

# 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-03131932v1

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

3

- 4 Amiot Christophe <sup>a g</sup>, Santos Cavalcante Cyntia <sup>a e j</sup>, Arvor Damien <sup>b</sup>, Bellón Beatriz <sup>a g k</sup>, Fritz
- 5 Hervé<sup>c g i</sup>, Harmange Clément <sup>a g</sup>, Holland Jeffrey D. <sup>d</sup>, Melo Isabel <sup>e</sup>, Metzger Jean Paul <sup>f</sup>, Renaud
- 6 Pierre-Cyril <sup>a g</sup>, Roque Fabio de Oliveira <sup>e h</sup>, Souza Franco Leandro <sup>e</sup>, Pays Olivier <sup>a g \*</sup>

7

- 8 <sup>a</sup> 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.
- d Department of Entomology, Purdue University, West Lafayette, Indiana, 47907 USA.
- 13 <sup>e</sup> Bioscience Institute, Federal University of Mato Grosso do Sul, Cidade Universitária, 79060-300
- 14 Campo Grande, MS, Brazil
- 15 <sup>f</sup> Universidade 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 <sup>i</sup> Sustainability Research Unit, Nelson Mandela University, Port Elizabeth, South Africa
- 21 <sup>j</sup> Wetlands International Brazil, Campo Grande, Mato Grosso do Sul, Brazil
- 22 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.

- 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	
45	

## 46 **Abstract**

63

64

65

66

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 four species of terrestrial mammal herbivores in the Brazilian savannas and tested whether those 50 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

effects. Thus, if we want to properly identify scales of effect for multiple species, it is necessary not

only to adopt a multi-scale approach, but also to use multiple definitions of habitat, considering

particularities of how each species interact with their environment.

## Introduction

68	
----	--

Identifying the spatial scale at which species respond to landscape structure, or the scale of effect (Jackson and Fahrig 2012) has become an important scientific challenge when investigating species-environment relationships (Fahrig et al. 2011; Redon et al. 2014; Miguet et al. 2016, 2017). 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 change on species (Levin 1992). This is particularly relevant in the understanding of mechanisms underlying the erosion of biodiversity, especially the mechanisms for which human activities shape land use and land cover changes (Newbold et al. 2015; Püttker et al. 2020). Knowing the scale at which species respond to landscapes should help decision-makers in designing management plan to maintain and restore biodiversity and their habitats (Haines-Young 2009).

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

Without knowing *a priori* what is the appropriate scale of study, landscape ecologists commonly assess landscape variables at multiple scales to select the scale that yields the best species—landscape relationship (e.g. Brennan et al. 2002; Holland et al. 2004; Holland and Yang 2016; Melo et al. 2017; Huais 2018). Different methods or statistical analyses have been proposed in the literature to test for the effect of landscape features on species attributes (e.g., richness, abundance, composition) at different spatial scales and to extract their goodness of fit through an 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

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.

Biological/ecological traits of species including for instance body size, dispersal distance, home range and habitat specialisation may affect the scale of effect (Miguet et al 2016). Those traits are particularly relevant when they are indicators of animal mobility, a key biological feature which determines the relationship between species attributes and their surrounding space (Bowman et al. 2002). Although the empirical studies are not all in agreement, we might expect that the scale of effect would be larger for larger-bodied species because they are more mobile with higher dispersal distance and home range than smaller-bodied species (Jackson and Fahrig 2012; Miguet et al 2016). Thus, when investigating the effect of landscape features on species occurrence, a general guideline for empirical studies is to consider a radius of a landscape to be 4-9 times the median dispersal distance of 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 spatial scale at which species responded to landscape variables. Finally, the scale of effect may differ between specialist and generalist species of similar body-size although this literature is sparse and not in agreement on whether habitat specialists would exhibit a larger or smaller scale of effect (Boscolo and Metzger 2009; Chaplin-Kramer et al. 2013). Comparisons of habitat generalists and 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 configuration, could also affect the scale of effect. Particularly, the type of maps of land use and land cover that are used as an input to derive patch-based landscape metrics may influence the assessments of scale of effect. This issue is especially relevant in tropical areas dominated by vegetation gradients (ranging from herbaceous layers with sparse trees to shrubland and forested areas, including degraded forests or regenerating forests) where the definition of landscape units can be arbitrary, and low thematic resolution in land cover maps (e.g., a small number of classes) may lead to an overly simplified representation of reality that may not adequately represent how species 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 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 25% of tree cover to map forest cover, Defries et al. (2001) defined forest cover as greater than 60% 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 subtropical zones of Africa, Australia and South America, Hirota et al. (2011) reported that percent tree cover in savannas was about 20% and contrasted with 80% in forested areas. While mammal species might respond differently to forest cover (Royo and Carson 2005; Young et al. 2013; Chamaillé-Jammes et al. 2016; Ferreguetti et al. 2017; Zimbres et al. 2018), the question whether the percent tree cover used to map forest cover affects the scale of effect of landscape metrics on the occurrence of species has never been studied. This might be critical when modelling the presence of forest dwelling species in complex mosaic of vegetation types such as in Brazil's Cerrado including 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 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 terrestris*, using camera traps in the Brazilian Cerrado. First, we studied how the occurrence of those species responded to three landscape metrics commonly used to describe landscape 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). These metrics have also been useful to explain mammal community distribution in Brazil (Mares et 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 detection of scale of effect of the landscape features on the occurrence of species. Finally, as the 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 biological/ecological traits.

We ran occupancy models for each species at 8 spatial extents from 0.5 to  $10 \, \mathrm{km}$  using 3 different maps of forest cover. We extracted the Akaike Information Criterion (AIC) for each model and examined whether species exhibited scale of effects. We hypothesized that mobility largely determines the relationship between landscape features and occurrence of the studied species. We 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 tapir should be larger than the brown brocket deer and the collared peccary, with the smallest scale of effect in the Azara's agouti. Regarding forest definition (that shaped both land use and land cover composition and spatial configuration), we predicted a stronger or clearer scale of effect in the two forest-dwelling species, the Azara's agouti and collared peccary, when considering high tree cover (e.g., 75%) compared to medium-low tree cover (50 – 25%).

# Methodology

172

204

205

173	
174	Study site
175	The Bodoquena Plateau region (20°25'29.28" to 21°44'19.72" S and 56°52'24.46"56° to 17'23.36"
176	W) is located in the state of Mato Grosso do Sul, Brazil (Fig. 1a). It is characterized by a mountain
177	chain (altitude $450-800\mathrm{m}$ ) with a tropical climate (annual temperature varies between $20\mathrm{and}$
178	22°C and rainfall between 1300 and 1700 mm). The study area is within the range of the Cerrado
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
189	project's products (Hansen et al. 2013). We used the "2000 percent tree cover" (TC) product, which
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

limited this to three variables in our study as (1) they are commonly used in the literature and (2)

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 not significantly correlated.

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

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)

Data from mammal species

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

Occupancy models

Creating large buffers around the locations of the 193 camera traps led to spatial overlaps and thus pseudoreplication. To tackle this issue, we used the Focus program (Holland et al. 2004) to conduct 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 combination of 57 independent locations allowing to increase buffer size up to 2 km (Fig. 1b). As 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 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.

