



Universidade Federal de Minas Gerais
Instituto de Ciências Biológicas
Departamento de Genética, Ecologia e Evolução



Programa de Pós-Graduação em Ecologia, Conservação e Manejo da Vida Silvestre

TESE DE DOUTORADO

Avaliação da ocupação e da adequabilidade ambiental para a preguiça-de-coleira (*Bradypus torquatus*, Illiger 1811): uma abordagem multi-espacial e multi-temporal com contribuições para a conservação da biodiversidade.

Paloma Marques Santos

Belo Horizonte – MG

2020

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Tese apresentada ao Instituto de Ciências
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Dr. Milton Cezar Ribeiro

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*À Maria luiza e à Maria Angélica,
por sempre me mostrarem
a força que há dentro de mim.*

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“A única coisa que separa as mulheres de cor de qualquer outra pessoa é a oportunidade.”

Viola Davis

“Os (...) entomologistas Bert Hölldobler e Ed Wilson fizeram uma afirmação intrigante no livro Journey to the ants. (...) há dois tipos de cientistas. Um é o teórico, que se interessa por uma questão específica e procura o melhor organismo para resolvê-la. (...). O outro tipo é o naturalista, que se interessa por uma classe específica de animais em si, percebendo que cada animal tem sua história para contar, que se revelará dotada de interesse teórico se for suficientemente estudada. Hölldobler e Wilson consideram-se pertencentes à segunda classe, assim como eu. “

Frans De Waal, no livro *“Eu, primata”*

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247 **Resumo**

248 De acordo com o Painel Intergovernamental sobre mudanças Climáticas (IPCC), áreas antrópicas
249 relacionadas à produção de alimentos são uma das grandes responsáveis pelas mudanças
250 climáticas. A conversão de grandes áreas naturais em áreas agropastoris cumpre uma função
251 fundamental no aumento da liberação do CO₂ no mundo. A crescente perda de habitat interrompe
252 fluxos gênicos e processos ecológicos importantes para a biodiversidade e assim, as mudanças no
253 uso da terra e seus efeitos nas mudanças climáticas correspondem a uma das principais causas de
254 extinção de espécies. Espécies arborícolas são as maiores atingidas pela perda de habitat, como é
255 o caso da Preguiça-de-Coleira (*Bradypus torquatus*), exclusiva da ameaçada Mata Atlântica.
256 Assim, a presente tese teve como objetivo principal entender a influência de fatores ambientais
257 na probabilidade de ocupação da preguiça-de-coleira e avaliar a adequabilidade ambiental atual e
258 futura para a espécie, além de contribuir com a geração de dados de biodiversidade de uso livre.
259 O primeiro capítulo buscou compilar e disponibilizar dados de ocorrência sobre os xenartros na
260 região neotropical, no intuito de facilitar análises e modelagens biológicas e ecológicas diversas.
261 O segundo capítulo objetivou identificar a influência da cobertura florestal e outras variáveis na
262 probabilidade de ocupação da preguiça-de-coleira. O terceiro capítulo investigou a
263 adequabilidade ambiental para a espécie, levando em consideração cenários de regeneração e
264 cenários de mudanças climáticas. Como principais resultados podemos citar: 1) a compilação de
265 mais 30 mil registros de xenartros – incluindo da preguiça-de-coleira, de diferentes instituições
266 de todo mundo e de diferentes fontes – dados estes que, em sua maioria, até então estavam
267 indisponíveis para o acesso público; 2) a descoberta de que preguiça-de-coleira responde a um
268 limiar de 35% de cobertura florestal, abaixo do qual a probabilidade de ocupação desta espécie
269 declina consideravelmente, chegando a quase zero em regiões com menos de 20% de floresta.
270 Além disso, verificamos que a porcentagem de áreas abertas, como pastos, pastos abandonados e
271 áreas de solo expostos, afetam negativamente a probabilidade de ocupação desta preguiça,
272 enquanto áreas com presença de árvores importantes para a sua dieta tem influência positiva; 3)
273 cenários de regeneração natural podem mitigar os efeitos das mudanças climáticas na
274 adequabilidade ambiental para a preguiça. De modo geral, essa tese reforça a importância de
275 áreas florestais como preditoras da ocorrência da espécie, sendo locais com baixa cobertura
276 florestal inadequadas para a sua presença. Além disso, a disponibilização de dados de ocorrência

277 dos xenartros irá facilitar o desenvolvimento de diversas pesquisas e avaliações
278 conservacionistas voltadas à este grupo relativamente ainda pouco estudado. Os produtos aqui
279 gerados irão contribuir com metas e ações estabelecidos em planos de ações nacionais, visando a
280 conservação a longo-prazo da espécie.

281

282 **Palavras-chave:** Mata Atlântica, desmatamento, biodiversidade, ecologia de paisagem,
283 conservação

284 **Abstract**

285 According to the Intergovernmental Panel on Climate Change (IPCC), food production is one of
286 the main responsible for climate change. The conversion of natural areas into large agricultural
287 areas plays a fundamental role in global warming. Additionally, the habitat loss may interrupt
288 gene flows and important ecological processes. Therefore, changes in land use and its effects on
289 climate change correspond to one of the main causes of species extinction. Arboreal species are
290 the most affected by habitat loss, due to their high dependence on forests, as is the case of the
291 maned sloth (*Bradypus torquatus*), exclusive of the threatened Atlantic Forest. Thus, the main
292 objective of this present thesis was to understand the influence of environmental factors in the
293 occupancy probability of *B.torquatus* and to evaluate the current and future environmental
294 suitability for the species, besides contributing to the generation of open biodiversity data. The
295 first chapter sought to compile and make available occurrence data on the xenarthrans, in order
296 to contribute with several analyzes and ecological modeling. The second chapter aimed to
297 identify the influence of forest cover and other variables on the occupancy probability of
298 *B.troquatus*. The third chapter evaluated the environmental suitability for the species,
299 considering regeneration and climate change scenarios. As a result: 1) We compiled over 30,000
300 xenarthran records, from different institutions around the world and from different sources -
301 including data that until then were unavailable for public access; 2) The occupancy probability of
302 the maned sloth is 0.97, but it decreases abruptly at 35% of forest cover, reaching zero in areas
303 with less than 20% of forest cover. In addition, open areas cover, such as pastures, abandoned
304 pastures and exposed soil, has a negative effect on *B.torquatus* occupancy probability, whereas
305 areas with the presence of important trees to their diet affect positively; 3) Natural regeneration
306 can mitigate the effects of climate change on environmental suitability for maned sloth. Overall,
307 this thesis reinforces that forest areas are important predictors for species, and their low coverage
308 implies in unsuitable areas for their presence. Additionally, the provision of xenarthran data will
309 favor the development of several research and conservationist assessments focused on the group.
310 The products generated here will contribute to goals and actions established in national action
311 plans, aiming the long-term conservation of the species.

312

313 **Keywords:** Atlantic Forest, deforestation, biodiversity, landscape ecology, conservation

314

315 **Introdução geral**

316 De acordo com o último relatório do IPCC (Painel Intergovernamental sobre Mudanças
317 Climáticas; em inglês *Intergovernmental Panel on Climate Change 2019*), cerca de 56% das
318 áreas terrestres mundiais são utilizadas para fins econômicos (pecuária, agricultura, silvicultura,
319 extração de madeira e infraestrutura), e apenas 16% são ocupadas por vegetação nativa, entre
320 florestas e áreas naturais não-florestais. Essa intensa conversão de ambientes naturais em
321 extensas áreas antrópicas cria ambientes cada vez mais hostis para diversas espécies, ameaçando
322 fortemente a conservação da biodiversidade mundial (BELLARD *et al.*, 2014; HADDAD *et al.*,
323 2015; KOUMARIS; FAHRIG, 2016).

324 A quantidade de habitat remanescente possui um papel fundamental na dinâmica de
325 populações e comunidades (FAHRIG, 2013). Alguns grupos reagem linearmente à perda de
326 habitat (BECA *et al.*, 2017), enquanto outros exibem uma resposta não-linear (BOESING;
327 NICHOLS; METZGER, 2018; MUYLAERT; STEVEN; RIBEIRO, 2016). Este último
328 relaciona-se aos limiares críticos (*critical thresholds*), e seu conceito sugere que pequenas
329 modificações na proporção de habitat disponível podem levar a mudanças abruptas nas respostas
330 ecológicas (SWIFT; HANNON, 2010). Além da quantidade de habitat, as espécies –
331 principalmente as florestais – necessitam de áreas naturais adequadas e bem estruturadas para
332 desempenhar suas atividades diárias. Habitats complexos englobam uma estratificação florestal
333 bem definida e formada por muitos estratos (AUGUST, 1983). Sendo assim, fatores como altura
334 do dossel, abertura da copa, composição florística, estratificação florestal e complexidade de
335 habitat são essenciais para manter a estrutura e a viabilidade de populações e comunidades em
336 uma mancha de habitat (GRELLE, 2003; PAIM, FERNANDA P. *et al.*, 2017; PAIM,
337 FERNANDA POZZAN *et al.*, 2018; SMITH *et al.*, 2019).

338 Áreas florestais também são grandes responsáveis por estocar grande parte do carbono
339 mundial, e transformá-las em áreas agropastoris - liberando o carbono armazenado – cumprem
340 papel fundamental no aquecimento global (IPCC, 2018). Portanto, diminuir – ou zerar – o
341 desmatamento em larga escala é uma das formas mais eficazes de mitigar os efeitos das
342 mudanças climáticas. Além disso, torna-se necessário o estímulo à regeneração e à restauração
343 florestal, uma vez que árvores jovens em crescimento são importantes sumidouros de carbono
344 (CASPERSEN *et al.*, 2000). Além de ajudar na regulação climática, o crescimento das áreas
345 florestais viabiliza a manutenção conservação da biodiversidade (METZGER *et al.*, 2019).

346 Alguns mecanismos conservacionistas se destacam para frear a perda de habitat e garantir
347 a preservação de áreas naturais, como por exemplo o estabelecimento de áreas protegidas. No
348 Brasil, as áreas protegidas são conhecidas por Unidades de Conservação e estão garantidas por
349 lei. O Sistema Nacional de Unidades de Conservação (SNUC – Lei 9,985/2000) foi instituído
350 como forma de potencializar o papel das diferentes categorias de UCs na conservação da
351 biodiversidade. Complementando as funções das UCs no Brasil, a Lei de Preservação da
352 Vegetação Nativa (LPVN, Lei 12,651/2012) estabelece a proteção de áreas ambientais áreas
353 sensíveis, como topos e encostas de morros, nascentes e margens de rios, lagos e lagoas – as
354 chamadas Áreas de Preservação Permanente (APPs), além da proteção de áreas de vegetação
355 nativa localizadas em propriedades particulares – Reservas Legais (RL). Tais mecanismos visam
356 a preservação de áreas não englobadas pelo SNUC, de forma a garantir diversos serviços
357 ecossistêmicos. Porém, de modo geral, algumas dessas áreas definidas e protegidas vêm sendo
358 constantemente ameaçadas por propostas de leis que flexibilizam a supressão vegetal para fins
359 agrícolas – no caso das RLs (Projeto de Lei n. 2362/2019, o qual já fora retirado de votação do
360 Senado Federal), ou seja por propostas de leis que permitem a abertura e uso de estradas dentro

361 de Parques Nacionais (Projetos de Lei 984/2019 - em trâmite - e 61/2013 - este último foi
362 barrado pelo Superior Tribunal Federal recentemente).

363 A Mata Atlântica brasileira ocupa hoje um território de 1,08 milhão km² (MUYLAERT
364 *et al.*, 2018). Com sua distribuição restrita a pouco mais de 28% da cobertura original
365 (REZENDE *et al.*, 2018), o bioma possui um alto déficit de Unidades de Conservação, APPs e
366 Reservas Legais (SOARES-FILHO *et al.*, 2014). Considerado um *hotspot* para a conservação da
367 biodiversidade (MYERS *et al.*, 2000), diversas espécies endêmicas e ameaçadas habitam a área
368 do bioma, e a deficiência das áreas legalmente protegidas pode agravar a situação de ameaça.
369 Dentre as espécies endêmicas, encontra-se a Preguiça-de-Coleira *Bradypus torquatus* (Pilosa:
370 Bradypodidae). Habitando quase que exclusivamente florestas ombrófilas densa, entre os estados
371 de Sergipe e Rio de Janeiro (HIRSCH; CHIARELLO, 2012), a Preguiça-de-Coleira se alimenta
372 basicamente folhas e brotos (CHIARELLO, 1998b; MONTGOMERY; SUNQUIST, 1975), fato
373 intrinsecamente relacionado à baixa taxa metabólica (NAGY; MONTGOMERY, 1980). Em
374 função de seu hábito estritamente arborícola e folívoro, a espécie é profundamente afetada pela
375 fragmentação, perda e destruição do habitat (CHIARELLO *et al.*, 2018; CHIARELLO;
376 MORAES-BARROS, 2014; SANTOS *et al.*, 2019). Tais fatores, associados a uma restrita e
377 descontínua distribuição tornam essa espécie ameaçada, e atualmente ela está classificada como
378 vulnerável tanto na lista nacional (CHIARELLO *et al.*, 2018), quanto na internacional
379 (CHIARELLO; MORAES-BARROS, 2014) de espécies ameaçadas. A preguiça-de-coleira está,
380 atualmente, no Plano de Ação Nacional para Conservação dos Primatas e Preguiça da Mata
381 Atlântica (PPMA), o qual institui algumas metas e ações para a conservação da preguiça. Essa
382 tese contribui diretamente na execução de importantes ações presentes no PPMA.

383 As preguiças pertencem à superordem Xenarthra juntamente com os tamanduás e
384 tamanduá – que compõe a Ordem Pilosa - e os tatus – da Ordem Cingulata (GIBB *et al.*, 2016).
385 Os xenartros é um dos grupos de mamíferos mais antigos existentes – sendo um dos quatro
386 clados basais dos mamíferos placentários (DELSUC *et al.*, 2002). Ademais, é o único grupo com
387 origem na América do Sul, onde sua história teve início, por volta de 65 milhões de anos atrás
388 (GIBB *et al.*, 2016; porém ver DELSUC; DOUZERY, 2008), e hoje conta com 13 gêneros
389 existentes – a grande maioria pertencentes aos tatus (SANTOS *et al.*, 2019).

390 Embora os xenartros possam prover enormes descobertas acerca da evolução dos
391 primeiros mamíferos placentários, de forma geral, muitos aspectos do grupo permanecem sem
392 respostas (SUPERINA; LOUGHRY, 2015). Os xenartros podem ser bastante críticos ao
393 ambiente em que vivem, tornando difíceis pesquisas que abordem parâmetros demográficos.
394 Felizmente, a ciência tem avançado de forma contínua, permitindo o desenvolvimento de
395 modelos estatísticos e matemáticos que possibilitam construir análises levando em consideração
396 a baixa detectabilidade das espécies em seu meio natural. A modelagem de ocupação é uma
397 alternativa para avaliar aspectos ecológicos e biológicos de espécies crípticas. Tal abordagem
398 estima a proporção da área, de fragmentos ou de unidades de amostra que é ocupada, corrigida
399 pela detecção imperfeita ou falsa ausência, i.e., a espécie pode estar lá, mas não foi possível
400 detectá-la, em consequência de diversos fatores, como comportamento das espécies ou
401 características do habitat (MACKENZIE *et al.*, 2006).

402 Adicionalmente, por meio da obtenção de dados já existentes em banco de dados e
403 bibliotecas virtuais, é possível desenvolver outros tipos de modelos que forneçam importantes
404 informações ecológicas acerca de um grupo. Aqui inclui-se a Modelagem de Distribuição de
405 Espécies (em inglês *Species Distribution Models* - SDM). Tal abordagem fundamenta-se

406 ecologicamente no conceito de nicho ecológico - o “*hiper-volume n-dimensional*”
407 (HUTCHINSON, 1957), ou o conjunto de condições e recursos que permitem à espécie suprir
408 suas necessidades mínimas para sobreviver (CHASE; LEIBOLD, 2003). Dessa forma, o SDM é
409 capaz de prever áreas potenciais para a distribuição de uma espécie ou avaliar áreas adequadas
410 para um grupo (FERRAZ *et al.*, 2012; GUISAN *et al.*, 2013; JOSE V; NAMEER, 2020) através
411 de dados de ocorrência e de variáveis ambientais, acessando a relação entre a distribuição de uma
412 determinada espécie e as condições ambientais (MARTÍNEZ-MEYER *et al.*, 2006; PETERSON,
413 2006), com amplo uso biológico (MORATO *et al.*, 2014; PORTUGAL *et al.*, 2019).

414 Diversas plataformas on-line, como *SpeciesLink* (<http://splink.cria.org.br/>), GBIF
415 (<https://www.gbif.org/>) e GIFT (<http://gift.uni-goettingen.de/home>), disponibilizam dados de
416 biodiversidade, os quais podem ser utilizados em variadas análises. Se somam a essas
417 plataformas os projetos de *datapapers* das séries ATLANTIC SERIES e NEOTROPICAL
418 SERIES, gerados e publicados nos três últimos anos. Ao compilar, integrar e disponibilizar
419 dados diversos de biodiversidade (muitos deles não publicados e presentes apenas em
420 dissertações, teses e relatórios de consultorias) de acesso gratuito e livre, os *datapapers* integra-
421 se ao *Open data movement* (tradução livre: movimento (ou iniciativa) por dados abertos) que
422 promove a disponibilização e uso de dados científicos de forma livre e irrestrita (BAACK, 2015).

423 Utilizando modelos de ocupação e modelos de distribuição de espécies, a presente tese
424 buscou entender a influência de fatores locais e espaciais sobre a probabilidade de ocupação da
425 preguiça-de-coleira e avaliar a adequabilidade ambiental presente e futura para a espécie ao
426 longo de sua distribuição. Essa tese buscou, ainda, a compilação e a disponibilização de dados
427 biológicos, se alinhando ao *open data movement*. A estrutura da tese se encontra dividida em três
428 capítulos. O primeiro capítulo – já publicado no periódico internacional *Ecology* - consistiu em

429 reunir, padronizar e disponibilizar dados de ocorrência da superordem Xenarthra em toda região
430 Neotropical (SANTOS *et al.*, 2019). Como resultado, compilamos mais de 35 mil registros de
431 ocorrência, distribuídos desde o Sul da Argentina/Chile, até a região Sudeste do Estados Unidos.
432 O *datapaper* contou com colaboração de 505 pesquisadores de 282 instituições de várias partes
433 do mundo que nos enviaram dados de xenartros de diversos, anos, tipo e locais. Além dos dados
434 enviados pelos colaboradores, foram compilados também dados de literatura – artigos, tese,
435 dissertação, relatórios. Constatou-se que dados de xenartros já publicados e disponíveis
436 correspondem a menos de 20% do *dataset*, e, portanto, o *datapaper* está sendo fundamental na
437 integração e disponibilização de dados exclusivos. O presente capítulo se junta a iniciativas
438 como ATLANTIC CAMTRAP (LIMA, F. *et al.*, 2017), ATLANTIC BATS (MUYLAERT *et*
439 *al.*, 2017) , ATLANTIC PRIMATES (CULOT *et al.*, 2018) e ATLANTIC SMALL MAMMALS
440 (BOVENDORP *et al.*, 2017), sendo o primeiro da série NEOTROPICAL a ser publicado
441 (SANTOS *et al.*, 2019).

442 O segundo capítulo - publicado no periódico internacional *Biological Conservation* -
443 buscou identificar e compreender como variáveis locais e espaciais influenciam a probabilidade
444 de ocupação da preguiça-de-coleira. Para isso, foram realizadas campanhas de campo a fim de se
445 detectar a espécie e coletar variáveis locais, em mais de 60 pontos amostrais localizados na
446 região Serrana do Espírito Santo. Por meio de ferramentas de SIG, realizamos análises espaciais
447 para calcular a porcentagem de floresta e de áreas abertas, utilizando a abordagem de multi-
448 escala. Por meio de seleção de modelos, selecionamos a escala de efeito mais apropriada e
449 utilizamos a modelagem de ocupação para identificar limiares críticos para a presença da
450 preguiça-de-coleira (SANTOS *et al.*, 2019).

451 Já o terceiro capítulo consiste na utilização de Modelos de Distribuição de Espécies e dos
452 dados compilados nos dois capítulos anteriores para avaliar a adequabilidade ambiental presente
453 e futura para a preguiça-de-coleira ao longo de sua distribuição. Para isso, levamos em
454 consideração um cenário atual e três cenários futuros relacionados à porcentagem de cobertura
455 florestal, e em cada cenário incluímos variáveis climáticas em diferentes cenários de mudanças
456 climáticas. Particularidades genéticas das preguiças-de-coleira foram consideradas, analisando as
457 duas Unidades Evolutivas Significativas separadamente.

458

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577 **Capítulo 1**

578 **NEOTROPICAL XENARTHANS: a dataset of occurrence of xenarthran species in the**
579 **Neotropics**

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598 **Introduction**

599 The superorder Xenarthra (*xenon* = stranger; *arthros* = articulations) is one of the four
600 major clades of placental mammals (Madsen et al. 2001, Delsuc et al. 2002, Delsuc and Douzery,
601 2008). This group, which includes the orders Cingulata (armadillos) and Pilosa (anteaters and
602 sloths), began its radiation in South America and later dispersed to Central and North America
603 through the establishment of land bridges (Vizcaíno and Loughry, 2008). Molecular dating
604 estimates Xenarthra split into Cingulata and Pilosa about 67 million years ago (Gibb et al. 2016).
605 However, molecular dating estimate that the group originated earlier, approximately 105 million
606 years ago (Springer 2003, Delsuc et al. 2004, Delsuc and Douzery 2008).

607 Xenarthrans encompass more than 150 described genera through Earth's history, including
608 recently extinct taxa such as glyptodonts and giant ground sloths (Fariña 1996). This number
609 decreased to only 13 extant genera, represented by 37 species (21 armadillos, six sloths, ten
610 anteaters) classified in six families: Dasypodidae (long-nosed armadillos), Chlamyphoridae (fairy,
611 hairy, three-banded, naked-tailed, and giant armadillos), Myrmecophagidae (lesser and giant
612 anteaters), Cyclopedidae (pygmy anteaters), Bradypodidae (three-toed sloths), and
613 Megalonychidae (two-toed sloths) (Abba et al. 2015, Gibb et al. 2016, Feijó and Cordeiro-Estrela
614 2016, Miranda et al. 2017, Feijó et al. 2018). They are distributed from the southeastern United
615 States of America to southern Patagonia, with more diversity in South America (Gardner 2008,
616 Vizcaíno and Loughry 2008, Taulman and Robbins 2014).

617 Each family has very distinctive morphological and ecological characteristics, making
618 Xenarthra a unique and morphologically diverse group. The bodies of armadillos are covered by a
619 carapace comprised of mobile articulated plates, which offers protection against predators
620 (McDonough and Loughry 2008). Anteaters possess a protrusible tongue and have a long and

621 tubular skull characterized by the complete loss of teeth, as well as large front claws used to feed
622 on ants and termites (Wilson and Mittermeier 2018). Three-toed sloths have three long digits on
623 each forelimb, whereas two-toed sloths only have two. Both are strictly arboreal and possess long
624 claws that they use to hang on to tree branches. Most mammals have seven cervical vertebrae;
625 however, the living sloths have a variable number: 5–8 in two-toed sloths (*Choloepus*) and 8–10
626 in three-toed sloths (*Bradypus*) (Buchholtz and Stepien 2009).

627 Besides their unique anatomical structures and ecology, the xenarthrans have an
628 exceptional physiology, with a lower metabolism than expected from similar-sized mammals,
629 and variable body temperature, characteristics that may be related to a low energy intake diet and
630 the ingestion of non-nutritive material during feeding (McNab 1984). Their diets are mainly
631 based on ants and termites (anteaters and armadillos), arthropods (armadillos), and leaves
632 (sloths). The living sloths are arboreal, feeding on leaves from trees, vines, and lianas, and their
633 low metabolism might assist in reducing the absorption of toxic leaves (Gilmore et al. 2001).
634 Armadillos use burrows as shelter and to buffer their body temperature from environmental
635 variation (Attias et al. 2018), as well as the Anteaters (Camilo-Alves and Mourão, 2006; Mourão
636 and Medri 2007). Additionally, xenarthrans play important ecological roles. Armadillos act as
637 ecosystem engineers with many species associated with their burrows (Desbiez and Kluyber
638 2013, Aya-Cuero et al. 2017). Armadillos and anteaters impact ant and termite populations
639 contributing, in turn, to the structure and diversity of plant communities (Rao 2000, Terborgh et
640 al. 2006). Sloths are important components of the arboreal vertebrate biomass, being top prey for
641 large raptors and contributing to the nutrient cycling of tropical forests. Their pelage is also the
642 substrate to a diverse array of microorganisms and invertebrates (Montgomery and Sunquist
643 1978; Higginbotham et al. 2014).

644 According to the International Union for Conservation of Nature (IUCN), five xenarthran
645 species are at some risk of extinction – *Bradypus pygmaeus*, *Bradypus torquatus*,
646 *Myrmecophaga tridactyla*, *Priodontes maximus*, and *Tolypeutes tricinctus*. Another four are near
647 threatened – *Cabassous chacoensis*, *Dasypus sabanicola*, *Tolypeutes matacus*, and *Zaedyus*
648 *pichiy* – all showing decreasing population trends. Additionally, because of recent taxonomic
649 revisions, three species of *Dasypus* and seven species of *Cyclopes* have pending conservation
650 status evaluations (Feijó and Cordeiro-Estrela 2016, Miranda et al. 2017, Feijó et al. 2018). For
651 all xenarthran species, the major threats are habitat loss resulting from fragmentation (Chiarello
652 and Moraes-Barros 2014, ICMBio 2015), domestic and feral dog attacks, roadkill (Chiarello and
653 Moraes-Barros 2014, Ascensão et al. 2017), subsistence hunting, illegal capture (ICMBio 2015),
654 and fires (Miranda et al. 2014).

655 Despite the increasing interest in living xenarthran species (Diniz and Brito 2012,
656 Superina et al. 2014), they remain a relatively understudied group when compared to other
657 groups of mammals (Superina and Loughry 2015). Therefore, many aspects of their ecology are
658 data deficient, making the correct assessment of population trends a difficult task. Conservation
659 of xenarthrans is becoming more pressing as this taxon faces increasing negative impacts from
660 anthropogenic actions. The Anteater, Sloth and Armadillo Specialist Group is a network of
661 mammalogists and conservationists within the IUCN. Its main goal is to comprehend and
662 provide information to aid in conservation action for those species. There are a growing number
663 of specific xenarthran conservation initiatives, including rehabilitation, education, monitoring
664 programs, and scientific research, supported by non-governmental organizations such as
665 Fundacion Aiunau (www.aiunau.org/), Colombia; Asociación Panamericana para la
666 Conservación (<http://www.appcpanama.org/>), Panama; Instituto de Pesquisa e Conservação de

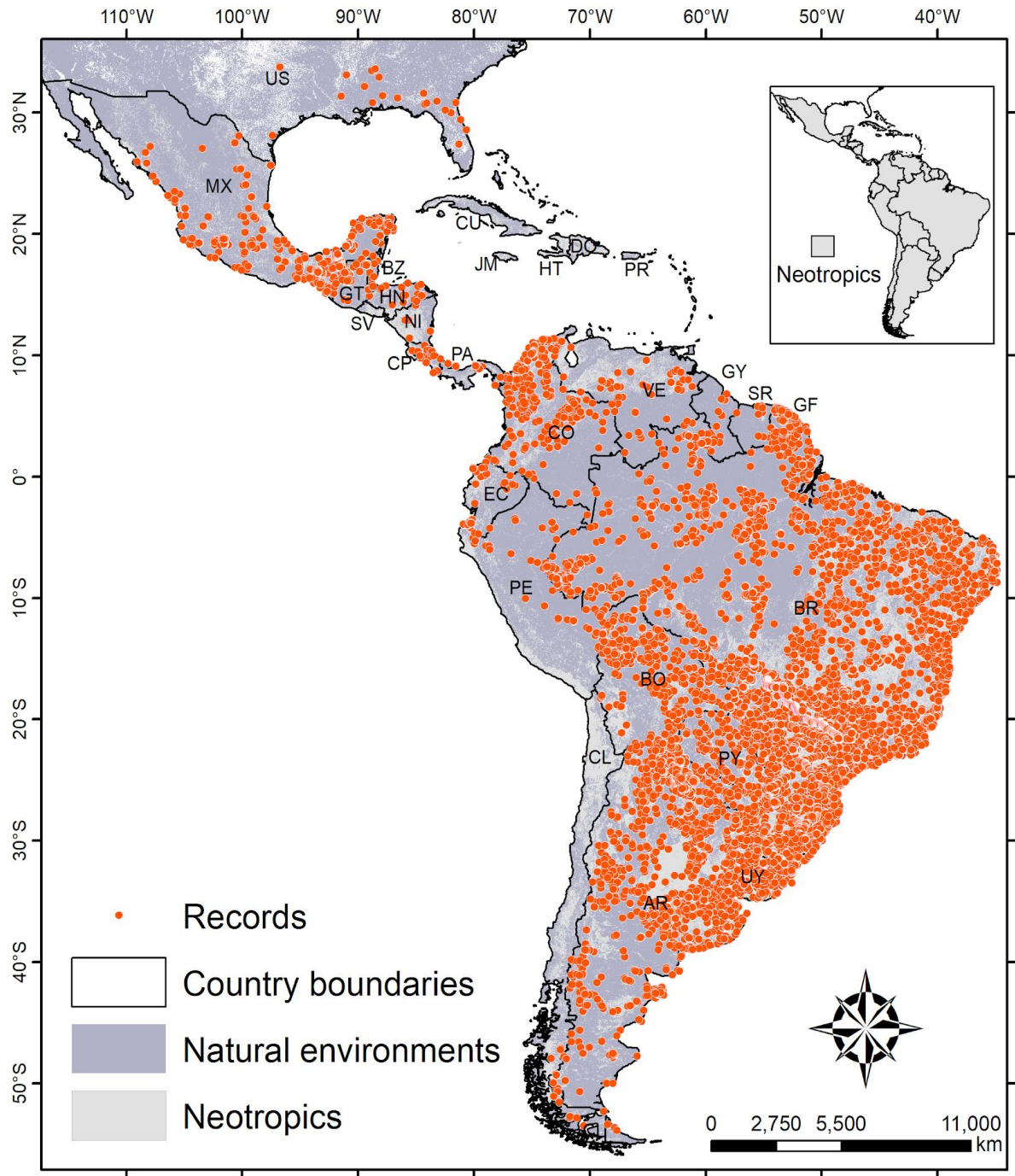
667 Tamanduás do Brazil; Projeto Tamanduá (<http://www.tamandua.org/>), Brazil; Projeto Bandeiras
668 e Rodovias (www.tamanduabandeira.org), Brazil; Proyecto Iberá - Oso Hormiguero,
669 (http://www.proyectoibera.org/especiesamenazadas_osohormiguero.htm), Argentina; Proyecto
670 Hormigueros (<https://www.cunaguaro.co/proyecto-hormigueros>), Colombia; Programa Tatu-bola
671 (<http://tatubola.org.br/>), Brazil; Armadillos de Chile (<http://www.armadilloschile.cl/>), Chile; The
672 Sloth Conservation Foundation (<https://slothconservation.com/>), United Kingdom; and Projeto
673 Tatu-canastra (www.icasconservation.org.br/o), Brazil. Nonetheless, relatively few species are
674 considered in those programs (such as *Myrmecophaga tridactyla*, *Priodontes maximus*, *Cyclopes*
675 sp. and *Tolypeutes tricinctus*).

676 Most xenarthran data are unpublished or have been published in the grey literature,
677 including reports, Master and PhD theses in different languages, with low accessibility to the
678 broad scientific community. In an effort to gather this dispersed and unavailable data in one
679 place, we compiled published and unpublished data on xenarthrans, from the years 1855 to 2018,
680 including occurrence and quantitative data. A total of 24,847 unique georeferenced records were
681 assembled, ranging from the northern frontier of the Neotropics (Mexico, southern USA, and
682 Caribbean regions), through to the austral south of South America (Chile, Argentina, Uruguay,
683 and Paraguay). Neotropical Xenarthrans is by far the largest dataset ever assembled with respect
684 to armadillos, sloths, and anteaters.

685 This work is part of the Neotropical, Atlantic, Brazil, and Amazonia series initiative, the
686 objective of which is to make data on the biodiversity of these regions publicly available. Until
687 now, the following data-papers of these series have been published: Atlantic Frugivory (Bello et
688 al. 2017), Atlantic Camtraps (Lima et al. 2017), Atlantic Small Mammals (Bovendorp et al.
689 2017), Atlantic Bats (Muylaert et al. 2017), Atlantic Birds (Hasui et al. 2018), Atlantic Mammal

690 Traits (Gonçalves et al. 2018), Atlantic Amphibians (Vancine et al. 2018), Non-volant mammals
691 from the Upper Paraná River Basin (Gonçalves et al. 2018), Atlantic Primates (Culot et al.
692 2018), Jaguar GPS movement (Morato et al. 2018), and Brazil roadkill (Grillo et al. 2018).

693



694
 695 **Fig. 1: Distribution of the Xenarthra records of the NEOTROPICAL XENARTHTRANS**
 696 **dataset in Neotropical realm.** Records are from years of 1855 to 2018; several populations may
 697 thus be extinct today.

