boscolo and Metzger 2009; Chaplin-Kramer et al. 2013). Comparisons of habitat generalists and 122 specialists have the added difficulty in consistently defining habitat for the different groups.

Landscape features, related to both land use and land cover composition and spatial 123 124 configuration, could also affect the scale of effect. Particularly, the type of maps of land use and 125 land cover that are used as an input to derive patch-based landscape metrics may influence the 126 assessments of scale of effect. This issue is especially relevant in tropical areas dominated by 127 vegetation gradients (ranging from herbaceous layers with sparse trees to shrubland and forested 128 areas, including degraded forests or regenerating forests) where the definition of landscape units can 129 be arbitrary, and low thematic resolution in land cover maps (e.g., a small number of classes) may 130 lead to an overly simplified representation of reality that may not adequately represent how species 131 perceive their environment (Comber et al. 2005; Rocchini et al. 2013). Moreover, the definition of what is a "forest," for instance, is vague, ambiguous and highly subjective (Bennett 2002) where 132 133 gradients in tree cover exist. Indeed, hundreds of different definitions of "forest" coexist (Chazdon et al. 2016), and obviously, none of these use a species perspective. While Hansen et al. (2010) used 134 25% of tree cover to map forest cover, Defries et al. (2001) defined forest cover as greater than 60% 135 136 tree cover. Although using different percent tree cover would thus select different amount of

canopy, it could also affect the type of vegetation that would be considered. In tropical and 137 subtropical zones of Africa, Australia and South America, Hirota et al. (2011) reported that percent 138 tree cover in savannas was about 20% and contrasted with 80% in forested areas. While mammal 139 species might respond differently to forest cover (Royo and Carson 2005; Young et al. 2013; 140 141 Chamaillé-Jammes et al. 2016; Ferreguetti et al. 2017; Zimbres et al. 2018), the question whether 142 the percent tree cover used to map forest cover affects the scale of effect of landscape metrics on the 143 occurrence of species has never been studied. This might be critical when modelling the presence of 144 forest dwelling species in complex mosaic of vegetation types such as in Brazil's Cerrado including 145 forests, savannas and grasslands (Bonanomi et al. 2019).

The aim of this study is to investigate the scale of effect in four common species of 146 147 terrestrial mammal herbivores, the Azara's agouti Dasyprocta azarae, the collared peccary Pecari tajacu, the brown brocket deer Mazama gouazoubira, and the south American tapir Tapirus 148 149 *terrestris*, using camera traps in the Brazilian Cerrado. First, we studied how the occurrence of 150 those species responded to three landscape metrics commonly used to describe landscape 151 composition (FC: percent of forest cover) and configuration (PD: patch density and ED: edge density) (Lidicker 1999; Bastian et al. 2006; Bennett et al. 2006; Lu et al. 2013; Lowicki 2017). 152 153 These metrics have also been useful to explain mammal community distribution in Brazil (Mares et 154 al. 1986; Arévalo-Sandi et al. 2018; Bovendorp et al. 2019; Püttker et al. 2020). Second, we investigated whether the definition of forest cover (i.e. 25, 50, or 75% of tree cover) affected the 155 detection of scale of effect of the landscape features on the occurrence of species. Finally, as the 156 157 four studied species differ in their body-size, dispersal and home range and forest specialisation (Table 1), we investigated whether the scale of effect varied among species with different 158 biological/ecological traits. 159

160 We ran occupancy models for each species at 8 spatial extents from 0.5 to 10 km using 3 161 different maps of forest cover. We extracted the Akaike Information Criterion (AIC) for each model 162 and examined whether species exhibited scale of effects. We hypothesized that mobility largely 163 determines the relationship between landscape features and occurrence of the studied species. We 164 therefore predicted that larger species with higher dispersal range and home range would exhibit a larger scale of effect than smaller species. In this context, the scale of effect of the South American 165 tapir should be larger than the brown brocket deer and the collared peccary, with the smallest scale 166 of effect in the Azara's agouti. Regarding forest definition (that shaped both land use and land cover 167 168 composition and spatial configuration), we predicted a stronger or clearer scale of effect in the two 169 forest-dwelling species, the Azara's agouti and collared peccary, when considering high tree cover 170 (e.g., 75%) compared to medium-low tree cover (50 - 25%).

172 Methodology

173

174 Study site

The Bodoguena Plateau region (20°25'29.28" to 21°44'19.72" S and 56°52'24.46"56° to 17'23.36" 175 W) is located in the state of Mato Grosso do Sul, Brazil (Fig. 1a). It is characterized by a mountain 176 177 chain (altitude 450 - 800 m) with a tropical climate (annual temperature varies between 20 and 22°C and rainfall between 1300 and 1700 mm). The study area is within the range of the Cerrado 178 179 biome (Brazilian savanna) in the region inside and around the Serra da Bodoquena National Park 180 (SBNP). The vegetation comprises a mix of alluvial semi-deciduous and submontane deciduous 181 forest (dry forest), wetlands, regenerating areas and agriculture including pastures and crops 182 (Oliveira and Marquis 2002; Klink and Machado 2005). SBNP represented an important protected 183 area due to acting as an ecological corridor of biodiversity and by maintaining remnants of Atlantic 184 Forest in the area. During the last decades, Cerrado has been experiencing a rapid agricultural 185 expansion (Rausch et al. 2019), leading to fragmentation (Strassburg et al. 2017). 186 187 Forest cover, landscape metrics and spatial extent

188 Landscape metrics were computed using forest maps derived from the Global Forest Change project's products (Hansen et al. 2013). We used the "2000 percent tree cover" (TC) product, which 189 190 corresponds to the proportion of trees per pixel, i.e. the canopy closure for all vegetation taller than 191 5 m in height. This global product is derived from the analysis of time series of Landsat images 192 with a 30 m spatial resolution. We additionally used the "Forest loss year" product from the Global 193 Forest Change project to remove all deforested areas between 2001 and 2017 (year of field 194 collection; see below) from the forest class in the TC, i.e., TC was set to 0 for deforested areas. 195 Based on this updated TC map, we then produced three forest maps for 2017 applying three

196 thresholds of percent tree cover (25, 50, and 75%) (Fig. 2).

197 Each of the three forest maps was then used to derive three landscape metrics: one metric of 198 landscape composition, forest cover (FC), and two additional metrics of landscape configuration, 199 patch density (PD) and the edge density (ED). FC is the proportion of pixels classified as forest in a 200 given area. PD measures the number of forest patches per km² and ED is the total length of all 201 edges of forest patches divided by the buffer area. These three metrics were calculated using the 202 ClassStat function from the SDMTools package (VanDerWal et al. 2014) in R 3.6.2 (R Development 203 Core Team 2019). Although a large range of landscape metrics could have been explored, we 204 limited this to three variables in our study as (1) they are commonly used in the literature and (2) 205 they allowed an assessment of forest cover and fragmentation in the Cerrado in 2017 when we

collected data in the field (Bonanomi et al. 2019). We checked that these three metrics were notsignificantly correlated.

208 We calculated ED, PD and FC at different scales, here considered as radii from the sampled 209 areas (i.e. buffer radius of 500, 750, 1000, 1500, 2000, 3000, 5000 and 10000 m). The range of 210 spatial extents allowed the investigation of local and dispersal movements of species (Miguet et al. 211 2016) and have been used to explain habitat selection in terrestrial herbivorous mammals (Johnson 212 1980; Danell et al. 2006). We chose to exclude two smaller buffer sizes, 100 and 250 m, as the 213 small number of pixels from Landsat images (30 m spatial resolution) did not provide enough 214 information to assess accurately PD, ED and FC (Strahler et al. 1986; Woodcock et al. 1987). 215 Finally, landscape variables showed scaling relations suggesting that they could be predicted

over a wide range of spatial extents in our study area and remained quite constant beyond a buffer
 radius of 3 km (Electronic Appendices Table S1 and Fig. S1)

218

219 Data from mammal species

220 We deployed a set of 193 camera traps (Reconyx-hyperfire HC500 and Bushnel-trophy Cam HD) in 221 a gradient of forest cover loss in the study area (Fig. 1b). Each camera trap was in the same location 222 on two different periods of 20 consecutive days between May 2016 and December 2017. They were 223 positioned 40 cm above the ground on the nearest tree of a computer-generated random point with a 224 10° inclination towards the ground. To increase detection of animal movement and maximise 225 identification of species, frame rate interval was set up at 3 seconds with 3 pictures per event. At the 226 end of the sampling period, SD cards were extracted and analysed visually using Wild.ID (TEAM 227 Network 2019). Each sampled picture was checked by two observers to reduce misidentification. In 228 this study, we focused on four species of terrestrial mammals typical from Cerrado: Azara's agouti, 229 collared peccary, brown brocket and South American tapir. Their ecological traits including body 230 size, home range, habitat use, diet and dispersal distance are presented in Table 1.