To investigate the effect of the landscape features (measured for the 8 buffer sizes and 3 percent threshold of tree cover) on the occurrence of the four mammal species, we run occupancy models using the *unmarked* R package (Fiske and Chandler 2011). For each of the 57 camera traps and focal species, we implemented a data set of 40 sampled days in which we indicated when the 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. 2015). Following this method, 13 camera traps (i.e., 23%) have detected the presence of Azara's agouti, 15 (26%) of collared peccary, 13 (23%) of brown brocket deer and 25 (44%) of South American tapir. Over the combination of 57 camera traps, there was a reasonable chance to detect the selected species. Our study assumed that we did have false detections and the absence of 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.

We used the species detection information to investigate the effects of spatial extent of habitat covariates on the probability of species occupancy. We ran occupancy models for which occupancy probability Ψ at each camera trap location and the detection probability p were modelled as a linear function of covariates x<sub>i</sub> using logit link functions (see Niedballa et al. 2015 for details). 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 starting with a model  $\Psi(.)p(.)$  where both occupancy and detection probabilities were constant (i.e., fixed as 1). As density of understory and weather could affect detection probability around the camera traps, we created two covariates called visibility (contrasting open vs. dense understory) and weather (contrasting rainy vs. dry season). Then we ran three other models  $\Psi(.)p(x_i)$  to test whether covariates x<sub>i</sub> including visibility around camera trap and weather, or both, affected detection probability p with occupancy probability \P constant. We compared these four models of detection  $(\Psi(.)p(.), \Psi(.)p(visibility), \Psi(.)p(weather), \Psi(.)p(visibility, weather)$  using AIC and selected the model with the lowest AIC and  $\triangle$ AIC higher than 2 with the closest competing model. Finally, from this selected (null) model of detection, we tested the effects of landscape metrics on occupancy probability  $\Psi$  at each of the 8 buffer sizes and three landscape metrics calculated from the three forest cover maps (25, 50 and 75%). Thus, for each landscape metric, we ran 8 models that we compared using AIC and null models. All models showing  $\triangle$ AIC < 2 from the best fitted models had similar statistical supports. Estimates ( $\pm$  SE) of landscape metrics with their P-values affecting

occupancy probability were extracted from the best candidate models. Statistical analyses were performed using R 3.6.2 (R Development Core Team 2019).

## **Results**

Performance of models run at different spatial extents highlighted scale of effects of landscape 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. 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 Table S2). For the other situations in the three species, we did not detect a scale of effects as null models including only covariates on detection probability p provided similar statistical models than candidate models including covariates affecting occupancy probability  $\Psi$  (Electronic appendix Table S2). The South American tapir did not exhibit any scale of effect for the three landscape metrics (Fig 3d, h, l).

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 different buffer sizes (Electronic appendices Table S2-4). The range of selected buffer sizes (i.e. the 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). Although a range of selected models emerged at 50% tree cover in PD in Azara's agouti, we did not detect any significant effect of this variable (i.e., P > 0.05, Table 2). Thus, we did not consider this result as robust. Altogether, our analyses suggest that the species exhibit a stronger scale of effect when considering high tree cover (e.g. 75%) compared to medium-low tree cover (50 – 25%).

Finally, regarding the estimate ( $\beta$ )  $\pm$  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).

#### **Discussion**

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 how we define forests.

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

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 effect on FC. These results might show that species that are more generalist in their habitat requirements might not be limited by forest cover using agricultural landscape with open areas to achieve their ecological requirements (e.g., food). A previous study in Brazilian Amazon reported that the lowland tapir was one of the species with a high ability to survive in more altered environments including regeneration areas with food opportunities (Teixeira-Santos et al. 2020). The lack of scale of effect on the occurrence of the South American tapir might also be observed if the response is not scale dependent (Martin and Fahrig 2012) or the landscape heterogeneity does not allow to investigate pattern at a larger scale in species with high dispersal range (> 3km for this species, Table 1). As in our study area, landscape variables remain quite fixed beyond a buffer 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.

Landscape features might affect the scale of effect on species occurrence. Lyra-Jorge et al. (2010) reported that two large and highly mobile carnivores, *Puma concolor* and *Chrysocyon brachyurus*, were best explained by edge density of the native vegetation at a coarse scale of 2 km. Our analyses showed that species that mainly use forest edges, the Azara's agouti and the collared peccary, exhibited a scale of effect on metrics associated to landscape configuration such as ED and PD. Regolin et al. (2017) reported that the richness of carnivores community increased with forest 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 neighbouring patch of remnant forest decreased. Here, as the studied species exhibited different patterns of scale of effect with the landscape metrics, our results supported previous studies 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. 2017). Unfortunately, among the four selected species, we do not have one mammal species that did not require forest.

While different percentages of tree cover have been used in the literature to map forest cover (Defries et al. 2001; Hansen et al. 2010), studies were reporting that occurrence of terrestrial 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 tree cover used to map forest cover affect the scale of effect of landscape metrics on the occurrence 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 detection of scale of effect varies with habitat definition. While the percent tree cover used to map forest cover shaped patch-based landscape metrics and thus the assessment of land use and land cover composition and spatial configuration, it also determines our ability to detect the correct scale of effects for species. This is a new and important result, which implies that before using multiple 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, particularly for species living in complex mosaic of vegetation types including forests, savannas and grasslands (Bonanomi et al. 2019) showing for instance environment gradients. Several percentages of tree cover have been used in the literature to map forest cover (Defries et al. 2001; Hansen et al. 2010). Although changing the percent tree cover allows to select different amount of

canopy, it also affects the type of vegetation and thus habitat for species (Hirota et al. 2011). These results are particularly important in the current landscape dynamics occurring in the Cerrado. In the last decades, agriculture expansion at the expense of native vegetation dramatically increased landscape fragmentation (Strassburg et al. 2017) and the need to investigate the mechanism 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