698

699 **METADATA**

700 **Class I - Data set descriptors**

701 **A. Data set identity**

702 **Title:** NEOTROPICAL XENARTHTRANS: A DATASET OF OCCURRENCE OF
703 XENARTHTRANS SPECIES IN THE NEOTROPICS

704

705 **B. Data set and metadata identification code**

706 **Suggested data set identity codes:**

707 NEOTROPICAL_XENARTHTRANS_QUANTITATIVE.csv

708 NEOTROPICAL_XENARTHTRANS_QUALITATIVE.csv

709 NEOTROPICAL_XENARTHTRANS_REFERENCES.csv

710

711 **C. Data set description**

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733 **Abstract:**

734 Xenarthrans – anteaters, sloths, and armadillos – have essential functions for ecosystem
735 maintenance, such as insect control and nutrient cycling, playing key roles as ecosystem
736 engineers. Because of habitat loss and fragmentation, hunting pressure, and conflicts with
737 domestic dogs, these species have been threatened locally, regionally, or even across their full
738 distribution ranges. The Neotropics harbor 21 species of armadillos, ten anteaters, and six sloths.
739 Our dataset includes the families Chlamyphoridae (13), Dasypodidae (7), Myrmecophagidae (3),
740 Bradypodidae (4), and Megalonychidae (2). We have no occurrence data on *Dasyus pilosus*

741 (Dasypodidae). Regarding Cyclopedidae, until recently, only one species was recognized, but
742 new genetic studies have revealed that the group is represented by seven species. In this data-
743 paper, we compiled a total of 42,528 records of 31 species, represented by occurrence and
744 quantitative data, totaling 24,847 unique georeferenced records. The geographic range is from
745 the south of the USA, Mexico, and Caribbean countries at the northern portion of the Neotropics,
746 to its austral distribution in Argentina, Paraguay, Chile, and Uruguay. Regarding anteaters,
747 *Myrmecophaga tridactyla* has the most records (n=5,941), and *Cyclopes* sp. has the fewest
748 (n=240). The armadillo species with the most data is *Dasypus novemcinctus* (n=11,588), and the
749 least recorded for *Calyptophractus retusus* (n=33). With regards to sloth species, *Bradypus*
750 *variegatus* has the most records (n=962), and *Bradypus pygmaeus* has the fewest (n=12). Our
751 main objective with Neotropical Xenarthrans is to make occurrence and quantitative data
752 available to facilitate more ecological research, particularly if we integrate the xenarthran data
753 with other datasets of Neotropical Series which will become available very soon (i.e. Neotropical
754 Carnivores, Neotropical Invasive Mammals, and Neotropical Hunters and Dogs). Therefore,
755 studies on trophic cascades, hunting pressure, habitat loss, fragmentation effects, species
756 invasion, and climate change effects will be possible with the Neotropical Xenarthrans dataset.

757

758 **D. Key words**

759 Xenarthra, Pilosa, Cingulata, Biodiversity Hotspot, Neotropical region, Neotropical mammals,
760 forest fragmentation, habitat loss

761

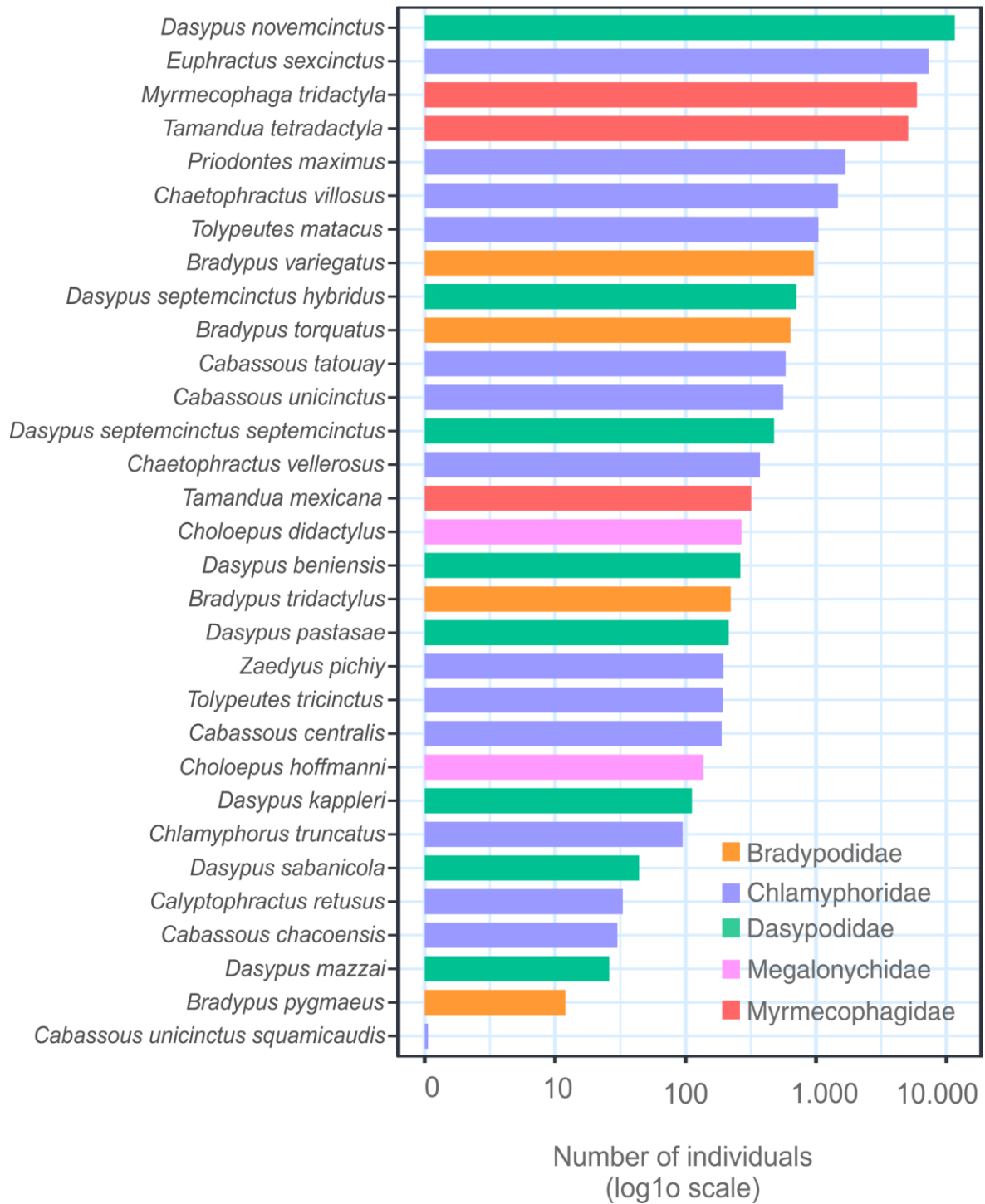
762 **E. Description**

763 To organize the dataset into the Neotropical realm, we selected all the countries and islands
764 indicated in the literature as the Neotropical area and merged all polygons into a single shapefile.
765 Our study extent includes the southern USA, Mexico, Central America, the Caribbean islands
766 and South America (Figure 1). The dataset is comprised of 42,528 records (Figure 1; Table 1),
767 including 40,794 (95.92%) with taxonomic certainty down to species-level identification, and
768 1,734 (4.08%) records of six genera with taxonomic uncertainty, with the majority of those
769 records belonging to *Dasypus* sp. (1,359 records, 78.37%). In some cases, we taxonomically
770 corrected some of those uncertain records, based on the distribution, on available literature, or on
771 evidence material sent to us by the authors and after consulting specialists (more details, Table
772 2). When correction was not possible, we opted to maintain only the genus. A special case
773 includes the genus *Cyclopes* sp. (240 records): at the beginning of this study, the genus was
774 monospecific, *Cyclopes didactylus*. In the middle of the process, the taxon was subjected to a
775 taxonomic review, which split the previously single species into seven (Miranda et al. 2017). To
776 avoid misinterpretation of the data, we decided to maintain *Cyclopes* sp.

777 The two armadillo families appear most frequently in terms of both the number of records and
778 species; Dasypodidae is responsible for the most records (n=14,790), followed by
779 Chlamyphoridae (n=13,887), and they are represented by seven (two subspecies) and 13 species,
780 respectively (Figures 3–5). Two of these species are the most common in the dataset (Figure 2):
781 *Dasypus novemcinctus* (Dasypodidae, n=11,588) and *Euphractus sexcinctus* (Chlamyphoridae,
782 n=7,325), both of which are species with wide distributions (Figures 3 and 4, respectively). Only
783 one species, *Cyclopes* sp., represents the family Cyclopedidae (Figure 9; see the recent
784 taxonomic review above). The other anteater species, Myrmecophagidae, is the third-most
785 numerous, (n=11,349), represented by its three species *Myrmecophaga didactyla* (n=5,941),

786 *Tamandua tetradactyla* (n=5,089), and *Tamandua mexicana* (n=319). The two sloth families are
787 the least numerous, mainly the Megalonychidae, with only 414 records (Figure 7). *Bradypus*
788 *variegatus*, from the family Bradypodidae, is the most-common sloth species, with 962 records
789 (Figure 8). *Bradypus pygmaeus*, also from the family Bradypodidae, is the least-common species
790 (n=12), both among sloth species and the dataset as a whole, reflecting its restricted insular
791 distribution (Figure 8).

792 Approximately 99.94% of the records are from native species, and only 0.06% are from
793 introduced (*Priodontes maximus*) and potentially introduced (*Bradypus* spp.) species. According
794 to the most-recent IUCN Red List classification (IUCN 2018) and considering taxonomically
795 confirmed species and the genus *Cyclopes*, 29,731 records (72.45%, 14 species and the genus
796 *Cyclopes*) are from Least Concern (LC) species, 2,020 (4.92%, five species) are from Near
797 Threatened (NT) species, 8,451 (20.6%, four species) are from Vulnerable (VU) species, and 12
798 records (0.03%, one species) are from Critically Endangered (CR) species. An additional 343
799 records (0.84%, four species) are from Data Deficient (DD) species, and another 477 records
800 (1.16%, two species) are from Not Evaluated (NE) species (Table 1).

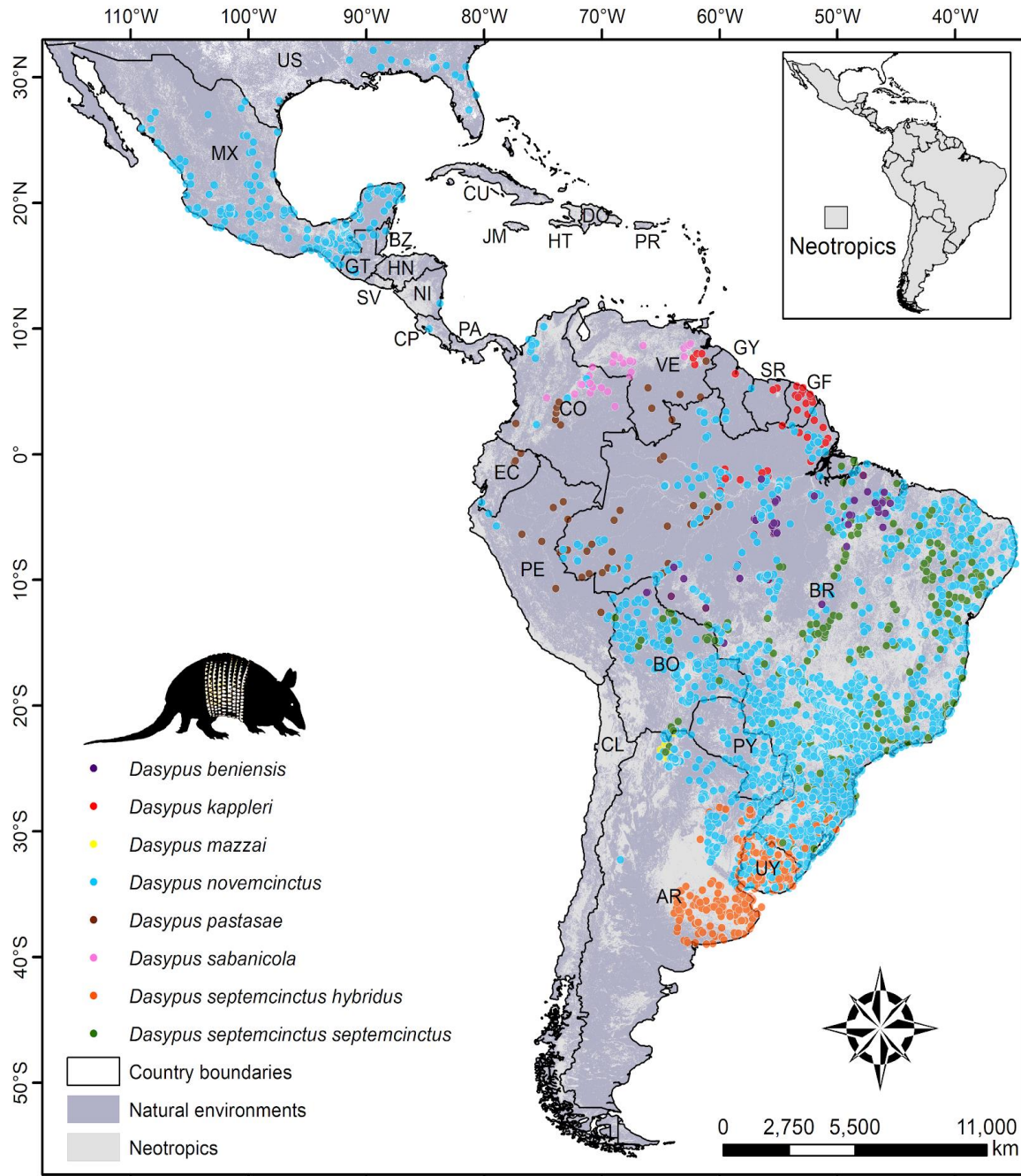


801

802 **Fig. 2: Number of records per species of armadillos, anteaters and sloths within**

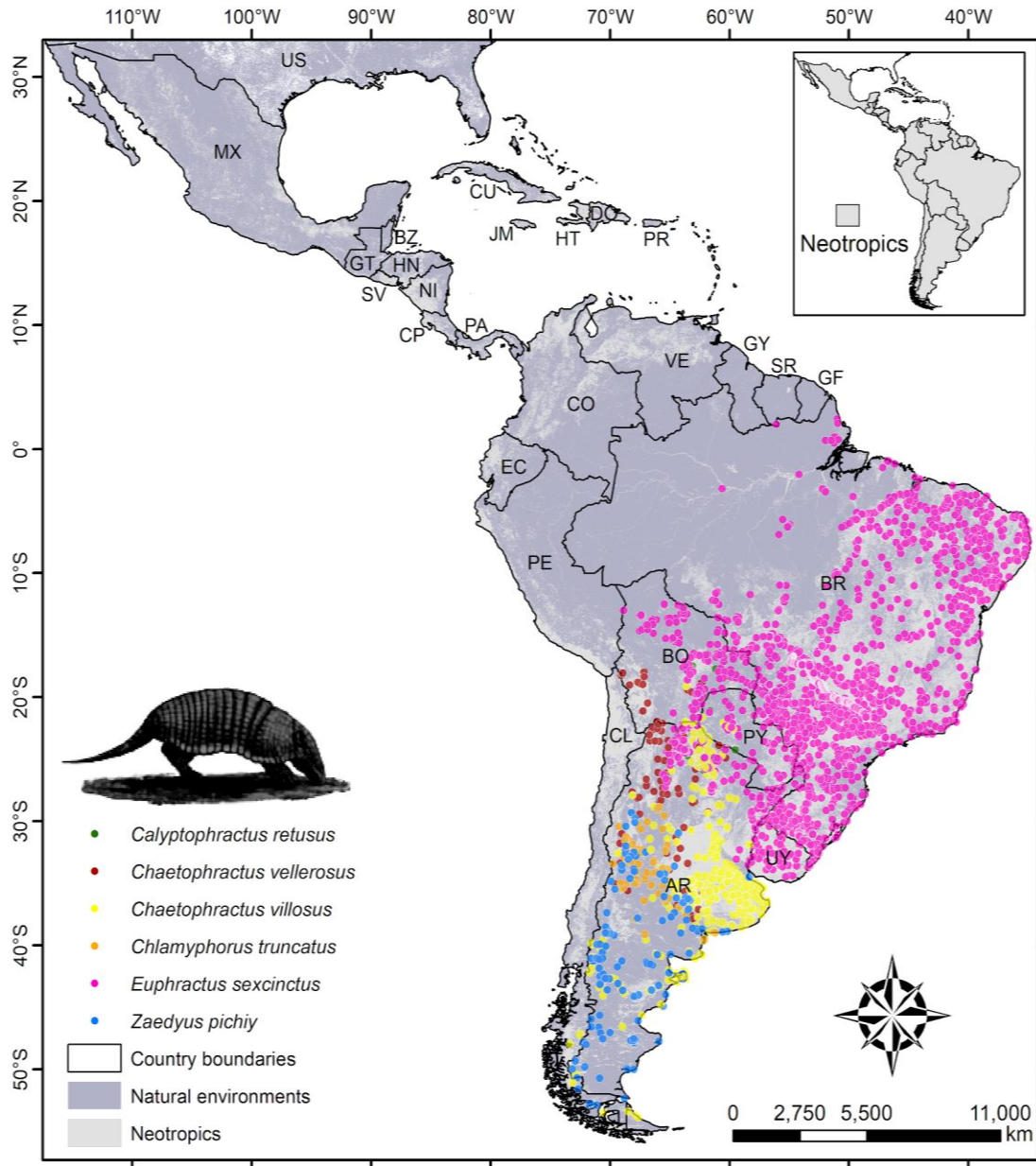
803 **NEOTROPICAL XENARTHANS dataset.** Species with unconfirmed identification are not

804 represented. Each color refers to a xenarthran family.



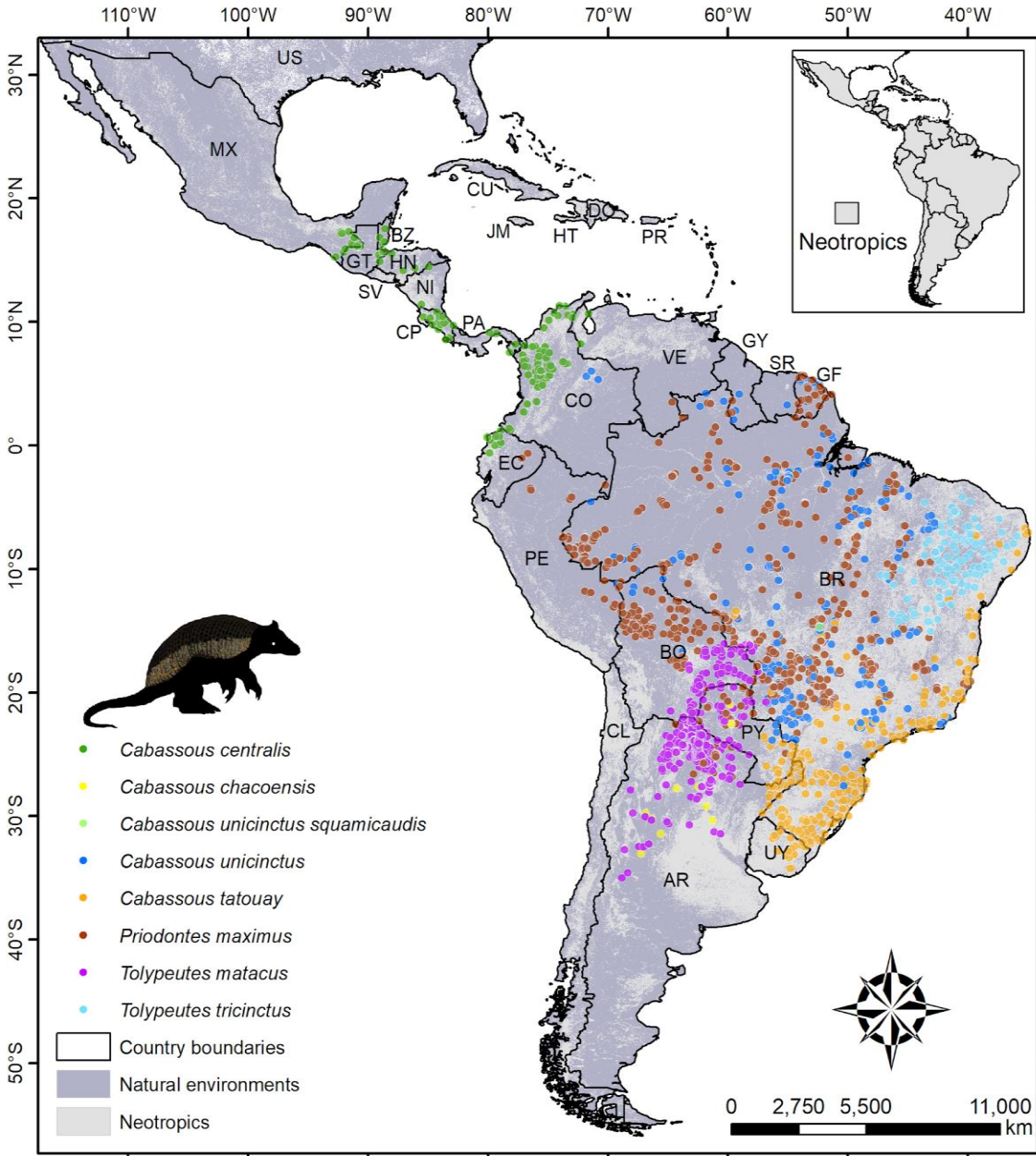
805

806 **Fig. 3: Distribution of the records of the family Dasypodidae of the NEOTROPICAL**
 807 **XENARTHANS dataset in Neotropical realm.** Records are from 1896 to 2018, thus several
 808 populations may now be extinct. Unconfirmed species with “sp.” are not included.



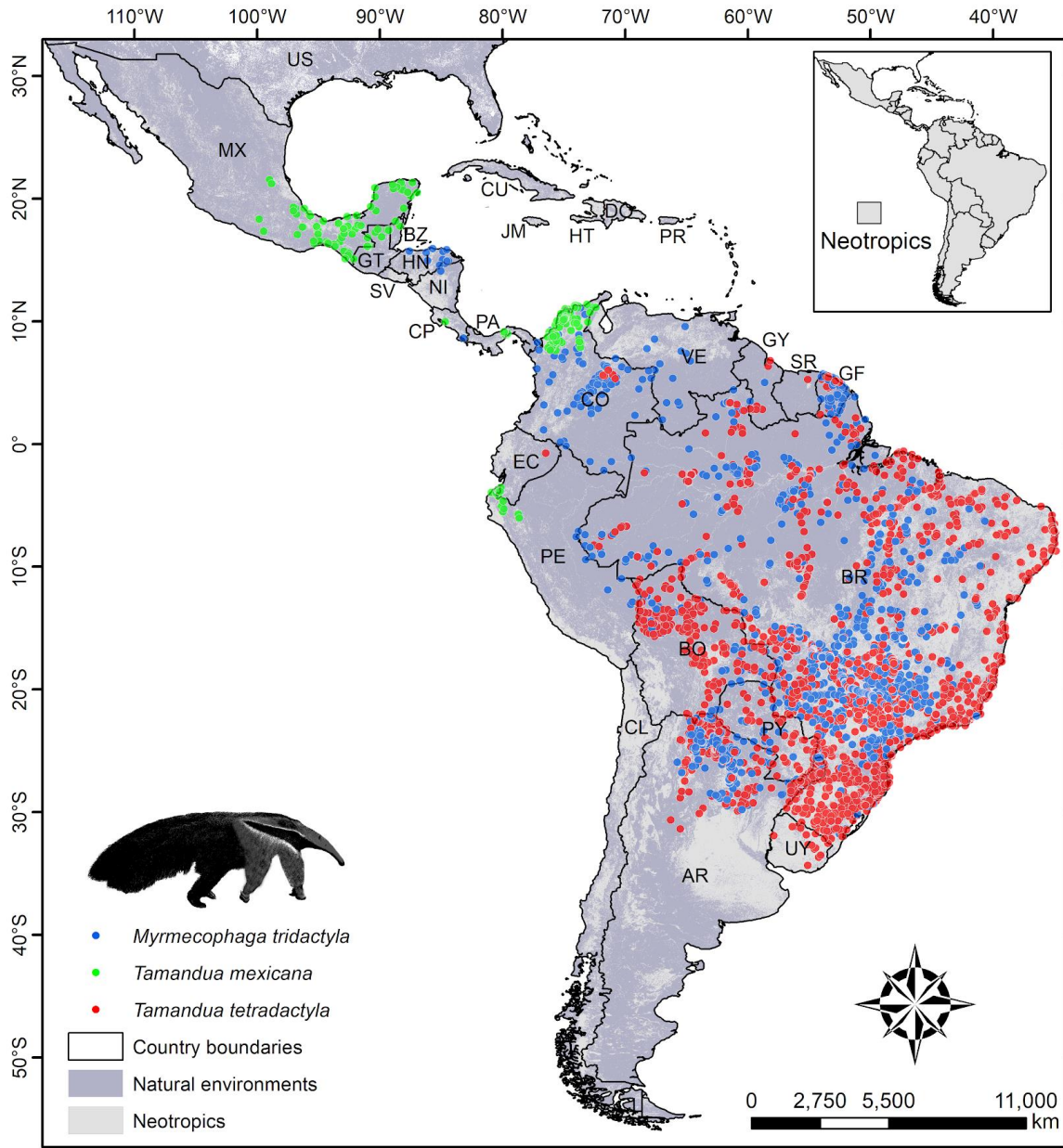
809

810 **Fig. 4: Distribution of the records of the family Chlamyphoridae (Subfamilies**
 811 **Chlamyphoniae and Euphractinae) of the NEOTROPICAL XENARTHTRANS dataset in**
 812 **Neotropical realm.** Records are from 1910 to 2018, thus several populations may now be
 813 extinct. Unconfirmed species with “sp.” are not included. Additional species in this family are
 814 shown in Figure 5.



815

816 **Fig. 5: Distribution of the records of the family Chlamyphoridae (Subfamily Tolypeutinae)**
 817 **of the NEOTROPICAL XENARTHTRANS dataset in Neotropical realm.** Records are from
 818 1855 to 2018, thus several populations may now thus be extinct. Unconfirmed species with “sp.”
 819 are not included.

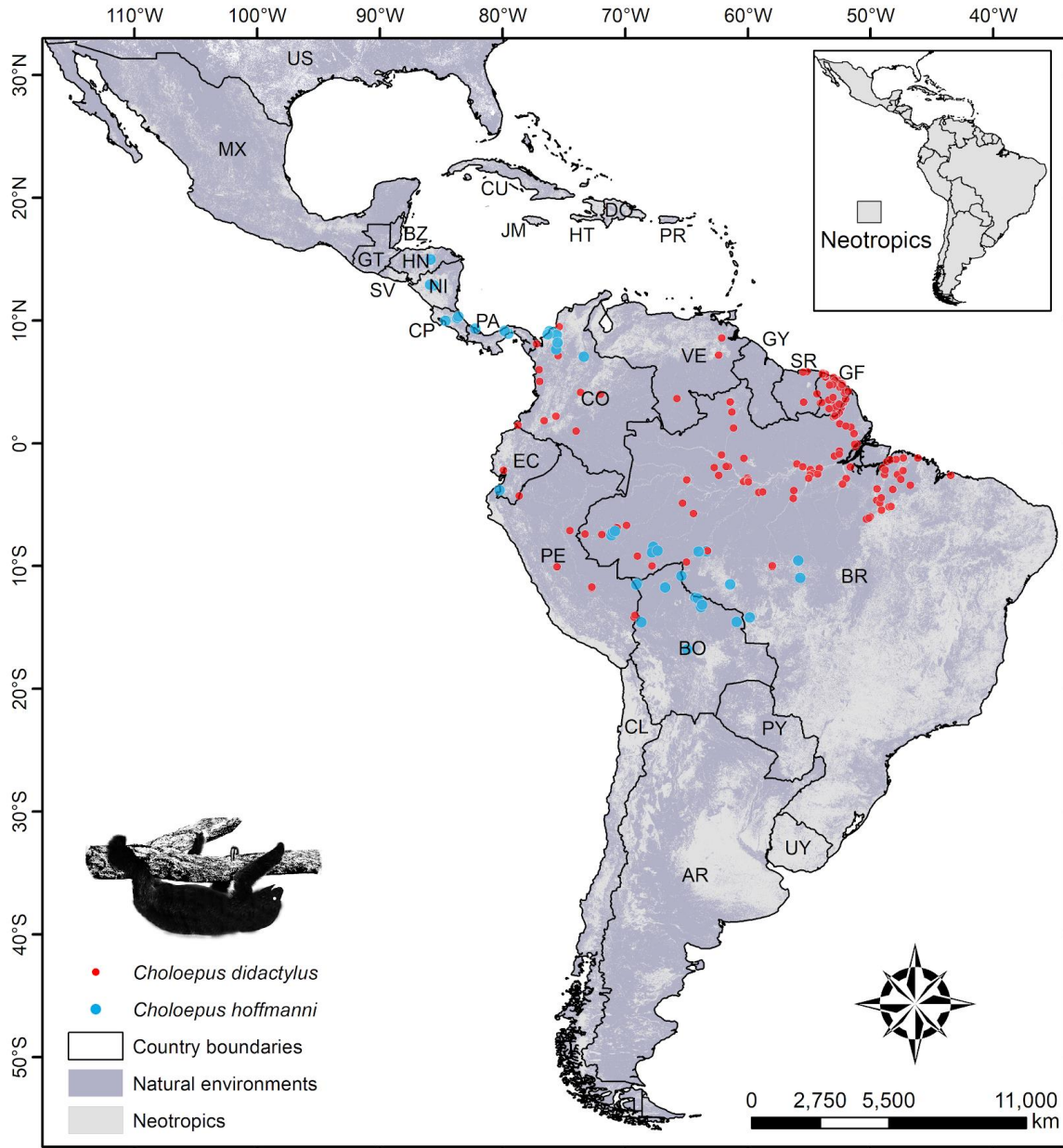


820

821 **Fig. 6: Distribution of the records of the family Myrmecophagidae of the NEOTROPICAL**

822 **XENARTHANS dataset in Neotropical realm.** Records are from 1899 to 2018, thus several

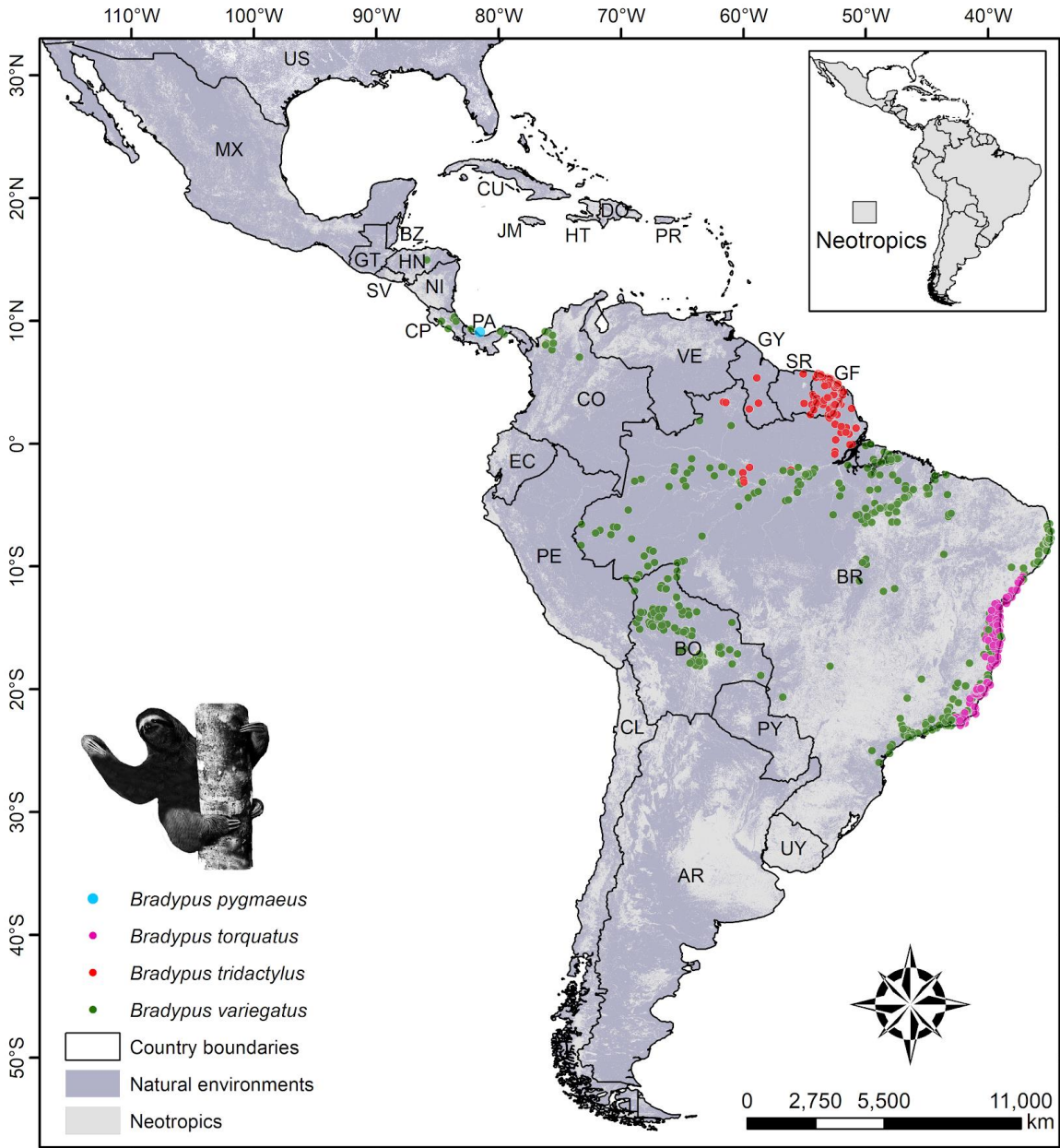
823 populations may now be extinct. Unconfirmed species with “sp.” are not included.