231

232 Occupancy models

233 Creating large buffers around the locations of the 193 camera traps led to spatial overlaps and thus 234 pseudoreplication. To tackle this issue, we used the Focus program (Holland et al. 2004) to conduct 235 a plot-buffer related sampling using 10 000 iterations on the 193 camera trap locations to select a set of spatially independent locations. Based on the retrials, Focus-2.0 randomly selected one 236 237 combination of 57 independent locations allowing to increase buffer size up to 2 km (Fig. 1b). As 238 the aim of our study was to understand the response of the occurrence of mammal species to the landscape metrics at wide range of different scales, we chose to use this combination to investigate 239 240 scaling relations beyond 2 km, i.e. 3, 5 and 10 km although some minimal overlapping existed

between several neighbouring buffers. Indeed, beyond 2 km, Focus-2.0 would have reduced the set of combinations to 30 camera traps for which occupancy models failed to converge. Thus, we kept the set of 57 selected locations and we then assessed each landscape metric described above at each buffer size and run occupancy models.

245 To investigate the effect of the landscape features (measured for the 8 buffer sizes and 3 246 percent threshold of tree cover) on the occurrence of the four mammal species, we run occupancy 247 models using the *unmarked* R package (Fiske and Chandler 2011). For each of the 57 camera traps 248 and focal species, we implemented a data set of 40 sampled days in which we indicated when the 249 studied species was detected (coded as 1) vs. non-detected (coded as 0). Then, from the 40 sampled days, we built a matrix of 8 sampling occasions of 5 consecutive days each (as in Niedballa et al. 250 251 2015). Following this method, 13 camera traps (i.e., 23%) have detected the presence of Azara's 252 agouti, 15 (26%) of collared peccary, 13 (23%) of brown brocket deer and 25 (44%) of South 253 American tapir. Over the combination of 57 camera traps, there was a reasonable chance to detect 254 the selected species. Our study assumed that we did have false detections and the absence of 255 detection of a focal species over one period indicated that sites were truly unoccupied during the sampling period or species were very scarce at the sampled sites during the survey. 256

257 We used the species detection information to investigate the effects of spatial extent of 258 habitat covariates on the probability of species occupancy. We ran occupancy models for which 259 occupancy probability Ψ at each camera trap location and the detection probability p were modelled 260 as a linear function of covariates x_i using logit link functions (see Niedballa et al. 2015 for details). 261 We ran models for each buffer size and species. First, we determined for each species how the probability of detection p should be implemented in null models. We used a stepwise approach 262 starting with a model $\Psi(.)p(.)$ where both occupancy and detection probabilities were constant (i.e., 263 264 fixed as 1). As density of understory and weather could affect detection probability around the 265 camera traps, we created two covariates called visibility (contrasting open vs. dense understory) and 266 weather (contrasting rainy vs. dry season). Then we ran three other models $\Psi(.)p(x_i)$ to test whether 267 covariates x_i including visibility around camera trap and weather, or both, affected detection 268 probability p with occupancy probability Ψ constant. We compared these four models of detection 269 $(\Psi(.)p(.), \Psi(.)p(visibility), \Psi(.)p(weather), \Psi(.)p(visibility, weather) using AIC and selected the$ 270 model with the lowest AIC and Δ AIC higher than 2 with the closest competing model. Finally, from 271 this selected (null) model of detection, we tested the effects of landscape metrics on occupancy 272 probability Ψ at each of the 8 buffer sizes and three landscape metrics calculated from the three 273 forest cover maps (25, 50 and 75%). Thus, for each landscape metric, we ran 8 models that we 274 compared using AIC and null models. All models showing $\Delta AIC < 2$ from the best fitted models 275 had similar statistical supports. Estimates $(\pm SE)$ of landscape metrics with their P-values affecting

276 occupancy probability were extracted from the best candidate models. Statistical analyses were

277 performed using R 3.6.2 (R Development Core Team 2019).

278

279 **Results**

280

281 Performance of models run at different spatial extents highlighted scale of effects of landscape 282 metrics on the occurrence of three species. Indeed, buffer sizes that showed the strongest statistical supports (i.e. $\Delta AIC < 2$) ranged from 500 to 1500 m in ED (Fig. 3a) and 500 to 2000 m in FC (Fig. 283 284 3i) in Azara's agouti, from 500 to 750 m in FC in collared peccary (Fig. 3j) and 750 m in PD in brown brocket deer (Fig. 3g) when percent tree cover was higher than 75% (Electronic appendix 285 Table S2). For the other situations in the three species, we did not detect a scale of effects as null 286 287 models including only covariates on detection probability p provided similar statistical models than 288 candidate models including covariates affecting occupancy probability Ψ (Electronic appendix 289 Table S2). The South American tapir did not exhibit any scale of effect for the three landscape 290 metrics (Fig 3d, h, l).

291 When the threshold of the percent tree cover used to map forest, cover decreased (from 75 to 50% and from 50 to 25%), we observed 3 different patterns from the performance of models at the 292 different buffer sizes (Electronic appendices Table S2-4). The range of selected buffer sizes (i.e. the 293 294 scale of effect) (1) remained similar as for instance in FC in collared peccary, (2) decreased as in Azara's agouti in ED, or (3) was no longer consistent in PD in brown brocket deer (Table 2). 295 296 Although a range of selected models emerged at 50% tree cover in PD in Azara's agouti, we did not 297 detect any significant effect of this variable (i.e., P > 0.05, Table 2). Thus, we did not consider this 298 result as robust. Altogether, our analyses suggest that the species exhibit a stronger scale of effect 299 when considering high tree cover (e.g. 75%) compared to medium-low tree cover (50 - 25%).

Finally, regarding the estimate (β) ± SE associated with the significant effect of each landscape metric extracted from the candidate models with the lowest AIC (buffer size indicated with * in Table 2), the occurrence of Azara's agouti and collared peccary increased with FC, brown brocket deer with PD, and Azara's agouti with ED when the threshold on percent tree cover was 75%. When this threshold decreased, the trend did not change in FC as in collared peccary although the strength of the effect felt to nearly significant value in Azara's agouti (Table 2).

306

307 **Discussion**

309 Scales of effects varied among mammal species from the Brazilian savannahs, and this variation

could be partially explained by species traits, but most importantly, was strongly influenced by howwe define forests.

The scale of effect of Azara's agouti ranged from 500 to 2000 m (for FC and ED), while for 312 313 collared peccary it ranged from 500 to 750 m (FC), and for brown brocket deer it was 750 m (PD). 314 The South American tapir did not exhibit any scale of effect on the three landscape metrics. Thus, 315 the spatial extents that have been detected suggest that ecological and behavioural mechanisms 316 underlying the scale of effects occurred at local scale and were associated with habitat selection and 317 home range as reported in medium and large-body-sized herbivore species (Johnson 1980; Danell et 318 al. 2006). These results also suggest that a range of spatial extents (from 500 to 2000 m in this 319 study) should be considered rather than a particular buffer size in the scale of effect of landscape metrics on species occurrence (Moraga et al. 2019). The ability of some species to live or use 320 321 fragmented sites depended on their biological traits such as body size (Cardillo et al. 2005), 322 ecological requirements (Miguet et al. 2016) or their ability to cross harsh habitats (Pires et al. 323 2002). We predicted that scale of effect would be larger for larger-bodied species because they are 324 more mobile, tend to be move farther and their home range are higher than smaller-bodied species 325 (Bowman et al. 2002; Jackson and Fahrig 2015). From performance of models with 75% tree cover, 326 we did not detect that scale of effect was larger in the largest studied species, the South American 327 tapir, and the scale of effect overlapped highly among the Azara's agouti, collared peccary and 328 brown brocket deer. Thus, body-size as indicator or animal mobility may not be a very strong 329 predictor of the scale of effect in our four studied tropical herbivorous species. The effect of the 330 dispersal distance on the scale of effect need to be tested on a wide range of species including 331 micromammals to address overall conclusions.

332 Contrary to the two studied generalist species, the South American tapir and brown brocket deer, the two forest-dwelling species, the Azara's agouti and collared peccary, exhibited a scale of 333 334 effect on FC. These results might show that species that are more generalist in their habitat 335 requirements might not be limited by forest cover using agricultural landscape with open areas to 336 achieve their ecological requirements (e.g., food). A previous study in Brazilian Amazon reported 337 that the lowland tapir was one of the species with a high ability to survive in more altered 338 environments including regeneration areas with food opportunities (Teixeira-Santos et al. 2020). 339 The lack of scale of effect on the occurrence of the South American tapir might also be observed if 340 the response is not scale dependent (Martin and Fahrig 2012) or the landscape heterogeneity does 341 not allow to investigate pattern at a larger scale in species with high dispersal range (> 3km for this 342 species, Table 1). As in our study area, landscape variables remain quite fixed beyond a buffer 343 radius of 3 km (Electronic Appendices Table S1 and Fig. S1), the opportunity to detect scale of

effect beyond this distance might be limited. Finally, as the effect of ecological traits might interact
to explain the scale of effect of forest cover on species occurrence (as for instance the studied
smaller bodied size species were the forest-dwelling ones), more studies are needed to clarify the
role of uncorrelated traits in a larger range of species.