388

389

390

391

392

393

394

395

396

397

398

399

400

401

402

403

404

405

406

407

408

409

Implications for conservation and land planning

Although the spatial extent (scale) at which landscape attributes are measured has a substantial 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 (Chamaillé-Jammes et al. 2016; Zimbres et al. 2018), the question whether forest cover (i.e., percent tree cover) has been appropriately selected to link landscape metrics on the occurrence of species 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 species. Although 75% tree cover seems to be a good proxy of forest cover to study the forestdwelling species in our study area, this might also be the case for other variables of landscape composition that are characterised by a continuous gradient such as agriculture or urbanisation. Moreover, from a conservation perspective, it is critical to test multiple scales at which species respond to the landscape variables. In the Cerrado, we recommend that initiatives examining spatial arrangement of landscape attributes of forest cover in relation to populations in terrestrial mammals (from small to large body sized species) should consider buffer sizes from 500 to 2000 m. However, 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 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 multiscale management based on the responses of multiple species (Neel et al. 2004). Thus, using multiple definitions of habitats and scales is critical as most initiatives of ecological zoning, biodiversity prioritizing or ecosystem restoration use landscape units that have a particular meaning in terms of land use dynamics and planning (usually justified by physical constraints) but not in 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
428	Aliaga-Rossel E, Kays R, Fragoso J (2008) Home-range use by the Central American agouti
429	(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
433	M, Igari AT, Martensen AC, Metzger JP (2014) Using ecological thresholds to evaluate the
434	costs and benefits of set-asides in a biodiversity hotspot. Science 345: 1041-1045
435	Bastian O, Krönert R, Lipský (2006) Landscape diagnosis on different space and time scales – a
436	challenge for landscape planning. Landsc Ecol 21: 359-374
437	Bhakti T, Goulart F, de Azevedo CS, Antonini Y (2018) Does scale matter? The influence of three-
438	level spatial scales on forest bird occurrence in a tropical landscape. PlosONE 13: e0198732
439	Bennett B (2002) What is a forest? on the vagueness of certain geographic concepts. Topoi 20:189-
440	201Bennett A, Radford JQ, Haslem A (2006) Properties of land mosaics: implications for
441	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
447	be the diversity patterns of medium- to large-bodied mammals if the fragmented Atlantic
448	Forest was a large metacommunity? Biol Conserv 211: 85-94
449	Bonanomi J, Tortato FR, Gomes FSR, Penha JM, Bueno AS, Peres CS (2019) Protecting forests at
450	the expense of native grasslands: Land-use policy encourages open-habitat loss in the
451	Brazilian cerrado biome. Perspect Ecol Conser 17: 26-31
452	Boscolo D, Metzger JP (2009) Is bird incidence in Atlantic forest fragments influenced by
453	landscape patterns at multiple scales? Landsc Ecol 24: 907-918
454	Bovendorp RS, Brum FT, McCleery RA, Baiser B, Loyola R, Cianciaruso MV, Galetti M (2019)
455	Defaunation and fragmentation erode small mammal diversity dimensions in tropical forests.
456	Ecography 42: 23-35
457	Bowman J, Jaeger JAG, Fahrig L (2002) Dispersal distance of mammals is proportional to home
458	range size. Ecology 83: 2049-2055
459	Brennan JM, Bender DJ, Contreras TA, Fahrig L (2002) Focal patch landscape studies for wildlife
460	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
466	Chaplin-Kramer R, de Valpine P, Mills NJ, Kremen C (2013) Detecting pest control services across
467	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/s13280-016-
476	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
495	Eisenberg JF, Redford KH (1999) Mammals of the Neotropics: volume 3: the central Neotropics.
496	University of Chicago Press, Chicago, Illinois.
497	Fahrig L, Baudry J, Brotons L, Burel FG, Crist TO, Fuller RJ, Martin JL (2011) Functional
498	landscape heterogeneity and animal biodiversity in agricultural landscapes. Ecol Lett 14: 101-
499	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
503	Fiske I, Chandler R (2011) unmarked: An R Package for Fitting Hierarchical Models of Wildlife
504	Occurrence and Abundance. J Stat Softw 43: 1-23
505	Fortin D, Courtois R, Etcheverry P, Dussault C, Gingras A (2008) Winter selection of landscapes by
506	woodland caribou: behavioural response to geographical gradients in habitat attributes. J Appl
507	Ecol 45:1392-1400
508	Fragoso JMS (1998) Home Range and movement patterns of white-lipped Peccary (Tayassu pecari)
509	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
518	Hansen MC, Stehman SV, Potapov PV (2010) Quantification of global gross forest cover loss. Proc
519	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
522	Holland J, Bert DG, Fahrig L (2004) Determining the spatial scale of species' response to habitat.
523	Bioscience 54: 227-233
524	Holland J, Yang S (2016) Multi-scale studies and the ecological neighbourhood. Curr Landsc Ecol
525	Rep 1: 135-145
526	Huais PY (2018) multifit: an R function for multi-scale analysis in landscape ecology. Landsc Ecol
527	33: 1023-1028
528	Jackson HB, Fahrig L (2012) What size is a biologically relevant landscape? Landsc Ecol 27: 929-
529	941
530	Jackson HB, Fahrig L (2015) Are ecologists conducting research at the optimal scale? Glob Ecol
531	Biogeogr 24: 52–63
532	Johnson DH (1980) The comparison of usage and availability measurements for evaluating resource
533	preference. Ecology 61: 65-71
534	Jansen PA, Hirsch BT, Emsens W-J, Zamora-Gutierrez V., Wikelski M, Kays R (2012) Thieving
535	rodents as substitute dispersers of megafaunal seeds. Proc Natl Acad Sci USA 109: 12610-
536	12615
537	Johnson DH (1980) The comparison of usage and availability measurements for evaluating resource
538	preference. Ecology 61: 65-71
539	Judas J, Henry O (1999) Seasonal variation of home range of collared peccary in tropical rain forest
540	of French Guiana. J Wildl Manag 63: 546-552
541	Keuroghlian A, Eaton DP, Longland WS (2004) Area use by white-lipped and collared peccaries
542	(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,
545	Law R, Tilman D, Loreau M, Gonzalez A (2004) The metacommunity concept: a framework
546	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
552	Lü Y, Feng X, Chen L, Fu B (2013) Scaling effects of landscape metrics: a comparison of two
553	methods. Phys Geog 34: 40-49
554	Lyra-Jorge MC, Ribeiro MC, Ciocheti C, Tambosi LR, Pivello VR (2010) Influence of multi-scale
555	landscape structure on the occurrence of carnivorous mammals in a human-modified savanna,
556	Brazil. Eur J Wildlife Res 56: 359-368
557	Mares MA, Ernest KA, Gettinger RD (1986) Small mammal community structure and composition
558	in the Cerrado province of Central Brazil. J Trop Ecol 2: 289-300
559	Martensen AC, Ribeiro MC, Banks-Leite C, Prado PI, Metzger JP (2012) Associations of Forest
560	Cover, Fragment Area, and Connectivity with Neotropical Understory Bird Species Richness
561	and Abundance. Conserv Biol 26: 1100-1111
562	Martin AE, Farhig L (2012) Measuring and selecting scales o effect for landscape predictors in
563	species-habitat models. Ecol Appl 22: 2277-2292
564	Mayor SJ, Schneider DC, Schaefer JA, Mahoney SP (2009) Habitat selection at multiple scales.
565	Ecoscience 16: 238-247
566	Melo GL, Sponchiado J, Cáceres N, Fahrig L (2017) Testing the habitat amount hypothesis for
567	South American small mammals. Biol Conser 209: 304-314
568	Michalski F, Peres CA (2007) Disturbance-mediated mammal persistence and abundance-area
569	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
574	Moraga AD, Martin AE, Fahrig L (2019) The scale of effect of landscape context varies with the
575	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
579	Newbold T, Hudson LN, Hill SLL et al (2015) Global effects of land use on local terrestrial
580	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.
587	Pardini R, de Buen, AA, Gardner TA, Prado PI, Metzger JP (2010) Beyond the fragmentation
588	threshold hypothesis: Regime shifts in biodiversity across fragmented landscapes. PLoS ONE
589	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
593	R, Vieira MV, Banks-Leite G, Fonseca CR et al. (2020) Indirect effects of habitat loss via
594	habitat fragmentation: Across-taxa analysis of forest-dependent species. Biol Cons 241:
595	108368
596	Rahbek C (2005) The role of spatial scale and the perception of large-scale species-richness
597	patterns. Ecol Lett 8: 224-239
598	Rausch LL, Gibbs HK, Schelly I, Brandão A, Morton DC, Filho AC, Strassburg B, Walker N,
599	Noojipady P, Barreto P, Meyer D (2019) Soy expansion in Brazil's Cerrado. Conserv Lett 12:
600	e12671Redon M, Bergès TC, Luque S (2014) Effects of increasing landscape heterogeneity
601	on local plant species richness: how much is enough? Landsc Ecol 29: 773-787
602	Regolin AL, Cherem JJ, Graipel ME, Bogoni JA, Ribeiro JW, Vancine MH, Tortato MA, Oliveira-
603	Santos LG, Fantacini FM, Luiz MR, de Castilho PV, Ribeiro MC, Cáceres NC (2017) Forest
604	cover influences occurrence of mammalian carnivores within Brazilian Atlantic Forest. J
605	Mammal 98: 1721–1731
606	Rocchini D, Foody GM, Nagendra H, Ricotta C, Anand M, He KS et al. (2013) Uncertainty in
607	ecosystem mapping by remote sensing. Comput Geosci 50:128–135.
608	Royo AA, Carson WP (2005) The herb community of a tropical forest in central Panamá: dynamics
609	and impact of mammalian herbivores. Oecologia 145: 66-75
610	Saab V (1999) Importance of spatial scale to habitat use by breeding birds in riparian forests: a
611	hierarchical analysis. Ecol Appl 9: 135-151
612	Schuster R Römer H, Germain RR (2013) Using multi-scale distribution and movement effects
613	along a montane highway to identify optimal crossing locations for a large-bodied mammal
614	community. PeerJ 1: e189
615	Strahler AH, Woodcock CE, Smith JA (1986) On the nature of models in remote sensing. Remote
616	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)
621	Environmental factors influencing the abundance of four species of threatened mammals in
622	degraded habitats in the eastern Brazilian Amazon. PLoS ONE 15: e0229459
623	Thornton DH, Fletcher RJ (2014) Body size and spatial scales in avian response to landscapes: a
624	meta-analysis. Ecography 37: 454-463
625	Vanderwal J, Falconi L, Januchowski S, Shoo L, Storlie C (2014) Package 'SDMTools'. Species
626	Distribution ModellingTools: Tools for processing data associated with species distribution
627	modelling exercises. R-package version 1.1.12
628	Villard M-A, Metzger JP (2014) Beyond the fragmentation debate: a conceptual model to predict
629	when habitat configuration really matters. J Appl Ecol 51: 309-318
630	Woodcock CE, Strahler AH (1987) The factor of scale in remote sensing. Remote Sens Environ 21:
631	311–332
632	Young HS, McCauley DJ, Helgen KM, Goheen JR, Otárola-Castillo E, Palmer TM, Pringle RM,
633	Young TP, Dirzo R (2013) Effects of mammalian herbivore declines on plant communities:
634	observations and experiments in an African savanna. J Ecol 101: 1030-1041
635	Zimbres B, Peres CA, Penido G, Machado RB (2018) Thresholds of riparian forest use by terrestrial
636	mammals in a fragmented Amazonian deforestation frontier. Biodivers Conserv 27: 2815-
637	2836