824

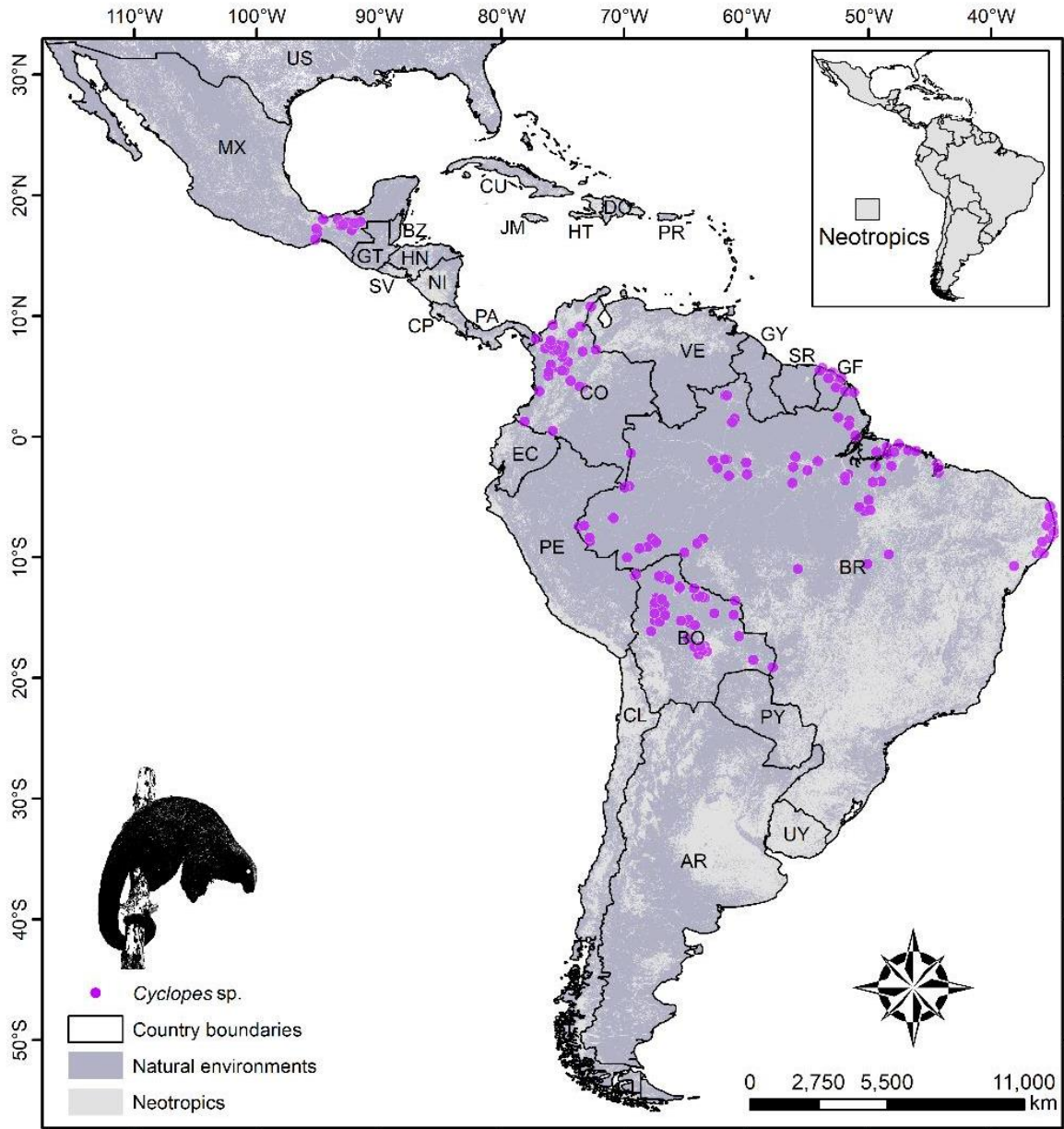
825 **Fig. 7: Distribution of the records of the family Megalonychidae of the NEOTROPICAL**
 826 **XENARTHANS dataset in Neotropical realm.** Records are from 1919 to 2017, thus several
 827 populations may now be extinct. Unconfirmed species with “sp.” are not included.

828



829

830 **Fig. 8: Distribution of the records of the family Bradypodidae of the NEOTROPICAL**
 831 **XENARTHANS dataset in Neotropical realm.** Records are from 1911 to 2018, thus several
 832 populations may now be extinct. Unconfirmed species with “sp.” are not included.



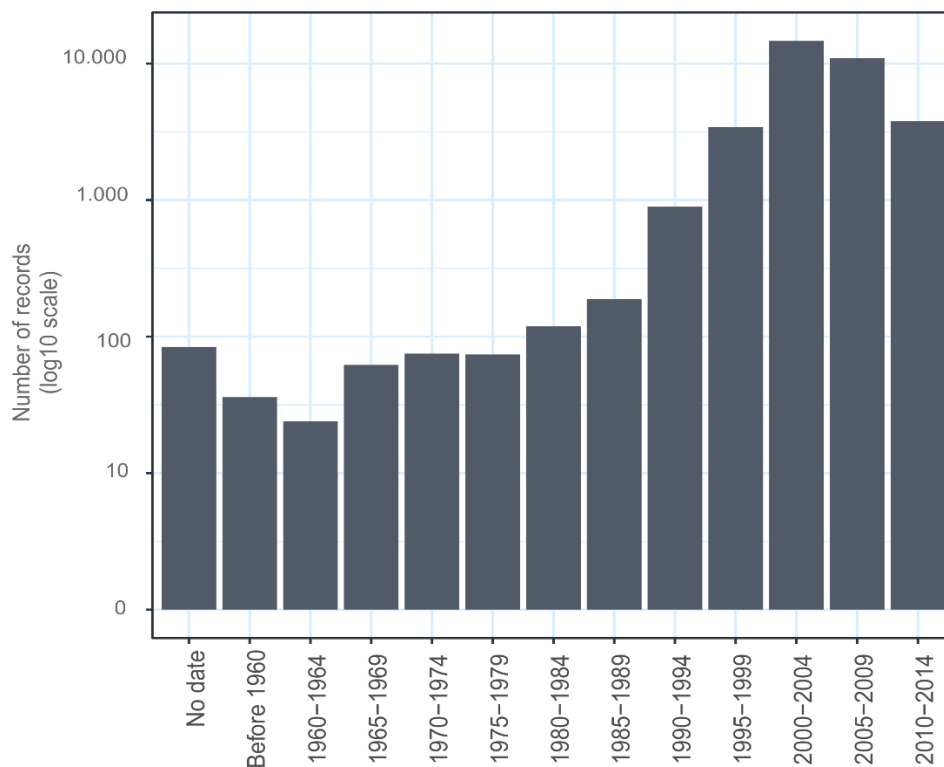
833

834 **Fig. 9: Distribution of the records of the family Cyclopedidae of the NEOTROPICAL**
 835 **XENARTHANS dataset in Neotropical realm.** Records are from 1931 to 2017, thus several
 836 populations may now be extinct.

837 Of the 19 different methods identified, camera trapping most-frequently documents xenarthran
 838 species, responsible for 13,866 of the records (38.42%), especially when used in conjunction

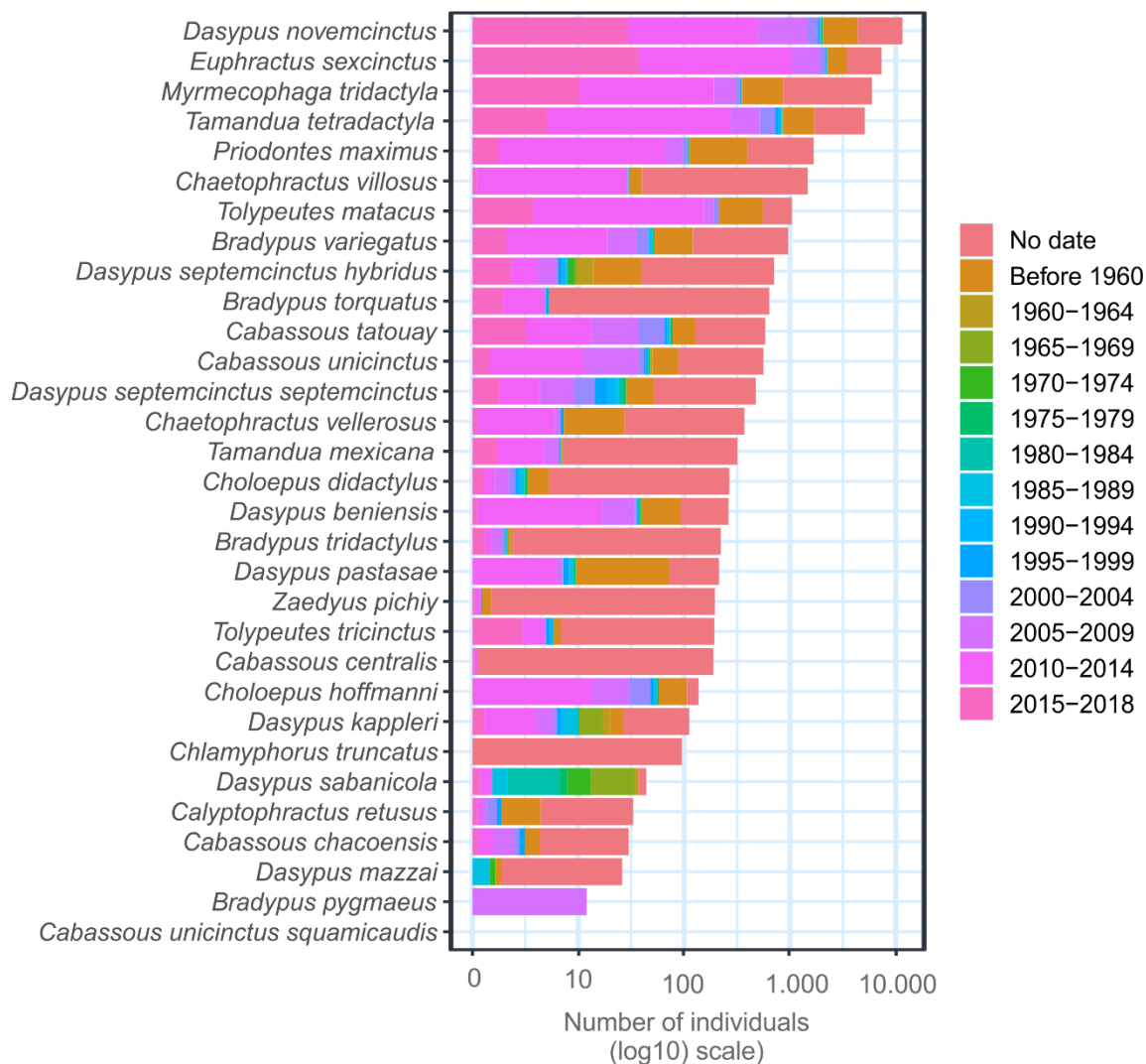
839 with other methods (Table 3). But even when used as the sole sampling method, camera trapping
840 is still the most-frequently method (32.4%). Roadkill is the second most-successful method,
841 accountable for 14.6% of the records. Active search is mostly used in combination with other
842 methods, followed by interview. Other methods include line transect, vestiges, museum
843 collections, telemetry, bibliographical survey, live trapping, sand plots, track plots, car
844 monitoring, and opportunistic encounters (Table 3). About 83.90% of the records correspond to
845 presence-only data, 10.02% to presence-absence data, and 6.08% to quantitative data.

846 According to our database, studies on *Xenarthra* have been increasing in recent years (Figure
847 10). Most of the data (< 10,000 records) were collected between 2010 and 2014, and this is the
848 major period for most species (Figure 11). There are less than 100 records from a less-studied
849 period from 1970 to 1974 (Figure 10) and 24 of the 31 species has records on this period (Figure
850 11).



851

852 **Fig.10: Number of xenarthran records over time.**



854 **Fig.11: Proportion of the records over time for each xenarthran species.**

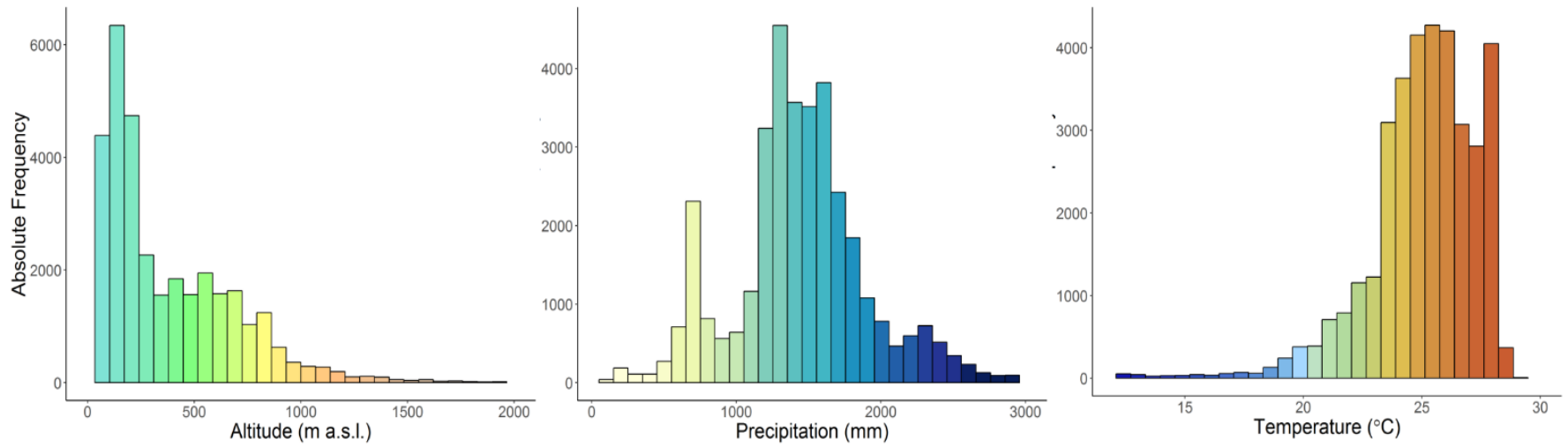
855 The records are concentrated in some regions along the Neotropical realm (Figure 1). On one hand,
 856 Bolivia, southeastern Brazil, northern Argentina, Mexico, and Guiana hold the most records.
 857 Alternatively, some regions, such as the Amazon - mainly due the low accessibility to this area
 858 (Figure 1) - and some countries in South and Central America such as Chile, Peru, Costa Rica, and

859 Panama, have few records - mainly due the lack of contact with researchers of these area; Figure
860 1; More details - Class II, Topic C: Data Limitations and Potential Enhancements).

861 To characterize the environmental conditions of the Neotropical area, we overlapped all data
862 locations with the environment maps, using the function “extract” of the raster package in the R
863 program. The raster files corresponding to temperature and precipitation values were obtained
864 from the WorldClim database 1.4 (<http://www.worldclim.org/version1>). The altitude of most
865 records is below 1,000 m, with a peak around lowland regions (between 200 and 300 m; Fig. 12).
866 The average annual temperature of the sampling sites varies mainly between 20 and 27° C, and
867 few sampling records show temperatures below this range (which is expected for the neotropical
868 region; Fig. 12). Lastly, annual precipitation varies mostly between 1,500 and 2,000 mm, and few
869 sampling sites show values above 2,000 mm. Elevation data were obtained from the digital
870 elevation map from the CGIARCSI database version v4.1 ([http://www.cgiar-csi.org/data/srtm-
871 90m-digital-elevation-database-v4-1](http://www.cgiar-csi.org/data/srtm-90m-digital-elevation-database-v4-1)). The digital elevation model has 1 km of spatial resolution
872 and is available in a unique file in “.tif” format.

873 Natural environment layers for the maps were obtained from the land-use map from the
874 http://due.esrin.esa.int/page_globcover.php. We used the description of Appendix III to identify
875 only the areas described as natural areas that overlapped with the Neotropics realm boundary
876 (http://due.esrin.esa.int/files/GLOBCOVER_Products_Description_Validation_Report_I2.1.1.pdf
877 f). The legend codes classified as natural environment areas were: 40, 50, 60, 70, 90, 100, 110,
878 120, 130, 140, 150, 160, 170, and 180.

879



880

881 **Fig. 12: Variation of altitudes, annual mean temperature, and annual precipitation of the study sites in the NEOTROPICAL**
 882 **XENARTHAN dataset.**

883 **Class II - Research Origin Descriptors**

884 **A. Overall project description**

885 **Identity**

886 A compilation of xenarthran (armadillos, anteaters, and sloths) occurrence and
887 quantitative data in Neotropical Regions.

888

889 **Originator (s)**

890 The Neotropical Xenarthrans project was coordinated by Paloma M. Santos at the
891 Universidade Federal de Minas Gerais (UFMG) and all authors contributed in the
892 assembly of the database. This research is part of the NEOTROPICAL, ATLANTIC,
893 BRAZIL, AMAZONIA, and NEOTROPICAL series, which is led by Mauro Galetti and
894 Milton Cezar Ribeiro, Universidade Estadual Paulista (UNESP).

895

896 **Period of the study**

897 Years of source publications range from 1855 – 2018

898

899 **Objectives**

900 With this paper, we aimed (i) to compile all raw and unpublished xenarthran data
901 collected in recent decades by researchers, projects, and consultants; (ii) to compile the
902 information available in the Portuguese, Spanish, and English literature regarding
903 xenarthran inventories in Neotropical regions, both occurrence only, population, and
904 community data, including the corresponding methods and sampling efforts.

905 **Abstract**

906 Same as above

907

908 **Source(s) of funding**

909 The compilation of this dataset was supported by grants, fellowships, and scholarships
910 from the Agence Nationale de la Recherche (National Research Agency from France)
911 CEBA, ref. ANR-10-LABX-25-01, Agencia Nacional de Promoción Científica y
912 Técnica de Argentina (National Agency for Scientific and Technical Promotion of
913 Argentina, AGENCIA), Alexander Koenig Society, Amazonas Distribuidora de Energia
914 S.A, Conselho Nacional de Desenvolvimento Científico e Tecnológico (Brazilian
915 Research Council, CNPq) numbers 307016/2015-3, 106186/79, 300982/82,
916 153423/2016-1, 301061/2007-6, 425746/2016-0, 207195/2014-5, 457434/2012-0,
917 563134/2010-0, 457497/2012-2, 457491/2012-4, 401887/2014-6, 459735/2014-4,
918 161089/2014-3, 301652/2015-5, 484350/2013-7, 503372/2014-5, 150123/2018-3,
919 472802/2010-0, 141057/2008-4, 158990/2014-5, 303006/2014-5, 141263/2016-4,
920 306392/2013-5, 140730/2010-9, 140039/2018-1, 216938/2014-7, 457458/2012-7,
921 141041/2008-0, 307781/2014-3, 307303/2017-9, 306695/2015-4 and 421361/2017-4,
922 Casadinho/PROCAD 552198/2011-0, Coordenação de Aperfeiçoamento de Pessoal de
923 Nível Superior (Coordination for the Improvement of Higher Education Personnel,
924 CAPES) numbers 1840/2015, 817737/2015, 1537137, 303757/2012-4, 080410/0,
925 004/2012 and 3261/2013, PNPd/CAPES number 283-60/2013, COFECUB/CAPES
926 numbers 88887.130202/201701 and 88887.143361/201700, American Society of
927 Mammalogists, Anglo American, Associação Comunidade Waimiri Atroari, Associação
928 de Defesa Etnoambiental Kanindé, BioFaces, Blue Moon, Brehm Foundation for
929 International Bird Conservation, BRVias, Cambuhy Agrícola Ltda., Chinese Academy

930 of Sciences President's International Fellowship Initiative number 2018PB0040,
931 Christian Rossell, Conservation International Brazil (CI Brazil), Cleveland Metropolitan
932 Zoo, Club de amigos del yagüareté, CNEC WorleyParsons resources and energy,
933 COFUSA Forestry company, Departamento Administrativo de Ciencia, Tecnología e
934 Innovación, Estudios de Doctorado en el Exterior (COLCIENCIAS), Comisión
935 Nacional Forestal, Secretaría de Medio Ambiente y Recursos Naturales, Comisión
936 Nacional de Áreas Naturales Protegidas-Reserva de la Biosfera Pantanos de Centla;
937 Área de Protección de Flora y Fauna Laguna de Términos, Companhia de Concessão
938 Rodoviária de Juiz de Fora – RJ (CONCER), Consejo Nacional de Investigaciones
939 Científicas y Técnicas (CONICET), Consejo Nacional de Ciencia y Tecnología de
940 México (CONACYT), Conservation International's Margot Marsh Primate Action Fund
941 numbers 1001257, 6002856, Conservation Leadership Programme number F02304217,
942 Conservation Program of Endangered Species (PROCER-Mexico) of the National,
943 Commission of Protected Areas (CONANP-Mexico), Conservation, Food and Health
944 Foundation, Consorcio del Mayab, Cota-Robles Graduate Fellowship, China Three
945 Gorges Corporation (CTG-Brazil), Darwin Initiative, DICE Small Grant Program,
946 Durrell Wildlife Conservation Trust, Earthwatch Institute, El Colegio de la Frontera
947 Sur, Unidad Campeche ECOSUR, Fauna and Flora International, Fazenda Barranco
948 Alto and Fazenda Vera Lúcia, Fundação para a Ciência e a Tecnologia (FCT)-Portugal
949 (SFRH/BD/40638/2007), Fibria Celulose S.A., Fundação de Amparo à Pesquisa do
950 Estado do Amazonas (FAPEAM), Fundação de Amparo à Pesquisa do Estado de Goiás
951 (FAPEG) number 11/2013, Fundação de Amparo à Pesquisa e ao Desenvolvimento
952 Científico e Tecnológico do Maranhão (FAPEMA), Fundação de Amparo à Pesquisa do
953 Estado de Minas Gerais (FAPEMIG) numbers CRA- RDP-00104-10; CRA 00357/07;
954 APQ 01145-10, APQ-03315-16 and APQ-00604-17, Fundação de Amparo à Pesquisa

955 do Estado do Rio Grande do Sul (FAPERGS), Fundação Carlos Chagas Filho de
956 Amparo à Pesquisa do Estado do Rio de Janeiro (FAPERJ) numbers E-
957 26/102.960/2012 and E26 /201.267/ 2014, Fundação de Amparo à Pesquisa e Inovação
958 do Espírito Santo (FAPES) numbers 0607/2015 and 0510/2016, Fundação de Amparo à
959 Pesquisa do Estado da Bahia (FAPESB), Fundação de Amparo à Pesquisa e Inovação
960 do Estado de Santa Catarina (FAPESC) numbers 06/2016 and 189/2016; Programa de
961 Apoio à Pesquisa (PAP/FAPESC) number 2017TR744, Fundação de Amparo à
962 Pesquisa do Estado de São Paulo (FAPESP) numbers 2013/04957-8, 2012/14245-2,
963 2013/18526-9, 2014/16320-7, 2013/50421-2, 2016/11595-3, 2014/01986-0,
964 2015/18381-6, 2014/23095-0, 2014/23132-2, 2013/50421-2, 2014/10192-7,
965 2015/25742-5, 2015/11521-7, 2011/22449-4, 2014/01986-0, 2013/10029-6,
966 2014/24921-0, 2016/16433-1, 2015/22844-1, 2017/12925-0, 2014/50434-0,
967 2014/09300-0, 2013/50421-2, 2015/10778 and 2014/16320-7, Fundação de Amparo à
968 Ciência e Tecnologia do Estado de Pernambuco (FACEPE), Fundação de Amparo ao
969 Ensino e Pesquisa (FAEP), Fundação de Tecnologia do Estado do Acre (FUNTAC),
970 Fundação Estadual do Meio Ambiente e Recursos Hídricos de Roraima number
971 01683/09-01, Fundação Grupo Boticário de Proteção à Natureza, Fundación Vida
972 Silvestre Argentina, Fundação de Desenvolvimento Científico e Cultural (FUNDECC) ,
973 Fundação de Apoio ao Desenvolvimento do Ensino, Ciência e Tecnologia do Estado de
974 Mato Grosso do Sul (FUNDECT) numbers 0223/2014 and 06/2016, Fundo Brasileiro
975 para a Biodiversidade – Tropical Forest Conservation Act agreement (FUNBio/TFCA),
976 Global Heritage Fund, Grupo de Ecología de Paisaje y Medio Ambiente de la
977 Universidad de Buenos Aires, Grupo Zema, Hamerton Zoo Park, Heinrich Hertz
978 Foundation, Hotel Belmond, Idea Wild, Instituto Estadual de Florestas (IEF/MG)
979 number 2101010400410, Ingenieros Civiles Asociados división Infraestructura (ICAi),

980 Instituto Brasileiro do Meio Ambiente e dos Recursos Naturais Renováveis (IBAMA),
981 Instituto Chico Mendes de Conservação da Biodiversidade (ICMBio), International
982 Foundation for Science, International Paper Co. of Brazil, Instituto de Pesquisas e
983 Estudos Florestais (IPEF), Kirschbaum Foundation, Lincoln Park Zoo, Liz Claiborne
984 and Art Ortenberg Jaguar Small Grants (Panthera Foundation), Lwarcel Celulose Ltda,
985 Max Planck Institute, Ministerio de Ecología y Recursos Naturales de la Provincia de
986 Misiones, Ministério Público de Minas Gerais, Ministry of Culture and Science of
987 North Rhine-Westphalia , Mohamed bin Zayed Species Conservation Fund numbers
988 12055114 and 0925815, National Council of Science and Technology of Mexico,
989 National Geographic Society grant number W314-14, National Institute for Science and
990 Technology in Wetlands (INAU / UFMT), National Science Foundation (NSF) Pre-
991 doctoral Fellowship grant number 0709598, Overbrook, Programa de auxílio ao
992 Pesquisador (PAPESQ/UNIFAP) number 015/2015, Pays de la Loire - Strategies
993 Internationales (CASEST), Pesquisas Ecológicas de Longa Duração (PELD/CNPQ)
994 numbers 88887.140649/2017-00 and 441278/2016-7 (JSS), PELD Parque Nacional da
995 Serra da Bodoquena (PELD/PNSB), People's Trust For Endangered Species (PTES),
996 President's International Fellowship Initiative grant number 2018PB0040, Primate
997 Conservation Inc. number 1158, Projeto de Conservação e Utilização Sustentável da
998 Diversidade Biológica Brasileira do Ministério do Meio Ambiente (PROBIO/MMA),
999 Programa Áreas Protegidas da Amazônia (ARPA), Programa Beca, Programa Clima,
1000 Naturaleza y Comunidades en Guatemala (CNCG), Programa das Nações Unidas para o
1001 Desenvolvimento (PNUD) , Programa de Concessão de Bolsa de Incentivo ao
1002 Pesquisador da Universidade Estadual de Goiás (PROBIP/UEG) number 009/2016,
1003 Programa Nacional de Incentivos a Investigadores, Projeto Estrada Ecológica and Novo
1004 Colégio, Projeto Onçafari, Pró-Reitoria de Pesquisa e Pós-Graduação da Universidade

1005 Estadual de Santa Cruz–(PROPP/UESC) number 00220.1100.1735, Pró - Reitoria de
1006 Assuntos Estudantis da Universidade Federal do Rio Grande do Sul (PRAE, UFRGS),
1007 Pró -Vida Brazil, Rainforest Alliance, Red de Macro Universidades de América Latina
1008 y el Caribe and Santander Bank , Resource Award, Ridgeway Trust , Royal Ontario
1009 Museum (ROM Governors), Rufford Foundation grants number 18442 -1, 12658 -1,
1010 20144 -2 and 21498 -1, San Diego Zoo Global, Segré Fondation, Secretaria do Meio
1011 Ambiente (SEMA), Secretaría de Ciencia y Técnica, Serviço Social do Comércio
1012 (SESC), Sistemas Estratégicos para la Gestión Ambiental (SEGA), Sistema Nacional de
1013 Pesquisa em Biodiversidade (SISBIOTA/CNPq) numbers 563134/2010 -0 and
1014 563216/2010 - 7, Smithsonian Tropical Research Institute, Sophie Danforth
1015 Conservation Biology Fund, SOS Mata Atlântica Foundation, Studienstiftung des
1016 Deutschen Volkes, Superintendência da Zona Franca de Manaus (SUFRAMA), Suzano
1017 Papel e Celulose S.A., The Ashoka Foundation, The Conservation, Food and Health
1018 Foundation, The Nature Conservancy, The Rolex Awards, The Scott Neotropical Fund
1019 of the Lincoln Park Zoological Society, The Species Survival Fund from Wildlife Trust,
1020 The Whitley Awards, The Woodland Park Zoo, Tijoá Energy, Tropical Ecology
1021 Assessment and Monitoring (TEAM), United States Agency for International
1022 Development (USAID), United States Department of the Interior (USDOI),
1023 Universidade Vila Velha, University of California, University of Puerto Rico, Unidad
1024 para el Cambio Rural (UCAR), Ministerio de Agroindustria, Argentina (PIA) numbers
1025 2011/10102 and 2014/14061, Universidad Nacional de San Luis, Facultad de Química,
1026 Bioquímica y Farmacia., Usina Caeté, Usina Coruripe LTDA, Wildlife Conservation
1027 Society (WCS) Bolivia, WCS Brazil, WCS Guatemala, WCS Jaguar Conservation
1028 Program, WCS Research Fellowship Program Grant , Wild Felid Legacy Scholarship,
1029 Wildlife Trust, World Wide Fund for Nature (WWF) United States of America, WWF

1030 Brazil, WWF EFN Scholarship Program, Zoological Gardens of Dortmund and
1031 Cologne, Zoological Society of London; EDGE of Existence (EDGE , ZSL),
1032 Zoological Society of San Diego, Consórcio Capim Branco Energia (CCBE). MCR
1033 thanks FAPESP (process #2013/50421-2), CNPq (processes # 312045/2013-1;
1034 #312292/2016-3), and PROCAD/CAPES (project # 88881.068425/2014-01) for their
1035 financial support.

1036

1037 **B. Specific Subproject description**

1038 **Site Description**

1039 The target area comprises all the Neotropical realm and part of North America (Mexico
1040 and southeastern United States). We included those areas to cover the entirety of
1041 Xenarthra's distribution. The Neotropical region is one of the eight biogeographic
1042 terrestrial realms (i.e., "ecozones") of the Earth's surface (Olson and Dinerstein 1998,
1043 Olson et al. 2001). It extends from parts of Mexico, southeast USA (southern Florida and
1044 coastal central Florida), Central America, and Caribbean countries, to the austral
1045 distribution of South America. Including the Central USA and the highlands of Mexico,
1046 the coordinates extend from 30°N to 50°S, and 110°W to 40°W, encompassing
1047 approximately 51% of the American continent territory – about 21 million km² of
1048 territorial extension - with most parts located in South America. The vegetation type is
1049 composed of different forests, savannas, mangroves, chaco, sandy vegetation, xeric
1050 vegetation, altitudinal open and rocky fields, cloud forest, floodplains, freshwater
1051 marshes, Andean and lowland environments, etc. The region includes more than half of
1052 the American continent population (approximately 55%), residing mainly in important
1053 and large metropolitan areas, such as São Paulo, Mexico City, Buenos Aires, and Rio de

1054 Janeiro (IBGE 2018, INEGI 2015, INDEC 2010). These figures suggest a strong and
1055 severe anthropogenic impact on natural areas, and consequently on biodiversity. The
1056 region has a high rate of endemism and a unique but threatened biodiversity. Therefore,
1057 the target area includes seven of the 35 world biodiversity hotspots: the Cerrado, the
1058 Atlantic Forest, Tropical Andes, Tumbes-Choco-Magdalena, Valdivian Forests,
1059 Mesoamerica and Caribbean Islands (Myers et al. 2000, Mittermeier et al. 2011), beyond
1060 the Amazon Forest, one of the largest and most important tropical forest in the world.

1061

1062 **Data Compilation**

1063 We created a collaborative network of researchers who had collected and published data
1064 on xenarthran occurrence and community composition in the Neotropics and invited
1065 everyone to contribute to this NEOTROPICAL XENARTHTRANS data-paper (i.e., their
1066 data would be added to our dataset). In addition, we collected occurrence data from the
1067 following museums and institutions: Museu Nacional/Universidade Federal do Rio de
1068 Janeiro, Rio de Janeiro, Brazil (MNRJ), Museu de Zoologia da Universidade de São
1069 Paulo, São Paulo, Brazil (MZUSP), Centro de Primatologia do Rio de Janeiro, Rio de
1070 Janeiro, Brazil (CPRJ), Museu da Fauna, Rio de Janeiro, Brazil (MFRJ), Museu Paraense
1071 "Emílio Goeldi", Pará, Brazil (MPEG), Fundação Zoobotânica do Rio Grande do Sul
1072 (FZBRS), Museu de História Natural do Capão da Imbuia, Paraná, Brazil (MHNCI),
1073 Centro Universitário Barra Mansa, Rio de Janeiro, Brazil (CUBM), Pontifícia
1074 Universidade Católica do Rio Grande do Sul, Rio Grande do Sul, Brazil (PUC-RS),
1075 Universidade Federal da Paraíba (UFPB), Paraíba, Brazil, Universidade Federal do Rio
1076 Grande do Norte (UFRN), Rio Grande do Norte, Brazil, Universidade Federal de Viçosa
1077 (UFV), Minas Gerais, Brazil, Universidade de Brasília (UnB), Brasília, Brazil,
1078 Universidade Federal de Santa Catarina (UFSC), Santa Catarina, Brazil, Museu

1079 Arquidiocesano Dom Joaquim, Santa Catarina, Brazil (MADJ), Museu do Colégio
1080 Agrícola de Camboriú Gert Hering, Santa Catarina, Brazil (MCAGH), Museu do
1081 Seminário Coração de Jesus, Santa Catarina, Brazil (MSCJ), Museu de Ciências Naturais,
1082 Rio Grande do Sul, Brazil (MCN), Museu de História Natural, Lisbon, Portugal (MHNL),
1083 Museu Bocage, Lisbon, Portugal (MBL), Museo de Historia Natural de Valparaíso, Chile
1084 (MHNV), Museu Argentino de Ciências Naturais Bernardino Rivadavia, Argentina
1085 (MACN), Museum Zoologicum Bogoriense (MZB), American Museum of Natural
1086 History, New York (AMNH), Field Museum of Natural History, Chicago (FMNH),
1087 British Museum (Natural History), London (BMNH), Smithsonian Institution National
1088 Museum of Natural History, Washington, D. C. (USNM), and Kansas University Natural
1089 History Museum, Lawrence, Kansas (KUNHM).