348 Landscape features might affect the scale of effect on species occurrence. Lyra-Jorge et al. 349 (2010) reported that two large and highly mobile carnivores, Puma concolor and Chrysocyon 350 *brachyurus*, were best explained by edge density of the native vegetation at a coarse scale of 2 km. 351 Our analyses showed that species that mainly use forest edges, the Azara's agouti and the collared 352 peccary, exhibited a scale of effect on metrics associated to landscape configuration such as ED and 353 PD. Regolin et al. (2017) reported that the richness of carnivores community increased with forest 354 cover and decreased with landscape fragmentation. Bogoni et al. (2018) showed that large herbivores decreased when native vegetation cover, forest fragment size, and the largest 355 356 neighbouring patch of remnant forest decreased. Here, as the studied species exhibited different 357 patterns of scale of effect with the landscape metrics, our results supported previous studies 358 showing that landscape composition and configuration are both important drivers in shaping occurrence of terrestrial mammals in native forests (Ochoa-Quintero et al. 2015; Regolin et al. 359 360 2017). Unfortunately, among the four selected species, we do not have one mammal species that did 361 not require forest.

362 While different percentages of tree cover have been used in the literature to map forest cover 363 (Defries et al. 2001; Hansen et al. 2010), studies were reporting that occurrence of terrestrial 364 mammal species was affected by forest cover (Chiarello 2000; Michalski and Peres 2007) and vegetation types (Lyra-Jorge et al. 2010). However, few studies have examined whether the percent 365 tree cover used to map forest cover affect the scale of effect of landscape metrics on the occurrence 366 367 of species. Overall, our analyses showed that species exhibited a stronger scale of effect when considering high tree cover (e.g., 75%) compared to medium-low tree cover (50 - 25%). Thus, the 368 369 detection of scale of effect varies with habitat definition. While the percent tree cover used to map 370 forest cover shaped patch-based landscape metrics and thus the assessment of land use and land 371 cover composition and spatial configuration, it also determines our ability to detect the correct scale 372 of effects for species. This is a new and important result, which implies that before using multiple 373 sizes of buffers to test the scale of effects, we need a previous and decisive step, which is to define correctly habitat types of species. However, defining properly what is an "habitat" is challenging, 374 375 particularly for species living in complex mosaic of vegetation types including forests, savannas 376 and grasslands (Bonanomi et al. 2019) showing for instance environment gradients. Several 377 percentages of tree cover have been used in the literature to map forest cover (Defries et al. 2001; 378 Hansen et al. 2010). Although changing the percent tree cover allows to select different amount of

379 canopy, it also affects the type of vegetation and thus habitat for species (Hirota et al. 2011). These

- 380 results are particularly important in the current landscape dynamics occurring in the Cerrado. In the
- 381 last decades, agriculture expansion at the expense of native vegetation dramatically increased
- 382 landscape fragmentation (Strassburg et al. 2017) and the need to investigate the mechanism
- 383 underlying the response of biodiversity to the erosion of native vegetation (e.g., forest cover) has
- been a critical topic for conservation purposes in Brazil (Pardini et al. 2010; Martensen et al. 2012;
- Banks-Leite et al. 2014; Ochoa-Quintero et al. 2015).
- 386

387 Implications for conservation and land planning

Although the spatial extent (scale) at which landscape attributes are measured has a substantial 388 389 impact on inferred species-landscape relationships, the way that habitats are defined has a critical impact on the scale of effect analysis. As mammal species might respond differently to forest cover 390 391 (Chamaillé-Jammes et al. 2016; Zimbres et al. 2018), the question whether forest cover (i.e., percent 392 tree cover) has been appropriately selected to link landscape metrics on the occurrence of species 393 appears as the first main concern to avoid failure in conservation initiatives. Here we recommend using and testing multiple definitions of habitats to assess the ecological requirements of studied 394 species. Although 75% tree cover seems to be a good proxy of forest cover to study the forest-395 dwelling species in our study area, this might also be the case for other variables of landscape 396 composition that are characterised by a continuous gradient such as agriculture or urbanisation. 397 398 Moreover, from a conservation perspective, it is critical to test multiple scales at which species 399 respond to the landscape variables. In the Cerrado, we recommend that initiatives examining spatial 400 arrangement of landscape attributes of forest cover in relation to populations in terrestrial mammals 401 (from small to large body sized species) should consider buffer sizes from 500 to 2000 m. However, 402 the puzzle is far more complicated with questions regarding communities. No management approach based on a single scale would benefit all species (Crouzeilles and Curran 2016; Bhakti et 403 404 al. 2018). It might be important to analyse a large range of species and set targets for each one using specific scales. This should be relevant in the context of systematic planning approaches and 405 406 multiscale management based on the responses of multiple species (Neel et al. 2004). Thus, using 407 multiple definitions of habitats and scales is critical as most initiatives of ecological zoning, 408 biodiversity prioritizing or ecosystem restoration use landscape units that have a particular meaning 409 in terms of land use dynamics and planning (usually justified by physical constraints) but not in 410 terms of functional responses of the organisms.

412 Acknowledgements

- 413 This work was funded by a CAPES (Coordenação de Aperfeiçoamento de Pessoal de Nível
- 414 Superior, Brazil) COFECUB (Comité Français d'Évaluation de la Coopération Universitaire et
- 415 *Scientifique* and Campus France) grant (French number: Sv 875-17 and Brazilian Process:
- 416 23038.001818/2020-70), the Institutional Program of Internationalization sponsored by the CAPES
- 417 (Finance Code 001; Capes-PrInt 41/2017 Process: 88881.311897/2018-01), the Conselho
- 418 Nacional de Desenvolvimento Científico e Tecnológico (CNPq Process: 301306/2018-4) and the
- 419 Pesquisa Ecológica de Longa Duração (PELD/ILTER) Planalto da Bodoquena and the University
- 420 of Angers (FragHerb project). We would like to thank the Programa de Pós Graduação em
- 421 Ecologia e Conservação (PPGEC) of the Universidade Federal de Mato Grosso do Sul (UFMS),
- 422 the CAPES, the COFECUB, Campus France, the Fundação de Apoio ao Desenvolvimento do
- 423 Ensino, Ciência e Tecnologia do Estado de Mato Grosso do Sul (FUNDECT) for their institutional
- 424 supports and Jacques Baudry for his helpful comments.
- 425
- 426

427 **References**

- Aliaga-Rossel E, Kays R, Fragoso J (2008) Home-range use by the Central American agouti
 (Dasyprocta punctata) on Barro Colorado Island, Panama. J Trop Eco 124: 367-374
- 430 Arévalo-Sandi A, Bobrowiec PED, Chuma VJUR, Norris D (2018) Diversity of terrestrial mammal
 431 seed dispersers along a lowland Amazon forest regrowth gradient. PloS one 13: e0193752.
- 432 Banks-Leite C, Pardini R, Tambosi LR, Pearse WD, Bueno AA, Bruscagin RT, Condez TH, Dixo
- M, Igari AT, Martensen AC, Metzger JP (2014) Using ecological thresholds to evaluate the
 costs and benefits of set-asides in a biodiversity hotspot. Science 345: 1041-1045
- Bastian O, Krönert R, Lipský (2006) Landscape diagnosis on different space and time scales a
 challenge for landscape planning. Landsc Ecol 21: 359-374
- Bhakti T, Goulart F, de Azevedo CS, Antonini Y (2018) Does scale matter? The influence of threelevel spatial scales on forest bird occurrence in a tropical landscape. PlosONE 13: e0198732
- Bennett B (2002) What is a forest? on the vagueness of certain geographic concepts. Topoi 20:189201Bennett A, Radford JQ, Haslem A (2006) Properties of land mosaics: implications for
- nature conservation in agricultural environments. Biol Conserv 133: 250-264
- 442 Bodmer RE (1989) Ungulate biomass in relation to feeding strategy within Amazonian forests.

443 Oecologia 81: 547-550

- 444 Bodmer RE (1991) Strategies of seed dispersal and seed predation in Amazonian ungulates.
- 445 Biotropica 23: 255-261

- 446 Bogoni JA, Graipel ME, Oliveira-Santos LGR, Cherem JJ, Giehl EL, Peroni N (2017) What would
- be the diversity patterns of medium- to large-bodied mammals if the fragmented Atlantic
 Forest was a large metacommunity? Biol Conserv 211: 85-94
- Bonanomi J, Tortato FR, Gomes FSR, Penha JM, Bueno AS, Peres CS (2019) Protecting forests at
 the expense of native grasslands: Land-use policy encourages open-habitat loss in the
 Brazilian cerrado biome. Perspect Ecol Conser 17: 26-31
- Boscolo D, Metzger JP (2009) Is bird incidence in Atlantic forest fragments influenced by
 landscape patterns at multiple scales? Landsc Ecol 24: 907-918
- Bovendorp RS, Brum FT, McCleery RA, Baiser B, Loyola R, Cianciaruso MV, Galetti M (2019)
 Defaunation and fragmentation erode small mammal diversity dimensions in tropical forests.
 Ecography 42: 23-35
- Bowman J, Jaeger JAG, Fahrig L (2002) Dispersal distance of mammals is proportional to home
 range size. Ecology 83: 2049-2055
- Brennan JM, Bender DJ, Contreras TA, Fahrig L (2002) Focal patch landscape studies for wildlife
 management: optimizing sampling effort across scales. In: Liu J, Taylor WW (eds.)
- 461 Integrating landscape ecology into natural resource management. Cambridge University
 462 Press, Cambridge, pp 68–91
- 463 Cardillo M, Mace GM, Jones KE, Bielby J, Bininda-Emonds ORP, Sechrest W, Orme CDL, Purvis
 464 A (2005) Multiple causes of high extinction risk in large mammal species. Science 309: 1239 465 1241
- Chaplin-Kramer R, de Valpine P, Mills NJ, Kremen C (2013) Detecting pest control services across
 spatial and temporal scales Agric Ecosyst Environ 181: 206-212
- 468 Chase JM, Leibold MA (2002) Spatial scale dictates the productivity-biodiversity relationship.
 469 Nature 416: 427-430
- 470 Chamaillé-Jammes S, Charbonnel A, Dray S, Madzikanda H, Fritz H (2016) Spatial Distribution of
- 471 a Large Herbivore Community at Waterholes: An Assessment of Its Stability over Years in
 472 Hwange National Park, Zimbabwe. PLoS ONE 11: e0153639
- 473 Chazdon RL, Pedro HS, Brancalion LL, Bennett-Curry A, Buckingham K, Kumar C, Moll-Rocek J,
- 474 Vieira ICG, Wilson SJ (2016) When is a forest a forest? Forest concepts and definitions in the
- 475 Era of forest and landscape restoration." Ambio 45 (5): 538–550. doi:10.1007/s13280-016476 0772-y
- 477 Chiarello AG (2000) Density and populations size of mammals in remnants of Brazilian Atlantic
 478 Forest. Conserv Biol 14:1649-1657
- 479 Cid B, Oliveira-Santos LGR, Mourão G (2013) Seasonal habitat use of Agoutis (Dasyprocta
- 480 *azarae*) is driven by the palm Attalea Phalerata in Brazilian Pantanal. Biotropica 45: 380-385