Table 1. Ecological traits of the studied mammal species

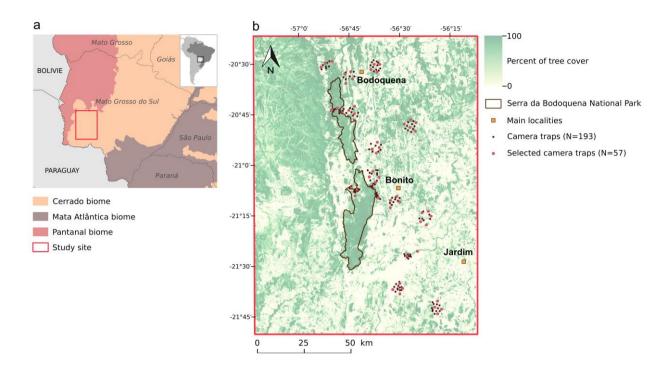
Studied species	Adult body mass (kg)	Home range (ha)	Dispersal range (m)	Habitat use	References
Azara's agouti	2.9	1.34 – 2.45	100 – 500	Forest/ edges	Aliaga-Rossel et al. 2008; Jansen et al. 2012; Cid et al. 2013
Brown brocket deer	21	2.7 – 348	750 – 1 000	Forest/ open area	Duarte et al. 2012 ; Pires 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) 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  $\Psi$  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. When several models showed similar statistical support (i.e.  $\Delta$ AIC < 2), estimate ( $\beta$ )  $\pm$  SE was extracted for the model with the lowest AIC indicated by the \* near the buffer size. Significant effects are indicated in bold.

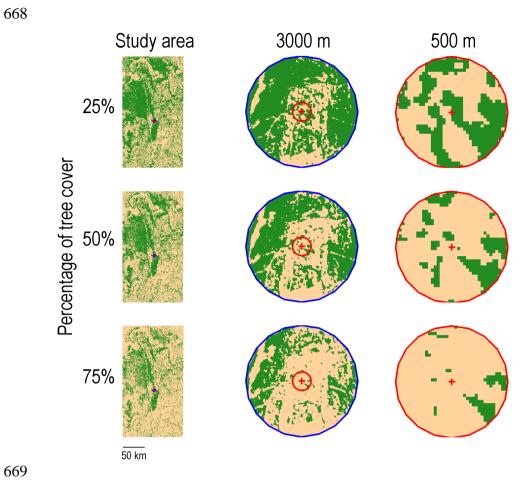
		Percent of tree cover per pixel									
Landscape metric	Model structure	Species	25%			50%			75%		
			Buffer size	$\beta \pm SE$	P	Buffer size	$\beta \pm SE$	P	Buffer size	$\beta \pm SE$	P
	p (weather+visibility) Ψ (ED)	Azara's agouti				500	$0.325 \pm 0.185$	0.079	500*-1500	$0.517 \pm 0.267$	0.053
ED	p (visibility) Ψ (ED)	Brown brocket deer									
LD	p (visibility) Ψ (ED)	Collared peccary									
	p (weather+visibility) Ψ (ED)	South American tapir									
	p (weather+visibility) Ψ (PD)	Azara's agouti				1500-2000*- 3000	-1.030 ± 0.675	0.128			
PD	p (visibility) Ψ (PD)	Brown brocket deer							750	$0.643 \pm 0.305$	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	$4.540 \pm 2.500$	0.070	500*-2000	$5.050 \pm 2.890$	0.081	500-1500*-2000	$9.000 \pm 4.450$	0.043
FC	p (visibility) Ψ (FC)	Brown brocket deer									
rc	p (visibility) Ψ (FC)	Collared peccary	500*-1000	$2.690 \pm 1.400$	0.029	500*-1000	$3.190 \pm 1.540$	0.039	500*-750	$4.030 \pm 1.890$	0.033
	p (weather+visibility) Ψ (FC)	South American tapir									