1090 After the first round of data compilation, we performed a systematic review of published
1091 literature on occurrence records and abundance of Xenarthra in the Neotropics using the
1092 following electronic databases: Web of Science (<http://scientific.thomson.com/isi/>),
1093 Science Direct (<https://www.sciencedirect.com/>), PubMed
1094 (<https://www.ncbi.nlm.nih.gov/pubmed/>), and Google Scholar
1095 (<https://scholar.google.com.br/>). We searched these databases for relevant articles using
1096 combinations of the following search terms within the title, abstract, and/or keywords:
1097 genus of Neotropical xenarthrans and location names (countries, states, and provinces or
1098 biomes and ecosystems) that lacked information in the first round of data compilation.
1099 Search terms were used in both English, Portuguese, and Spanish.

1100

1101 **Research Methods**

1102 All studies that report location coordinates of occurrence or abundance of xenarthrans
1103 were included in NEOTROPICAL XENARTHTRANS, independent of sampling method

1104 or effort. All the recorded coordinates are in decimal degrees, and the datum is WGS 84.
1105 Although most of such data includes the precise location of species records, a small
1106 portion of the data refers to municipalities, roads, protected areas, or farms (i.e., the
1107 location is not precise). However, we have a PRECISION attribute on the database that
1108 allows users to track the geographic coordinate precision and identify whether the level
1109 of precision satisfies their requirements.

1110 The focus of our data-paper is Neotropical regions, and we delineated this boundary
1111 considering the literature description of the realm boundary, encompassing Central
1112 America including Mexico, the Caribbean islands, and South America. For the complete
1113 database, however, we also allowed contributors to send data for the transitional zone
1114 between the Neotropics and neighboring regions, both at northern and southern parts of
1115 the Neotropical range. Therefore, if users are interested in the stricto sensu Neotropical
1116 regions, we suggest that they use their boundaries of interest, overlap that data, and
1117 extract which data they want to use.

1118 We organized the entire dataset into two different sub-datasets, separated according to
1119 the data type: quantitative information (i.e., more than one record per point) and
1120 qualitative information (i.e., presence and absence and presence-only data; only one
1121 record per point). The quantitative information (Table 4) contains all quantitative data
1122 (i.e. number of records per point) obtained by different methods, such as linear transect,
1123 telemetry, and camera trapping. Qualitative data (Table 5) includes all records of
1124 occurrence data (individual and occasional records of xenarthran species, independent
1125 of the method used), including interviews, museum collections, line transects, camera
1126 traps, roadkill, active searches, vestiges, and opportunistic visualization. All the missing
1127 information was labeled as “NA” in the dataset.

1128 Data were compiled from Darwin 1842, Bard 1855, Lahille 1899, Miller 1899, Allen
1129 1905, Thomas 1919, Yepes 1938a, 1938b, 1939, 1944, Minoprio 1945, Giai 1950,
1130 Vieira 1953, Zeballos 1960, Roig 1965, Rood 1970, Crespo 1974, 1982, Olrog 1976,
1131 1979, Orcés and Albuja 1976, Wetzel and Avila-Pires 1980, Daciuk et al. 1981, Mares
1132 et al. 1981, 1996, McCarthy 1982, Massoia and Chebez 1985, Montgomery 1986,
1133 Navone 1987, 1988, George et al. 1988, Mascarenhas and Puerto 1988, Cardich 1988,
1134 Ojeda and Mares 1989, Cuaron et al. 1989, Stallings et al. 1990, Janzen 1991, Márquez
1135 et al. 1991, Carrillo-Jiménez and Wong-Reyes 1992, Del Vitto et al. 1993,
1136 Administración de Parques Nacionales 1993, Parker et al. 1993, Heinonen and Bosso
1137 1994, Juliá et al. 1994, Barnnet and Cunha 1994, Vidoz and González 1994, NEA 1995,
1138 Pujalte et al. 1995, Brooks 1995, Bolkovi et al. 1995, Rodríguez and Chinchilla 1996,
1139 Capllonch et al. 1996, Cherem and Perez 1996, Heinonen Fortabat and Chebez 1997,
1140 Gaia 1998a, 1998b, 1998c, INEFAN/GEF 1998, BIOLAW 1998a, 1998b, Suare et al.
1141 1998, Cadena et al. 1998, Díaz 1999, Acosta and Murúa 1999, Toledo et al. 1999,
1142 Bolkovic 1999, Calouro 1999, Chiarello 1999, Cuervo-Maya et al. 2000, Cullen et al.
1143 2000, Diaz 2000, Lopes and Ferrari 2000, Sánchez 2000, Tabeni 2000, 2001, Wallauer
1144 et al. 2000, Balabusic and Ruiz 2001, McCain 2001, Printes et al. 2001, Vizcaíno and
1145 Giallombardo 2001, Bowen-Jones 2001, Caro et al. 2001, Graipel et al. 2001, Instituto
1146 Ambiental do Paraná 2002a, 2002b, 2004, 2006a, 2006b, 2015a, 2015b, 2015c,
1147 Marques-Aguiar et al. 2002, Perovic 2002, Sanabria and Quiroga 2002, Soares and
1148 Carneiro 2002, Trucco and Gato 2002, Ceballos 2002, Chalukian et al. 2002, Fernandez
1149 2003, Gil and Heinonen 2003, Jerez and Halloy 2003, Medri et al. 2003, Moreno-
1150 Bejarano and Álvarez-León 2003, Moura 2003, Notarnicola and Navone 2003, Prado et
1151 al. 2003, 2008, Rodrigues et al. 2003, Agüero et al. 2003, Sanderson and Silveira 2003,
1152 Trolle 2003, Agüero et al. 2005, Braga 2003, 2010, Peres et al. 2003, Castaño et al.

1153 2003, Amorim et al. 2004, Cherem et al. 2004, 2007, 2008, 2011, Cunha 2004, Fischer
1154 et al. 2004, Hervas et al. 2004, ICMBio 2004, 2005, 2006, 2007, 2008a, 2008b, 2008c,
1155 2009, 2010, 2011a, 2011b, 2012a, 2012b, 2013a, 2013b, 2013c, 2016, Iwanaga 2004,
1156 Jones and Young 2004, Barbarán 2004, Naranjo et al. 2004, Pérez-Jimeno and Amaya
1157 2004, Prada 2004, Sánchez et al. 2004, Superina et al. 2004, Unidad Administrativa
1158 Especial de Parques Nacionales Naturales de Colombia 2004, 2005, Mendes Pontes
1159 2004, Brito et al. 2004, Arroyo Cabrales et al. 2005, Azurduy et al. 2005, Instituto de
1160 Biología 2005, Mendes Pontes et al. 2005, 2016, Boada and Román 2005, Vaz 2005,
1161 Carrizo et al. 2005, CATIE et al. 2005, Alves and Andriolo 2005, Corredor et al. 2005,
1162 Cherem 2005, Srbek-Araujo and Chiarello 2005, 2013, Cisneros 2006, Correa and Ortiz
1163 2006, Escobar Ocampo et al. 2006, Huertas and Donegan 2006, Barahona Cáceres
1164 2006, Lorenzutti and Almeida 2006, Monguillot 2006, Mora 2006, Oliveira et al. 2006,
1165 2013b, 2013a, 2015, 2016, Pavé and Calderón 2006, Reis et al. 2006, Ruiz 2006,
1166 Bordignon et al. 2006, Chassot et al. 2006, Chavez-Leon 2006, Rocha and Dalponte
1167 2006, Aldana et al. 2006, Lara-Ruiz and Srbek-Araujo 2006, Trinca et al. 2006,
1168 Corredor-Carrillo and Muñoz-Saba 2007, FATMA 2007, 2009, 2010a, 2010b, 2016,
1169 Wetzel et al. 2007, Bovendorp and Galetti 2007, Cáceres et al. 2007, Kasper et al. 2007,
1170 Díaz-N and Sánchez-Giraldo 2008, Gardner 2008, Landmann et al. 2008, Barros 2008,
1171 Modesto et al. 2008a, 2008b, Moreira et al. 2008, Moreira-Ramírez et al. 2008, Penter
1172 et al. 2008, Pereira et al. 2008, 2013, Rodríguez-Mahecha et al. 2008, Silva 2008, Silva
1173 and Mendes Pontes 2008, Smith 2008, Tavares and Koenemann 2008, Vizcaíno and
1174 Loughry 2008, Bezerra et al. 2008, Santos et al. 2008, 2016, Araújo et al. 2008,
1175 Miranda et al. 2008, 2009, Abreu-Júnior and Köhler 2009, Chiquito and Percequillo
1176 2009, INEA 2009, Ballesteros et al. 2009, INEA 2010, 2011a, 2011b, 2015a, 2015b,
1177 Koenemann 2009, Lima 2009, López and Hernández 2009, Mejía-Correa and Díaz-

1178 Martínez 2009, Moreira De Lima et al. 2009, Pautasso et al. 2009, Pérez Jimeno and
1179 Llarín Amaya 2009, Sáenz-Bolaños and Carrillo-Jiménez 2009, Timm et al. 2009,
1180 Torres et al. 2009, Cassano and Kierulff 2009, Di Bitetti et al. 2009, Goulart et al. 2009,
1181 Rocha and Silva 2009, Srbek-Araujo et al. 2009, Pessôa et al. 2009, Santos-Filho and
1182 Silva 2009, Kreutz et al. 2009, CONABIO 2010, Araújo Xavier et al. 2010a, 2010b,
1183 Galván-Guevara 2010, Gomez and Bayly 2010, Balaguera-Reina et al. 2010, Instituto
1184 de Investigaciones Ambientales del Pacifico (IIAP) 2010, Jayat and Ortiz 2010, Macedo
1185 et al. 2010, Muñoz-Saba 2010, 2012, Paviolo 2010, Rezini 2010, Botello et al. 2010,
1186 Bruna et al. 2010, Castaño and Corrales 2010, Braga et al. 2010, 2014, 2018, Rosa et al.
1187 2010, Dantas-Torres et al. 2010, Miranda and Superina 2010, Moraes-Barros et al.
1188 2010, Ramírez-Chaves and Noguera-Urbano 2010, Wallace et al. 2010, 2013,
1189 Corpogujaira et al. 2011, Cruz-Rodríguez et al. 2011, Arcos Delgado 2011, Fabian et al.
1190 2011, Galvis-Penuela et al. 2011, González-Zamora et al. 2011, Marques et al. 2011,
1191 Nascimento and Campos 2011, Tortato et al. 2011, 2014, Diniz and Brito 2012, Hack
1192 2012a, 2012b, 2014, 2017, Manzatti and Franco 2012, Martins et al. 2012, 2016,
1193 Massoia et al. 2012, Padilla and Pérez 2012, Bertassoni et al. 2012, Penido and Zanzini
1194 2012, Bertassoni et al. 2017, Salvador 2012, Sandoval-Gómez et al. 2012, SPVS 2012,
1195 Bonjorne 2012, Cabrera 2012, Hirsch and CHIARELLO 2012, Brocardo and Cândido
1196 Júnior 2012, Delciellos et al. 2012, Santos and Mendes-Oliveira 2012, Chacón Pacheco
1197 et al. 2012, 2013, 2017, Rocha et al. 2012, 2013, 2015, Botelho et al. 2012, Di Blanco et
1198 al. 2012, Catzefflis and Thoisy 2012, Monterrubio et al. 2012, Escobar-Lasso et al. 2013,
1199 Feijó and Langguth 2013, Fundación Jatun Sacha 2013, Geraldo Junior et al. 2013,
1200 Pazio 2013, Quiroga 2013, Rocha 2013, Rojano et al. 2013, 2016, Silva et al. 2013a,
1201 2013b, 2013c, 2017, Weber et al. 2013, Bogoni et al. 2013, 2016a, 2016b, Nunes et al.
1202 2013, Vásquez et al. 2013, Corrêa et al. 2013, Garmendia et al. 2013, Hack and Krüger

1203 2013, Bauni et al. 2013, Costa 2014, Díaz-Pulido et al. 2014, Hack and Santos 2014,
1204 Juraszek et al. 2014, Miranda Júnior and Bertassoni 2014, Miretzki and Braga 2014,
1205 Pereira and Bazilio 2014, Racero-Casarrubia and González-Maya 2014, BirdLife
1206 International 2014, Bogoni 2014, Canon and Trujillo 2014, Carvalho et al. 2014,
1207 Clozato et al. 2014, 2015, Freitas et al. 2014, Borges et al. 2014, 2015, Brown et al.
1208 2014, Magioli et al. 2014, 2016, Feijó and Cordeiro-Estrela 2014, 2016, Hannibal 2014,
1209 Dias and Bocchiglieri 2015a, 2015b, Aximoff et al. 2015b, 2015a, Hannibal et al. 2015,
1210 2018, Baltensperger and Brown 2015, Jiménez-Alvarado et al. 2015, Batista 2015,
1211 Reyes-Amaya et al. 2015, Cherem and Althoff 2015, Feijó et al. 2015, 2018, Ramírez-
1212 Mejía and Sánchez 2015, Hannibal and Neves-Godoi 2015, Hendges et al. 2015,
1213 Fernandes-Ferreira et al. 2015, Benchimol and Peres 2015, De La Ossa Lacayo and De
1214 La Ossa 2015, Zanin et al. 2015, Falconi et al. 2015, Giné et al. 2015, Melo et al. 2015,
1215 David Lopez 2016, De La Ossa and Galván-Guevara 2016, Ecofuturo 2016, Figueroa-
1216 De-León et al. 2016, Bernardo 2016, Preuss et al. 2016, Tinoco-Sotomayor et al. 2016,
1217 Xavier 2016, Deffaci et al. 2016, Cuyckens et al. 2016, Porras et al. 2016, Paviolo et al.
1218 2016, Abreu-Júnior et al. 2017, Cuarón Orozco and Ramos Rivera 2017, Dias et al.
1219 2017b, 2017a, Dirzo Minjarez and Ramos Rivera 2017, Esselstyn 2017, Attias 2017,
1220 González Cózatl and Ramos Rivera 2017, Gual Díaz and Ramos Rivera 2017, Horváth
1221 and Ramos Rivera 2017, León Cortés and Ramos Rivera 2017, León Paniagua and
1222 Ramos Rivera 2017a, 2017b, López González and Ramos Rivera 2017, López Wilchis
1223 and Ramos Rivera 2017, Massocato et al. 2017, Meave del Castillo and Ramos Rivera
1224 2017, Millen and Lim 2017, Monterrubio and Ramos Rivera 2017, Morales Pérez and
1225 Ramos Rivera 2017, Moreira-Ramírez 2017, Muñoz Alonso and Ramos Rivera 2017,
1226 Navarro Singüenza and Ramos Rivera 2017, Oliveira and Hannibal 2017, Polisar et al.
1227 2017, Pozo de la Tijera and Ramos Rivera 2017a, 2017b, Ressler and Ramos Rivera

1228 2017, Sánchez Cordero Dávila and Ramos Rivera 2017, South Australian Museum
1229 2017, Velázquez Montes and Ramos Rivera 2017a, 2017b, Altamirano González Ortega
1230 and Ramos Rivera 2017, Ceballos González and Ramos Rivera 2017, Cepeda-D and
1231 Chacón Pacheco 2017, Cervantes Reza and Ramos Rivera 2017, Zimbres et al. 2017,
1232 Fornitano et al. 2017, Eaton et al. 2017, Laurindo et al. 2017, Cabral et al. 2017, Beca et
1233 al. 2017, Rosa and Souza 2017, Passos et al. 2017, Arevalo et al. 2017, Luna et al. 2017,
1234 Juárez-López et al. 2017, Lima et al. 2017, Bôlla et al. 2017, Feeney 2018, Gall 2018,
1235 Iezzi et al. 2018, Leuchtenberger et al. 2018, UNIBIO and IBUNAM 2018, University
1236 of Minnesota Bell Museum of Natural History 2018, Bradley 2018, Braun and King
1237 2018, Campos et al. 2018a, 2018b, de la Torre et al. 2018, Gonçalves et al. 2018a,
1238 2018b, Xavier da Silva et al. 2018, Lemos et al. in press, Sampaio et al. in press, 2010,
1239 Andrade-Núñez and Aide 2010, Arimoro et al. 2017, Bartrina et al. 2010, D`Bastiani et
1240 al. 2018, DPNVS 1999, Esquivel 2001, FMB 2005, Hurtado and Pacheco 2015, Juárez-
1241 López et al. 2017, Mourthé 2013, Mourthé and Barnett 2014, Núñez-Regueiro et al.
1242 2015, Osto et al. 2004, Queirolo 2016, Ubaid et al. 2010, Wallace et al. 2013, Texera
1243 1973, Atalah 1975, Johnson et al. 1992, Mccarthy et al. 1999, Timock and Vaughan
1244 2002, Vaughan et al. 2007, Suutari et al. 2010, Peery and Pauli 2014, Ruiz-Aravena
1245 2012, Pauli et al. 2012, 2014, Peery and Pauli 2012, Sierpe et al. 2013, Arriagada et al.
1246 2017, Garcés-Restrepo et al. 2017, 2018, Sáenz-Bolaños et al. 2018, and Fountain et al.
1247 2018.

1248

1249 **Taxonomic data**

1250 Taxonomic identity was checked by experts of each family:

1251 a) **Chlamyphoridae and Dasypodidae:** A. Desbiez, G. Mourão, A. Feijó, A.

1252 Bocchiglieri, N. Attias.

1253 b) **Myrmecophagidae and Cyclopedidae:** F. Miranda, V. Tavares, F. Rodrigues,
1254 A. Paglia, A. Bertassoni.

1255 c) **Bradypodidae and Megalonychidae:** A. Chiarello, P. Santos, G. Guiné, S.
1256 Silva.

1257

1258 All data were therefore changed with respect to the identity of most recent nomenclature
1259 and distribution, following Gibb et al. (2016), Miranda et al. (2017), Feijó et al. (2018),
1260 and other available literature (Table 2). The taxonomic uncertainties were maintained,
1261 using sp.

1262

1263 **Validation**

1264 Specialists checked the localities for each xenarthran species and excluded the points
1265 that were erroneous (Table 2). Due to changes in taxonomic classification, several
1266 records of *Cyclopes didactylus* in North and South America were registered as *Cyclopes*
1267 sp., following Miranda et al. (2017).

1268

1269 **C. Data Limitations and Potential Enhancements**

1270 The extant xenarthrans are a fascinating study group of interest, partly due to their
1271 ecology, biology, and taxonomy. Some species have a nocturnal lifestyle, others only
1272 inhabit tree canopies, and, in addition, they also show a great ability to camouflage
1273 themselves in the environment in which they live. This ability reflects directly in the
1274 number of studies; the super order Xenarthra has much less research than other mammal
1275 groups and therefore, gathering data from the various species is a significant challenge.

1276 Nevertheless, we recognize the many efforts that researchers made to collect and
1277 identify the xenarthran data compiled here, given the cryptic habits of several species.

1278 The xenarthran taxonomy, particularly for armadillos, is unusual and usually requires
1279 detailed examination for a correct classification. Therefore, some of the species are
1280 prone to misidentification, mainly due the similar morphological characteristics and
1281 similar common names in different regions. The misidentification and the uncertainty
1282 apply notably when it comes to data from interviews, or when the data are very old. In
1283 all cases, we resorted to the latest taxonomic classification and known distribution
1284 (Abba et al. 2015; Gibb et al 2016; Feijó and Cordeiro-Estrela, 2016, Miranda et al.
1285 2017, Feijó et al. 2018; Table 2), besides an extensive consultation with xenarthran
1286 specialists (for more details, see item 4, section B, Class II and Table 2). Some of the
1287 records, mainly those from museums or interviews, were excluded due to a lack of
1288 certainty regarding occurrences. Two special cases should be highlighted. The recent
1289 taxonomic review of the genus *Cyclopes* (Miranda *et al.* 2017) led to uncertainty
1290 regarding the data that we received. In this case, after consulting key specialists on these
1291 species, we decided to adopt *Cyclopes* sp., thus avoiding errors in the current
1292 distribution of the species. The same approach applies to the genus *Dasypus*. We
1293 followed a recent review of the taxonomic classification, which splits the species
1294 *Dasypus kappleri* into two other species, *D. beniensis* and *D. pastasae* (Feijó *et al.*
1295 2018).

1296 The geographical accuracy of data is another important topic which deserves attention.
1297 We received several types of coordinates, some of them referring to the municipality or
1298 cities, while others referred to the fragment or point sampled. This difference could
1299 represent a bias in our dataset. Therefore, we strongly recommended that co-authors add

1300 the precision (in meters). In case of lack of clarity, we sent a protocol created by us to
1301 assist in estimating precision.

1302 We made significant efforts to compile data from all different regions of the Neotropics
1303 and indeed, data from the different biomes and regions were assembled. There was,
1304 however, a concentration of records in some areas (e.g., the southeastern region of
1305 Brazil) in contrast to regions with few records (e.g., the Amazon, some countries of
1306 Central and South America). To fill this gap, we focused the literature search on those
1307 regions (see 2. Data Compilation section B, Class II), even though there are some areas
1308 with few or no studies. This pattern was found in other data-papers (such as Lima et al.
1309 2017 and Vancine et al. 2017) and seems to reflect the frequency with which the
1310 different natural areas are being studied, often related to accessibility, mainly in the
1311 Amazon region. Additionally, we were unable to contact researchers who are
1312 developing (or have developed) xenarthran research in some neotropical countries, such
1313 as Chile, Peru, Costa Rica, and Panama. This gap is unrelated to lack of studies, but is
1314 related to lack of contact with local researchers, since the literature reports xenarthran
1315 data on those areas (Suutari 2010, Peery and Pauli 2012, Ruiz-Aravena and 2012),

1316 We highlight that the data come from different studies, and consequently, the records
1317 have different sampling efforts, methods, taxonomic scope, and collection date
1318 (including some records without the collection date). Therefore, we strongly
1319 recommend the use of the data to be carried out with a confiable selection criteria. The
1320 lack of quantitative data and the prevalence of occurrence (presence-absence and
1321 presence-only) data reflect the challenges involved when working with this group. A
1322 xenarthran survey may be complex and require alternative methods and significant
1323 efforts. In some cases, it may be necessary to use two or three different methods if

1324 questions aside from the presence of the species (e.g., abundance or habitat use) are to
1325 be investigated.

1326 Despite these limitations, we compiled the largest and most-complete dataset of
1327 xenarthran species of the Neotropical region to date. The merit belongs to the
1328 researchers who have provided us with a great deal of information from different kinds
1329 of studies. With this data-paper, we are providing data often hidden in the gray
1330 literature, or in databases inaccessible to the public. We hope that the public who will
1331 use the database will be able to answer questions, such as: 1) general patterns of
1332 distribution, 2) gaps in xenarthran studies and which species deserve more attention, 3)
1333 poorly studied regions and where to focus more surveys.

1334 **Class III - Data set Status and Accessibility**

1335 **A. Status**

1336 **Latest update**

1337 December 2018.

1338

1339 **Latest Archive date**

1340 December 2018.

1341

1342 **Metadata Status**

1343 Latest update December 2018, which refers to the submitted version of revision process

1344

1345 **B. Accessibility**

1346 Original NEOTROPICAL XENARTHANS dataset can be accessed on the
1347 ECOLOGY repository. All the data – in its updated version and complementary
1348 material – are fully available for both public use and research purposes.

1349

1350 **Storage location and medium:**

1351 The dataset can be accessed on the GitHub Inc. repository
1352 (https://github.com/LEEClab/Neotropical_Xenarthans) in .CSV format. A mirror of this
1353 repository will also be available at https://github.com/LEEClab/Neotropical_series,
1354 where all the other data-papers of NEOTROPICAL SERIES will be available.

1355

1356 **Contact persons:**

1357 Paloma Marques Santos, Universidade Federal de Minas Gerais (UFMG), Instituto de
1358 Ciências Biológicas, Departamento de Biologia Geral, Belo Horizonte (MG), 312700-
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1361 Ecologia, Rio Claro (SP), 13506-900, Brazil. E-mail: miltinho.astronauta@gmail.com.

1362

1363 **Copyright restrictions:**

1364 None

1365

1366 **Proprietary restrictions:**

1367 Please cite this data-paper when using its data in publications. We also request that
1368 researchers and teachers inform us of how they are using this data.

1369

1370 **Costs:**

1371 None

1372 **Class IV - Data Structural Descriptors**

1373 **A. Data set File**

1374 **Identity:**

1375 NEOTROPICAL_XENARTHTRANS_QUANTITATIVE.csv;

1376 NEOTROPICAL_XENARTHTRANS_QUALITATIVE.csv;

1377 NEOTROPICAL_XENARTHTRANS_REFERENCES.csv.

1378 **Size:**

1379 NEOTROPICAL_XENARTHTRANS_QUANTITATIVE.csv, 2,585 records, 1,

1380 169KB;

1381 NEOTROPICAL_XENARTHTRANS_QUALITATIVE.csv, 39,943 records,

1382 14,631 KB;

1383 NEOTROPICAL_XENARTHTRANS_REFERENCES.csv, 1,669 references, 302

1384 KB.

1385

1386 **Format and storage mode:**

1387 comma-separated values (.csv).

1388

1389 **Header Information:**

1390 See column descriptors in section B.

1391

1392 **Alphanumeric attributes:**

1393 Mixed.

1394

1395 **Data anomalies:**

1396 If no information is available for a given record, this is indicated as 'NA'.

1397

1398 **B. Variable Information**

1399 **1) Table 4. Reference Information**

1400 **2) Table 5. Quantitative information**

1401 **3) Table 6. Qualitative information**

1402

1403 **Class V - Supplemental Descriptors**

1404 **A. Data acquisition**

1405 **1. Data request history: None**

1406 **2. Data set updates history: None**

1407 **3. Data entry/verification procedures**

1408

1409 **B. History of dataset usage**

1410

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1462 2016. EncicloVida (Online database: <http://www.enciclovida.mx>). CONABIO. Mexico.
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1470 science not only in Brazil, but for the entire Neotropic realm and for the whole world.

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1477 **TABLES**

1478 **Table 1. Species information.** Family, IUCN conservation status and number of
 1479 records of the armadillos, anteaters and sloth species reported in the NEOTROPICAL
 1480 XENARTHTRANS dataset. Species without confirmed identification (sp.; excluding
 1481 *Cyclopes* sp.) were not included in the species counting. DD=Data Deficient, NE=Not
 1482 Evaluated, LC=Least Concern, NT=Near Threatened, VU=Vulnerable, CR=Critically
 1483 Endangered.

Group	Family	Species*	IUCN	
			Status	Number of records
Armadillos	Chlamyphoridae	<i>Cabassous centralis</i>	DD	189
		<i>Cabassous chacoensis</i>	NT	30
		<i>Cabassous tatouay</i>	LC	585
		<i>Cabassous unicinctus*</i>	LC	562
		<i>Calyptophractus retusus</i>	DD	33
		<i>Chaetophractus vellerosus</i>	LC	372
		<i>Chaetophractus villosus</i>	LC	1,473
		<i>Chlamyphorus truncatus</i>	DD	95
		<i>Euphractus sexcinctus</i>	LC	7,325
		<i>Priodontes maximus</i>	VU	1,678
		<i>Tolypeutes matacus</i>	NT	1,044

		<i>Tolypeutes tricinctus</i>	VU	194
		<i>Zaedyus pichiy</i>	NT	195
	Dasypodidae	<i>Dasypus beniensis</i>	NE	278
		<i>Dasypus kappleri</i> **	LC	94
		<i>Dasypus mazzai</i> ***	DD	26
		<i>Dasypus novemcinctus</i>	LC	11,588
		<i>Dasypus pastasae</i>	NE	217
		<i>Dasypus sabanicola</i>	NT	44
		<i>Dasypus septemcinctus hybridus</i> **	NT	707
		<i>Dasypus septemcinctus septemcinctus</i> **	LC	477
Anteaters	Cyclopedidae	<i>Cyclopes</i> sp.**	NT*	240
	Myrmecophagidae	<i>Myrmecophaga tridactyla</i>	VU	5,941
		<i>Tamandua mexicana</i>	LC	319
		<i>Tamandua tetradactyla</i>	LC	5,089
Sloths	Bradypodidae	<i>Bradypus pygmaeus</i>	CR	12
		<i>Bradypus torquatus</i>	VU	638
		<i>Bradypus tridactylus</i>	LC	222
		<i>Bradypus variegatus</i>	LC	962
	Megalonychidae	<i>Choloepus didactylus</i>	LC	268
		<i>Choloepus hoffmanni</i>	LC	137

1484 *Including the registers of the subspecies.

1485 **A re-evaluation is required because of recent taxonomic changes.

1486 *** Taxonomic synonym *Dasytus yepesi*.

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1489 **Table 2:** List of species registered in the data paper, the synonym (If applicable),
 1490 reference we use to confirm the record and justification of why we corrected the species
 1491 (If applicable). - means “no information”.

Species	Synonyms	Reference	Justification
<i>Bradypus pygmaeus</i>	-	Anderson and Handley 2001, Pauli 2018	-
<i>Bradypus torquatus</i>	-	Hirsch and Chiarello 2012, Gardner 2008, Pauli 2018	-
<i>Bradypus tridactylus</i>	-	Moraes-Barros et al 2010, Gardner 2008, Pauli 2019	Some records of <i>Bradypus tridactylus</i> were corrected to <i>B.variegatus</i> , since <i>B.tridactylus</i> occurs only north of the Amazon River and east of the Rio Negro, and some points were well outside the known distribution for the species
<i>Bradypus variegatus</i>	-	Moraes-Barros et al 2010, Gardner 2008, Pauli 2020	-
<i>Cabassous centralis</i>	-	Abba and Superina 2010	-
<i>Cabassous chacoensis</i>	-	Abba and Superina 2010	-

<i>Cabassous tatouay</i>	-	Abba and Superina 2010, Feijó and Langguth 2013	-
<i>Cabassous unicinctus</i>	-	Abba and Superina 2010, Feijó and Langguth 2013	The genus <i>Cabassus</i> is represented by <i>C. tatouay</i> in northeastern Brazil. Historically these records were erroneously applied to <i>C.unicinctus</i> . Thus, all the records from northeastern Brazil were classified as <i>C. tatouay</i>
<i>Calyptopractus retusus</i>	<i>Chlamyphorus retusus</i> ; <i>Burmeisteria retusa</i>	Abba and Superina 2010, Delsuc 2009	-
<i>Chaetopractus vellerosus</i>	<i>Chaetopractus nationi</i> ; <i>Dasypus vellerosus</i>	Abba and Superina 2010, Abba et al. 2015	All the records of <i>Chaetopractus nationi</i> were considered as <i>C. vellerosus</i>
<i>Chaetopractus villosus</i>	<i>Dasypus villosus</i>	Abba and Superina 2010	-
<i>Chlamyphorus truncatus</i>	-	Abba and Superina 2010	-
<i>Choloepus didactylus</i>	-	Chiarello and Plese 2014, Moraes-Barros 2018	-
<i>Choloepus hoffmanni</i>	-	Plese and Chiarello 2014, Moraes-Barros 2018	-

<i>Cyclopes</i> sp.	<i>Cyclopes didactylus</i>	Miranda et al. 2017	All the records were considered as <i>Cyclopes</i> sp., due to the recent taxonomic review
<i>Dasypus beniensis</i>	<i>Dasypus kappleri</i>	Feijó and Cordeiro-Estrela 2016	The records south of the Amazon River were considered <i>Dasypus beniensis</i>
<i>Dasypus kappleri</i>	-	Feijó and Cordeiro-Estrela 2017	<i>Dasypus kappleri</i> was divided into three species. With the new arrangement, <i>D. kappleri</i> is restricted to the region of the Guiana shield. The records south of the Amazon River refer to <i>D. beniensis</i> , and to the west of the Guiana shield and north of the Amazon River refer to <i>D. pastasae</i> .
<i>Dasypus mazzai</i>	<i>Dasypus yepesi</i>	Feijó and Cordeiro-Estrela 2014	-
<i>Dasypus novemcinctus</i>	-	Abba and Superina 2010, Feijó et al. 2018	-
<i>Dasypus pastasae</i>	<i>Dasypus kappleri</i>	Feijó and Cordeiro-Estrela 2016	The records of west Guiana shield and north Amazon River were considered <i>Dasypus pastasae</i>
<i>Dasypus sabanicola</i>	-	Abba and Superina 2010, Feijó et al. 2018	-

<i>Dasypus septemcinctus hybridus</i>	<i>Dasypus hybridus</i>	Abba and Superina 2010, Feijó et al. 2018	All the records of <i>Dasypus hybridus</i> were re-classified as <i>D. septemcinctus hybridus</i> due to recent taxonomic changes, which considered <i>D. hybridus</i> as a subspecies of <i>D. septemcinctus</i>
<i>Dasypus septemcinctus septemcinctus</i>	<i>Dasypus septemcinctus</i>	Abba and Superina 2010, Feijó et al. 2018	For all records of <i>D. septemcinctus</i> we added the subspecies <i>D. septemcinctus septemcinctus</i> , due to the recent taxonomic review
<i>Euphractus sexcinctus</i>	-	Abba and Superina 2010	-
<i>Myrmecophaga tridactyla</i>	-	Miranda et al. 2014, Bertassoni 2018	-
<i>Priodontes maximus</i>	<i>Priodontes giganteus</i>	Abba and Superina 2010	All the records of <i>Priodontes giganteus</i> were considered as <i>P. maximus</i> , since the name <i>P. giganteus</i> is not used anymore, according to specialists.
<i>Tamandua mexicana</i>	-	Ortega et al. 2014, Bertassoni 2018	-
<i>Tamandua tetradactyla</i>	-	Miranda et al. 2014. Bertassoni 2018	-
<i>Tolypeutes matacus</i>	-	Abba and Superina 2010, IUCN 2018	-

<i>Tolypeutes tricinctus</i>	-	Abba and Superina 2010, Feijó et al. 2015	-
<i>Zaedyus pichiy</i>	-	Abba and Superina 2010	-

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Table 3. Method information. All the methods identified, their absolute

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frequency and percentage.