- 481 Comber A, Fisher P, Wadsworth R (2005) You know what land cover is but does anyone else?... an
- 482 investigation into semantic and ontological confusion. Int J Remote Sens 26: 223-228
- 483 Crawley MJ, Harral JE (2001) Scale dependence in plant biodiversity. Science 291: 864-868

484 Crouzeilles R, Curran M (2016) Which landscape size best predicts the influence of forest cover on
 485 restoration success? A global meta-analysis on the scale of effect. J Appl Ecol 53: 440-448

- 486 Danell K, Bergström R, Duncan P, Pastor J (2006) Large herbivores and ecosystem functioning. –
 487 Cambridge University Press, Cambridge
- 488 Defries RS, Hansen MC, Townshend JRG, Janetos AC, Loveland TR (2001) A new global 1-km
 489 dataset of percentage tree cover derived from remote sensing. Global Change Biol 6: 247-254.
- 490 Delsol R, Loreau M, Haegeman B (2018) The relationship between the spatial scaling of
 491 biodiversity and ecosystem stability. Glob Ecol Biogeogr 27: 439-449
- 492 Duarte JMB, Vogliotti A, Zanetti EV, Oliveira ML, Tiepolo LM, Rodrigues LF, Almeida, LB
- 493 (2012) Avaliação do risco de extinção do veado-catingueiro *Mazama gouazoubira* G. Fischer
 494 [von Waldheim], 1814, no Brasil. Biodiversi Brasil, 3: 50-58
- Eisenberg JF, Redford KH (1999) Mammals of the Neotropics: volume 3: the central Neotropics.
 University of Chicago Press, Chicago, Illinois.
- Fahrig L, Baudry J, Brotons L, Burel FG, Crist TO, Fuller RJ, Martin JL (2011) Functional
 landscape heterogeneity and animal biodiversity in agricultural landscapes. Ecol Lett 14: 101–
 112.
- 500 Ferreguetti AC, Tomas WM, Fergallo HG (2017) Differences in the mammalian habitat use in a
- 501 mosaic of vegetation types of an Atlantic rain-forest Reserve, Brazil. Mastozool Neotrop 24:
 502 355-364
- Fiske I, Chandler R (2011) unmarked: An R Package for Fitting Hierarchical Models of Wildlife
 Occurrence and Abundance. J Stat Softw 43: 1-23
- Fortin D, Courtois R, Etcheverry P, Dussault C, Gingras A (2008) Winter selection of landscapes by
 woodland caribou: behavioural response to geographical gradients in habitat attributes. J Appl
 Ecol 45:1392-1400
- Fragoso JMS (1998) Home Range and movement patterns of white-lipped Peccary (Tayassu pecari)
 Herds in the Northern Brazilian Amazon. Biotropica 30: 458-469
- 510 Fryxell JM, Hazell M, Börger L, Dalziel BD, Haydon DT, Morales JM, McIntosh T, Rosatte RC
- 511 (2008) Multiple movement modes by large herbivores at multiple spatiotemporal scales. Proc
 512 Natl Acad Sci USA 105: 19114-19119
- 513 Gabriel D, Sait SM, Hodgson JA, Schmutz U, Kunin WE, Benton TG (2010) Scale matters: the
- 514 impact of organic farming on biodiversity at different spatial scales. Ecol Lett 13: 858-869

- 515 Hansen MC, Potapov PV, Moore R, Hancher M, Turubanova SA, Tyukavina A, Thau D, Stehman
- 516 SV, Goetz SJ, Loveland TR, Kommareddy A (2013) High-resolution global maps of 21 st-517 century forest cover change. Science 342: 850-853
- Hansen MC, Stehman SV, Potapov PV (2010) Quantification of global gross forest cover loss. Proc
 Natl Acad Sci USA 107: 8650–8655
- 520 Hirota M, Holmgren M, Van Nes EH, Scheffer M (2011) Global Resilience of Tropical Forest and
 521 Savanna to Critical Transitions. Science 334: 232-235
- Holland J, Bert DG, Fahrig L (2004) Determining the spatial scale of species' response to habitat.
 Bioscience 54: 227-233
- Holland J, Yang S (2016) Multi-scale studies and the ecological neighbourhood. Curr Landsc Ecol
 Rep 1: 135-145
- Huais PY (2018) multifit: an R function for multi-scale analysis in landscape ecology. Landsc Ecol
 33: 1023-1028
- Jackson HB, Fahrig L (2012) What size is a biologically relevant landscape? Landsc Ecol 27: 929–
 941
- Jackson HB, Fahrig L (2015) Are ecologists conducting research at the optimal scale? Glob Ecol
 Biogeogr 24: 52–63
- Johnson DH (1980) The comparison of usage and availability measurements for evaluating resource
 preference. Ecology 61: 65-71
- Jansen PA, Hirsch BT, Emsens W-J, Zamora-Gutierrez V., Wikelski M, Kays R (2012) Thieving
 rodents as substitute dispersers of megafaunal seeds. Proc Natl Acad Sci USA 109: 12610 12615
- Johnson DH (1980) The comparison of usage and availability measurements for evaluating resource
 preference. Ecology 61: 65-71
- Judas J, Henry O (1999) Seasonal variation of home range of collared peccary in tropical rain forest
 of French Guiana. J Wildl Manag 63: 546-552
- Keuroghlian A, Eaton DP, Longland WS (2004) Area use by white-lipped and collared peccaries
 (*Tayassu pecari* and *Tayassu tajacu*) in a tropical forest fragment. Biol Conserv 120:411–25

543 Klink CA, Machado RB (2005) Conservation of the Brazilian Cerrado. Conserv Biol 19: 707-713

- 544 Leibold MA, Holyoak M, Mouquet N, Amarasekare P, Chase JM, Hoopes MF, Holt RD, Shurin JB,
- Law R, Tilman D, Loreau M, Gonzalez A (2004) The metacommunity concept: a framework
 for multi-scale community ecology. Ecol Lett 7: 601-613
- 547 Levin SA (1992) The problem of pattern and scale in Ecology. Ecology 73: 1943-1967
- 548 Lidicker Jr WZ (1999) Responses of mammals to habitat edges: a landscape perspective. Landsc
- 549 Ecol 14: 331