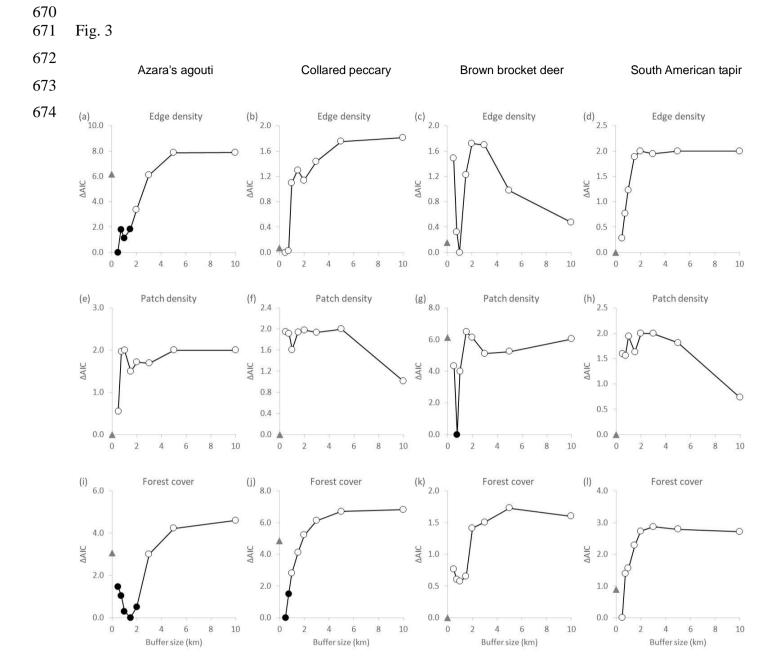
649	Figure captions
650	
651	Fig. 1 Map of the study site, Bodoquena, state of Mato Grosso do Sul, Brazil (a) and forest cover
652	including the location of camera traps (b). Camera traps selected to run occupancy models (see
653	methods) are highlighted in red.
654	
655	Fig. 2 Example of forest cover 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	$\Delta AIC \le 2$ included the smallest AIC without the null model (black dots).





667 Fig. 2





## **Electronic appendices**

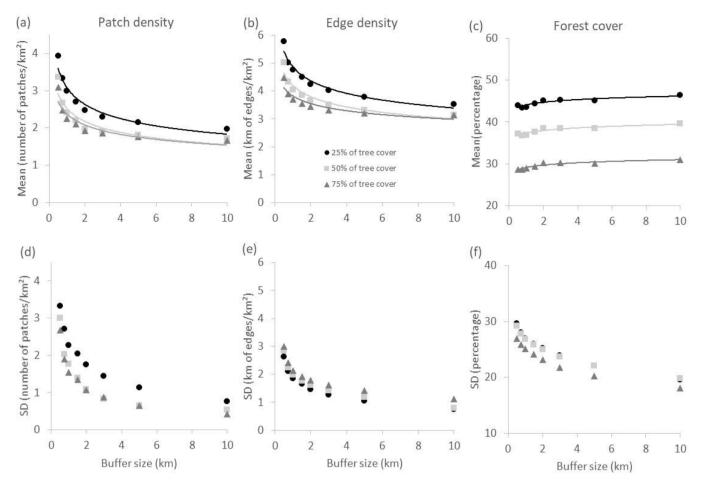


Fig. S1 Scaling relations with spatial extent (buffer size) of the mean values of landscape metrics, patch density (a), edge density (b), forest cover (c) and their standard deviation (SD) (d, e, f) assessed around our camera traps at three different percent tree cover per pixel (25, 50 and 75%) (See 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 <sup>2</sup>
	25%	Power***	y=3.082x <sup>-0.226</sup>	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 <sup>-0.157</sup>	0.96
Edge Density	50%	Power***	$y=4.196x^{-0.145}$	0.92
	75%	Power***	$y=3.823x^{-0.108}$	0.86
	25%	Logarithm***	$y=0.917\ln(x)+44.058$	0.81
Forest Cover	50%	Logarithm***	$y=0.915\ln(x)+37.312$	0.86
	75%	Logarithm***	$y=0.804\ln(x)+29.143$	0.89

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 *Mazama gouazoubira*, and South American tapir *Tapirus terrestris*,) in the Brazilian Cerrado hotspot across buffer size (from 500 to 10000 m) considering 75% as the threshold of tree cover. Model structure mentions how occupancy probability  $\Psi$  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 agouti						d peccary		Brown	brocket deer			South American tapir				
Landscape metric	Buffer size	Model structure	AIC	ΔΑΙС	Buffer size	Model structure	AIC	ΔΑΙС	Buffer size	Model structure	AIC	ΔΑΙС	Buffer size	Model structure	AIC	ΔΑΙΟ
	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) p (weather +	149.63	3.38	2000	p (visibility) Ψ(ED) p (visibility)	174.61	1.13	2000	p (visibility) Ψ(ED) p (visibility)	147.12	1.72	2000	p (weather + visibility) Ψ(ED) p (weather +	233.04	2.00
	3000	visibility) Ψ(ED) p (weather +	152.34	6.09	3000	Ψ(ED) p (visibility)	174.91	1.44	3000	Ψ(ED) p (visibility)	147.10	1.70	3000	visibility) Ψ(ED) p (weather +	232.99	1.95
	5000	visibility) Ψ(ED) p (weather +		7.87	5000	Ψ(ED) p (visibility)	175.22	1.75	5000	Ψ(ED) p (visibility)	146.38	0.98	5000	visibility) Ψ(ED) p (weather +	233.04	2.00
	10000	visibility) Ψ(ED)	154.13	7.88	10000	Ψ(ED)	175.28	1.81	10000	Ψ(ED)	145.88	0.47	10000	visibility) Ψ(ED)	233.04	2.00
	Null	p (weather + visibility) Ψ(1)	152.43	0.00	Null	p (visibility) Ψ(1)	173.54	0.00	Null	p (visibility) Ψ(1)	145.56	6.13	Null	p (weather + visibility) Ψ(1)	231.04	0.00
	500	p (weather + visibility) Ψ(PD)	152.98	0.54	500	p (visibility) Ψ(PD)	175.49	1.95	500	p (visibility) Ψ(PD)	143.78	4.35	500	p (weather + visibility) Ψ(PD)	232.63	1.59
	750	p (weather + visibility) Ψ(PD)	154.40	1.97	750	p (visibility) Ψ(PD)	175.46	1.91	750	p (visibility) Ψ(PD)	139.42	0.00	750	p (weather + visibility) Ψ(PD)	232.60	1.56
	1000	p (weather + visibility) Ψ(PD)	154.43	2.00	1000	p (visibility) Ψ(PD)	175.15	1.61	1000	p (visibility) Ψ(PD)	143.41	3.99	1000	p (weather + visibility) Ψ(PD)	232.98	1.94
PD	1500	p (weather + visibility) Ψ(PD)	153.93	1.50	1500	p (visibility) Ψ(PD)	175.49	1.94	1500	p (visibility) Ψ(PD)	145.92	6.49	1500	p (weather + visibility) Ψ(PD)	232.67	1.63
	2000	p (weather + visibility) Ψ(PD)	154.15	1.71	2000	p (visibility) Ψ(PD)	175.52	1.98	2000	p (visibility) Ψ(PD)	145.54	6.12	2000	p (weather + visibility) Ψ(PD)	233.04	2.00
	3000	p (weather + visibility) Ψ(PD)	154.12	1.69	3000	p (visibility) Ψ(PD)	175.48	1.93	3000	p (visibility) Ψ(PD)	144.53	5.11	3000	p (weather + visibility) Ψ(PD)	233.04	1.99
	5000	p (weather + visibility) Ψ(PD)	154.43	2.00	5000	p (visibility) Ψ(PD)	175.54	2.00	5000	p (visibility) Ψ(PD)	144.66	5.24	5000	p (weather + visibility) Ψ(PD)	232.85	1.81
	10000	p (weather + visibility) Ψ(PD)	154.43	2.00	10000	p (visibility) Ψ(PD)	174.55	1.01	10000	p (visibility) Ψ(PD)	145.46	6.03	10000	p (weather + 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) Ψ(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
C	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