Method	Frequency	Percentage
Camera trap	13,783	32.4
Road kill	6,225	14.6
Opportunistic	4,339	10.2
Line transect	2,551	6
Active search and interview	2,141	5.03
Bibliographical survey	1,609	3.78
Monitoring	1,358	3.19
Museum scientific collection	780	1.83
Vestige	673	1.58
Active search	624	1.47
Interview	523	1.23
Telemetry	343	0.81
Active search and vestige	243	0.57
Capture	238	0.56
Tracks plot	99	0.23
Car monitoring	74	0.17

Line transect and camera trap	74	0.17
Line transect and live trap	74	0.17
Data base	65	0.15
Burrows sampling	60	0.14
Apprehension	46	0.11
Active search and opportunistic	41	0.1
Live trap	39	0.09
Interview and vestige	21	0.05
Active search, vestige and interview	11	0.03
Live trap and vestige	11	0.03
Interview and bibliographical survey	9	0.02
Active search and roadkill	4	0.01
Camera trap and vestige	4	0.01
Interview, museum scientific collection and bibliographical survey	4	0.01
Vestige and road kill	4	0.01
Active search and camera trap	3	0.01
Active search and live trap	3	0.01
Line transect and vestige	3	0.01

Active search, vestige and camera trap	2	0
Capture and interview	2	0
Active search and bibliographical survey	1	0
Active search and roadkill collection	1	0
Active search, bibliographical survey and interview	1	0
Interview and museum scientific collection	1	0
Interview, museum scientific collection and vestige	1	0
Sand plot	1	0
Vestige and museum scientific collection	1	0
NA*	6,438	15.1
<hr/>		
Total	42,528	100
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*No information

Table 4. Reference information. Description of the fields related to references.

Type of information	Field	Description	Levels	Example
REFERENCE INFORMATION	REF_ID	Reference identification that links the references to a specific record in the dataset.	NEOXEN_1001 – NEOXEN_2669	NEOXEN_1703
	REF_TYPE	Type of the publication: “Thesis” refers to theses, dissertations and monographs ; “Technical document” refer to reports and wildlife management programs.	Abstract Book Book chapter Database In prep In press Management plan Personnal Report Scientific paper Submitted Thesis Unpublished	Scientific paper

	REFERENCE	Study reference in <i>Ecology</i> style.	1554 references	Santos, P. M., A. G. Chiarello, M. C. Ribeiro, J. W. Ribeiro, and A. P. Paglia. 2016. Local and landscape influences on the habitat occupancy of the endangered maned sloth <i>Bradypus torquatus</i> within fragmented landscapes. <i>Mammalian Biology - Zeitschrift für Säugetierkunde</i> 81:447–454.
	Ref_STATUS	Located and unlocated references	Located Not Located	Located
	CITATION_in_text	Citation format in text: author and year		Santos et al. 2016

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1502 **Table 5. Quantitative information.** Description of the fields related to

1503 NEOTROPICAL XENARTHANS abundance data.

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Field	Description	Levels	Example
ORDEMBD	Identification code of each species record. Each code is exclusive and represents the record in the sampling area and the dataset of origin. Records made in the same area in different studies received different codes.	ABBA_00001 - ABBA_02128; ACSA_00001 - ACSA_00060; PAMS_00001 - PAMS_00033	ACSA0001
REFERENCE	Study reference for the published records.		Carrillo-Bilbao G, Cuircama V, Ruíz R, Martin-Solano S (2016) Área de vida, dieta, preferênci a de estrato vertical y uso del tiempo de <i>Bradypus variegatus</i> , liberados
STUDY_AIM	Information about the objective of the study.		ABD-COMM
DATA_TYPE	Type of collected data.	Presence only Presence-absence Abundance	Abundance
SPECIES	The name of the registered species according to the current taxonomy.	See Table 1	<i>Bradypus torquatus</i>

GENUS	The name of the genus according to the current taxonomy.		<i>Bradypus</i>
FAMILY	The name of the family according to the current taxonomy.		Bradypodidae
ORDER	The name of the ORDER according to the current taxonomy.		Pilosa
IUCN_STATUS	Information about the threatened status according to IUCN red list.	NE DD LC NT VU EN CR EW EX	VU
SP_ORIGIN	Information about the origin of that species in that sampling area.	Native Potentially introduced Introduced	Native
SITE	Name of the sampling area.		La Estación Científica Agroecológica Fátima
AREA_HA	Area of the sampling site in hectares.	0.05 - 5.400.000	3096
MUNICIPALITY	Municipality of the sampling site.		Fatima

STATE	State or province of the sampling site.		Pastaza
COUNTRY	Country of the sampling site.		Ecuador
LONG_X	Longitude corrected and transformed into decimal degrees (datum WGS84).	Decimal Degree	-56.11345
LAT_Y	Latitude corrected and transformed into decimal degrees (datum WGS84).	Decimal Degree	-19.566064
PRECISION	Coordinate precision, in meters, of the sampling site.	3 - 290000	30
UC	Information about the protection of the sampling area. Yes: if the area is a conservation unit or is within a conservation unit; No: If the area is outside a conservation unit.	Yes - No	Yes
VEG_TYPE	Vegetation type of the sampling area location, as described in the reference paper, or described by the data collector.		Amazon forest
ANNUAL_RAIN	Annual rainfall WorldClim v. 1.4., in mm, available in http://www.worldclim.org/version1 . Access on August 2nd, 2018.		164
ANNUAL_TEMP	Average annual temperature WorldClim v. 1.4., in Celsius degrees, available in http://www.worldclim.org/version1 . Access on August 2nd, 2018.		25.1

ALTITUDE	Altitude in meters above sea level, from the Hydro-1K dataset (United States Geological Survey – USGS, 2001. HYDRO 1K: Elevation Derivative Database. Available from: < http://edc.usgs.gov/products/elevation/gtopo30/hydro/namerica.html >). Access on August 2nd, 2018.		469
Olsoneconame	ECO_NAME column of the shapefile wwf_terr_ecos available in WWF website (Olson et al. 2001).		Caatinga
OlsonG200r	Olson bioregion (Olson et al. 2001), represented by G200_REGIO column of the shapefile wwf_terr_ecos		Cerrado Woodlands and Savannas
METHOD	Sampling method described in the reference paper or by the data collector.		Line transect
N_POINTS	Number of sampling points.	1 - 1401	500
D_POINTS	Distance, in kilometers, between the sampling points.	0 - 1.6	1.2
LTR_EFF	Effort in kilometers walked on transects in the sampling area.	0.5 – 5000	618
LTR_ABD_10KM	Abundance of individuals registered on transects. Calculated as the number of individuals/10km.	0 – 25.71	0.32

CAM_EFF	Survey effort in the sampling area using camera traps. Measured in hours.	1.03 - 36000	3600
CAM_ABD	Number of records registered on camera traps.	0 - 94	25
SURV_EFF	Survey effort in the sampling area. Measured in days, hours and kilometers.	8.8h – 24.3h	24.1h
SURV_ABD	Number of records registered on survey. Calculated as the number of counts/day.	0 - 85	33
COL_STRT_MO	Month when data collection started	January - December	May
COL_STRT_YR	Year when data collection started.	1896- 2018	1996
COL_END_MO	Month when data collection ended.	January - December	December
COL_END_YR	Year when data collection ended.	1855- 2018	1996

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1508 **Table 6. Qualitative information.** Description of the fields related to NEOTROPICAL

1509 XENARTHANS occurrence data.

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Field	Description	Levels	Example
ORDEMBD	Identification code of each species record. Each code is exclusive and represents the record in the sampling area and the dataset of origin. Records made in the same area in different studies received different codes.	ABBA_00001 - ABBA_02128; ACSA_00001 - ACSA_00060; PAMS_00001 - PAMS_00033	PAMS_0001
REFERENCE	Study reference for the published records.		Carrillo-Bilbao G, Cuircama V, Ruíz R, Martin-Solano S (2016) Área de vida, dieta, preferência de estrato vertical y uso del tiempo de <i>Bradypus variegatus</i> , liberados
STUDY_AIM	Information about the objective of the study.		Occur
DATA_TYPE	Type of collected data.	Presence only Presence-absence	Presence-absence
SPECIES	The name of the registered species according to the current taxonomy.	See Table 1	<i>Bradypus variegatus</i>
GENUS	The name of the genus according to the current taxonomy.		<i>Bradypus</i>

FAMILY	The name of the family according to the current taxonomy.		Bradypodidae
ORDER	The name of the ORDER according to the current taxonomy.		Pilosa
IUCN_STATUS	Information about the threatened status according to IUCN red list.	NE DD LC NT VU EN CR EW EX	LC
SP_ORIGIN	Information about the origin of that species in that sampling area.	Native Potentially introduced	Native
SITE	Name of the sampling area.		La Estación Científica Agroecológica Fátima
AREA_HA	Area of the sampling site in hectares.	0.05 - 5.400.000	3096
MUNICIPALITY	Municipality of the sampling site.		Fatima
STATE	State or province of the sampling site.		Pastaza
COUNTRY	Country of the sampling site.		Ecuador
LONG_X	Longitude corrected and transformed into decimal degrees (datum WGS84).	Decimal Degree	-56.11345

LAT_Y	Latitude corrected and transformed into decimal degrees (datum WGS84).	Decimal Degree	-19.566064
PRECISION	Coordinate precision, in meters, of the sampling site.	3 - 290000	30
UC	Information about the protection of the sampling area. Yes: if the area is a conservation unit or is within a conservation unit; No: If the area is outside a conservation unit.	Yes - No	Yes
VEG_TYPE	Vegetation type of the sampling area location, as described in the reference paper, or described by the data collector.		Amazon forest
ANNUAL_RAIN	Annual rainfall WorldClim v. 1.4., in mm, available in http://www.worldclim.org/version1 . Access on August 2nd, 2018.		164
ANNUAL_TEMP	Average annual temperature WorldClim v. 1.4., in Celsius degrees, available in http://www.worldclim.org/version1 . Access on August 2nd, 2018.		25.1
ALTITUDE	Altitude in meters above sea level, from the Hydro-1K dataset (United States Geological Survey – USGS, 2001. HYDRO 1K: Elevation Derivative Database. Available from: http://edc.usgs.gov/products/elevation/gtopo30/hydro/america.html) Access on August 2nd, 2018.		469

Olsoneconame	ECO_NAME column of the shapefile wwf_terr_ecos available in WWF website (Olson et al. 2001).		Uruguayan savanna
OlsonG200r	Olson bioregion (Olson et al. 2001), represented by G200_REGIO column of the shapefile wwf_terr_ecos		Cerrado Woodlands and Savannas
METHOD	Sampling method described in the reference paper or by the data collector.		Line transect
N_POINTS	Number of sampling points.	1 - 1401	500
D_POINTS	Distance, in kilometers, between the sampling points.	0 - 1.6	1.2
LTR_EFF	Effort in kilometers walked on transects in the sampling area.	0.5 – 5000	618
CAM_EFF	Survey effort in the sampling area using camera traps. Measured in hours.	1.03 - 36000	3600
SURV_EFF	Survey effort in the sampling area. Measured in days, hours and kilometers.	8.8h – 24.3h	24.1h
COL_STRT_MO	Month when data collection started	January - December	May
COL_STRT_YR	Year when data collection started.	1896- 2018	1996
COL_END_MO	Month when data collection ended.	January - December	December

COL_END_YR	Year when data collection ended.	1855- 2018	1996
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3213

3214 **Capítulo 2**

3215 **Living on the Edge: Forest cover threshold effect on endangered maned sloth**
3216 **occurrence in Atlantic Forest**

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3237 **Abstract**

3238 Habitat loss and the isolation of remaining habitats are undoubtedly the two greatest
3239 threats to biodiversity conservation, especially for the maned sloth, due to its ecological
3240 restrictions. In this study, we identified a critical threshold of forest cover for maned
3241 sloth occurrence and explored the effects of other local and landscape variables. We
3242 sampled 68 sites, where we searched for the maned sloth and collected local habitat
3243 variables. We calculated the percentage of forest cover and open areas, assessing the
3244 appropriated scale through model selection. We used occupancy models and model
3245 selection methods to identify the threshold and assess occupancy and detection
3246 probabilities. The occupancy probability of the maned sloth is 0.97, but it decreases
3247 abruptly at 35% of forest cover, reaching zero in areas with less than 20% of forest
3248 cover. The two landscape variables are the most important predictors of sloth
3249 occupancy, based on the cumulative weight of evidence, were: Forest cover (78%) and
3250 Open areas cover (46%); the latter influencing negatively maned sloth occupancy. This
3251 is the first attempt to identify the habitat requirements of the threatened maned sloth in a
3252 fragmented area using landscape and local variables. Our results imply that conservation
3253 of maned sloth will benefit from an increase in the amount of native forest at the
3254 landscape scale. Given difficulties in the creation of new public protected areas, this
3255 improvement could be achieved via restoration of areas located in private properties
3256 which are protected by the Brazilian Forest Code.

3257 **Keywords:** *Bradypus torquatus*; Pilosa; Threatened; Threshold; Atlantic Forest;

3258 Habitat loss

3259

3260

3261 **1. Introduction**

3262 Human-dominated areas have expanded throughout the world and are primarily
3263 responsible for increased deforestation (LEBLOIS *et al.*, 2017). Habitat fragmentation
3264 and loss leads to patch size reduction, increases habitat isolation and edge effects, and
3265 have several negative consequences for biodiversity (FAHRIG, 2003). Worldwide, most
3266 of Earth's native ecosystems are within 1 km of the habitat edge (HADDAD *et al.*,
3267 2015), embedded in anthropogenic matrix that is often inhospitable to biodiversity
3268 (CASTELLÓN; SIEVING, 2006). These anthropogenic matrices can act as a filter,
3269 restricting dynamic processes such as dispersion and movement (MORAES *et al.*, 2018;
3270 MORATO *et al.*, 2018), gene flow (DIXO *et al.*, 2009), and pollination (PAVAGEAU
3271 *et al.*, 2017). Additionally, deforested areas have limited shelter or refuge for native
3272 fauna, exposing species to opportunist predation by domestic dogs (DOHERTY *et al.*,
3273 2017), poaching (FERREGUETTI *et al.*, 2018) and roadkill (ASCENSAO *et al.*, 2017;
3274 ASSIS; GIACOMINI; RIBEIRO, 2019). All these threats are commonly related with
3275 human activities in a non-habitat matrix areas (DOHERTY *et al.*, 2017;
3276 FERREGUETTI *et al.*, 2018).

3277 Among the various attributes intrinsic to anthropized environments, the amount
3278 of remaining habitat is perhaps one of the most influential in the dynamics of
3279 populations and communities (FAHRIG, 2013). When habitat loss increases and
3280 reaches a critical threshold, small modifications in the remaining habitat can produce
3281 more severe changes in biological responses (WITH; KING, 1999; Huggett, 2005).
3282 Theory predicts that structural and functional connectivity suffers when the amount of
3283 remaining habitat drops below this critical threshold, and a species distribution becomes
3284 more dependent on patch size and isolation (LANDE, 1987; FAHRIG, 2003). The
3285 threshold value will vary among species, following the species' habitat requirements

3286 (Muylaert et al., 2016), patch size (MAGIOLI *et al.*, 2015), matrix composition
3287 (BOESING; NICHOLS; METZGER, 2018), and biome (Melo et al., 2018).

3288 As the proportion of suitable habitat increases (*i.e.* above a given threshold),
3289 other variables may drive the ecological dynamics, besides the landscape context, such
3290 as habitat quality. The local habitat structure and composition plays an important role in
3291 occurrence and persistence of animal communities (August, 1983; Santos et al., 2016).
3292 Increasing habitat complexity creates a safe and dynamic environment, providing
3293 different resources and conditions necessary for reproduction (SHENBROT *et al.*,
3294 2002) and foraging success (JACOB; MATTER; CAMERON, 2017). In addition, a
3295 more complex habitat structure may increase the alpha and beta diversities by providing
3296 suitable habitats for a higher number of species (CAMARGO; SANO; VIEIRA, 2018;
3297 GRELLE, 2003).

3298 The Atlantic Forest is a biodiversity hotspot (ZACHOS; HABEL, 2011) and one
3299 of the most threatened biomes in the world and its area has been reduced to 28% of its
3300 original cover (REZENDE *et al.*, 2018). Most remaining forest patches are smaller than
3301 50 ha and about half of the remaining forest is within 100 meters of any forest edge
3302 (RIBEIRO *et al.*, 2009). Landscape composition and configuration pose a threat to
3303 several species, including the endemic maned sloth (*Bradypus torquatus*;
3304 Bradypodidae). According to Santos *et al.* (2019), in a data paper that compiled the
3305 largest data set on abundance and occurrence of Neotropical Xenarthrans, maned sloth
3306 (with 638 records) are distributed along Atlantic Forest coast from Sergipe to Rio
3307 Janeiro, Brazil. The species inhabits dense ombrophyllous forests (HIRSCH;
3308 CHIARELLO, 2012) and feeds mainly on leaves and shoots (MONTGOMERY;
3309 SUNQUIST, 1975; CHIARELLO, 1998b). The species rarely descends to the ground,
3310 moving mostly through the forest canopy (MONTGOMERY; SUNQUIST, 1975). Due

3311 to its strictly arboreal habit and its low capacity to move in non-forested areas, the
3312 species is highly affected by the habitat loss (CHIARELLO; MORAES-BARROS,
3313 2014) Its restricted distribution and the dependence on forest led the International Union
3314 for Conservation of Nature (IUCN) to classify the maned sloth as Vulnerable (VU) in
3315 2014 (CHIARELLO; MORAES-BARROS, 2014).

3316 There is strong evidence that habitat loss and fragmentation negatively influence
3317 maned sloth occurrence (CHIARELLO; MORAES-BARROS, 2014). Intuitively,
3318 increasing the connection between fragments would be a logical strategy to improve the
3319 conservation of the species. Nevertheless, what exactly does “improve connection”
3320 mean for the maned sloth? What are the minimum habitat amount requirements to
3321 maintain this species? Is there a critical threshold, below which landscape variables
3322 would most likely influence the species occurrence? These questions are relevant to
3323 decision-makers responsible for applying resources to direct conservation efforts; these
3324 efforts are led by the National Center for Research and Conservation of Brazilian
3325 Primates (CPB) and linked to Chico Mendes Institute for Biodiversity Conservation
3326 (ICMBio), a Brazilian federal agency responsible for the National Action Plans.

3327 In previously study, Santos *et al.* (2016) found a strong positive relationship
3328 between local variables (e.g., canopy height and proportion of important trees) and
3329 maned sloth occurrence – these local variables were more important than factors related
3330 to habitat fragmentation (e.g., patch size and isolation metrics). However, the studied
3331 area encompassed a region with >36% of forest cover, a level that exceeds forest
3332 thresholds found in other ecosystems with different taxa (thresholds ~30 – 35% forest
3333 cover; ANDRÉN, 1994; ESTAVILLO *et al.*, 2013). Motivated by our previous
3334 experience, here, we expanded the study region to encompass a broader area covering
3335 the full gradient of available habitat (5 to 100% forest cover) to identify and assess

3336 larger-scale variables for maned sloth occurrence. In this study, we aimed to better
3337 understand what affects maned sloth occupancy probability at the landscape level and in
3338 a local context to improve the subsidies for landscape and local management,
3339 respectively. We also contribute to the understanding of maned sloth habitat
3340 requirements, which are still unclear. We hypothesize there is a threshold relationship
3341 between the forest cover (%) and maned sloth occurrence: above this threshold maned
3342 sloth occurrence should be high and governed primarily by local variables and below
3343 this threshold maned sloth occupancy should decline rapidly regardless of local
3344 variables.

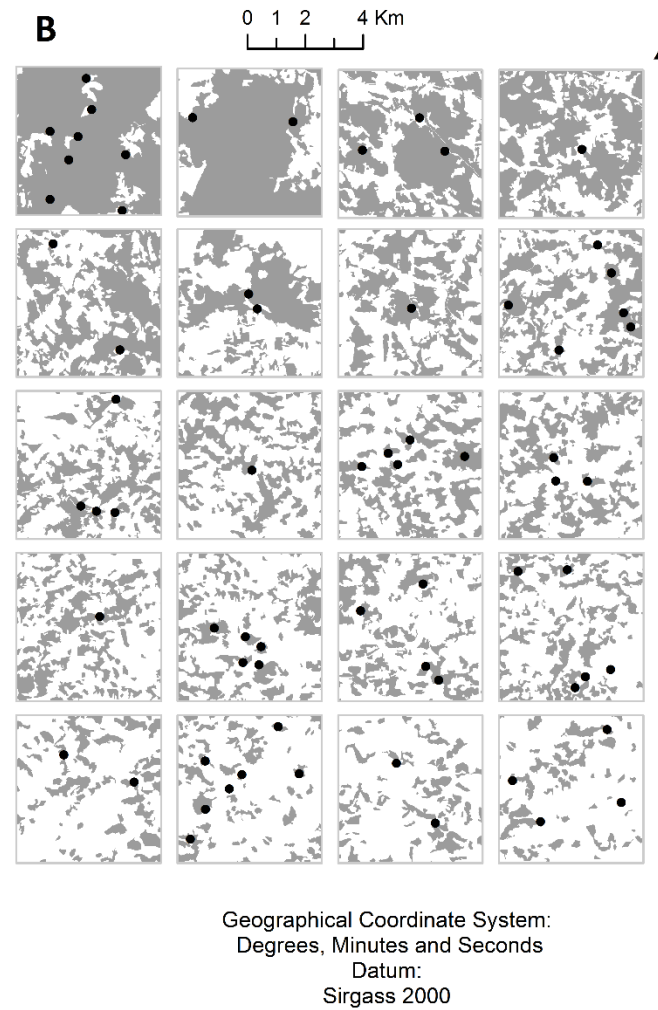
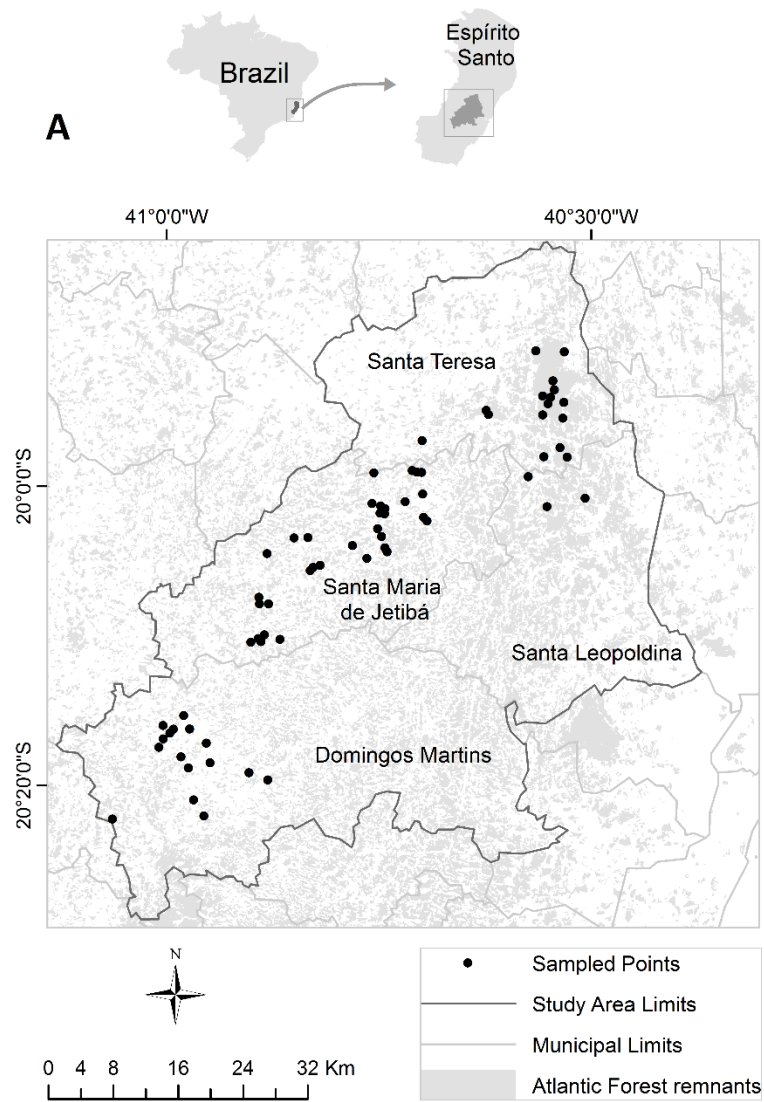
3345 **2. Material and Methods**

3346 *2.1. Study area*

3347 We conducted this study in the mountainous region of Espírito Santo State in
3348 southeast Brazil (Fig. 1), which historically consisted entirely of the Atlantic Forest
3349 (MUYLAERT *et al.*, 2018). The study region encompasses an area of approximately
3350 187,000 ha (Fig. 1), composed primarily of dense lower-montane and montane
3351 ombrophyllous forests (INSTITUTO BRASILEIRO DE GEOGRAFIA E
3352 ESTATÍSTICA, 2012). The elevation ranges from 600 to 1200 m.a.s.l. and
3353 temperatures are mild throughout the year (mean annual temperature = 22.1°C). The
3354 annual mean precipitation is 1232 mm, with most rain concentrated between November
3355 and March (INCAPER, 2013a, b, d, c).

3356 The study region has 34% of native forests, concentrated (> 85%) in the
3357 northeast, where the largest protected areas are located, while the southern part has less
3358 than 20% of forest cover (Fig. 1). Most forest areas are privately owned (97.2%),
3359 mainly by descendents from European settlers, who arrived at the region in the early

3360 19th century, and established small agricultural properties and a family farming system –
3361 which remains the economic base today (INCAPER, 2013a, b, d, c).



3363 **Fig. 1:** Study area with the location of the 68 sampled points for recording maned sloth within Atlantic Forest remnants, Espírito Santo, Brazil

3364 (A). Example of the forest cover gradient throughout the region, in which the sampled points are inserted (B).

3365 2.2. *Sampling design and covariates selection*

3366 We selected 68 sampling points (i.e., sites) using GrassGis 7.4.1 software
3367 (GRASS Development Team, 2018) across a range of forest cover (5-100%) within the
3368 study region. Specifically, we select 19 sites on northeast (higher percentage of forest
3369 cover), 16 sites in the south (lower percentage of forest cover), and 33 sites from the
3370 central Santa Maria de Jetibá region (medium percentage of forest cover). To minimize
3371 spatial autocorrelation, sites were at least 500 m away from each other, a distance that is
3372 large enough to accommodate the home ranges of multiple individuals – the average
3373 sloth home range was estimated at 4.5 ha in previous studies conducted in this same
3374 area (CHIARELLO, 1998a). Sloth surveys were conducted during the dry season to
3375 avoid effects of rain on species detection. Specifically, we surveyed 33 sites in the
3376 central Santa Maria de Jetibá region from April to August in 2014 (Santos et al., 2016),
3377 and 35 different sites from April to October in 2017 and from March to July in 2018.
3378 The sites surveyed in 2014 were not surveyed in 2017/2018 and vice-versa. Surveys
3379 within a given dry season were spaced at approximately 1-month intervals such that
3380 each site was surveyed six times. During each survey, we established two perpendicular
3381 200-m transects, forming a cross (Figure A.1). We walked the resulting four 100-m long
3382 segments at a speed of about 0.24 km/h, searching the canopies for maned sloths. To
3383 maximize detection, we walked the transects between 7 am - 5 pm, when the species is
3384 most active (CHIARELLO, 1998a; GINÉ *et al.*, 2015). We measured two local
3385 variables, canopy height (m) and the proportion of important trees (hereafter important
3386 trees) at five locations, including the center point (intersection of the two perpendicular
3387 transects) and at the halfway point of each 100-m transects (Figure A.1). At each
3388 location, we measured the canopy height with a rangefinder (Bushnell model Sport 850,
3389 4 x 20 mm) and we recorded the presence/absence of important tree species

3390 (*Micropholis venulosa*, *Cupania furfuraceae*, *Creecopia* spp., and *Ficus* spp.) within a 5
3391 m radius of each location. Collectively, these species represent more than 40% of the
3392 maned sloth diet (CHIARELLO, 1998b). We combined the measurements at the five
3393 locations for each site to determine the site-specific mean canopy height and the
3394 proportion of locations with at least one of the important trees.

3395 We also calculated two compositional landscape structure variables that we felt
3396 may influence maned sloth probability of occurrence: forest cover (%) and open areas
3397 cover (%). We expected each covariate of interest to influence maned sloth ecology and
3398 biology differently (Table 1). For example, forest cover is important to reproduction,
3399 foraging success, and other aspects of maned sloth ecology and behavior. Therefore, we
3400 expected a positive relationship between the forest cover and maned sloth occurrence.
3401 Conversely, open areas cover may deter or hamper the species' dispersal, since moving
3402 through these areas may expose sloths to predation, starvation, temperature stress, and
3403 other hazards. As the open areas cover increases, we expect maned sloth occupancy to
3404 decrease. At the local level, important trees are a necessary food source and
3405 consequently, there should be more maned sloth in areas with a high proportion of those
3406 trees, leading to higher species occurrence and detection probabilities. The relationship
3407 with detection should also be positive since it may be easier to detect the species if
3408 individuals spend more time in or around these food sources. Many trees also serve as
3409 shelter where sloths can safely rest and thermoregulate appropriately, which is usually
3410 done in the upper strata of the forest canopy. Accordingly, we expect occupancy
3411 probability to increase with increasing canopy height. However, detection is more
3412 difficult the higher the sloth is in the canopy since vegetation density also increase with
3413 canopy height, obstructing our viewing capacity.

3414 **Table 1:** Local and landscape level variables thought to influence occupancy
 3415 probability (Ψ) or detection probability (p). Hypothesized positive (+) effect or negative
 3416 (-) effect on occupancy probability or detection probability.

Type of Variable	Variable	Level	Ecological Importance	Parameter	Effect on Ψ	Effect on p
Class	% Forest cover	Landscape	Breeding Foraging success Shelter	Ψ	+	
Class	% Open Areas cover	Landscape	Dispersal Exposure to threats	Ψ	-	
Local	Important trees	Local	Food source	Ψ/p	+	+
Local	Canopy Height (m)	Local	Shelter Thermoregulation	Ψ/p	+	-

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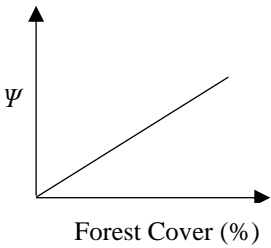
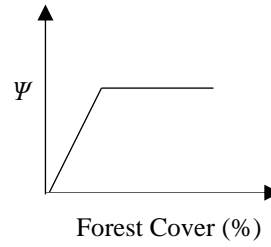
3419 To calculate landscape variables for each sampled site, we used a 2015 land use
 3420 and land cover map with 10-m resolution, georeferenced in the UTM Projection
 3421 System, SIRGAS 2000 Datum, Zone 24s, provided by Geobases (IEMA-ES,
 3422 <https://geobases.es.gov.br/>). We ran the landscape analyses with Fragstats 4.2
 3423 (McGarigal et al., 2012) and considered six buffer sizes (i.e. scale of effect or spatial
 3424 extension; Miguët et al., 2015) for each sampled sites: 100 m, 200 m, 300 m, 400 m,
 3425 500 m and 600 m. The usage of multiple spatial extents when evaluating the scale of
 3426 effect of landscape structure influence on species occurrence, abundance or species
 3427 richness is of utmost importance in landscape ecology studies (GESTICH *et al.*, 2019).
 3428 Within each buffer, we calculated the percentage of forest cover and the percentage of
 3429 open areas cover – the latter including existing and abandoned pasture and exposed soil.

3430 *2.3. Occupancy and detection probabilities*

3431 We used a step-wise approach (four steps) to evaluate factors influencing maned
3432 sloth occupancy and detection probabilities. First, we determined the scale of effect
3433 (buffer size) for each of our landscape covariates: percentage of forest cover and
3434 percentage of open areas cover. Using a general model structure where both occupancy
3435 and detection probabilities are modeled as additive functions of local variables
3436 (important trees and canopy height), we fitted 36 models, where sloth occupancy was
3437 modeled as a linear function (on the logit scale) of all possible combinations of the six
3438 spatial extents for percent forest cover and percent open areas cover (Table A.1). We
3439 used the best-supported scale of effect for each of the landscape variables in subsequent
3440 analyses. We also performed a correlation analysis using the best-supported scale of
3441 effect for forest cover (500 m) and open area cover (600 m) and our local covariates to
3442 ensure that none of our covariates were highly correlated ($|r| < 0.6$; Table A.2).

3443 Second, we explored if there was evidence of a threshold relationship between
3444 the probability of sloth occurrence and the percent of forest cover, using a general
3445 detection probability structure and an occupancy structure that accounted for potential
3446 variation associated with our other covariates. Specifically, we built eight occupancy
3447 structures: one corresponding to the linear relation (on the logit scale) between the sloth
3448 occupancy probability and percent forest cover and the other seven models reflected
3449 threshold effects at specified values of forest cover (e.g., 20%, 25%, 30%,..., 50%
3450 forest cover; Tables 2 and A.4). For example, a model with a specified threshold of 30%
3451 forest cover suggests that sloth occupancy starts to decline when the forest cover falls
3452 below 30%, but sloth occupancy is approximately equal at sites above the threshold (see
3453 Table 2 for a graphical representation of these relationships). We modelled thresholds
3454 from 20% to 50% forest cover (increasing in 5% increments) and used a model selection

3455 criteria (AICc) to determine which threshold level was best supported by our data
 3456 (Table A.4).
 3457 **Table 2:** Eight competing global models with different threshold relations between
 3458 forest cover (%) and occupancy probability (Ψ) of maned sloth within Atlantic Forest,
 3459 Espírito Santo, Brazil. The “X” in “ Ψ Forest Cover_X%” represents the different
 3460 values of percentage of forest cover threshold.

Model	Relationship	Graphical Representation
Ψ Forest Cover % + Open Area Cover %+ Important trees + Canopy Height, <i>p</i> Canopy Height + Important Trees	Linear	
Ψ Forest Cover_X%* + Open Area Cover %+ Important trees + Canopy Height, <i>p</i> Canopy Height + Important Trees	Threshold at X% of Forest Cover	

3461 *Forest cover ranges 20 to 50%, 5 by 5%.