550 Lowicki D (2017) Landscape metrics as an indicators of landscape value. Probl Landsc Ecol Vol.

 551
 44: 99-108

- Lü Y, Feng X, Chen L, Fu B (2013) Scaling effects of landscape metrics: a comparison of two
 methods. Phys Geog 34: 40-49
- Lyra-Jorge MC, Ribeiro MC, Ciocheti C, Tambosi LR, Pivello VR (2010) Influence of multi-scale
 landscape structure on the occurrence of carnivorous mammals in a human-modified savanna,
 Brazil. Eur J Wildlife Res 56: 359-368
- Mares MA, Ernest KA, Gettinger RD (1986) Small mammal community structure and composition
 in the Cerrado province of Central Brazil. J Trop Ecol 2: 289-300
- Martensen AC, Ribeiro MC, Banks-Leite C, Prado PI, Metzger JP (2012) Associations of Forest
 Cover, Fragment Area, and Connectivity with Neotropical Understory Bird Species Richness
 and Abundance. Conserv Biol 26: 1100-1111
- Martin AE, Farhig L (2012) Measuring and selecting scales o effect for landscape predictors in
 species-habitat models. Ecol Appl 22: 2277-2292
- Mayor SJ, Schneider DC, Schaefer JA, Mahoney SP (2009) Habitat selection at multiple scales.
 Ecoscience 16: 238-247
- Melo GL, Sponchiado J, Cáceres N, Fahrig L (2017) Testing the habitat amount hypothesis for
 South American small mammals. Biol Conser 209: 304-314
- Michalski F, Peres CA (2007) Disturbance-mediated mammal persistence and abundance-area
 relationships in Amazonian forest fragments. Conserv Biol 21:1626-1640
- 570 Miguet P, Jackson HB, Jackson ND, Martin AE, Fahrig L (2016) What determines the spatial extent
 571 of landscape effects on species? Landsc Ecol 31: 1177–1194
- 572 Miguet P, Fahrig L, Lavigne C (2017) How to quantify a distance-dependent landscape effect on a
 573 biological response. Methods Ecol Evol 8: 1717–1724
- Moraga AD, Martin AE, Fahrig L (2019) The scale of effect of landscape context varies with the
 species' response variable measured. Landsc Ecol 34: 703-715
- 576 Haines-Young R (2009) Land use and biodiversity relationships. Land Use Policy 26: 178–186
- 577 Neel MC, McGarigal K, Cushman SA. (2004) Behavior of class-level landscape metrics across
 578 gradients of class aggregation and area. Landsc Ecol 19: 435–455
- Newbold T, Hudson LN, Hill SLL et al (2015) Global effects of land use on local terrestrial
 biodiversity. Nature 520: 45–50
- 581 Niedballa J, Sollmann R, bin Mohamed A, Bender J (2015) Defining habitat covariates in camera 582 trap based occupancy studies. Sci Rep 5: 17041
- 583 Ochoa-Quintero JM, Gardner TA, Rosa I, de Barros Ferraz SF, Sutherland WJ (2015) Thresholds of
- 584 species loss in Amazonian deforestation frontier landscapes. Conserv Biol 29: 440-451

- 585 Oliveira PS, Marquis RJ (2002) The Cerrados of Brazil: Ecology and Natural History of a
- 586 Neotropical Savanna, Columbia University Press, New York/USA. 373p.
- Pardini R, de Buen, AA, Gardner TA, Prado PI, Metzger JP (2010) Beyond the fragmentation
 threshold hypothesis: Regime shifts in biodiversity across fragmented landscapes. PLoS ONE
 5: e13666
- 590 Pires AS, Lira PK, Fernandez FAS, Schittini GM, Oliveira LC (2002) Frequency of movements of
 591 small mammals among Atlantic Coastal Forest fragments in Brazil. Biol Cons 108: 229-237
- 592 Püttker T, Crouzeilles R, Almeida-Gomes M, Schmoeller M, Maurenza D, Alves-Pinto H, Pardini
- R, Vieira MV, Banks-Leite G, Fonseca CR et al. (2020) Indirect effects of habitat loss via
 habitat fragmentation: Across-taxa analysis of forest-dependent species. Biol Cons 241:
 108368
- Rahbek C (2005) The role of spatial scale and the perception of large-scale species-richness
 patterns. Ecol Lett 8: 224-239
- Rausch LL, Gibbs HK, Schelly I, Brandão A, Morton DC, Filho AC, Strassburg B, Walker N,
 Noojipady P, Barreto P, Meyer D (2019) Soy expansion in Brazil's Cerrado. Conserv Lett 12:
 e12671Redon M, Bergès TC, Luque S (2014) Effects of increasing landscape heterogeneity
 on local plant species richness: how much is enough? Landsc Ecol 29: 773-787
- Regolin AL, Cherem JJ, Graipel ME, Bogoni JA, Ribeiro JW, Vancine MH, Tortato MA, OliveiraSantos LG, Fantacini FM, Luiz MR, de Castilho PV, Ribeiro MC, Cáceres NC (2017) Forest
 cover influences occurrence of mammalian carnivores within Brazilian Atlantic Forest. J
- 605 Mammal 98: 1721–1731
- Rocchini D, Foody GM, Nagendra H, Ricotta C, Anand M, He KS et al. (2013) Uncertainty in
 ecosystem mapping by remote sensing. Comput Geosci 50:128–135.
- Royo AA, Carson WP (2005) The herb community of a tropical forest in central Panamá: dynamics
 and impact of mammalian herbivores. Oecologia 145: 66-75
- Saab V (1999) Importance of spatial scale to habitat use by breeding birds in riparian forests: a
 hierarchical analysis. Ecol Appl 9: 135-151
- 612 Schuster R Römer H, Germain RR (2013) Using multi-scale distribution and movement effects
- along a montane highway to identify optimal crossing locations for a large-bodied mammal
 community. PeerJ 1: e189
- Strahler AH, Woodcock CE, Smith JA (1986) On the nature of models in remote sensing. Remote
 Sens Environ 20: 121–139
- 617 Strassburg BBN, Brooks T, Feltran-Barbieri R, Iribarrem A, Crouzeilles R, Loyola R, Latawiec AE,
- 618 Oliveira Filho FJB, Scaramuzza CAM, Scarano FR, Soares-Filho B, Balmford A (2017)
- 619 Moment of truth for the Cerrado hotspot. Nat Ecol Evol 1: 99

- 620 Teixeira-Santos J, Ribeiro ACdC, Wiig Ø, Pinto NS, Cantanhêde LG, Sena L, et al. (2020)
- Environmental factors influencing the abundance of four species of threatened mammals in
 degraded habitats in the eastern Brazilian Amazon. PLoS ONE 15: e0229459
- Thornton DH, Fletcher RJ (2014) Body size and spatial scales in avian response to landscapes: a
 meta-analysis. Ecography 37: 454-463
- Vanderwal J, Falconi L, Januchowski S, Shoo L, Storlie C (2014) Package 'SDMTools'. Species
 Distribution ModellingTools: Tools for processing data associated with species distribution
 modelling exercises. R-package version 1.1.12
- Villard M-A, Metzger JP (2014) Beyond the fragmentation debate: a conceptual model to predict
 when habitat configuration really matters. J Appl Ecol 51: 309-318
- Woodcock CE, Strahler AH (1987) The factor of scale in remote sensing. Remote Sens Environ 21:
 311–332
- 632 Young HS, McCauley DJ, Helgen KM, Goheen JR, Otárola-Castillo E, Palmer TM, Pringle RM,
- Young TP, Dirzo R (2013) Effects of mammalian herbivore declines on plant communities:
 observations and experiments in an African savanna. J Ecol 101: 1030-1041
- Zimbres B, Peres CA, Penido G, Machado RB (2018) Thresholds of riparian forest use by terrestrial
 mammals in a fragmented Amazonian deforestation frontier. Biodivers Conserv 27: 28152836

638639 Table 1. Ecological traits of the studied mammal species

Studied species	Adult body mass (kg)	Home range (ha)	Dispersal range (m)	Habitat use	References
Azara's agouti	2.9	1.34 - 2.45	100 - 500	Forest/ edges	Aliaga-Rossel et al. 2008 ; Jansen et al. 2012 ; Cid et al. 2013
Brown brocket deer	21	2.7 - 348	750 - 1 000	Forest/ open area	Duarte et al. 2012 ; Pires et al. 2018
Collared peccary	22	24 - 800	500 - 1 000	Forest/ edges	Bodmer 1991; Fragoso 1998; Judas and Henry 1999; Keuroghlian et al 2004
South American tapir	225	0.1 - 100	> 3 000	Forest/ open area	Bodmer 1989; Emmons and Feer 1997; Pires et al. 2018

Table 2. Results of occupancy models predicting the occurrence of four species of terrestrial mammals across buffer size (from 500 to 10 000m)

641 considering three threshold of percent tree cover per pixel (25, 50 and 75%). This table addressed the selected range of models (and thus buffer size)

that have been selected from AIC model selection (See Electronic Appendix Table S1 for details). Model structure refers to how occupancy probability

643 Ψ at the camera trap i and the detection probability p were implemented. Analyses were carried out separately for each species and landscape metric

644 including edge density, ED, patch density, PD, and the percentage of forest cover, FC. When several models showed similar statistical support (i.e.

645 ΔAIC < 2), estimate (β) ± SE was extracted for the model with the lowest AIC indicated by the * near the buffer size. Significant effects are indicated 646 in bold.

- 647
- 648

			Percent of tree cover per pixel												
Landscape metric	Model structure	Species		25%			50%								
			Buffer size	$\beta\pm SE$	Р	Buffer size	$\beta\pm SE$	Р	Buffer size	$\beta\pm SE$	Р				
	p (weather+visibility) Ψ (ED)	Azara's agouti				500	0.325 ± 0.185	0.079	500*-1500	0.517 ± 0.267	0.053				
ED	p (visibility) Ψ (ED)	Brown brocket deer													
	p (visibility) Ψ (ED)	Collared peccary													
	p (weather+visibility) Ψ (ED)	South American tapir													
						1500-2000*-									
	p (weather+visibility) Ψ (PD)	Azara's agouti				3000	-1.030 ± 0.675	0.128							
PD	p (visibility) Ψ (PD)	Brown brocket deer							750	0.643 ± 0.305	0.035				
	p (visibility) Ψ (PD)	Collared peccary													
	p (weather+visibility) Ψ (PD)	South American tapir													
	p (weather+visibility) Ψ (FC)	Azara's agouti	500*-1500	4.540 ± 2.500	0.070	500*-2000	5.050 ± 2.890	0.081	500-1500*-2000	9.000 ± 4.450	0.043				
FC	p (visibility) Ψ (FC)	Brown brocket deer													
	p (visibility) Ψ (FC)	Collared peccary	500*-1000	2.690 ± 1.400	0.029	500*-1000	3.190 ± 1.540	0.039	500*-750	4.030 ± 1.890	0.033				
	p (weather+visibility) Ψ (FC)	South American tapir													

649	Figure captions
650	
651	Fig. 1 Map of the study site, Bodoquena, state of Mato Grosso do Sul, Brazil (a) and forest cover
652	including the location of camera traps (b). Camera traps selected to run occupancy models (see
653	methods) are highlighted in red.
654	
655	Fig. 2 Example of forest cover assessed for different threshold of "forest" definition (minimum tree
656	cover per pixel of 25, 50 and 75%) around one camera trap at two buffers sizes, 3000 m (blue disc)
657	and 500 m (red disc). Forest pixels are represented in green.
658	
659	Fig. 3 Comparison of model performance for the effect of landscape metric for forests defined with
660	a 75% of tree cover per pixel. The grey triangle is the null model in which occupancy probability is
661	fixed as 1. The scale of effect is detected from spatial extent (i.e. the range of buffer sizes) for which
662	$\Delta AIC \leq 2$ included the smallest AIC without the null model (black dots).