Table S3. 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 50% as the threshold of tree cover. Model structure mentions how occupancy probability  $\Psi$  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.

Procedure	Azara's agou	uti		Collared	d peccary		Brown b	brocket deer			South American tapir						
Nall   vishility   Vish   124   289   Nall   Vish   173.54   1.88   500   Vish   Vis			Model structure	AIC	ΔΑΙС			AIC	ΔΑΙС			AIC	ΔΑΙС		Model structure	AIC	ΔΑΙС
Fig.   Properties   1,000						-											
Soo		Null		152.43	2.89	Null		173.54	0.00	Null		145.56	1.43	Null		231.04	0.00
Process				110.51												222.22	
Procedure		500		149.54	0.00	500		175.12	1.58	500		145.48	1.35	500		232.32	1.27
Powelfare		750		151.63	2.08	750		175.08	1.54	750		144.80	0.67	750		232.20	1.25
ED   1500   visibility) \( \text{P(ED)} \)   15.15   2.61   1000   \( \text{P(ED)} \)   175.51   198   1000   \( \text{P(ED)} \)   14.15   14.55   1.44   1500   \( \text{visibility} \) \( \text{P(ED)} \)   1500   \( \text{visibility} \) \( \text{P(ED)} \)   15.05   1.07   1500   \( \text{P(ED)} \)   14.55   1.44   1500   \( \text{visibility} \) \( \text{P(ED)} \)   15.05   1.07   1500   \( \text{P(ED)} \)   14.55   1.44   1500   \( \text{visibility} \) \( \text{P(ED)} \)   15.05   1.07   1500   \( \text{P(ED)} \)   14.15   1.07   1.000   \( \text{visibility} \) \( \text{P(ED)} \)   15.05   1.07   1.000   \( \text{P(ED)} \)   17.55   1.07   1.000   \( \text{P(ED)} \)   17.55   1.000   \( \text{P(ED)} \)   17.55   1.000   \( \text{P(ED)} \)   17.55   1.000   1.000   \( \text{P(ED)} \)   17.55   1.000   \( \text{P(ED)} \)   14.13   1.000   \( \text{P(ED)} \)   1.000   \( \text{P(ED)} \)   14.13   1.000   \( \text{P(ED)} \)   1.000   \( \text{P(ED)} \)   14.13   1.000   \( \text{P(ED)} \)   1.000   \( \text{P(ED)} \)   17.55   1.000		750		131.03	2.00	750		175.00	1.54	750		144.00	0.07	750		232.29	1.23
ED		1000		152.15	2.61	1000		175.53	1.98	1000		144.50	0.37	1000		232.89	1.85
For   Part   P	ED																
Property   Property   Property   Property   Property   Program		1500	visibility) Ψ(ED)	151.83	2.29	1500	Ψ(ED)	175.51	1.97	1500	Ψ(ED)	145.58	1.44	1500	visibility) Ψ(ED)	233.00	1.95
Process																	
Solution		2000		152.35	2.80	2000		175.30	1.76	2000		146.10	1.97	2000		233.04	2.00
Process																	
Solidary   Y(ED)   154.09   4.55   5000   Y(ED)   175.18   1.64   10000   Y(ED)   144.87   0.74   5000   visibility) P(ED)   233.00   232.41   23		3000		153.49	3.95	3000		175.43	1.89	3000		145.26	1.13	3000		232.96	1.92
Process		5000		154.00	155	5000		175 42	1 90	5000		144 97	0.74	5000		222.00	1.96
Null		3000		134.09	4.55	2000		173.43	1.07	2000		144.07	0.74	2000		233.00	1.70
Nul		10000		154.25	4.71	10000		175.18	1.64	10000		144.13	0.00	10000		232.41	1.37
Null visibility) \( \text{V(I)} \)			• , , ,												• , , .		
Null			n (weather +				n (visibility)				n (visibility)				n (weather +		
P		Null		152.43	2.05	Null		173.54	0.00	Null		145.56	1.83	Null		231.04	0.00
Property						- 14				- 1							
Process   Proc		500	visibility) Ψ(PD)	154.27	3.88	500	Ψ(PD)	175.43	1.88	500	Ψ(PD)	143.73	0.00	500	visibility) Ψ(PD)	233.03	1.99
PD   000   0																	
PD   1000   visibility) \(V(PD)   152.87   2.49   1000   \(V(PD)   174.45   0.91   1000   \(V(PD)   146.62   2.89   1000   visibility) \(V(PD)   232.17 \)   1500   visibility) \(V(PD)   152.24   1.85   1500   \(V(PD)   174.35   0.80   1500   \(V(PD)   147.06   3.33   1500   \(V(PD)   150.08   1500   Visibility) \(V(PD)   232.55   1200   Visibility) \(V(PD)   150.08   1500   V(PD)   175.07   1.53   1200   V(PD)   146.83   3.10   2000   Visibility) \(V(PD)   232.67 \)   1500   Visibility) \(V(PD)   151.52   1.13   3000   V(PD)   175.07   1.53   1.81   3000   V(PD)   146.83   3.10   2000   Visibility) \(V(PD)   232.87 \)   1500   Visibility) \(V(PD)   151.52   1.13   3000   V(PD)   175.35   1.81   3000   V(PD)   146.92   3.19   3000   Visibility) \(V(PD)   232.87 \)   1000   Visibility) \(V(PD)   153.40   3.01   5000   V(PD)   175.30   1.75   5000   V(PD)   146.82   2.55   5000   Visibility) \(V(PD)   232.75 \)   1000   Visibility) \(V(PD)   153.43   3.05   10000   V(PD)   175.54   2.00   10000   V(PD)   147.44   3.71   10000   Visibility) \(V(PD)   231.83   1000   V(PD)   175.40   1.000   V(PD)   175.40   1.000   V(PD)   147.44   3.71   10000   Visibility) \(V(PD)   231.83   V(PD)   1000		750		153.96	3.58	750		174.24	0.70	750		144.95	1.22	750		233.01	1.97
PD   1500		1000		452.05	2.10	4000			0.04	1000			2.00	4000		222.15	
1500   visibility   vivio   vivio   visibility   vivio   visibility   vivio   visibility   vi		1000		152.87	2.49	1000		174.45	0.91	1000		146.62	2.89	1000		232.17	1.12
P (weather +   P (wisibility)   P (weather +   P (wisibility)   P (weather +	PD	1500		152.24	1 85	1500		174 35	0.80	1500		147.06	3 33	1500		232.55	1.51
2000		1500		132.24	1.05	1300		174.33	0.00	1300		147.00	3.33	1300		232.33	1.51
Probability   Property   Proper		2000		150.38	0.00	2000		175.07	1.53	2000		146.83	3.10	2000		233.04	2.00