3462

3463 Third, using the best supported threshold occupancy structure, we explored local
 3464 factors that could influence sloth detection probability. Specifically, we fitted four
 3465 detection probability structures that included all additive combinations of canopy height
 3466 and important tree (Table 3). Finally, we considered occupancy structures to explore
 3467 additive (16 models) and interactive effects (6 models) of our local and landscape
 3468 variables, fitting models with up to 7 parameters to avoid the overparameterization. We
 3469 explored interactive relationships between forest cover and the other occupancy
 3470 covariates because the effect of some variables (e.g., important trees and open areas
 3471 cover) may change as the forest cover increases.

3472 All models were fitted using the single-season occupancy model (MACKENZIE
3473 *et al.*, 2006) incorporated in program MARK (WHITE; BURHNAM 1999). Using our
3474 global model, we conducted a goodness-of-fit test and estimated overdispersion (\hat{c})
3475 using the parametric bootstrap procedure (MACKENZIE; BAILEY, 2004) implemented
3476 in program PRESENCE (software version 2.12.20; HINES, 2006). We evaluated model
3477 selection using second-order Akaike Information Criterion (AICc) or Quasi-Akaike
3478 Information Criterion (QAICc; Burnham and Anderson, 2002), selecting only those
3479 models with $\Delta\text{AICc} < 2$ (MACKENZIE *et al.*, 2006). In addition, we calculated
3480 cumulative model weights (w_+) for each explanatory variable in our balanced set of
3481 models associated with each of the four steps described above (BURNHAM;
3482 ANDERSON, 2002). Then, we used estimated effects from our best-supported model to
3483 generate predict maps for the maned sloth distribution at the study area.

3484 **3. Results**

3485 We detected the maned sloth 53 times at 32 out 68 sampled sites (naïve
3486 occupancy estimate = 47%). Among the sites with sloth detections, 53% had only one
3487 detection, 34% had two detections, and 6% had three or four detections. The forest
3488 cover around the sampled sites range from 5% to 100%, open areas cover ranges from 0
3489 to 69%, the proportion of important trees range from 0 to 1 and canopy height range
3490 from 8.9 to 30.5 m (Table A.3).

3491 The evaluation of the scale of effect (buffer size) for our two landscape variables
3492 suggested that a model that included forest cover at 500m and open area cover at 600m
3493 was approximately twice as likely as any other model in candidate set (Table A.1).
3494 These scales of effects were used in all subsequent modeling. The parametric bootstrap
3495 goodness of fit revealed some evidence of overdispersion ($\hat{c}=2.59$), so we used Quasi-
3496 AICc (BURNHAM; ANDERSON, 2002) for model selection procedures. Our best-

3497 supported forest threshold relationship suggested that maned sloth occupancy declined
 3498 when forest cover was below 30-35% (Tables 3A, A.4). We found some evidence that
 3499 detection probability was positively affected by canopy height ($w_+ = 0.32$; $\hat{\beta} = 0.09$,
 3500 $SE(\hat{\beta}) = 0.07$; Fig. 2), but a model with constant detection probability was more
 3501 parsimonious (model weight = 0.44, Table 3B). To ensure less bias in our occupancy
 3502 estimates, we used the more general structure where detection could vary among sites
 3503 with different canopy height in our evaluation of factors influencing sloth occupancy.
 3504

3505 **Table 3:** Summary of the model selection results for models of: (1) Scale off effect, (2)
 3506 thresholds effects of Forest Cover, (3) detection probability structure and (4) occupancy
 3507 probability structure. Reported statistics include QAICc = Quasi-AICc; w = model
 3508 weight; $-2\text{Log}(L) = -2 \text{Log Likelihood}$; K = number of parameters.

Model	QAICc	ΔQAICc	w	$-2\text{Log}(L)$	K
<i>(1) Threshold effect</i>					
$\Psi \sim$ Forest Cover_35% + Open Area Cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	117.32	0	0.21	250.07	8
$\Psi \sim$ Forest Cover + Open Area Cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	119.57	2.24	0.07	255.75	8
<i>(2) Detection probability</i>					
$\Psi \sim$ Forest Cover_35% + Open Area Cover + Important trees + Canopy Height, $p \sim$ null	114.30	0	0.44	255.22	6

$\Psi \sim$ Forest Cover_35% + Open Area Cover + Important trees + Canopy Height, $p \sim$ Canopy Height	114.98	0.68	0.32	250.64	7
$\Psi \sim$ Forest Cover_35% + Open Area Cover + Important trees + Canopy Height, $p \sim$ Important trees	116.56	2.26	0.14	254.63	7
$\Psi \sim$ Forest Cover_35% + Open Area Cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	117.33	3.027	0.01	250.07	8

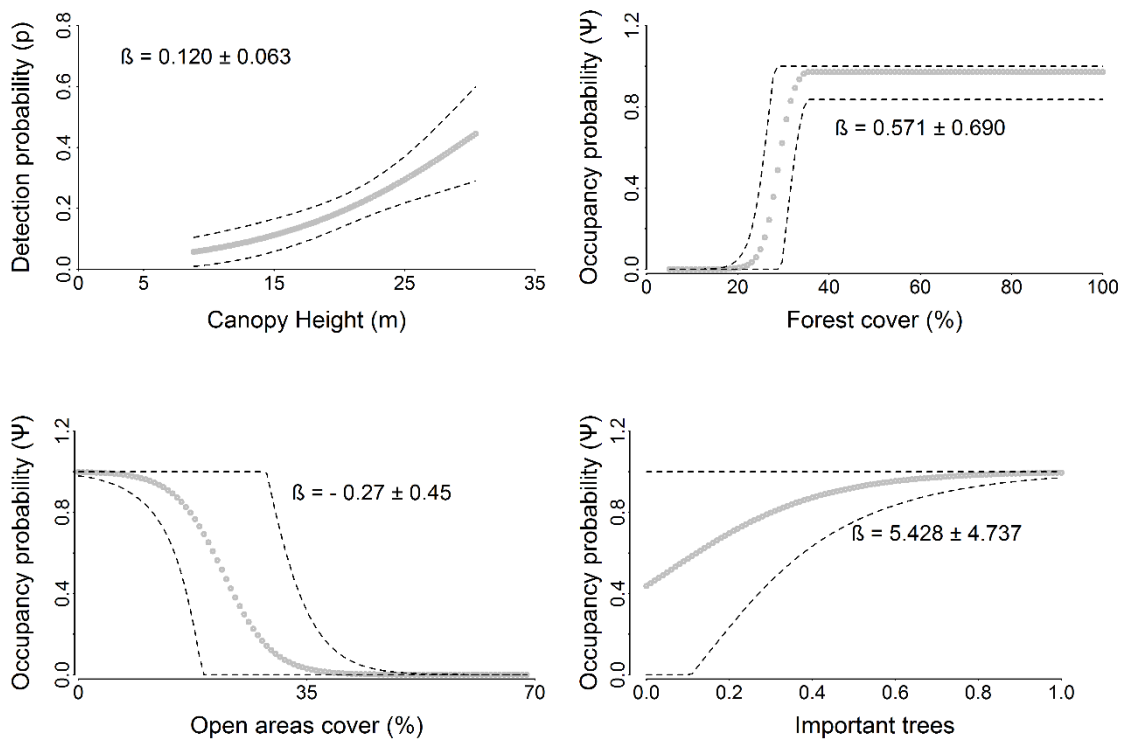
(3) Occupancy probability

$\Psi \sim$ Forest Cover_35%	110.24	0	0.20	255.92	4
$\Psi \sim$ Forest Cover_35% + Open Area Cover	111.09	0.86	0.13	252.11	5
$\Psi \sim$ Forest Cover_35% + Important trees	111.35	1.12	0.12	252.748	5
Global model	114.98	4.74	0.02	249.42	7
Null model	115.01	4.77	0.02	273.98	3

3509

3510 Among competing occupancy models, three had the $\Delta\text{QAICc} < 2$ (Tables 3C,
3511 A.5). Our best-supported model suggests that sloth occurrence is extremely high (near
3512 1.0) when there is over 35% forest cover surrounding a site, but sloth occurrence drops
3513 rapidly when forest cover is below this threshold (Fig. 2). Our data also suggested that
3514 sloths do not occur at sites with $\leq 20\%$ of forest cover (Fig 2). As expected, the
3515 proportion of important trees showed a positive relationship with occupancy probability
3516 and the percentage of open areas cover had a negative influence on sloth occurrence

3517 (Fig. 2). We found no evidence of interactive relationship between the forest cover and
 3518 the other three variables (Table A.5), possibly due to our distribution of sampling sites.
 3519 Landscape variables have the most pronounced importance: forest cover had the highest
 3520 cumulative weight ($w_+ = 0.78$), followed by the open area cover ($w_+ = 0.46$). The two
 3521 local variables had the lowest cumulative weights $w_+ < 0.40$ (Fig. 3).

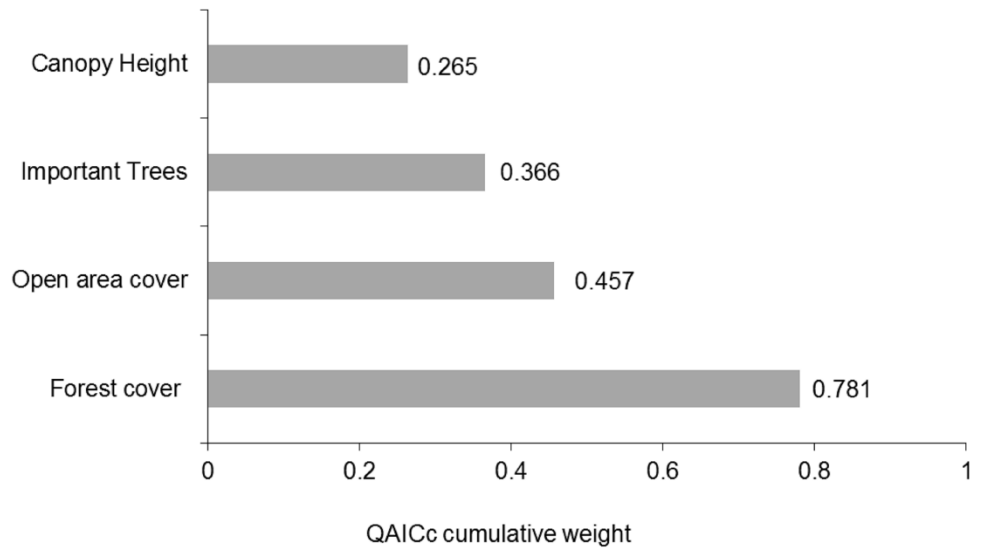


3522

3523 **Fig. 2:** Maned sloth detection and occupancy probabilities at 68 sampled sites within
 3524 Atlantic Forest, Espírito Santo, Brazil, predicted by the three best-supported models.

3525 The slope parameters (betas) estimated for the covariates “forest cover”, “open
 3526 areas cover” (second ranked model) and “important trees” (third ranked model) are also
 3527 shown. The gray line indicates the estimated covariate relationship and the black dash
 3528 indicate the standard error. The relationships between open areas and important trees are
 3529 graphed using the mean forest cover (%).

3530

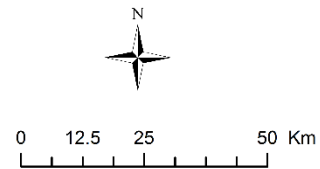
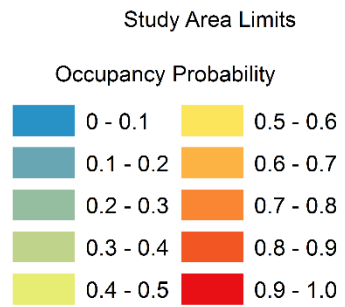
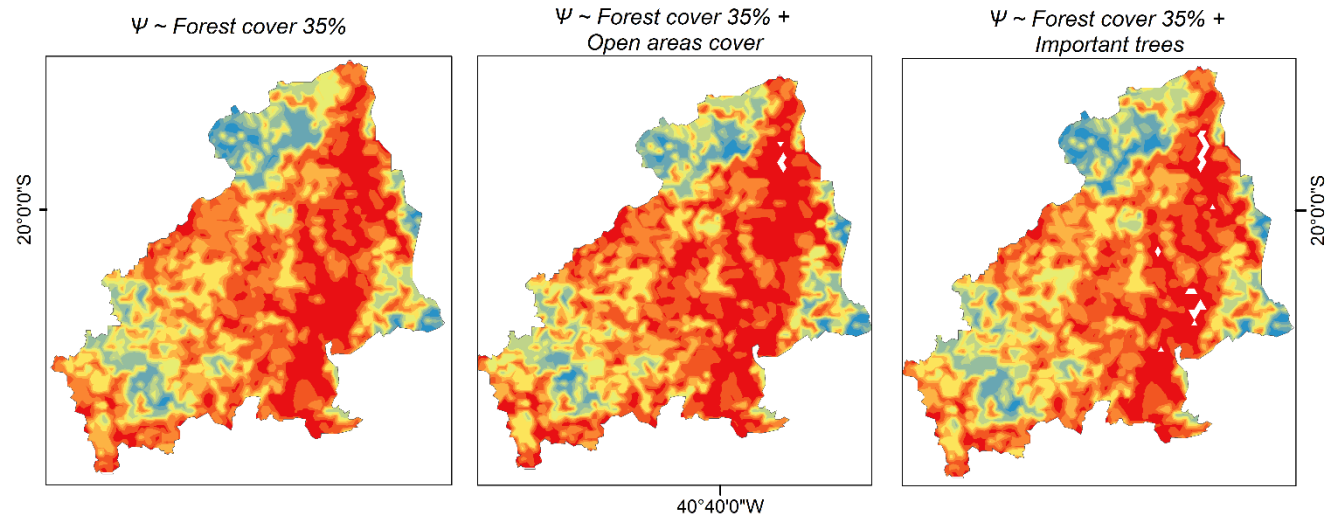


3531

3532 **Fig. 3:** Cumulative weight of evidence (QAIcC cumulative weight) for each covariate
 3533 used to model maned sloth occupancy within Atlantic Forest, Espírito Santo, Brazil.

3534

3535 Our predictive maps generated from the top three occupancy models show the
 3536 maned sloth occurring with higher probabilities in a relatively wide band that stretches
 3537 approximately north-southward in the study region (Fig. 4). In all three predictions, the
 3538 maned sloth has a high occupancy probability in the northeast and central part of the
 3539 study area. Conversely, the sloth is almost absent in the areas in the south, north and
 3540 east edge – which corresponds to the areas with low forest cover.



Geographical Coordinate System:
 Degrees, Minutes and Seconds
 Datum:
 Sirgass 2000

3542 **Fig. 4:** Predicted estimates of maned sloth occupancy probability in the study area within Atlantic Forest, Espírito Santo, Brazil; the three best-
3543 supported occupancy models were used.

3544 4. Discussion

3545 We found a threshold effect for the occupancy probability of the threatened and
3546 cryptic maned sloth. In contrast to previous study (SANTOS *et al.*, 2016), our results
3547 suggest that both forest cover and open areas cover strongly affect maned sloth
3548 occurrence across forest fragmentation gradients. The effect of local variables on
3549 detection probability at occupied sites was similar to our previous study, with canopy
3550 height positively influencing the maned sloth's detection. This positive relationship was
3551 contrary to our expectation and may be related to the higher abundance of the maned
3552 sloth in taller forests which, apparently, overcompensates the difficulties in finding
3553 sloths high up in the canopy.

3554 4.1 – Thresholds effects on occupancy probability

3555 The best-supported model predicts that areas with more than 35% of forest cover
3556 (65% of the study area) are likely to support maned sloth ($\hat{\psi} = 0.97$, $SE(\hat{\psi}) = 0.13$), but
3557 occupancy estimates decrease abruptly in areas below that threshold (Fig. 2), which
3558 corresponds to 35% of the study area (Fig. 4). Other habitat specialist groups exhibit
3559 similar responses to changes in habitat amount, including small to medium-sized
3560 terrestrial mammals (ESTAVILLO; PARDINI; DA ROCHA, 2013; OCHOA-
3561 QUINTERO *et al.*, 2015), woody plants (LIMA; MARIANO-NETO, 2014), and birds
3562 (ZUCKERBERG; PORTER, 2010). We also observed the absence of the maned sloth in
3563 areas with less than 20% of forest cover (15% of the study area; Fig.4). Forest loss in
3564 the region is already severe (originally 100% forest cover) and additional forest loss will
3565 increase the isolation of remaining forest patches and decrease patch size. These
3566 processes are likely to lead to the local extinction of sloth in some patches and reduce
3567 sloth movement between patches making recolonization difficult.

3568 As a forest specialist, habitat loss represents one of the major threats for the
3569 maned sloth. Landscapes with high concentration of natural habitats have smaller inter-
3570 patch distances and large numbers of patches (number of patches peaks at 30%, see
3571 GUSTAFSON; PARKER, 1992). This configuration allows the maned sloth – a strictly
3572 arboreal species – to move easily between forest patches. As the forest cover decreases,
3573 patch isolation and matrix resistance increase, followed by a decrease in patch size
3574 (ANDRÉN, 1994; SWIFT; HANNON, 2010). Isolated and small fragments might no
3575 longer support viable populations of maned sloth. In general, community composition
3576 may exhibit a shift (PARDINI *et al.*, 2010; BANKS-LEITE *et al.*, 2014; MAGIOLI *et*
3577 *al.*, 2015), holding mainly generalists species, since they can exploit different types of
3578 habitat to complement their diet and behavior or resource needs (ESTAVILLO;
3579 PARDINI; DA ROCHA, 2013; LIMA; MARIANO-NETO, 2014).

3580 *4.2 – Interactions between Forest Cover and local/landscape levels*

3581 Contrary to our expectations, we found no evidence of interaction between the
3582 forest cover and the local variables, though our findings may be influenced by our
3583 limited number of sites with forest cover values between 20-35%, where sloth
3584 occupancy estimates are greater than 0, but less than 1 (Fig 2). Forest structure and
3585 composition play an unquestionable role in sustaining high biodiversity (AUGUST,
3586 1983). Habitat loss may cause important changes in the habitat structure of remaining
3587 patches, with an increase of light demanding arboreal species (early successional), and
3588 canopy openness, and a decrease of overall basal area (ROCHA-SANTOS *et al.*, 2016;
3589 BENCHIMOL *et al.*, 2017). Species richness may decrease within patches (ANDRADE
3590 *et al.*, 2015; MAGIOLI *et al.*, 2015), and the compositional dissimilarity of plants
3591 between fragments may increase (ARROYO-RODRÍGUEZ *et al.*, 2013; BENCHIMOL

3592 *et al.*, 2017). This simplification of habitat composition and structure can limit food
3593 resource for the maned sloth, since in general, the sloths have a very restrictive diet,
3594 feeding on a few arboreal species (CHIARELLO, 1998b; MONTGOMERY;
3595 SUNQUIST, 1975), including some shade tolerant species (CHIARELLO, 1998b). The
3596 reduction of canopy height and increase of canopy openness can also influence sloths,
3597 since they spend most of their time on high canopy strata resting, feeding, breeding, and
3598 executing thermoregulatory behavior (MONTGOMERY; SUNQUIST, 1975; PAULI *et*
3599 *al.*, 2016; GARCÉS-RESTREPO *et al.*, 2017).

3600 Our second-best model suggests that sloth occupancy is influenced by both the
3601 open areas cover and forest cover, with a strong negative effect of open areas (Fig 2).
3602 Therefore, open areas may limit maned sloth movements, and the species may avoid it –
3603 a pattern also found previously (FALCONI *et al.*, 2015). Sloths can survive in small
3604 fragments of forest surrounded by an intermediate forest cover (20-35%) due to their
3605 relatively small home range (CHIARELLO, 1998b; FALCONI *et al.*, 2015). They can
3606 exploit living fences (*e.g.* narrow linear strips of planted trees; LEÓN; HARVEY,
3607 2006), isolated trees, forestry systems, and agriculture crops if these habitat patches
3608 provide food resources or are located between potentially accessible forest
3609 fragments(CASSANO; KIERULFF; CHIARELLO, 2011; PEERY; PAULI, 2014;
3610 VAUGHAN *et al.*, 2007). This is also true for other arboreal mammal, such as *Alouatta*
3611 *palliata mexicana* (ASENSIO *et al.*, 2009), *Colobus anglicus palliatus* (ANDERSON;
3612 ROWCLIFFE; COWLISHAW, 2007), and *Callicebus personatus* found in our study
3613 area (Pers. Observation). The use of living fences, isolated trees, forestry systems and
3614 agriculture crops – habitats with poor resources and conditions – by animals that are
3615 considered strictly arboreal provides evidence of the species adaptive capacity to
3616 maximize resource consumption in more anthropogenic landscapes due to the lower

3617 quality of existing forest fragments. However, the use of open areas may represent a
3618 severe predation hazard since the sloths, and other arboreal species, are frequently
3619 exposed to attacks by domestic and feral dogs (VAUGHAN *et al.*, 2007; OLIVEIRA *et*
3620 *al.*, 2008; GARCÉS-RESTREPO *et al.*, 2018). Similarly, since disturbed forests are of
3621 low quality, the sloths would be foraging on a lower and more open canopy, becoming
3622 more exposed to predation by ocelot (*Leopardus pardalis* - DELIBES *et al.*, 2011) and
3623 tayra (*Eira Barbara* - SÁENZ-BOLAÑOS *et al.*, 2018). In addition, the energy
3624 expenditure to cross larger open areas may be higher than its putative resource gains,
3625 since the sloth anatomy is not adapted to movements on the ground (GOFFART, 1971).

3626 4.3 – Implications for conservation

3627 We provide the first assessment of the maned sloth habitat requirements in a
3628 broader landscape context, representing the full spectrum of forest cover gradient, and
3629 with a multi-scale perspective covering a wide range of scale of effects. This
3630 understanding can lead to better targeted conservation actions, including a more
3631 accurate assessment of the species conservation status and threats. The use of thresholds
3632 in conservation and management actions might be debated, mainly because the
3633 threshold value may exclude species that require more contiguous habitat (JOHNSON,
3634 2013; VAN DER HOEK; ZUCKERBERG; MANNE, 2015). Additionally, when habitat
3635 loss is severe, biodiversity recovery may require different conservation approaches and
3636 the previous conservation state may never be obtained (JOHNSON, 2013; VAN DER
3637 HOEK; ZUCKERBERG; MANNE, 2015). Still, maintaining 30% to 35% of forest
3638 cover in an area can be important for a wide diversity of species and may conserve
3639 several ecological process essential to ecosystem function (BANKS-LEITE *et al.*, 2014;
3640 MARTENSEN *et al.*, 2012; MUYLAERT; STEVEN; RIBEIRO, 2016).

3641 To improve maned sloth conservation, our results support the need for an
3642 increase in forest cover, which will also increase habitat connectivity, by: 1) restoring
3643 and increasing riparian vegetation and 2) increasing matrix permeability with
3644 agroforestry systems and living fences. The Atlantic Forest has lost much of its riparian
3645 vegetation, and if this vegetation could be restored, the forest cover could increase to
3646 30% (REZENDE *et al.*, 2018), approaching the threshold level found in our study.
3647 Sloths typically use riparian forest for foraging and dispersal (RAMIREZ *et al.*, 2011;
3648 GARCÉS-RESTREPO *et al.*, 2018) and an early study observed maned sloths using
3649 swamp forests in Poço das Antas Reserve (PINDER, 1985). These areas are also useful
3650 for birds (TREMBLAY; ST. CLAIR, 2011) butterflies (VAN HALDER *et al.*, 2015),
3651 and bats (MUYLAERT; STEVEN; RIBEIRO, 2016; WORDLEY *et al.*, 2015).
3652 Additionally, riparian vegetation has a positive impact on water quality by reducing the
3653 nutrient and sediment load in streams (DOSSKEY *et al.*, 2010).

3654 Still, conserving and restoring riparian vegetation may not be enough to ensure
3655 the conservation of maned sloth. According to the Brazilian Native Vegetation
3656 Protection Law (Federal Law 12,727/2012), 20% of forest areas in rural properties must
3657 be set aside as legal forest reserves. However, this value is far below our threshold of
3658 35%, and we found that maned sloth is unlikely to exist in areas with less than 20% of
3659 forest cover (Fig. 2). Thereby, we reinforce that areas with less than 20% forest cover
3660 are unlikely to sustain viable populations of sloths. Additionally, the current law
3661 incorporates riparian vegetation in the calculation of the legal reserves, resulting in
3662 future deficits in the overall vegetation (SOARES-FILHO *et al.*, 2014), including our
3663 study area where 97.3% of forest cover are privately owned. To guarantee more
3664 consistent conservation, the restoration of riparian forest must be followed by forest
3665 reforestation in other areas.

3666 By mixing native vegetation with agricultural production, the agroforestry
3667 systems may be a sustainable way of reducing the contrast between the forest cover and
3668 the matrix, along with increasing forest area (SCHROTH; HARVEY, 2007; SANTOS
3669 *et al.*, 2019). When combined with other forested areas, these systems can offer more
3670 suitable habitat for maned sloths than the traditional agricultural system (CASSANO;
3671 KIERULFF; CHIARELLO, 2011) Moreover, providing different trees in the form of
3672 living fences yield new resource areas, resting refugia, and can act as movement
3673 corridors between habitat patches for the sloths, enhancing the landscape connectivity
3674 and reducing the pressure to traverse the anthropogenic open matrix (CASTELLÓN;
3675 SIEVING, 2006; LEÓN; HARVEY, 2006). Together, the restoration and increase of
3676 riparian vegetation and the creation of living fences and biodiverse agroforestry system
3677 may provide useful corridors and habitats for the maned sloth and for a wide range of
3678 species, thus allowing a long-term biodiversity conservation.

3679 In this paper, we found a threshold relationship between maned sloth occurrence
3680 and forest cover using an occupancy modeling framework. This framework is
3681 recommended for species with low detection probabilities, such as sloth species. Sloth
3682 occurrence declined quickly in areas with less than 35% forest cover and sloth were
3683 unlikely to occur in areas with less than 20% forest cover. Future studies aiming at
3684 understanding the potential interaction between landscape and local variables for this
3685 species may concentrate effort within this forest cover range and could provide valuable
3686 information for conservation efforts in degraded forest areas. In addition, we strongly
3687 recommend long-term monitoring of sloth in our study region – as well as in other key
3688 distribution areas of the species –incorporating movement monitoring, to estimate
3689 changes in occupancy over time and determine sloth movements and resource use
3690 within and between forest patches surrounded by different types of non-habitat and

3691 poor-habitat matrix. Such studies would make it possible to better infer how the maned
3692 sloth uses and selects their habitats in a heterogeneous and anthropogenic landscape,
3693 improving the management and conservation of this lesser known species.

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3707

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3970 **Appendix A. Supplementary data**

3971 **Table A.1:** Model selection results for supported models of scale of effect. The same general
 3972 model structure was used for sloth occupancy, Ψ (Canopy Height + ImpTrees + Forest cover
 3973 *scale of effect* + Open areas cover *scale of effect*), varying the scale (buffer, m) of each
 3974 landscape variable. Detection probability was modeled as a function of both local covariates,
 3975 p (Canopy Height + ImpTrees). For each model we report: AICc = Akaike information
 3976 criteria corrected for small samples; w = Model Weight; -2Log(L) = Measurements of model
 3977 fit; K = number of parameters. The bolded model represents the most parsimonious model in
 3978 the candidate model set.

Model	AICc	ΔAICc	w	K	-2log(L)
General structure: Ψ (Canopy Height + ImpTrees + Forest cover <i>scale of effect</i> + Open areas cover <i>scale of effect</i>), p (Canopy Height + ImpTrees)					
Forest cover 500 + Open areas cover 600	271.02	0.00	0.09	8	252.58
Forest cover 600 + Open areas cover 600	272.33	1.31	0.05	8	253.89
Forest cover 500 + Open areas cover 400	272.47	1.45	0.04	8	254.03
Forest cover 600 + Open areas cover 400	272.47	1.45	0.04	8	254.03
Forest cover 100 + Open areas cover 400	272.56	1.54	0.04	8	254.12
Forest cover 400 + Open areas cover 400	272.59	1.57	0.04	8	254.15
Forest cover 600 + Open areas cover 500	272.72	1.69	0.04	8	254.28
Forest cover 300 + Open areas cover 400	272.74	1.72	0.04	8	254.30
Forest cover 500 + Open areas cover 500	272.79	1.77	0.04	8	254.35
Forest cover 100 + Open areas cover 300	272.79	1.77	0.04	8	254.35
Forest cover 200 + Open areas cover 400	272.90	1.87	0.04	8	254.45
Forest cover 500 + Open areas cover 300	273.08	2.06	0.03	8	254.64
Forest cover 600 + Open areas cover 300	273.09	2.07	0.03	8	254.65
Forest cover 100 + Open areas cover 200	273.10	2.07	0.03	8	254.66
Forest cover 400 + Open areas cover 600	273.10	2.08	0.03	8	254.66
Forest cover 400 + Open areas cover 300	273.19	2.16	0.03	8	254.75
Forest cover 300 + Open areas cover 300	273.23	2.21	0.03	8	254.79
Forest cover 400 + Open areas cover 500	273.24	2.22	0.03	8	254.80
Forest cover 200 + Open areas cover 300	273.28	2.26	0.03	8	254.84
Forest cover 100 + Open areas cover 500	273.43	2.41	0.03	8	254.99
Forest cover 300 + Open areas cover 600	273.50	2.48	0.03	8	255.06
Forest cover 300 + Open areas cover 500	273.60	2.58	0.02	8	255.16
Forest cover 100 + Open areas cover 600	273.81	2.78	0.02	8	255.37

Forest cover 200 + Open areas cover 500	273.81	2.79	0.02	8	255.37
Forest cover 300 + Open areas cover 200	273.81	2.79	0.02	8	255.37
Forest cover 400 + Open areas cover 200	273.83	2.81	0.02	8	255.39
Forest cover 600 + Open areas cover 200	273.86	2.83	0.02	8	255.42
Forest cover 200 + Open areas cover 200	273.86	2.83	0.02	8	255.42
Forest cover 500 + Open areas cover 200	273.86	2.83	0.02	8	255.42
Forest cover 200 + Open areas cover 600	273.95	2.93	0.02	8	255.51
Forest cover 100 + Open areas cover 100	278.60	7.58	0.00	8	260.16
Forest cover 300 + Open areas cover 100	279.27	8.25	0.00	8	260.83
Forest cover 200 + Open areas cover 100	279.48	8.46	0.00	8	261.04
Forest cover 500 + Open areas cover 100	279.63	8.61	0.00	8	261.19
Forest cover 400 + Open areas cover 100	280.82	9.80	0.00	8	262.38
Forest cover 600 + Open areas cover 100	280.85	9.83	0.00	8	262.41

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3981 **Table A.2:** Pearson Correlation Matrix for selected covariates.

	Canopy Height (m)	Important Trees	Open areas cover (%) in 600 m	Forest Cover (%) in 500 m
Canopy Height (m)	1			
Important Trees	0.39	1		
Open areas cover (%) in 600 m	-0.46	-0.32	1	
Forest Cover (%) in 500 m	0.25	0.27	-0.50	1

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3984 **Table A.3:** General characteristics for the sampled sites in study area. Where: Total mean –
 3985 mean of the respective variable in al sampled sites; Non-detected = mean of the respective
 3986 variable in all sites which the maned sloth were not detected; Detected = mean of the
 3987 respective variable in all sites which the maned sloth were detected; m = meters; SD =
 3988 Standard Deviation.

Variable	Total Mean (SD)	Non-detected (SD)	Detected (SD)
Forest cover (%)	48.72 (±21.05)	41.47 (±10.49)	56.87 (±20.96)
Open areas cover (%)	10.46 (±11.75)	16.59 (±17.29)	3.56 (±3.47)
Important Trees (0-1)	0.63 (±0.27)	0.53 (±0.31)	0.73 (±0.20)
Canopy Height (m)	19.59 (±3.34)	18 (±3.12)	21.38 (±3.41)
Total of the sampled Sites	68	36	32

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3991 **Table A.4:** Model selection results for eight models exploring thresholds effect between
3992 forest cover and the Maned sloth occupancy probability. All models include a general
3993 detection probability structure, p (Canopy Height+ ImpTrees), and an occupancy structure
3994 that accounted for potential variation associated with our other covariates, Ψ (Open areas
3995 cover + ImpTrees + Canopy Height + *Threshold Relationship*) . For each model we report:
3996 QAICc = Quasi-AICc; w = Model Weight; $-2\text{Log}(L)$ = Measurements of model fit; K =
3997 number of parameters.

Model	QAICc	Δ QAICc	w	$-2\text{log}(L)$	K
$\Psi \sim$ Forest cover_35% + Open areas cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	117.33	0	0.213	250.07	8
$\Psi \sim$ Forest cover_30% + Open areas cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	117.7	0.37	0.18	251.02	8
$\Psi \sim$ Forest cover_40% + Open areas cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	118.09	0.77	0.15	252.02	8
$\Psi \sim$ Forest cover_45% + Open areas cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	118.35	1.02	0.13	252.65	8
$\Psi \sim$ Forest cover_50% + Open areas cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	118.51	1.20	0.12	253.12	8
$\Psi \sim$ Forest cover_25% + Open areas cover + Important trees + Canopy Height, $p \sim$ Canopy Height + Important Trees	118.93	1.61	0.1	254.13	8
$\Psi \sim$ Forest cover + Open areas cover + Important trees + Canopy Height,	119.57	2.24	0.07	255.74	8

$p \sim$ Canopy Height + Important Trees

$\Psi \sim$ Forest cover_20% + Open areas cover 119.99 2.66 0.06 256.81 8

+ Important trees + Canopy Height,

$p \sim$ Canopy Height + Important Trees

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4001 **Table A.5:** Model selection results for occupancy structures that included additive (16
4002 models) and interactive effects (6 models) of our local and landscape variables on Maned
4003 sloth occupancy probability. All models include the supported detection probability structure,
4004 p (Canopy Height). For each model we report: QAICc = Quasi-AICc; w = Model Weight; -
4005 $2\text{Log}(L) = -2 \text{Log Likelihood}$; K = number of parameters; + = Additive models; * =
4006 Interactive models. The bolded model represents the three better models in the candidate
4007 model set ($\Delta\text{QAICc} < 2$).