Electronic appendices





patch density (a), edge density (b), forest cover (c) and their standard deviation (SD) (d, e, f)

assessed around our camera traps at three different percent tree cover per pixel (25, 50 and 75%)

(See Table S1 for the statistics of the relationships in a, b and c).

- Table S1. Shape of the relationship between buffer size (x) and landscape metrics (y) including
- patch density PD, edge density ED and the percent forest cover (FC) assessed at three percent tree
 cover per pixel (25, 50 and 75%). *** indicates that P value < 0.001.

Landscape metric	Percent tree cover per pixel	Relationship	Equation	R ²
	25%	Power***	y=3.082x ^{-0.226}	0.94
Patch Density	50%	Power***	y=2.520x ^{-0.211}	0.87
	75%	Power***	y=2.372x ^{-0.189}	0.87
	25%	Power***	y=4.868x ^{-0.157}	0.96
Edge Density	50%	Power***	y=4.196x ^{-0.145}	0.92
	75%	Power***	y=3.823x ^{-0.108}	0.86
	25%	Logarithm***	y=0.917ln(x)+44.058	0.81
Forest Cover	50%	Logarithm***	y=0.915ln(x)+37.312	0.86
	75%	Logarithm***	y=0.804ln(x)+29.143	0.89

691 Table S2. Performance of occupancy models predicting the occurrence of four species of terrestrial mammals (Azara's agouti Dasyprocta azarae, collared peccary Pecari tajacu, brown brocket deer 692 Mazama gouazoubira, and South American tapir Tapirus terrestris,) in the Brazilian Cerrado 693 694 hotspot across buffer size (from 500 to 10000 m) considering 75% as the threshold of tree cover. Model structure mentions how occupancy probability Ψ at the camera trap i and the detection 695 probability p were implemented. Analyses were carried out separately for each species and 696 697 landscape metric including edge density, ED, patch density, PD, and the percentage of forest cover, 698 FC. Buffer size is indicated in m. AIC is the Akaike criterion. Models of similar statistical supports 699 with $\Delta AIC < 2$ from the minimum model are in bold.

700

Azara's agouti					Collared peccary				Brown brocket deer				South American tapir				
Landscape metric	Buffer size	Model structure	AIC	ΔΑΙC	Buffer size	Model structure	AIC	ΔAIC	Buffer size	Model structure	AIC	ΔΑΙC	Buffer size	Model structure	AIC	ΔAIC	
	Null	p (weather + visibility) Ψ(1)	152.43	6.18	Null	p (visibility) Ψ(1)	173.54	0.07	Null	p (visibility) Ψ(1)	145.56	0.16	Null	p (weather + visibility) Ψ(1)	231.04	0.00	
	500	p (weather + visibility) Ψ(ED)	146.25	0.00	500	p (visibility) Ψ(ED)	173.47	0.00	500	p (visibility) Ψ(ED)	146.89	1.49	500	p (weather + visibility) Ψ(ED)	231.32	0.28	
	750	p (weather + visibility) Ψ(ED)	148.06	1.80	750	p (visibility) Ψ(ED)	173.50	0.02	750	p (visibility) Ψ(ED)	145.73	0.33	750	p (weather + visibility) Ψ(ED)	231.81	0.77	
	1000	p (weather + visibility) Ψ(ED)	147.39	1.13	1000	p (visibility) Ψ(ED)	174.57	1.10	1000	p (visibility) Ψ(ED)	145.40	0.00	1000	p (weather + visibility) Ψ(ED)	232.27	1.23	
ED	1500	p (weather + visibility) Ψ(ED)	148.08	1.83	1500	p (visibility) Ψ(ED)	174.77	1.30	1500	p (visibility) Ψ(ED)	146.63	1.23	1500	p (weather + visibility) Ψ(ED)	232.93	1.89	
	2000	p (weather + visibility) Ψ(ED)	149.63	3.38	2000	p (visibility) Ψ(ED)	174.61	1.13	2000	p (visibility) Ψ(ED)	147.12	1.72	2000	p (weather + visibility) Ψ(ED)	233.04	2.00	
	3000	p (weather + visibility) Ψ(ED)	152.34	6.09	3000	p (visibility) Ψ(ED)	174.91	1.44	3000	p (visibility) Ψ(ED)	147.10	1.70	3000	p (weather + visibility) Ψ(ED)	232.99	1.95	
	5000	p (weather + visibility) Ψ(ED)	154.12	7.87	5000	p (visibility) Ψ(ED)	175.22	1.75	5000	p (visibility) Ψ(ED)	146.38	0.98	5000	p (weather + visibility) Ψ(ED)	233.04	2.00	
	10000	p (weather + visibility) Ψ(ED)	154.13	7.88	10000	p (visibility) Ψ(ED)	175.28	1.81	10000	p (visibility) Ψ(ED)	145.88	0.47	10000	p (weather + visibility) Ψ(ED)	233.04	2.00	
	Null	p (weather +	152.43	0.00	Null	p (visibility)	173 54	0.00	Null	p (visibility)	145 56	613	Null	p (weather +	231.04	0.00	
	500	visibility) Ψ(1) p (weather +	152.45	0.54	500	Ψ(1) p (visibility)	175.04	1.05	500	Ψ(1) p (visibility)	143.50	4.35	500	visibility) Ψ(1) p (weather +	231.04	1 59	
	750	visibility) Ψ(PD) p (weather +	154.40	1.07	750	Ψ(PD) p (visibility)	175.49	1.95	750	Ψ(PD) p (visibility)	145.76	4.55	750	visibility) Ψ(PD) p (weather +	232.03	1.59	
	1000	visibility) Ψ(PD) p (weather +	154.40	2.00	1000	Ψ(PD) p (visibility)	175.40	1.61	1000	Ψ(PD) p (visibility)	143.41	3.00	1000	visibility) Ψ(PD) p (weather +	232.00	1.04	
PD	1500	visibility) Ψ(PD) p (weather +	152.02	1.50	1500	Ψ(PD) p (visibility)	175.40	1.01	1500	Ψ(PD) p (visibility)	145.02	6.40	1500	visibility) Ψ(PD) p (weather +	232.90	1.04	
ID	2000	visibility) Ψ(PD) p (weather +	155.95	1.50	2000	Ψ(PD) p (visibility)	175.50	1.94	2000	Ψ(PD) p (visibility)	145.52	6.12	2000	visibility) Ψ(PD) p (weather +	232.07	2.00	
	2000	visibility) Ψ(PD) p (weather +	154.15	1./1	2000	Ψ(PD) p (visibility)	175.52	1.98	2000	Ψ(PD) p (visibility)	145.54	6.12	2000	visibility) Ψ(PD) p (weather +	255.04	2.00	
	3000	visibility) Ψ(PD) p (weather +	154.12	1.69	3000	Ψ(PD) p (visibility)	1/5.48	1.93	3000	Ψ(PD) p (visibility)	144.55	5.11	3000	visibility) Ψ(PD) p (weather +	233.04	1.99	
	5000	visibility) Ψ(PD) p (weather +	154.43	2.00	5000	Ψ(PD) p (visibility)	175.54	2.00	5000	Ψ(PD) p (visibility)	144.66	5.24	5000	visibility) Ψ(PD) p (weather +	232.85	1.81	
	10000	visibility) Ψ(PD)	154.43	2.00	10000	Ψ(PD)	174.55	1.01	10000	$\Psi(PD)$	145.46	6.03	10000	visibility) Ψ(PD)	231.77	0.73	
	Null	p (weather + visibility) Ψ(1)	152.43	3.07	Null	p (visibility) Ψ(1)	173.54	4.84	Null	p (visibility) Ψ(1)	145.56	0.00	Null	p (weather + visibility) $\Psi(1)$	231.04	0.89	
	500	p (weather + visibility) Ψ(FC)	150.83	1.46	500	p (visibility) Ψ(FC)	168.70	0.00	500	p (visibility) Ψ(FC)	146.33	0.77	500	p (weather + visibility) Ψ(FC)	230.15	0.00	
	750	p (weather + visibility) Ψ(FC)	150.40	1.03	750	p (visibility) Ψ(FC)	170.20	1.49	750	p (visibility) Ψ(FC)	146.16	0.60	750	p (weather + visibility) Ψ(FC)	231.54	1.39	
	1000	p (weather + visibility) Ψ(FC)	149.66	0.30	1000	p (visibility) Ψ(FC)	171.50	2.80	1000	p (visibility) Ψ(FC)	146.13	0.57	1000	p (weather + visibility) Ψ(FC)	231.71	1.56	
FC	1500	p (weather + visibility) Ψ(FC)	149.37	0.00	1500	p (visibility) Ψ(FC)	172.83	4.13	1500	p (visibility) Ψ(FC)	146.21	0.65	1500	p (weather + visibility) Ψ(FC)	232.44	2.28	
	2000	p (weather + visibility) Ψ(FC)	149.87	0.50	2000	p (visibility) Ψ(FC)	173.92	5.22	2000	p (visibility) Ψ(FC)	146.97	1.41	2000	p (weather + visibility) Ψ(FC)	232.88	2.73	
	3000	p (weather + visibility) Ψ(FC)	152.37	3.00	3000	p (visibility) Ψ(FC)	174.83	6.13	3000	p (visibility) Ψ(FC)	147.06	1.51	3000	p (weather + visibility) Ψ(FC)	233.02	2.87	
	5000	p (weather + visibility) Ψ(FC)	153.60	4.23	5000	p (visibility) Ψ(FC)	175.41	6.71	5000	p (visibility) Ψ(FC)	147.29	1.73	5000	p (weather + visibility) Ψ(FC)	232.94	2.79	
	10000	p (weather + visibility) Ψ(FC)	153.97	4.61	10000	p (visibility) Ψ(FC)	175.52	6.82	10000	p (visibility) Ψ(FC)	147.16	1.60	10000	p (weather + visibility) Ψ(FC)	232.87	2.72	