3000 visibility) \( \text{VPD} \)   151.52   1.13   3000 \( \text{VPD} \)   175.35   1.81   3000 \( \text{VPD} \)   146.92   3.19   3000 \( \text{visibility} \text{VPD} \)   232.87 \\   5000 visibility) \( \text{VPD} \)   153.40   3.01   5000 \( \text{VPD} \)   175.30   1.75   5000 \( \text{VPD} \)   146.28   2.55   5000 \( \text{visibility} \) \( \text{VPD} \)   232.75 \\   1000 visibility) \( \text{VPD} \)   153.43   3.05   10000 \( \text{VPD} \)   175.54   2.00   10000 \( \text{VPD} \)   146.28   2.55   5000 \( \text{visibility} \) \( \text{VPD} \)   232.75 \\   1000 visibility) \( \text{VPD} \)   153.43   3.05   10000 \( \text{VPD} \)   175.54   2.00   10000 \( \text{VPD} \)   147.44   3.71   10000 \( \text{visibility} \) \( \text{VPD} \)   231.83 \\   1000 visibility) \( \text{VPD} \)   152.43   3.28   Null \( \text{VPI} \)   173.54   3.45   Null \( \text{VPI} \)   145.56   0.00   Null \( \text{visibility} \) \( \text{VPD} \)   231.04 \\   1000 visibility) \( \text{VFC} \)   149.15   0.00   500 \( \text{VFC} \)   170.09   0.00   500 \( \text{VFC} \)   145.80   0.25   500 \( \text{visibility} \) \( \text{VPC} \)   231.63 \\   1000 visibility) \( \text{VFC} \)   150.26   1.11   750 \( \text{VFC} \)   170.80   0.71   750 \( \text{VFC} \)   145.80   0.25   500 \( \text{visibility} \) \( \text{VFC} \)   231.54 \\   1000 visibility) \( \text{VFC} \)   150.26   1.11   750 \( \text{VFC} \)   171.95   1.86   1000 \( \text{VFC} \)   145.80   0.25   500 \( \text{visibility} \) \( \text{VFC} \)   231.88 \\   1000 visibility) \( \text{VFC} \)   149.44   0.29   1500 \( \text{VFC} \)   171.95   1.86   1000 \( \text{VFC} \)   145.80   0.23   1500 \( \text{visibility} \) \( \text{VFC} \)   232.60 \\   1500 visibility) \( \text{VFC} \)   150.31   1.16   2000 \( \text{VFC} \)   173.87   3.78   2000 \( \text{VFC} \)   146.62   1.07   2000 \( \text{visibility} \) \( \text{VFC} \)   232.95 \\   1500 visibility) \( \text{VFC} \)   152.47   3.32   3000 \( \text{VFC} \)   174.78   4.69   3000 \( \text{VFC} \)   146.72   117   30																	
Solidary   Feature   Fea		3000	visibility) Ψ(PD)	151.52	1.13	3000		175.35	1.81	3000		146.92	3.19	3000		232.87	1.83
P (weather +																	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		5000		153.40	3.01	5000		175.30	1.75	5000		146.28	2.55	5000		232.75	1.71
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		10000		152.42	2.05	10000		175.54	2.00	10000		147.44	2.71	10000		221.02	0.70
$ FC \begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		10000	visibility) Ψ(PD)	153.43	3.05	10000	Ψ(PD)	175.54	2.00	10000	Ψ(PD)	147.44	3.71	10000	visibility) Ψ(PD)	231.83	0.79
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$																	
Procedure   Proc		NT 11		150.40	2.20	NT 11		172.54	2.45	N: 11		145.56	0.00	N. 11		221.04	0.41
$ FC \begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		Null		132.43	3.28	Null	. ,	1/3.34	5.45	Null		145.50	0.00	Null		231.04	0.41
$FC \begin{tabular}{l l l l l l l l l l l l l l l l l l l $		500		149 15	0.00	500		170.09	0.00	500		145.80	0.25	500		230.63	0.00
FC		300	• / / /	149.13	0.00	300		170.09	0.00	300		145.00	0.23	300	•	230.03	0.00
FC $\begin{pmatrix} 1500 \\ 20$		750		150.26	1.11	750		170.80	0.71	750		145.68	0.12	750		231.54	0.91
FC $\begin{pmatrix} 1000 & visibility \ V(FC) & 149.31 & 0.16 & 1000 & V(FC) & 171.95 & 1.86 & 1000 & V(FC) & 145.82 & 0.26 & 1000 & visibility \ V(FC) & 231.88 \\ p (weather + p (weather + p (visibility) & p (weather + p (visibil$		,,,	•			700	. ,		0.7.2	700			0.12	700	•		0.52
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		1000		149.31	0.16	1000		171.95	1.86	1000		145.82	0.26	1000		231.88	1.25
1500   visibility  v(FC)   149,44   0.29   1500   v(FC)   173,02   2.93   1500   v(FC)   143,79   0.23   1500   visibility) v(FC)   232,60	FC		p (weather +				p (visibility)				p (visibility)				p (weather +		
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		1500		149.44	0.29	1500	Ψ(FC)	173.02	2.93	1500	Ψ(FC)	145.79	0.23	1500		232.60	1.97
$\begin{array}{cccccccccccccccccccccccccccccccccccc$																	
3000 visibility) $\Psi$ (FC) 152.47 3.32 3000 $\Psi$ (FC) 174.78 4.69 <b>3000</b> $\Psi$ (FC) 146.72 <b>1.17</b> 3000 visibility) $\Psi$ (FC) 233.02 p (weather + p (visibility) p (visibility) p (weather +		2000		150.31	1.16	2000	. ,	173.87	3.78	2000		146.62	1.07	2000		232.95	2.31
p (weather + p (visibility) p (visibility) p (weather +		2000		150.45	2.22	2000		17470	4.60	2000		146.72	1.15	2000		222.02	2.20
		3000	• / / /	152.47	3.32	3000		174.78	4.69	3000		146.72	1.17	3000	•	233.02	2.38
		5000		152.62	4.40	5000		175.20	5 21	5000		146.00	1.44	5000		222.05	2 22
5000 Visibility) T(C) 155.05 4.46 5000 T(C) 175.39 5.51 5000 T(C) 140.39 1.44 5000 Visibility) T(C) 252.55 p (weather + p (visibility) p (visibility) p (visibility) p (weather +		2000		155.63	4.48	5000		1/5.39	3.31	5000		146.99	1.44	5000		232.95	2.32
		10000		154.00	4.85	10000		175 50	5.42	10000		146 96	1.40	10000		232.88	2 25