Model	QAICc	ΔQAICc	w	$-2\text{Log}(L)$	K
Ψ Forest cover_35%	110.23	0	0.20	255.92	4
Ψ Forest cover_35% + Open areas cover	111.10	0.86	0.13	252.11	5
Ψ Forest cover_35% + Important trees	111.35	1.12	0.12	252.74	5
Ψ Forest Cover_35% + Canopy height	112.36	2.12	0.07	255.36	5
Ψ Open areas cover	112.46	2.22	0.07	261.58	4
Ψ Forest Cover_35% + Open areas cover + Important trees	112.66	2.43	0.06	249.91	6
Ψ Forest Cover_35% + Open areas cover+ Canopy height	113.40	3.17	0.04	251.83	6
Ψ Forest Cover_35% + Canopy Height + Important trees	113.47	3.23	0.04	251.96	6

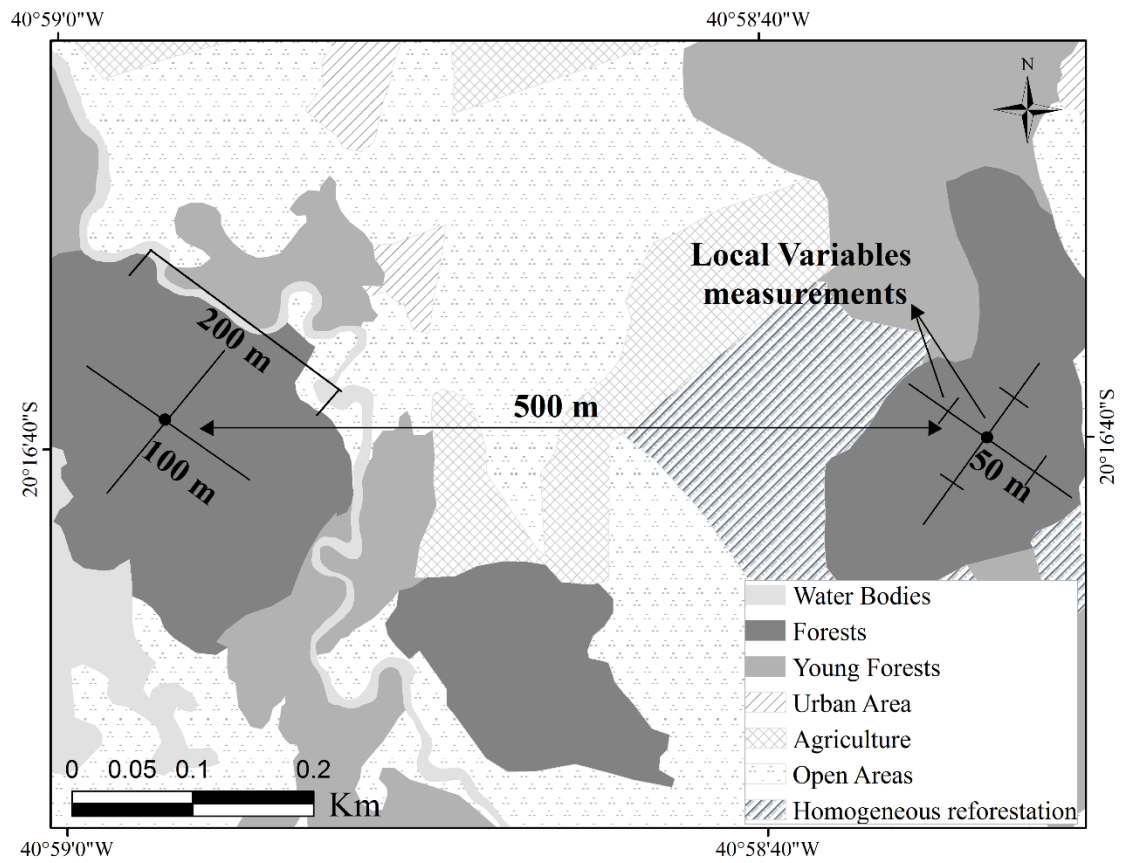
Ψ Forest Cover_35% *	113.49	3.25	0.04	252.08	6
Open areas cover					
Ψ Open areas cover+	113.639	3.40	0.04	258.56	5
Important trees					
Ψ Forest Cover_35% *	113.71	3.48	0.04	252.61	6
Important trees					
Ψ Open areas cover +	114.62	4.39	0.02	261.09	5
Canopy height					
Ψ Forest Cover_35% *	114.68	4.44	0.022	255.14	6
Canopy height					
Global model	115.00	4.74	0.02	249.42	7
Null model	115.01	4.77	0.019	273.98	3
Ψ Open areas cover+	115.36	5.12	0.02	256.74	6
Canopy height +					
Important trees					
Ψ Open areas cover*	115.72	5.48	0.01	257.73	6
Important trees					
Ψ Canopy height +	115.75	5.517	0.01	263.91	5
Important trees					
Ψ Important trees	115.94	5.71	0.01	270.44	4
Ψ Canopy height	116.56	6.33	0.01	260.10	4

Ψ Open areas cover*	116.62	6.38	0.01	272.12	6
Canopy height					
Ψ Canopy height	118.13	7.89	0.00	263.80	6
*Important trees					

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4012 **Fig. A.1:** Schematic representation of sample design.

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Capítulo 3

Natural regeneration can mitigate the climate change effects on suitable areas for the endangered maned sloth in Atlantic Forest, Brazil.

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4032 **Abstract**

4033 Climate changes and habitat loss are two of the major factors responsible for the species
4034 extinction. Urgent conservation measures are needed to stop the spread of these global
4035 threats, such as encouraging forest regeneration. This article sought to evaluate the
4036 environmental suitability for the endangered maned sloth (*Bradypus torquatus*) in its entire
4037 distribution, using SDM tools, and modeling the two evolutionary units separately – ESU01
4038 (North) and ESU02 (South). We also used Maxent to model future scenarios, taking into
4039 account climate change in 2050 and regeneration scenarios – BAU, scenario 02 less optimistic
4040 and scenario 03 – more optimistic. We used climate variables from WorldCim V.1 and we
4041 calculated two landscape variables – forest cover and pasture cover. We analysed the
4042 generated models in face of the Brazilian legislation, to understand how the suitable areas are
4043 in relation to their legal conservation. Suitable areas correspond to 17.71 % of the delimited
4044 area for ESU01, and 7.0% for ESU02. ESU01 exhibits a decrease of suitable areas in
4045 scenarios BAU and 02, but a stability in scenario 03. On the other hand, ESU02 exhibits a
4046 increase of suitable areas in all scenarios. For both populations, forest cover plays an
4047 important role in the current environmental suitability, but this pattern may change in future
4048 scenarios, mainly in ESU02, in which the climate variable gains importance in future
4049 scenarios. Most part of suitable areas are outside from any legal protection, and this pattern
4050 continues in future scenarios. The future conservation of *B.torquatus* will depend on landscape
4051 planning, in order to include different types of forest regeneration.

4052

4053 **Keywords:** Conservation, Forest regeneration, Landscape ecology, Global warming, Pilosa

4054

4055 **1. Introduction**

4056 Successful biodiversity conservation is intrinsically linked to the our capacity to
4057 foresee how of if a given habitat will support the presence of different species and viable
4058 populations in the future. Climate change will act as the primary actor in the following years,
4059 with severe negative impacts on biodiversity distribution (GILANI *et al.*, 2020), species
4060 richness and composition (GOUVEIA *et al.*, 2016a), and suitable areas for viable populations
4061 (ZHANG *et al.*, 2020). Human activities, such as agriculture, forestry, and other land uses are
4062 responsible for 23% of global greenhouse gas emissions (IPCC, 2018). In addition, several
4063 biomes are experiencing high rates of habitat loss and decreasing biodiversity, since small
4064 and isolated habitats may be inadequate to sustain viable populations (FAHRIG, 2003;
4065 SILVA *et al.*, 2015).

4066 Globally, forest areas play a fundamental role in regulating local and global climate,
4067 mitigating the effects caused by climate (PREVEDELLO *et al.*, 2019; SILVÉRIO *et al.*,
4068 2015); thus, reducing deforestation is imperative and urgent. Additionally, the regeneration
4069 of natural habitats is crucial to reduce the impacts of global warming (BAPTISTA; RUDEL,
4070 2006; RUDEL *et al.*, 2005; RUDEL; BATES; MACHINGUIASHI, 2002), since growing
4071 young forests are excellent carbon sinks (CASPERSEN *et al.*, 2000; STRASSBURG *et al.*,
4072 2014). Deforested areas can be recovered in two ways: active restoration and passive — or
4073 natural — regeneration (CHAZDON, 2012). The latter is considered the cheapest form of
4074 forest recovery, since it is simply a natural process of ecological succession (CHAZDON,
4075 2012). The former involves human labor for planting seedlings and seeding directly and is
4076 therefore costlier, especially for small landowners (LAMB; ERSKINE; PARROTTA, 2005;
4077 but see BANKS-LEITE *et al.*, 2014). Some conservations initiatives are fundamental in order
4078 to ensure the long-term preservation of natural environments and the species that inhabit
4079 them, such as the establishment of a protected areas system (NAUGHTON-TREVES;

4080 HOLLAND; BRANDON, 2005) and the conservation of natural areas in private lands
4081 (BOWERS, 1999; METZGER *et al.*, 2019).

4082 The Brazilian Atlantic Forest has been reduced to just over 28% of its original size
4083 (REZENDE *et al.*, 2018), and is composed mainly of fragments smaller than 50 ha that are
4084 immersed in a heterogeneous matrix (RIBEIRO *et al.*, 2009), and highly vulnerable to
4085 climate change (BELLARD *et al.*, 2014; SOBRAL-SOUZA *et al.*, 2018). Still, the biome has
4086 been experiencing a decrease in deforestation rates, with the lowest deforestation rate in 30
4087 years recorded between 2017 and 2018 (FUNDAÇÃO SOS MATA ATLÂNTICA, 2018).
4088 Furthermore, some regions report an increase in forest cover, mainly due to natural
4089 succession processes (BAPTISTA; RUDEL, 2006; DE REZENDE *et al.*, 2015; TEIXEIRA *et*
4090 *al.*, 2009).

4091 Even with its long history of deforestation, the biome is still home to the endemic
4092 maned sloth (*Bradypus torquatus*), a threatened species according to the national red list
4093 (ICMBio. CHIARELLO *et al.*, 2018) and by the international list (IUCN. CHIARELLO ;
4094 MORAES-BARROS, 2014). Due to their strictly arboreal and folivorous habit
4095 (CHIARELLO, 1998a, b), maned sloths need sizeable forest cover for their survival and are
4096 practically absent in areas with less than 20% forest cover (SANTOS *et al.*, 2019).
4097 Additionally, the presence of large open areas such as abandoned pasture lands has a
4098 detrimental effect on them (FALCONI *et al.*, 2015; SANTOS *et al.*, 2019). Since they are
4099 highly adapted and at the same time, restricted to the arboreal environment, a very high
4100 energy expenditure is required to cross these open areas and doing so exposes them to
4101 opportunistic predation events (PEERY; PAULI, 2014; VAUGHAN *et al.*, 2007).
4102 Additionally, they are very sensitive to climatic conditions, especially to changes in
4103 temperature, since their low metabolic rate causes their body temperature to vary according to
4104 the environment (CHIARELLO, 1998a; GINÉ *et al.*, 2015).

4105 *B. torquatus* has a restricted and discontinuous distribution in the Atlantic Coast
4106 forest, (HIRSCH; CHIARELLO, 2012; SANTOS *et al.*, 2019), inhabiting fragments in four
4107 Brazilian states—Sergipe, Bahia, Espírito Santo, and Rio de Janeiro (HIRSCH;
4108 CHIARELLO, 2012; SANTOS *et al.*, 2019). Overall, populations of *B. torquatus* are
4109 genetically structured, with northern lineages (Sergipe and Bahia) being divergent from
4110 southern ones (Espírito Santo and Rio de Janeiro) to the point of being considered two
4111 distincts Evolutionary Significant Units (ESUs) (SCHETINO; COIMBRA; SANTOS, 2017).
4112 This diversification occurred during the Miocene-Pliocene transition, probably as a result of
4113 climate events in Atlantic Forest (SCHETINO; COIMBRA; SANTOS, 2017), and the current
4114 natural boundaries between the two lineages is located in northern Espírito Santo (from the
4115 limit with Bahia, to the north of Rio Doce). Consequently, it is extremely important to treat
4116 these two lineages as if they were different species (MOREIRA *et al.*, 2014; SCHETINO;
4117 COIMBRA; SANTOS, 2017).

4118 Species Distribution Models (SDM) are an important tool given its capacity for
4119 predicting areas where the species in question is likely to be found (ELITH; LEATHWICK,
4120 2009). Through occurrence data and environmental variables, this tool allows researchers to
4121 understand the relationship between the distribution of a given species and the environmental
4122 conditions, and thus to map environmentally suitable areas for the species (FERRAZ *et al.*,
4123 2012). The approach has been widely used to identify priority areas for species conservation
4124 (CARRASCO *et al.*, 2020; PORTUGAL *et al.*, 2019; RODRIGUES *et al.*, 2004), define
4125 areas for regeneration through ecological corridors (MORATO *et al.*, 2014), predict potential
4126 hybridization zones between native and invasive species (MORAES *et al.*, 2019), and
4127 identify possible effects of habitat regeneration on species conservation (ANGELIERI *et al.*,
4128 2016), among other uses.

4129 Climatic stability and large forest areas act positively for the occurrence of *B.*
4130 *torquatus* (CHIARELLO 1998a; GINÉ *et al.*, 2015; MOREIRA *et al.*, 2014; SANTOS *et al.*,
4131 2019), yet, it is uncertain how climate and landscape changes will interact in the future.
4132 Using the Species Distribution Models (SDM) and considering the two evolutionary units,
4133 this novel study aimed to 1) Evaluate the environmental suitability for *B. torquatus*
4134 throughout its distribution; 2) Assess environmental suitability in the future, using natural
4135 forest regeneration and climate change scenarios; 3) Analyze the resulting scenarios - present
4136 and future - in accordance with the Brazilian environmental legislation. Thus, we evaluated
4137 whether suitable areas are legally protected and the likely gains from suitable areas will be
4138 protected in the future.

4139 **2. Material and Methods**

4140 *2.1 Study area*

4141 The Atlantic Forest covers part of the Brazilian east coast, extending to more inland areas
4142 in its southern distribution (MUYLAERT *et al.*, 2018). Throughout its extent, the biome
4143 encompasses different vegetation types, and the ombrophilous formations - preferred by
4144 *B.torquatus* - are located near coastal areas and in the eastern slope of Serra do Mar
4145 (OLIVEIRA-FILHO; FONTES, 2000). We delimited the study area using the WWF
4146 terrestrial ecoregions database (OLSON *et al.*, 2001), selecting only the vegetation types
4147 present in the current known distribution of *B.torquatus* (HIRSCH; CHIARELLO, 2012). We
4148 also included ecoregions present in the most northerly areas of ecoregions, comprising the
4149 states of Pernambuco (PE), Paraiba (PB), and Rio Grande do Norte (RN, Fig. 1). Such
4150 coverage is justified by the fact that there are reports of the species in such areas, although
4151 this has never been confirmed (HIRSCH; CHIARELLO, 2012). The defined area is 237,935

4152 km², encompassing coastal areas in eight states and consisting mainly of perennial vegetation,
4153 sandbanks, and mangroves (Fig. 1).

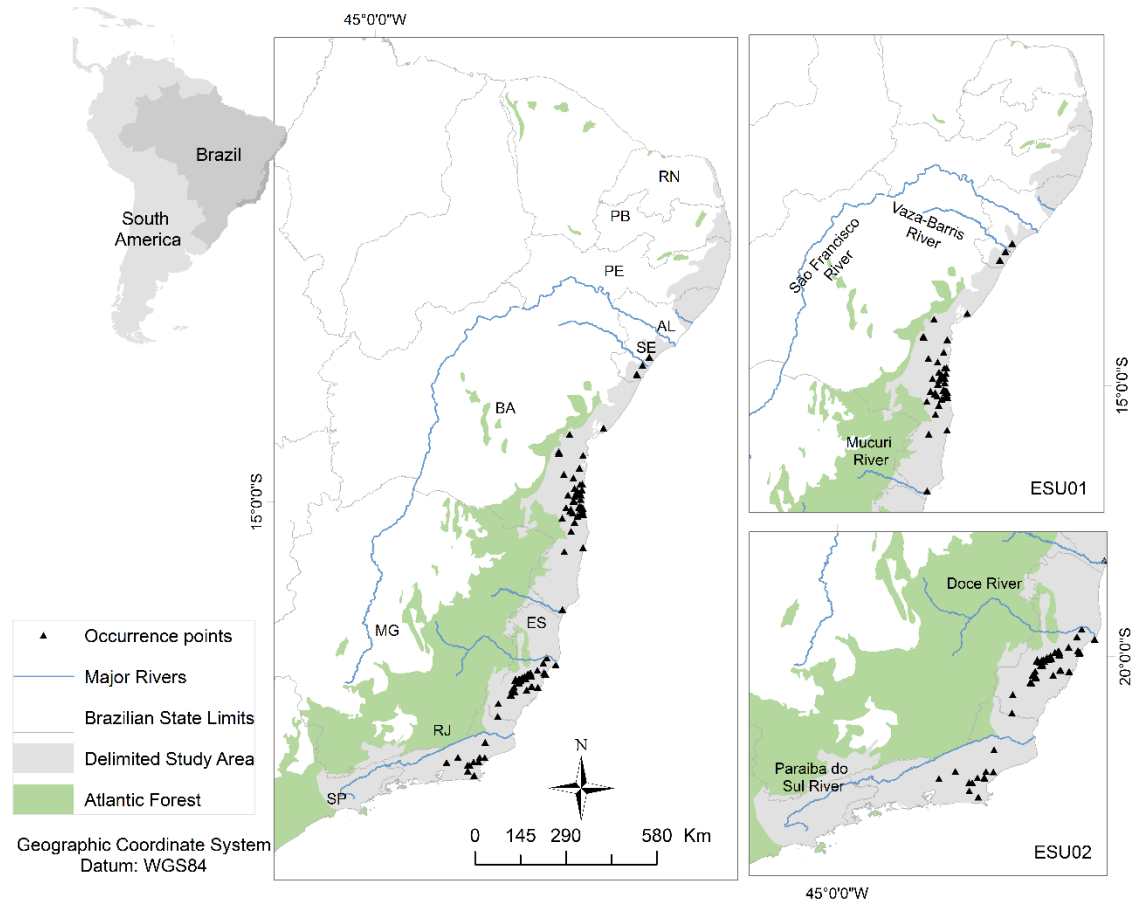
4154 *2.2 Modeling framework*

4155 In this study, we opted to model current and future suitable areas separately for each of
4156 the Evolutionary Significant Units (ESU) of *B. torquatus* (SCHETINO; COIMBRA;
4157 SANTOS, 2017). We used ESU01 and ESU02 to refer to the two modeled areas (Fig. 1):
4158 ESU01 corresponding to the northern population, located in Sergipe and Bahia, and ESU02
4159 to the southern population, present in Espírito Santo and Rio de Janeiro (SCHETINO;
4160 COIMBRA; SANTOS, 2017).

4161 First, we modelled the current scenario, using current climate data and current landscape
4162 data. Then, we modeled the future environmental suitability surfaces in three scenarios: 1)
4163 Scenario 01, Business as Usual (BAU), projected future climate, but without changes in
4164 landscape. In this case we analyzed the losses/gains of suitable areas only from the
4165 perspective of climate change. The other two scenarios correspond to the analysis of
4166 losses/gains of suitable areas from the perspective of climate change and of regeneration
4167 scenarios: 2) Scenario 02, less optimistic, projected future climate and changes in landscape,
4168 with a small rate of regeneration, only enough to maintain the species, i.e., 35% of forest
4169 cover, a threshold level of forest cover we detected previously (Santos et al., 2019); and 3)
4170 Scenario 03, more optimistic, projected future climate and changes in landscape with high
4171 rate of regeneration, beyond the minimum amount to maintain the species. We used the same
4172 variables for present and for future scenarios.

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4175

4176 **Fig. 1** Study area delimited by WWF Ecorregions (Olson et al., 2001) and distribution of
 4177 occurrence points in each Evolutionary Significant Units (ESU).

4178 2.3 Occurrence points and sampling bias

4179 To construct the habitat suitability models we compiled *B. torquatus* occurrence data
 4180 available from Neotropical Xenarthrans data papers (SANTOS *et al.*, 2019). This database is
 4181 a vast compilation of data from various sources and researchers, including data that were
 4182 previously unknown and inaccessible, totaling 638 records. Subsequently, we filtered all
 4183 occurrence points in five steps. First, all points with coordinate accuracy above 100 meters
 4184 were eliminated. The data paper included the information about coordinate precision,
 4185 allowing us select only location points, and not locality. Second, all points related to
 4186 apprehension, interview, and roadkill were eliminated, since these are often not representative
 4187 of the real location of the species and are often inaccurate. Third, only data obtained from

4188 2009 to present were selected, to be temporally consistent with the environmental data used.
4189 Fourth, duplicate points were eliminated, using the program RStudio 3.5.1 (R CORE TEAM,
4190 2018), *vegan* package (Okansen et al., 2007). Fifth, we maintain all points with a minimum
4191 distance of 500 m to avoid spatial correlation (SANTOS, *et al.*, 2016, 2019). At the end, a
4192 database with 174 points was generated, consisting of 94 points from ESU01 and 70 points
4193 from ESU02 (Fig. 01). Finally, we developed a sampling probability surface, called bias grid,
4194 to reduce the sampling bias in areas more intensely sampled than others (PHILLIPS *et al.*,
4195 2009) by decreasing the weight of these locations. The bias grid was created by calculating
4196 the Gaussian Kernel density of the sampling locations with a distance of 30 km. This value
4197 best represents the two large clusters of sampling points - in the south of Bahia and in the
4198 mountain region of Espirito Santo.

4199 *2.4 Environmental variables and future scenarios*

4200 We obtained the bioclimatic data through the WorldClim data base (WordClim V.1)
4201 and elevation data through USGS (United States Geological Service) (Table A.1). Since the
4202 bioclimatic variables are generally correlated with each other, a factor analysis with
4203 maximum variation rotation was performed, separately for each ESU, to check for
4204 multicollinearity. We selected only 2 bioclimatic variables for the ESU: Precipitation of
4205 Wettest Month (Bio13) and Precipitation Seasonality (Bio15) for ESU01 and Precipitation of
4206 Driest Month (Bio14) and Temperature Seasonality (Bio04), for ESU02. For future climate
4207 scenarios, we selected the bioclimatic variables from two Representative Concentration
4208 Pathways (RCPs), related to different scenarios of greenhouse gas emissions for 2050. The
4209 RCP 4.5 represent the less optimistic scenario, and RCP 8.5 represents the more pessimistic
4210 climate scenario (IPCC, 2014). For each RCP, we considered three Global Climate Model
4211 (GCM) – GCMS4, HadGEM2-ES and MIROC-ESM. These models have performed very
4212 well in other studies of primates in the Atlantic Forest (GOUVEIA *et al.*, 2016a).

4213 The two landscape variables chosen for modeling was forest cover and pasture cover,
4214 which are important predictors for the presence of *B. torquatus* (SANTOS, PALOMA
4215 MARQUES *et al.*, 2019). We used the spatial data available from The Brazilian Foundation
4216 for Sustainable Development (In Portuguese Fundação Brasileira para o Desenvolvimento
4217 Sustentável - FBDS) and the Laboratory of Processing Images and Geoprocessing (In
4218 Portuguese Laboratório de Processamento de Imagens e Geoprocessamento - LAPIG (Table
4219 A.1). The software GrassGis 7.4.4 (Grass, 2018) was used to perform all landscape analyses.

4220 To create the regeneration scenarios, we used a map of pixels with values of natural
4221 regeneration potential in pastures areas, ranging from 0 to 100, where higher values
4222 represented greater potential for regeneration (NIEBUHR *et al.*, 2017). First, we calculated
4223 how much area would need to be regenerated to ensure the presence of *B. torquatus* in the
4224 delimited area, based on Santos *et al.*, (2019a), which reported a critical threshold on habitat
4225 amount to *B.torquatus* presence: the occupancy probability start to decline in areas with less
4226 than 35% of forest cover. Currently, only 25.45% of the area for maned sloth in ESU01 is
4227 covered by the Atlantic Forest remnants, and 27.61% in ESU02. To guarantee future
4228 populations of the species, it would be necessary to restore 9.55% (11,794.67 km²) and 7.39%
4229 (8,455.20 km²), respectively, to reach 35% of forest cover. So, for each ESU, we selected the
4230 pixels necessary to reach this goal (Table 01).

4231 Second, we selected the pixels in the regeneration map to create the different scenario
4232 – except for Scenario 01 (BAU). For Scenario 02, we selected all pixels with regeneration
4233 potential higher than 10 for ESU01 and selected all pixels with regeneration potential higher
4234 than 40 for ESU02 (Table 01). For Scenario 03, we selected all pixels with some regeneration
4235 potential for both ESUs (Table 1). The third step consisted of mosaicking the raster created
4236 with the current forest cover raster (Table A.1), creating forest cover maps for future
4237 scenarios. The reverse process was done with the pasture cover, in which we subtract from

4238 the current raster pasture areas that will be potentially regenerated, creating the future pasture
 4239 cover maps. The fourth step was to calculate the percentage of forest and pasture in each
 4240 pixel of 1km.

4241 **Table 1** Accumulated area of pixels in the regeneration map for both ESU (central and left
 4242 column). The bold lines correspond to the minimum pixel values selected to create the
 4243 Scenario 02 map.

Pixel Value	Accumulated area (Km ²) of pixels -	
	ESU01	ESU02
90	393.55	600.81
80	1227	1976.47
70	2041.61	3364.14
60	3210.95	5372.95
50	4600.42	7795
40	6226.5	10606.72
30	8007.82	13743.87
20	10382.57	17815.78
10	13216.83	22714.19
0	22800.23	39661.32

4244

4245 To rule out possible correlations between all selected variables, we performed a
 4246 correlation analysis using a Pearson correlation matrix, for which the results were $p > 0.7$
 4247 indicating that none of the variables were correlated. For the modeling, the chosen
 4248 environmental variables were all standardized to a spatial resolution of 0.0083 decimal
 4249 degrees (approximately 1 km).

4250 2.5 Species Distribution Models (SDM)

4251 We use the Maximum Entropy algorithm to generate the distribution models through
4252 the Maxent 3.4.1 software (PHILLIPS; DUDI, 2008). Maxent is a modeling technique with
4253 high accuracy and better performance than other methods (MORAES *et al.*, 2019; ROURA-
4254 PASCUAL *et al.*, 2009). MaxEnt uses only presence points and background points – also
4255 namely as pseudo-absence points, environmental variables and sampling points to calculates
4256 an environmental suitability index, indicating where the species is most likely to occur, with
4257 values ranging from 0 to 1.

4258 We calibrate the current and future models using bootstrapping methods with 10
4259 random partitions, in which 70% of the data set went for training and 30% went for testing.
4260 All models were generated with a convergence threshold of 1.0^{E-5} with 500 interactions and
4261 10,000 background points. We use a Jackknife test to measure the relative importance of each
4262 variable in model performance. To evaluate the performance of resulting models, we
4263 calculate the Area Under the Curve (AUC) of the Receiving Operator Characteristic (ROC),
4264 in which values above 0.75 correspond with great discriminatory capacity between the
4265 generated models and random models.

4266 The 20 final models (10 models for each ESU) for current (2 models) and future
4267 scenarios (18 models – 9 from each RCP) were converted into binary maps to discriminate
4268 between suitable and unsuitable maps, applying a threshold rule defined by the 10th
4269 percentile of training presence points. This threshold removes suitability values that are less
4270 than the 10% higher suitability values (MUKHERJEE *et al.*, 2020). Binomial probability and
4271 omission error were also evaluated. We create consensus models to facilitate the presentation
4272 on the results, through the extension BioDinamica, from DinamicaEgo software (SOARES-

4273 FILHO; RODRIGUES; COSTA, 2009). The results of each RCP are available in the
4274 supplementary material (Fig S1).

4275 *2.6 Suitable areas conservation*

4276 After applying the threshold in all resulting maps, we analyzed the different scenarios
4277 according to Brazilian Native Protection Conservation Law (LPVN. Law 12,727/2012) and to
4278 the National System of Conservation Units (In portuguese Sistema Nacional de Unidades de
4279 Conservação – SNUC. Law 9,985/2000). The LPVN established the Legal Reserve (RL, from
4280 Reserva Legal, in portuguese) of 20% of native vegetation in private properties and the
4281 Permanent Protection Area (APP from Área de Preservação Permanente, in Portuguese)
4282 focused on preservation of river springs, river banks, lakes, lagoons, hilltops and steep slopes.
4283 The SNUC establish two types of protected areas – Strictly protected areas and Protected
4284 areas of sustainable use. We obtained a GIS database available from different official sources
4285 to calculate the percentage of suitable areas in three different conservation categories:
4286 riparian APP, Strictly Protected Area and Protected Area of Sustainable Use (Table A.1).

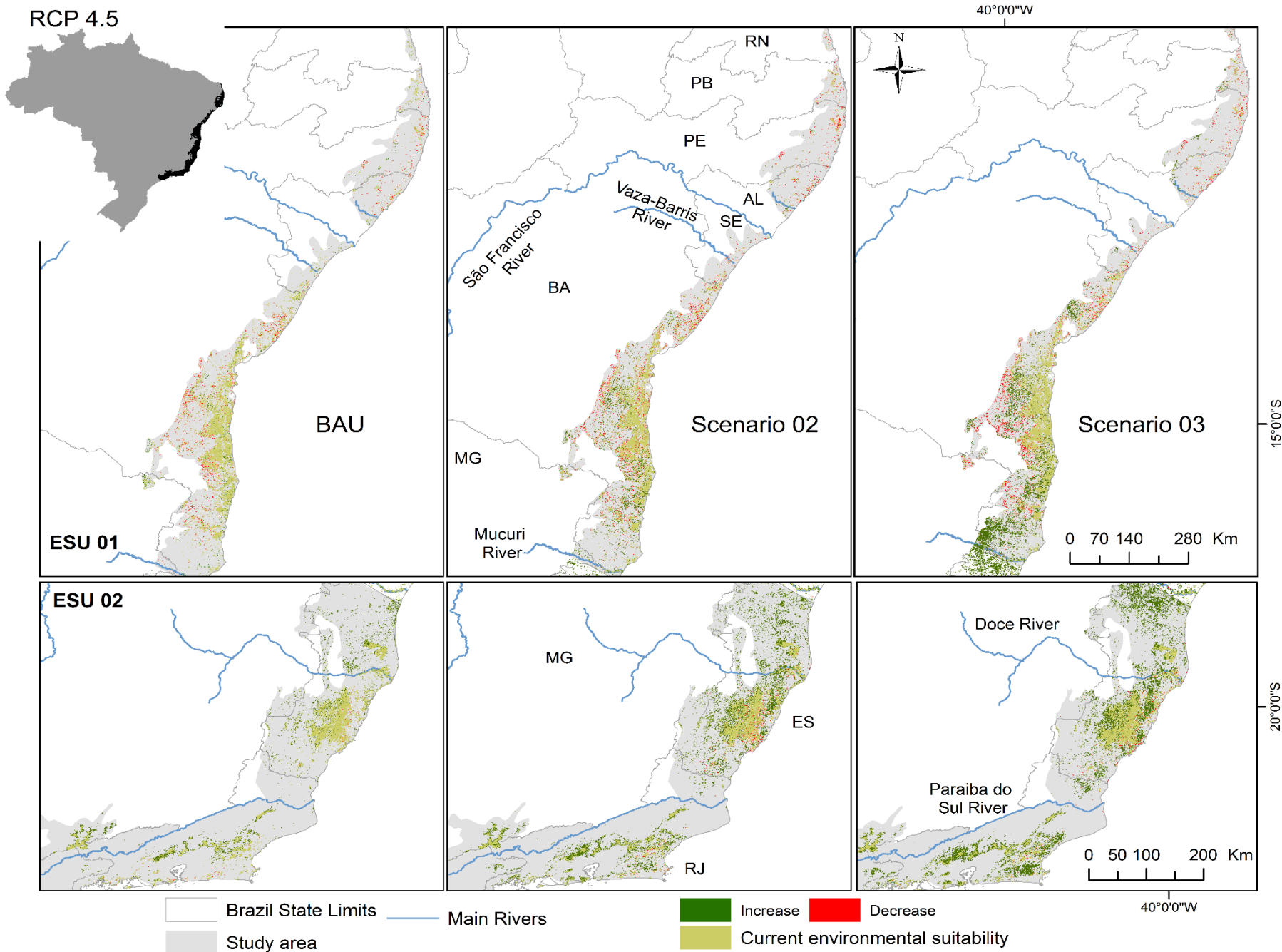
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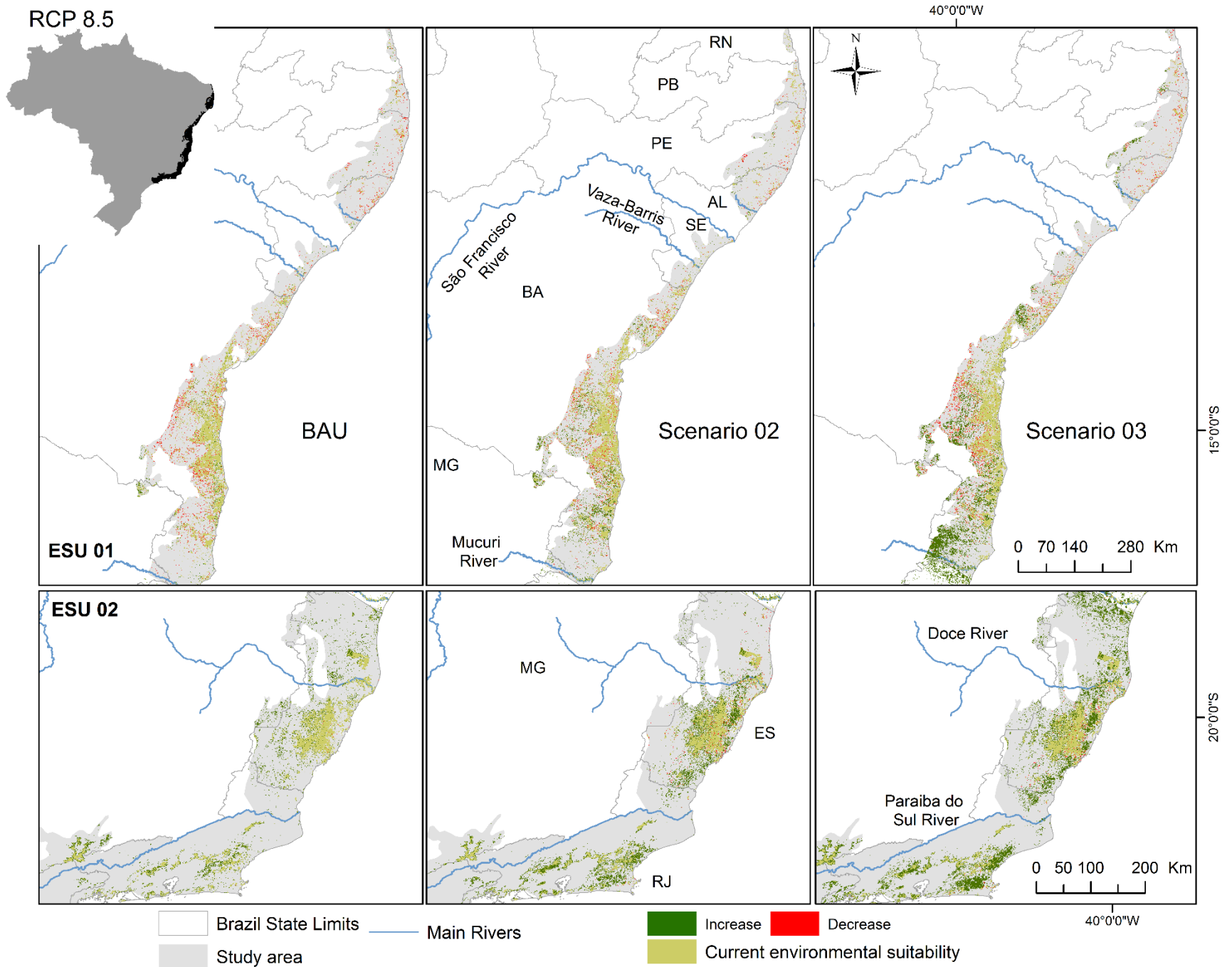
4288

4289 **3. Results**

4290 All resulting models had good discriminatory power according to AUC values,
4291 ranging from 0.8774 to 0.9613 (Table A.2). The current model predicts that 29,886.07 km² -
4292 only 4.15% of the study area – are suitable for *B.torquatus*. ESU01 was found to have higher
4293 cover of suitable areas (17.71 % or - 21,872.38 km²) then ESU02 (7.0% or- 8,013.69 km²).
4294 Two areas of the range of *B. torquatus* stand out for their high suitability: South Bahia and
4295 Center-South Espírito Santo (Fig. 2).