702 Table S3. Performance of occupancy models predicting the occurrence of four species of terrestrial 703 mammals (Azara's agouti Dasyprocta azarae, collared peccary Pecari tajacu, brown brocket deer 704 Mazama gouazoubira, and South American tapir Tapirus terrestris,) in the Brazilian Cerrado 705 hotspot across buffer size (from 500 to 10000 m) considering 50% as the threshold of tree cover. Model structure mentions how occupancy probability Ψ at the camera trap i and the detection 706 probability p were implemented. Analyses were carried out separately for each species and 707 708 landscape metric including edge density, ED, patch density, PD, and the percentage of forest cover, 709 FC. Buffer size is indicated in m. AIC is the Akaike criterion. Models of similar statistical supports 710 with $\Delta AIC < 2$ from the minimum model are in bold.

711

Azara's agou	ıti			Collare	d peccary	Brown	brocket deer			South American tapir						
Landscape metric	Buffer size	Model structure	AIC	ΔΑΙΟ	Buffer size	Model structure	AIC	ΔΑΙΟ	Buffer size	Model structure	AIC	ΔΑΙΟ	Buffer size	Model structure	AIC	ΔΑΙC
	Null	p (weather + visibility) Ψ(1)	152.43	2.89	Null	p (visibility) Ψ(1)	173.54	0.00	Null	p (visibility) Ψ(1)	145.56	1.43	Null	p (weather + visibility) Ψ(1)	231.04	0.00
	500	p (weather + visibility) Ψ(ED)	149.54	0.00	500	p (visibility) Ψ(ED)	175.12	1.58	500	p (visibility) Ψ(ED)	145.48	1.35	500	p (weather + visibility) Ψ(ED)	232.32	1.27
	750	p (weather + visibility) Ψ(ED) p (weather +	151.63	2.08	750	p (visibility) Ψ(ED) p (visibility)	175.08	1.54	750	p (visibility) Ψ(ED) p (visibility)	144.80	0.67	750	p (weather + visibility) Ψ(ED) p (weather +	232.29	1.25
	1000	p (weather + visibility) Ψ(ED) p (weather +	152.15	2.61	1000	$\Psi(ED)$ p (visibility)	175.53	1.98	1000	$\Psi(ED)$ p (visibility)	144.50	0.37	1000	p (weather + visibility) Ψ(ED) p (weather +	232.89	1.85
ED	1500	visibility) Ψ(ED) p (weather +	151.83	2.29	1500	Ψ(ED) p (visibility)	175.51	1.97	1500	Ψ(ED) p (visibility)	145.58	1.44	1500	visibility) Ψ(ED) p (weather +	233.00	1.95
	2000	visibility) Ψ(ED) p (weather +	152.35	2.80	2000	Ψ(ED) p (visibility)	175.30	1.76	2000	Ψ(ED) p (visibility)	146.10	1.97	2000	visibility) Ψ(ED) p (weather +	233.04	2.00
	3000	visibility) Ψ(ED) p (weather +	153.49	3.95	3000	Ψ(ED) p (visibility)	175.43	1.89	3000	Ψ(ED) p (visibility)	145.26	1.13	3000	visibility) Ψ(ED) p (weather +	232.96	1.92
	10000	p (weather + visibility) $\Psi(ED)$	154.09	4.55	10000	Ψ(ED) p (visibility) Ψ(ED)	175.43	1.89	10000	Ψ(ED) p (visibility) Ψ(ED)	144.87	0.74	10000	p (weather + visibility) Ψ(ED)	233.00	1.96
	10000	visionity) 1(ED)	134.23	4.71	10000	I(LD)	175.18	1.04	10000	I(LD)	144.15	0.00	10000	visionity) 1(ED)	252.41	1.57
DD	Null	p (weather + visibility) Ψ(1)	152.43	2.05	Null	p (visibility) $\Psi(1)$	173.54	0.00	Null	p (visibility) $\Psi(1)$	145.56	1.83	Null	p (weather + visibility) $\Psi(1)$	231.04	0.00
	500	p (weather + visibility) $\Psi(PD)$ p (weather +	154.27	3.88	500	Ψ(PD) n (visibility)	175.43	1.88	500	Ψ(PD) n (visibility)	143.73	0.00	500	p (weather + visibility) $\Psi(PD)$ p (weather +	233.03	1.99
	750	visibility) Ψ(PD) p (weather +	153.96	3.58	750	Ψ(PD) p (visibility)	174.24	0.70	750	Ψ(PD) p (visibility)	144.95	1.22	750	visibility) Ψ(PD) p (weather +	233.01	1.97
	1000	visibility) Ψ(PD) p (weather +	152.87	2.49	1000	Ψ(PD) p (visibility)	174.45	0.91	1000	Ψ(PD) p (visibility)	146.62	2.89	1000	visibility) Ψ(PD) p (weather +	232.17	1.12
10	1500	visibility) Ψ(PD) p (weather +	152.24	1.85	1500	Ψ(PD) p (visibility)	174.35	0.80	1500	Ψ(PD) p (visibility)	147.06	3.33	1500	visibility) Ψ(PD) p (weather +	232.55	1.51
	2000	visibility) Ψ(PD) p (weather +	150.38	0.00	2000	Ψ(PD) p (visibility) W(PD)	175.07	1.53	2000	Ψ(PD) p (visibility)	146.83	3.10	2000	visibility) Ψ(PD) p (weather +	233.04	2.00
	5000	p (weather + visibility) $\Psi(PD)$	151.52	3.01	5000	Ψ(PD) p (visibility) Ψ(PD)	175.30	1.81	5000	Ψ(PD) p (visibility) Ψ(PD)	146.92	2.55	5000	p (weather + visibility) Ψ(PD)	232.87	1.65
	10000	p (weather + visibility) Ψ(PD)	153.43	3.05	10000	p (visibility) Ψ(PD)	175.54	2.00	10000	p (visibility) Ψ(PD)	147.44	3.71	10000	p (weather + visibility) Ψ(PD)	231.83	0.79
		n (weather +				n (visibility)				n (visibility)				n (weather +		
	Null	visibility) $\Psi(1)$ p (weather +	152.43	3.28	Null	$\Psi(1)$	173.54	3.45	Null	$\Psi(1)$ p (visibility)	145.56	0.00	Null	visibility) $\Psi(1)$ p (weather +	231.04	0.41
	500	visibility) $\Psi(FC)$ p (weather +	149.15	0.00	500	Ψ(FC) p (visibility)	170.09	0.00	500	Ψ(FC) p (visibility)	145.80	0.25	500	visibility) $\Psi(FC)$ p (weather +	230.63	0.00
	750	visibility) Ψ(FC) p (weather +	150.26	1.11	750	Ψ(FC) p (visibility)	170.80	0.71	750	Ψ(FC) p (visibility)	145.68	0.12	750	visibility) Ψ(FC) p (weather +	231.54	0.91
FC	1000	visibility) Ψ(FC) p (weather +	149.31	0.16	1000	Ψ(FC) p (visibility)	171.95	1.86	1000	Ψ(FC) p (visibility)	145.82	0.26	1000	visibility) Ψ(FC) p (weather +	231.88	1.25
10	1500	visibility) Ψ(FC) p (weather +	149.44	0.29	1500	Ψ(FC) p (visibility)	173.02	2.93	1500	Ψ(FC) p (visibility)	145.79	0.23	1500	visibility) Ψ(FC) p (weather +	232.60	1.97
	2000	visibility) Ψ(FC) p (weather +	150.31	1.16	2000	Ψ(FC) p (visibility)	173.87	3.78	2000	Ψ(FC) p (visibility)	146.62	1.07	2000	visibility) Ψ(FC) p (weather +	232.95	2.31
	5000	p (weather +	152.47	3.32	5000	r(FC) p (visibility)	175.20	4.69	5000	r(FC) p (visibility)	146.72	1.17	5000	p (weather +	233.02	2.38
	10000	p (weather + visibility) $\Psi(FC)$	155.05	4.85	10000	p (visibility) Ψ(FC)	175.50	5.42	10000	p (visibility) Ψ(FC)	140.99	1.44	10000	p (weather + visibility) $\Psi(FC)$	232.95	2.32