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  $\Psi$  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 metric	Buffer size	Model structure	AIC	ΔΑΙС	Buffer size	Model structure	AIC	ΔΑΙС	Buffer size	Model structure	AIC	ΔΑΙС	Buffer size	Model structure	AIC	ΔΑΙС
	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)	152.81	1.24	750	p (visibility) Ψ(ED)	175.54	1.99	750	p (visibility) Ψ(ED)	143.64	0.00	750	p (weather + visibility) Ψ(ED)	232.79	1.75
	1000	p (weather + visibility) Ψ(ED) p (weather +	153.75	2.18	1000	p (visibility) Ψ(ED) p (visibility)	175.36	1.82	1000	p (visibility) Ψ(ED) p (visibility)	144.16	0.53	1000	p (weather + visibility) Ψ(ED) p (weather +	233.04	2.00
ED	1500	p (weather + visibility) Ψ(ED) p (weather +	153.89	2.32	1500	P(ED) p (visibility)	175.39	1.85	1500	P (VISIBILITY) Ψ(ED) p (visibility)	146.32	2.68	1500	yisibility) 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 +	154.27	2.70	3000	Ψ(ED) p (visibility)	175.31	1.77	3000	Ψ(ED) p (visibility)	146.82	3.19	3000	visibility) Ψ(ED) p (weather +	232.98	1.94
	5000	visibility) Ψ(ED) p (weather +	154.41	2.85	5000	Ψ(ED) p (visibility)	175.46	1.92	5000	Ψ(ED) p (visibility)	146.21	2.58	5000	visibility) Ψ(ED) p (weather +	232.92	1.88
	10000	visibility) Ψ(ED)	154.43	2.86	10000	Ψ(ED)	175.41	1.87	10000	Ψ(ED)	146.23	2.60	10000	visibility) Ψ(ED)	231.68	0.63
		p (weather +				p (visibility)				p (visibility)				p (weather +		
	Null	visibility) Ψ(1) p (weather +	152.43		Null	Ψ(1) p (visibility)	173.54		Null	Ψ(1) p (visibility)	145.56		Null	visibility) Ψ(1) p (weather +	231.04	0.00
	500	visibility) Ψ(PD) p (weather +	154.21		500	Ψ(PD) p (visibility)	175.17	2.18	500	Ψ(PD) p (visibility)	147.01		500	visibility) Ψ(PD) p (weather +	233.03	1.99
	750 1000	visibility) Ψ(PD) p (weather +	154.39	1.65	750 1000	Ψ(PD) p (visibility)	173.04 174.04	0.06 1.06	750 1000	Ψ(PD) p (visibility)	147.30 147.54		750 1000	visibility) Ψ(PD) p (weather +	232.69	1.65
PD	1500	visibility) Ψ(PD) p (weather + visibility) Ψ(PD)	154.09 152.91	0.48	1500	Ψ(PD) p (visibility) Ψ(PD)	174.04	0.00	1500	Ψ(PD) p (visibility) Ψ(PD)	147.54		1500	visibility) Ψ(PD) p (weather + visibility) Ψ(PD)	233.04	2.35 2.00
	2000	p (weather + visibility) Ψ(PD)	152.71		2000	p (visibility) Ψ(PD)	172.98	0.12	2000	p (visibility) Ψ(PD)	147.55		2000	p (weather + visibility) Ψ(PD)		
	3000	p (weather + visibility) Ψ(PD)	153.32		3000	p (visibility) Ψ(PD)	174.65	1.67	3000	p (visibility) Ψ(PD)	147.56		3000	p (weather + visibility) Ψ(PD)	231.96	0.92
	5000	p (weather + visibility) Ψ(PD)	153.62		5000	p (visibility) Ψ(PD)	174.72		5000	p (visibility) Ψ(PD)	147.41		5000	p (weather + visibility) Ψ(PD)	231.46	
	10000	p (weather + visibility) Ψ(PD)	153.82		10000	p (visibility) Ψ(PD)	175.46	2.47	10000	p (visibility) Ψ(PD)	146.94		10000	p (weather + visibility) Ψ(PD)	232.47	1.43
	10000	3, , ,	100.02	1107	10000		175.10	2	10000	, ,	1.0.7.	1.00	10000		232.17	
	Null	p (weather + visibility) Ψ(1)	152.43	3.88	Null	p (visibility) Ψ(1)	173.54	2.41	Null	p (visibility) Ψ(1)	145.56	0.00	Null	p (weather + visibility) Ψ(1)	231.04	0.00
	500	p (weather + visibility) Ψ(FC)	148.55	0.00	500	p (visibility) Ψ(FC)	171.13	0.00	500	p (visibility) Ψ(FC)	145.76	0.21	500	p (weather + visibility) Ψ(FC)	231.49	0.44
	750	p (weather + visibility) Ψ(FC)	150.32	1.76	750	p (visibility) Ψ(FC)	171.57	0.44	750	p (visibility) Ψ(FC)	145.58	0.03	750	p (weather + visibility) Ψ(FC)	232.01	0.97
	1000	p (weather + visibility) Ψ(FC)	149.82	1.27	1000	p (visibility) Ψ(FC)	172.33	1.20	1000	p (visibility) Ψ(FC)	145.62	0.06	1000	p (weather + visibility) Ψ(FC)	232.26	1.22
FC	1500	p (weather + visibility) Ψ(FC)	150.08	1.53	1500	p (visibility) Ψ(FC)	173.21	2.08	1500	p (visibility) Ψ(FC)	145.58	0.02	1500	p (weather + visibility) Ψ(FC)	232.88	1.83
	2000	p (weather + visibility) Ψ(FC)	150.91	2.35	2000	p (visibility) Ψ(FC)	173.94	2.81	2000	p (visibility) Ψ(FC)	146.44	0.88	2000	p (weather + visibility) Ψ(FC)	233.03	1.98
	3000	p (weather + visibility) Ψ(FC)	152.55	4.00	3000	p (visibility) Ψ(FC)	174.82	3.69	3000	p (visibility) Ψ(FC)	146.45	0.89	3000	p (weather + visibility) Ψ(FC)	233.04	1.99
	5000	p (weather + visibility) Ψ(FC)	153.57	5.02	5000	p (visibility) Ψ(FC)	175.41	4.28	5000	p (visibility) Ψ(FC)	146.73	1.17	5000	p (weather + visibility) Ψ(FC)	232.91	1.87
	10000	p (weather + visibility) Ψ(FC)	153.98	5.42	10000	p (visibility) Ψ(FC)	175.51	4.38	10000	p (visibility) Ψ(FC)	146.84	1.28	10000	p (weather + visibility) Ψ(FC)	232.85	1.81