712

Table S4. Performance of occupancy models predicting the occurrence of four species of terrestrial mammals (Azara's agouti Dasyprocta azarae, collared peccary Pecari tajacu, brown brocket deer Mazama gouazoubira, and South American tapir Tapirus terrestris,) in the Brazilian Cerrado hotspot across buffer size (from 500 to 10000 m) considering 25% as the threshold of tree cover. Model structure mentions how occupancy probability Ψ at the camera trap i and the detection probability p were implemented. Analyses were carried out separately for each species and landscape metric including edge density, ED, patch density, PD, and the percentage of forest cover, FC. Buffer size is indicated in m. AIC is the Akaike criterion. Models of similar statistical supports with $\Delta AIC < 2$ from the minimum model are in bold.

Azara's agou	ıti			Collare	d peccary	Brown	brocket deer			South American tapir						
Landscape	Buffer	Model structure	AIC	ΔAIC	Buffer size	Model structure	AIC	ΔΑΙΟ	Buffer	Model structure	AIC	ΔΑΙΟ	Buffer	Model structure	AIC	ΔΑΙΟ
ED	Null	p (weather + visibility) Ψ(1)	152.43	0.87	Null	p (visibility) Ψ(1)	173.54	0.00	Null	p (visibility) Ψ(1)	145.56	1.92	Null	p (weather + visibility) Ψ(1)	231.04	0.00
	500	p (weather + visibility) Ψ(ED)	151.57	0.00	500	p (visibility) Ψ(ED)	175.50	1.96	500	p (visibility) Ψ(ED)	145.42	1.78	500	p (weather + visibility) Ψ(ED)	232.87	1.83
	750	p (weather + visibility) Ψ(ED) p (weather +	152.81	1.24	750	$\Psi(ED)$ p (visibility)	175.54	1.99	750	p (visibility) Ψ(ED) p (visibility)	143.64	0.00	750	p (weather + visibility) Ψ(ED) p (weather +	232.79	1.75
	1000	visibility) Ψ(ED) p (weather +	153.75	2.18	1000	Ψ(ED) p (visibility)	175.36	1.82	1000	Ψ(ED) p (visibility)	144.16	0.53	1000	visibility) Ψ(ED) p (weather +	233.04	2.00
ED	1500	visibility) Ψ(ED) p (weather +	153.89	2.32	1500	Ψ(ED) p (visibility)	175.39	1.85	1500	Ψ(ED) p (visibility)	146.32	2.68	1500	visibility) Ψ(ED) p (weather +	232.68	1.64
	2000	visibility) Ψ(ED) p (weather +	153.95	2.38	2000	Ψ(ED) p (visibility)	175.49	1.95	2000	Ψ(ED) p (visibility)	147.01	3.37	2000	visibility) Ψ(ED) p (weather +	232.99	1.95
	3000	visibility) Ψ(ED) p (weather + visibility) Ψ(ED)	154.27	2.70	3000	Ψ(ED) p (visibility) Ψ(ED)	175.31	1.77	3000	Ψ(ED) p (visibility) Ψ(ED)	146.82	3.19	3000	visibility) Ψ(ED) p (weather + visibility) Ψ(ED)	232.98	1.94
	10000	p (weather + visibility) $\Psi(ED)$	154.41	2.85	10000	Ψ(ED) p (visibility) Ψ(ED)	175.40	1.92	10000	P(ED) p (visibility) Ψ(ED)	146.21	2.58	10000	p (weather + visibility) $\Psi(ED)$	232.92	0.63
	10000	(15)011(22)	101110	2.00	10000	1(22)	175.11	107	10000	(LD)	110.25	2.00	10000	(LD)	201100	0100
	Null	p (weather + visibility) Ψ(1)	152.43	0.00	Null	p (visibility) Ψ(1)	173.54	0.56	Null	p (visibility) Ψ(1)	145.56	0.00	Null	p (weather + visibility) $\Psi(1)$	231.04	0.00
	500	p (weather + visibility) Ψ(PD)	154.21	1.78	500	p (visibility) Ψ(PD)	175.17	2.18	500	p (visibility) Ψ(PD)	147.01	1.46	500	p (weather + visibility) Ψ(PD)	233.03	1.99
	750	visibility) $\Psi(PD)$	154.39	1.95	750	$\Psi(PD)$	173.04	0.06	750	$\Psi(PD)$	147.30	1.74	750	visibility) $\Psi(PD)$	232.69	1.65
	1000	visibility) Ψ(PD) p (weather +	154.09	1.65	1000	Ψ(PD) p (visibility)	174.04	1.06	1000	Ψ(PD) p (visibility)	147.54	1.99	1000	p (weather + visibility) Ψ(PD) p (weather +	233.39	2.35
PD	1500	visibility) Ψ(PD) p (weather +	152.91	0.48	1500	Ψ(PD) p (visibility)	172.98	0.00	1500	Ψ(PD) p (visibility)	147.53	1.97	1500	visibility) Ψ(PD) p (weather +	233.04	2.00
	2000	visibility) Ψ(PD) p (weather +	152.71	0.28	2000	Ψ(PD) p (visibility)	173.10	0.12	2000	Ψ(PD) p (visibility)	147.55	2.00	2000	visibility) Ψ(PD) p (weather +	232.52	1.48
	3000	p (weather +	153.32	0.89	3000	Ψ(PD) p (visibility) W(PD)	174.65	1.67	3000	Ψ(PD) p (visibility)	147.56	2.00	3000	p (weather +	231.96	0.92
	10000	p (weather + visibility) $\Psi(PD)$	153.62	1.19	10000	Ψ(PD) p (visibility) Ψ(PD)	175.46	2.47	10000	P(PD) p (visibility) Ψ(PD)	147.41	1.38	10000	p (weather + visibility) $\Psi(PD)$	231.40	1.43
	10000	(15)011(12)	155.62	103	10000	1(12)	175.10	2,	10000	1(12)	110.71	100	10000	(15)0111(12)	232.17	
	Null	p (weather + visibility) $\Psi(1)$	152.43	3.88	Null	p (visibility) Ψ(1)	173.54	2.41	Null	p (visibility) $\Psi(1)$	145.56	0.00	Null	p (weather + visibility) $\Psi(1)$	231.04	0.00
	500	p (weather + visibility) Ψ(FC) p (weather +	148.55	0.00	500	$\Psi(FC)$	171.13	0.00	500	$\Psi(FC)$	145.76	0.21	500	p (weather + visibility) Ψ(FC) p (weather +	231.49	0.44
	750	visibility) Ψ(FC) p (weather +	150.32	1.76	750	Ψ(FC) p (visibility)	171.57	0.44	750	Ψ(FC) p (visibility)	145.58	0.03	750	visibility) $\Psi(FC)$ p (weather +	232.01	0.97
FC	1000	visibility) Ψ(FC) p (weather +	149.82	1.27	1000	Ψ(FC) p (visibility)	172.33	1.20	1000	Ψ(FC) p (visibility)	145.62	0.06	1000	visibility) Ψ(FC) p (weather +	232.26	1.22
re	1500	visibility) Ψ(FC) p (weather +	150.08	1.53	1500	Ψ(FC) p (visibility)	173.21	2.08	1500	Ψ(FC) p (visibility)	145.58	0.02	1500	visibility) Ψ(FC) p (weather +	232.88	1.83
	2000	visibility) Ψ(FC) p (weather +	150.91	2.35	2000	Ψ(FC) p (visibility)	173.94	2.81	2000	Ψ(FC) p (visibility)	146.44	0.88	2000	visibility) Ψ(FC) p (weather +	233.03	1.98
	3000	visibility) Ψ(FC) p (weather +	152.55	4.00	3000	Ψ(FC) p (visibility)	174.82	3.69	3000	Ψ(FC) p (visibility)	146.45	0.89	3000	visibility) Ψ(FC) p (weather +	233.04	1.99
	5000	visibility) Ψ(FC) p (weather + visibility) Ψ(FC)	153.57	5.02	5000	Y(FC) p (visibility)	175.41	4.28	5000	P(FC) p (visibility)	146.73	1.17	5000	visibility) Ψ(FC) p (weather + visibility) Ψ(FC)	232.91	1.87
	10000	visibility) $\Upsilon(FC)$	155.98	3.42	10000	1(FC)	1/3.31	4.38	10000	T(FC)	140.84	1.28	10000	visibility) $\tau(rC)$	232.83	1.81