4296 In future scenarios, both Esu01 and ESU02 show differences in future land cover
4297 changes (Table 02, Fig. 02). The ESU01 would experience a loss of suitable areas, even in
4298 the regeneration scenarios, mainly further inland and more northern areas (Table 02; Fig. 02).
4299 In the BAU scenario, the suitable areas would decrease more abruptly, reaching 4033.02 km²
4300 (4.0%) in losses by 2050 (Table 02). The loss would be subtler in Scenario 02, and in
4301 Scenario 03, a stability in the suitable areas would take place only in RCP 4.5 (Table 2). On
4302 the other hand, ESU02 would experience an increase in the amount of suitable areas,
4303 including in the scenario without regeneration (Table 2; Fig 02).





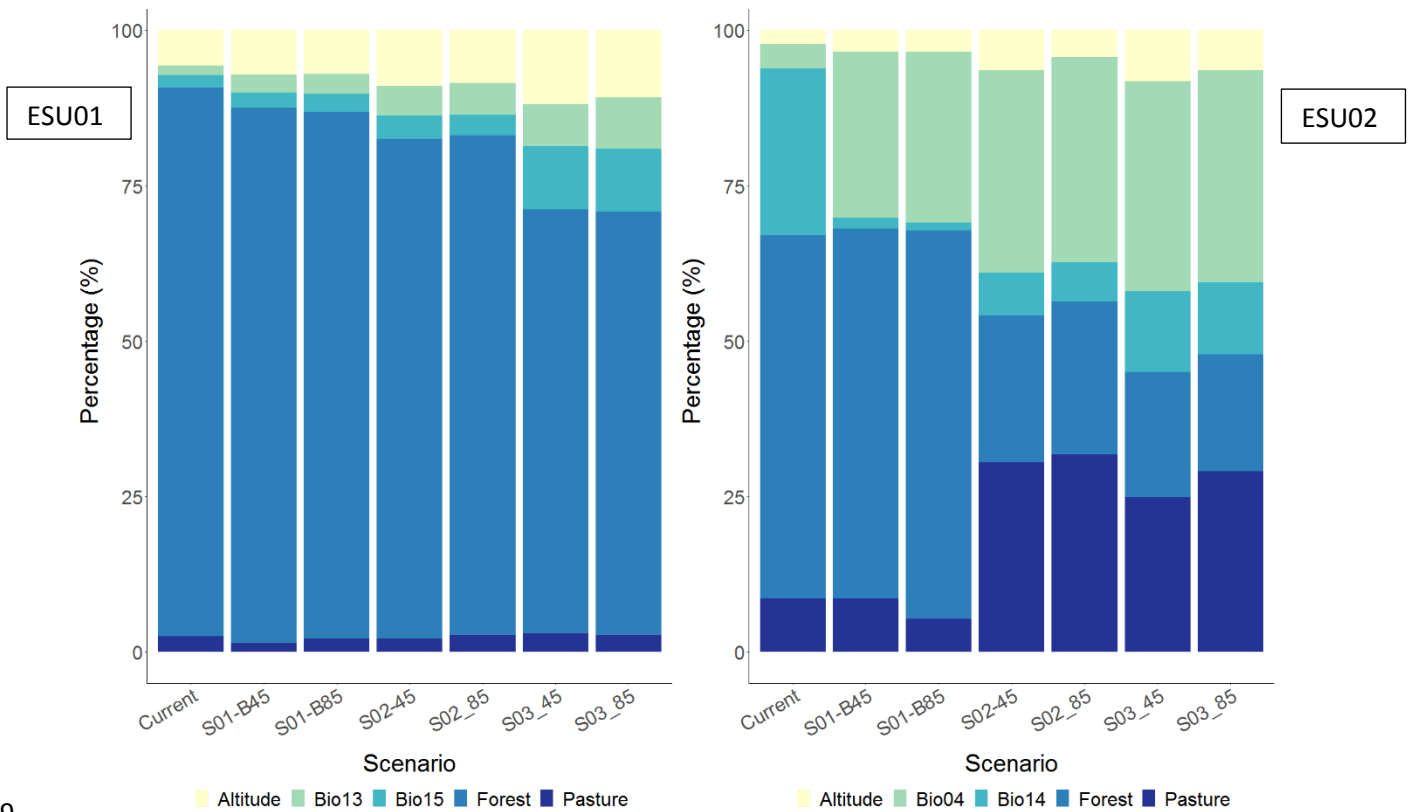
4306 **Fig. 02** Increase and decrease of suitable areas in relation to current environmental suitability for both populations of *Bradypus torquatus*,
4307 considering regeneration scenarios and climate changes. BAU – Business As Usual; RN – Rio Grande do Norte; PB – Paraíba; PE –
4308 Pernambuco; AL – Alagoas; SE – Sergipe; BA – Bahia; MG – Minas Gerais; ES – Espírito Santo; RJ – Rio de Janeiro.

4309 **Table 02** Suitable habitat in km² for the two populations, in the different scenarios. S01 –
 4310 Scenario 01 (BAU), S02 - Scenario 02, S03 - Scenario 03. 45 and 85 – RCP 4.5 and 8.5,
 4311 respectively

Scenario	ESU01	ESU02	Total area
Current	21872.38	8013.69	29886.07
S01-45	16939.36	9450.87	26390.23
S01-85	16130.52	10366.02	26496.53
S02-45	18595.45	12525.23	31120.68
S02-85	19382.81	13002.29	32385.10
S03-45	22030.83	14055.82	36086.65
S03-85	21115.40	14270.71	35386.11

4312

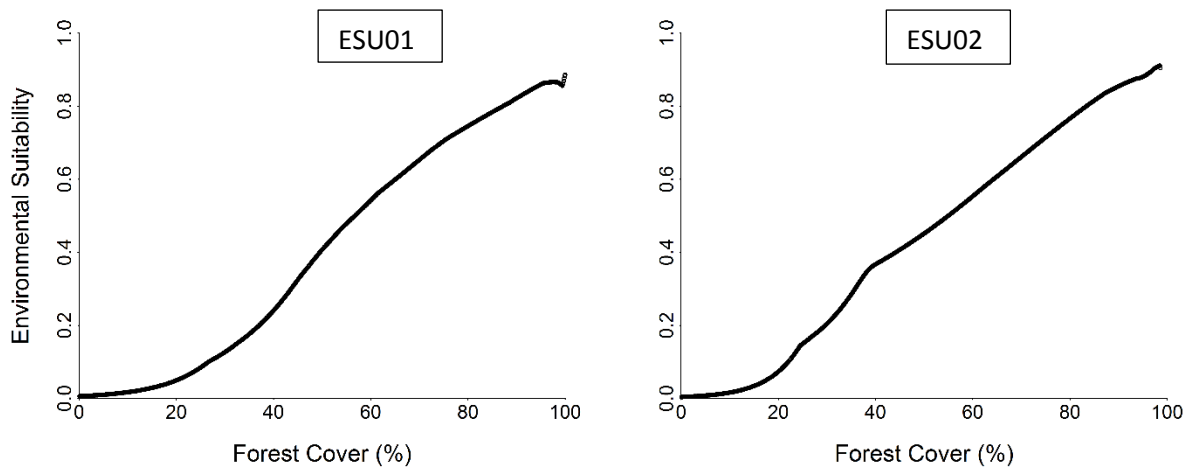
4313 According to the Jackknife test, the environmental parameter with the highest
 4314 influence on environmental suitability was forest cover for ESU01 (Fig.3), which positively
 4315 influenced the suitability (Fig.04). However, ESU02 showed a different pattern: the forest
 4316 cover was only important in scenarios without regeneration (Fig.3). In both regeneration
 4317 scenarios, Bio 04 (Temperature Seasonality) and pasture cover had more influence on
 4318 environmental suitability.



4319

4320 **Fig. 03** Results in percentage for the Jackknife test for variable of environmental variables
 4321 in the development of Maxent model.

4322



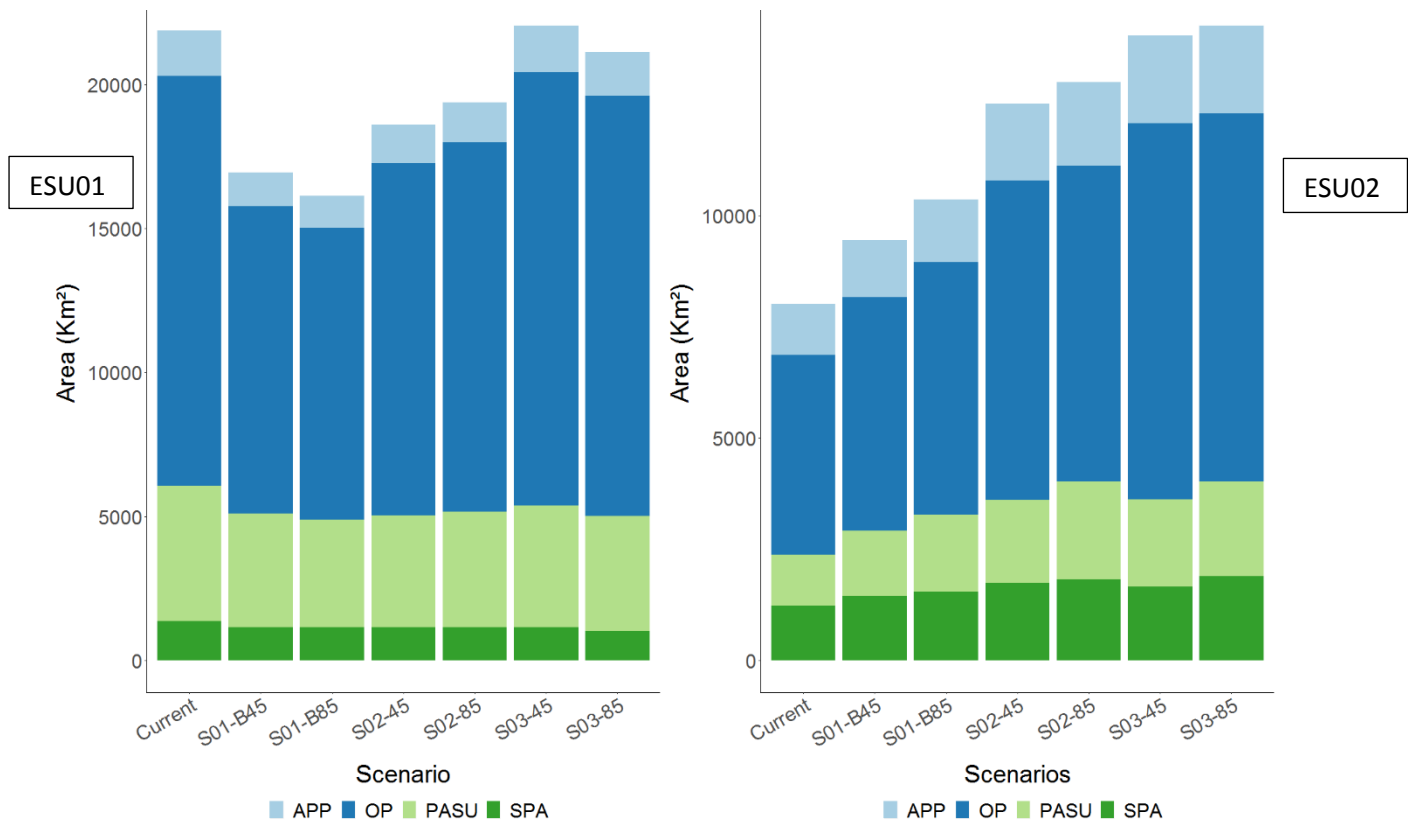
4323

4324 **Fig. 04** Response curves of the most important variable and the environmental suitability.

4325 Currently, most part of these suitable areas in both ESUs are excluded from any legal
 4326 protection (ESU01 – 65.03% and ESU02 – 56%; Fig.5). Among the protected regions, areas

4327 of Sustainable use protection encompass more suitable areas in ESU01 (21.4%), followed by
 4328 APP (7.27%) and Strictly protected areas (6.30%). For ESU02, suitable areas are present
 4329 almost in same proportion in the three conservation categories (15.32%, 14.44%, and 14.25,
 4330 respectively - Fig.5). The same trend for both ESUs remains throughout the future scenarios,
 4331 with the majority of the suitable areas remaining unprotected (Fig.5).

4332



4333

4334 **Fig. 5** Distribution of Suitable Areas in different conservation categories, according to
 4335 conservation Categories, where: APP - Permanent Protection Areas; PASU – Protected
 4336 Areas of Sustainable Use; SPA – Strictly Protected Areas/ OP – Outside Protected Areas.

4337

4338 **4. Discussion**

4339 *4.1 Overview of environmental Suitability models*

4340 The strong influence of climate change on the species distribution and the size of
4341 suitable areas is undeniable (GOUVEIA *et al.*, 2016b; STRUEBIG *et al.*, 2015). However, as
4342 we demonstrate here, and in accordance with other works (PREVEDELLO *et al.*, 2019;
4343 SILVÉRIO *et al.*, 2015), forest regeneration can mitigate the negative influence of climate
4344 change. We also highlight the importance of considering the genetic structure of species in
4345 ecological analyses. Genetically distinct populations of the same species present some
4346 particularities and distinct relations within the inhabited environment. Additionally, the
4347 inclusion of other variables besides climate data (e.g. landscape variables) allowed us to
4348 draw different conclusions from what was previously published (Moreira *et al.*, 2014).

4349 South Bahia and Center-South of Espírito Santo represents two important strongholds
4350 for *B.torquatus* (Fig 02). Both locations present well-preserved forest cover and include
4351 important protected areas, such as Una Biological Reserve and UNA Wildlife Refuge (BA)
4352 and Augusto Ruschi Biological Reserve (ES). Our models correctly predict the current low
4353 environmental suitability already indicated by gaps in the distribution of *B. torquatus*, in
4354 North Espírito Santo and North Rio de Janeiro (Fig.2). These areas present a low forest
4355 cover, which reduces the environmental suitability for the species. Yet, two large blocks of
4356 forest in the distribution gap between Bahia and Espírito Santo have been classified as
4357 suitable, which leads us to believe the existence of other undetected influences. Other studies
4358 highlight past epirogenic and climatic events which resulted in forest shrinkage,
4359 fragmentation and vegetation change (SCHETINO; COIMBRA; SANTOS, 2017;
4360 VASCONCELOS *et al.*, 1992). This gap region is distinct from the rest of the ombrophilous
4361 Atlantic Forest, with high seasonality and the presence of a semi-deciduous forest (JESUS;

4362 ROLIM, 2005). Similarly, previous studies carried out in North of Rio de Janeiro state
4363 highlights the low tree diversity of the semi-deciduous forests existing there (OLIVEIRA-
4364 FILHO; FONTES, 2000), linked to a strongly seasonal climate (HIRSCH; CHIARELLO,
4365 2012; OLIVEIRA-FILHO *et al.*, 2005), which likely acts as a limiting factor for the presence
4366 of strictly folivorous species.

4367 The northernmost part of *B. torquatus*' range, the regions of Pernambuco (PE),
4368 Paraiba (PB) and Rio Grande do Norte (RN - Fig. 02), present low environmental suitability.
4369 Previous research indicates similar result (MOREIRA *et al.*, 2014), evidencing the current
4370 absence of the species in the region. In contrast, Sergipe (SE), which represent the known
4371 northern limit of *B. torquatus*' range, still presents important though isolated areas suitable
4372 for the species, especially near the border of this state with Bahia (Fig. 2, Fig. S1). These
4373 populations deserve special attention; since the confirmation of this species in the region in
4374 2009 (CHAGAS *et al.*, 2009), only one unpublished study has been conducted on this
4375 population (SANTOS, P.M.; BOCCHIGLIERI; *et al.*, 2019). Thus, several ecological
4376 aspects remain unknown. Furthermore, the genetics of this population have not yet been
4377 studied (SCHETINO; COIMBRA; SANTOS, 2017), and it has not been included in any
4378 national conservation plan.

4379 *4.2 Present and future scenarios*

4380 The ESU01 concentrates large and continuous suitable areas for *B. torquatus*, with a
4381 stability trend only in the most positive scenarios (Table 02). Additionally, most current and
4382 future suitable areas in ESU01 represent one of the most climatically stable regions of the
4383 Atlantic Forest (CARNAVAL *et al.*, 2009; SOBRAL-SOUZA *et al.*, 2018). Overall, the
4384 Atlantic forest presents an increase in precipitation seasonality from east to west, and also
4385 from south to north (OLIVEIRA-FILHO; FONTES, 2000), and this trend will become

4386 stronger with climate changes (IPCC, 2018). The increase of precipitation seasonality and
4387 the decrease in precipitation of the wettest month, combined with a low forest cover, might
4388 explain why the environmental suitability declined mainly further inland and in northern
4389 areas - more vulnerable to climate changes (CARNAVAL *et al.*, 2009). Fluctuations in the
4390 precipitation regimes, may affect the production of young leaves, a preferred food item that
4391 becomes more abundant during the wet season (CHIARELLO, 1998b).

4392 The BAU scenario – which represents the climate changes effects without the forest
4393 regeneration - presents a sharp decrease in suitable areas. This trend decreased in the less
4394 optimistic scenario (Scenario 02 - Fig 02; Table 02), ratifying the importance of forest
4395 regeneration in mitigating the effects of climate change. Even though, the decrease in
4396 suitable areas in practically all scenarios (Scenario 02 – Fig 02, Table 02) lights a warning
4397 for the species conservation, since the region was one of the most deforested between 2017
4398 and 2018, counteracting the national trend of decreasing and stabilizing deforestation in the
4399 Atlantic Forest (FUNDAÇÃO SOS MATA ATLÂNTICA, 2018).

4400 In the ESU02 region, the environmental suitability depends on other factors, besides
4401 the percentage of forest cover, such as the presence of pasture and climate conditions. A
4402 slight improvement in climatic conditions and a large increase in forest cover may result in a
4403 continuous increase on suitable areas, even in the BAU scenario (Fig 2, Table 02). The
4404 southeast Atlantic Forest is considered climatically unstable (CARNAVAL *et al.*, 2009),
4405 presenting a strong seasonality in precipitation and in temperature throughout the year
4406 (OLIVERA - FILHO; FONTES, 2000). With global warming, the temperature seasonality
4407 and the precipitation of the driest month tends to decrease (Table S3). More constant
4408 temperatures might benefits the species, since their body temperature varies with the
4409 environment and activity decreases during cold periods (CHIARELLO, 1998a; GINÉ *et al.*,
4410 2015). Differently, the decrease of precipitation of the driest month can be harmful to the

4411 species, since it may exacerbate water deficit during the driest months, affecting leaf
4412 production in the dry season (LIEBERMAN; LIEBERMAN, 1984; REICH, 1995). The
4413 maned sloths from ESU02 are able to find and consume young leaves throughout the year
4414 (CHIARELLO, 1998b), despite the seasonal variation that is typical of this region today, but
4415 this flexibility might be compromised if precipitation seasonally increases. As we highlighted
4416 above, the two distribution gaps of this species have stronger seasonality in precipitation than
4417 non-gap areas.

4418 Regarding the forest remnants, the increase in suitable areas is concentrated in
4419 specific locations, around the largest forest areas (Fig. 02). The region englobes a few large
4420 forest patches, as well as regions with medium-sized fragments, that are functionally
4421 connected (SANTOS; JÚNIOR; EUGENIO, 2012; SANTOS *et al.*, 2016), and also a high
4422 incidence of pastures. This combination creates ideal conditions for increasing forest cover,
4423 since larger fragments encompass a greater diversity of species – including important
4424 animals seed dispersal (WUNDERLE, 1997) - and therefore, their matrices are more
4425 exposed to seed rain (CHAZDON, 2012; CROUZEILLES *et al.*, 2016; TAMBOSI *et al.*,
4426 2014). Additionally, the region experiment lower rates of habitat loss, presenting the lowest
4427 deforestation rate in the last 30 years (FUNDAÇÃO SOS MATA ATLÂNTICA, 2018).

4428 *4.3 Conservation Status of Suitable areas*

4429 In both ESUs, the percentage of suitable areas legally protected in the three categories
4430 we analyzed is consistently low. Part of this unprotected area is somewhat protected by the
4431 Atlantic Forest Law (Law 11. 428/2006), which prohibits the suppression of primary and late
4432 secondary forest areas, except in rare cases, in which environmental compensation is given.
4433 However, it allows the suppression of early secondary areas and of the initial stage,
4434 threatening, for example, the incidence of the natural regeneration of an area

4435 (VARJABEDIAN, 2010). Thus, it's quite necessary to guarantee the conservation of these
4436 areas, in order to ensure a substantial increase in forest cover, and consequently the
4437 environmental suitability for several forest species endemic to the Atlantic Forest.

4438 The other areas lying outside of the aforementioned conservation categories may
4439 belong to areas of Legal Reserves. According to LPVN, Legal Reserves is an instrument for
4440 preserving areas of natural vegetation in private areas, and in the Atlantic Forest this area
4441 occupies 20% of private property. The Legal Reserves system, however, has been constantly
4442 threatened with the argument that it is necessary to increase agricultural production areas
4443 (METZGER *et al.*, 2019), and its presence in the future is unknown. Legal Reserves have
4444 been neglected since the approval of the LPVN in 2012, which ruled that APP areas also
4445 must enter into the calculation of the RL areas. This was, in addition to forgiving the debt of
4446 small landowners who deforested their RL areas before 2008, a decision that cost the
4447 regeneration of thousands of hectares (BRANCALION *et al.*, 2016; SOARES-FILHO *et al.*,
4448 2014). Legal Reserves have fundamental ecological roles, including improving regional and
4449 global climate regulation (METZGER *et al.*, 2019). They compose the landscape and along
4450 with APPs and can increase the area of regional forest, facilitating the flow of organisms
4451 between protected areas and maintaining the genetic viability of populations (METZGER *et*
4452 *al.*, 2019; PADOVEZI *et al.*, 2018).

4453 **5. Conclusions**

4454 Our models evaluate the environmental suitability for *B. torquatus*, which does not
4455 necessarily indicate that the species is currently present in the area. This approach is highly
4456 useful for conservation and management purposes (MUKHERJEE *et al.*, 2020; PORTUGAL
4457 *et al.*, 2019; PRÉAU *et al.*, 2020), and in particular, helps to identify and prioritize areas for
4458 reintroduction (DANKS; KLEIN, 2002; MARTÍNEZ-MEYER *et al.*, 2006; PANT *et al.*,

4459 2020; PETERSON, 2006) , and to identify areas to create potential wildlife corridors to link
4460 suitable yet noncontiguous areas (LIU *et al.*, 2018; MORATO *et al.*, 2014).

4461 Natural regeneration is a reality in Atlantic Forest, and as we show here, it can
4462 guarantee a decrease in local effects of climate change effects on environmental suitability
4463 for a forest dependent species. The majority of pasture lands are economically unproductive
4464 (STRASSBURG *et al.*, 2014), and therefore present an opportunity for forest recovery
4465 (LATAWIEC *et al.*, 2015). Our models will contribute to several conservation actions for *B.*
4466 *torquatus* and are especially useful for our different analyses for the two ESUs. Even though
4467 its conservation status is threatened, mainly due to habitat loss, there remain adequate areas
4468 to sustain viable populations.

4469 In this sense, it is extremely important to ensure the maintenance of areas that are
4470 already protected, mainly those in South Bahia, Espírito Santo and Rio de Janeiro, by
4471 decreasing external pressures. Promoting compliance with environmental law in the entire
4472 analyzed area will also be necessary to preserve APPs and Legal Reserves. Lastly, the
4473 natural regeneration of abandoned areas should be encouraged, especially in the more
4474 northerly areas (Sergipe and North Bahia) that currently contain suitable but isolated areas.
4475 In this case, it is also necessary to collaborate for their protection, to minimize the
4476 suppression of regrowth, and to monitor the forests' successional stages, interfering when
4477 necessary to increase its quality.

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4725 **Appendix A. Supplementary data**4726 **Table S1** Characteristics of variables used to delineate environmental suitability for *Bradypus torquatus*.

Use	Variable	Description	Original Resolution	Year	Source	ESU 01	ESU 02
Environmental Suitability Models	Forest Cover Percentage	Forest Cover percentage in 1x1km pixel	5 x 5 m	2018	Brazilian Foundation for Sustainable Development (In Portuguese Fundação Brasileira para o Desenvolvimento Sustentável - FBDS) Source: https://www.fbds.org.br	X	X
	Pasture cover Percentage	Pasture Cover percentage in 1x1km pixel	30x30m	2015	Image Processing and Geoprocessing Lab (In portuguese: Laboratório Processamento de Imagens e Geoprocessamento - LAPIG). Source: www.pastagem.org	X	X
	Elevation	Global elevation data	30 arc-second (1x1km)	2004	NASA Shuttle Radar Topography Mission Source: https://www2.jpl.nasa.gov/srtm/	X	X
	Bioclimatic Variables	Bio 04 = Temperature Seasonality (standard deviation *100)	30 arc-second (1x1km)	2005	Worldclim global climate variables V1. Source: https://www.worldclim.org/		X

		Bio 13 = Precipitation of Wettest Month				X	
		Bio 14 = Precipitation of Driest Month					X
		Bio 15 = Precipitation Seasonality (Coefficient of Variation)				X	
	hydic APPs	Areas of water APPs, which include rivers, water springs, lakes, lagoons and artificial reservoirs	5 x 5 m	2018	Brazilian Foundation for Sustainable Development (In Portuguese Fundação Brasileira para o Desenvolvimento Sustentável - FBDS) Source: https://www.fbds.org.br	X	X
Conservation Analyses	Strictly Protected areas (SPA) and Protected Areas of Sustainable use (PASU)	Shapefile with the distribution of protected areas of integral conservation and sustainable use			Ministry of the Environment (In Portuguese: Ministério do Meio Ambiente - MMA). Source: http://mapas.mma.gov.br/i3geo/datadownload.htm	X	X

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4729 **Table S2** AUC scores, test omission, and variable contribution for each scenario. All the scenarios, $p < 0.05$.

Scenarios	Model Evaluation				Variable Contribution				
	Test AUC	AUC SD	10 percentil	Omission test	Altitude	Bio13	Bio15	Forest cover	Pasture cover
ESU01									
Current Scenario	0.929	0.017	0.367	0.094	5.701	1.563	2.032	88.177	2.527
BAU_RCP45_CCSM4	0.913	0.020	0.408	0.206	4.387	1.241	4.885	88.103	1.385
BAU_RCP45_HadGCM2-ES	0.927	0.016	0.447	0.212	7.772	5.505	1.122	83.621	1.981
BAU_RCP45_MIROC-ESM	0.913	0.020	0.364	0.171	9.338	1.985	1.204	86.375	1.097
BAU_RCP85_CCSM4	0.910	0.021	0.467	0.282	7.186	1.375	5.133	83.575	2.730
BAU_RCP85_HadGCM2-ES	0.934	0.017	0.398	0.088	6.456	5.340	2.055	84.459	1.690
BAU_RCP85_MIROC-ESM	0.928	0.016	0.424	0.176	7.434	2.960	1.440	86.301	1.865
Scenario02_RCP45_CCSM4	0.898	0.024	0.411	0.212	7.036	2.540	6.047	81.776	2.601
Scenario02_RCP45_HadGCM2-ES	0.920	0.017	0.411	0.177	7.948	9.630	2.370	78.928	1.125
Scenario02_RCP45_MIROC-ESM	0.901	0.020	0.414	0.200	12.047	1.927	2.852	80.534	2.641
Scenario02_RCP85_CCSM4	0.918	0.021	0.419	0.135	7.271	1.090	5.797	82.929	2.914
Scenario02_RCP85_HadGCM2-ES	0.924	0.022	0.357	0.147	8.880	10.103	1.037	76.972	3.008
Scenario02_RCP85_MIROC-ESM	0.925	0.018	0.422	0.153	9.236	4.348	2.928	81.323	2.165
Scenario03_RCP45_CCSM4	0.911	0.024	0.409	0.124	12.919	3.580	15.051	65.247	3.203
Scenario03_RCP45_HadGCM2-ES	0.915	0.021	0.429	0.147	7.874	13.495	8.270	68.739	1.623
Scenario03_RCP45_MIROC-ESM	0.881	0.031	0.413	0.247	14.790	3.333	7.038	70.477	4.362
Scenario03_RCP85_CCSM4	0.877	0.029	0.387	0.218	12.305	2.018	15.929	65.945	3.803
Scenario03_RCP85_HadGCM2-ES	0.931	0.019	0.416	0.147	8.441	16.522	3.520	69.660	1.856
Scenario03_RCP85_MIROC-ESM	0.903	0.024	0.430	0.200	11.738	6.034	11.220	68.419	2.589
ESU02									
Current Scenario	0.961	0.009	0.407	0.171	2.168	26.888	3.959	58.351	8.634
BAU_RCP45_CCSM4	0.958	0.010	0.360	0.143	2.400	0.980	27.780	57.820	11.020

BAU_RCP45_HadGCM2-ES	0.958	0.011	0.444	0.171	1.988	1.712	28.348	60.901	7.052
BAU_RCP45_MIROC-ESM	0.957	0.009	0.327	0.124	5.882	2.464	24.092	59.607	7.955
BAU_RCP85_CCSM4	0.958	0.009	0.321	0.119	3.637	1.208	27.208	63.208	4.739
BAU_RCP85_HadGCM2-ES	0.957	0.010	0.380	0.152	3.263	2.063	28.305	59.181	7.187
BAU_RCP85_MIROC-ESM	0.947	0.011	0.380	0.210	3.512	0.556	26.853	65.104	3.975
Scenario02_RCP45_CCSM4	0.939	0.015	0.322	0.195	4.761	3.817	35.359	22.331	33.733
Scenario02_RCP45_HadGCM2-ES	0.957	0.013	0.443	0.176	4.972	7.331	34.382	22.525	30.791
Scenario02_RCP45_MIROC-ESM	0.932	0.020	0.326	0.186	9.562	9.339	27.906	26.211	26.982
Scenario02_RCP85_CCSM4	0.957	0.011	0.309	0.081	4.395	4.811	35.060	24.801	30.933
Scenario02_RCP85_HadGCM2-ES	0.947	0.018	0.351	0.148	2.585	9.614	33.919	23.968	29.914
Scenario02_RCP85_MIROC-ESM	0.931	0.020	0.311	0.176	6.037	4.382	30.008	24.977	34.596
Scenario03_RCP45_CCSM4	0.945	0.017	0.350	0.129	9.576	11.100	37.400	19.186	22.738
Scenario03_RCP45_HadGCM2-ES	0.938	0.020	0.317	0.157	4.506	12.906	34.907	20.090	27.591
Scenario03_RCP45_MIROC-ESM	0.940	0.017	0.354	0.138	10.354	14.881	29.340	21.175	24.251
Scenario03_RCP85_CCSM4	0.932	0.020	0.369	0.162	6.149	11.919	35.391	17.890	28.651
Scenario03_RCP85_HadGCM2-ES	0.941	0.018	0.316	0.105	3.943	13.433	32.701	18.893	31.031
Scenario03_RCP85_MIROC-ESM	0.925	0.022	0.379	0.162	9.322	9.434	34.090	19.735	27.419

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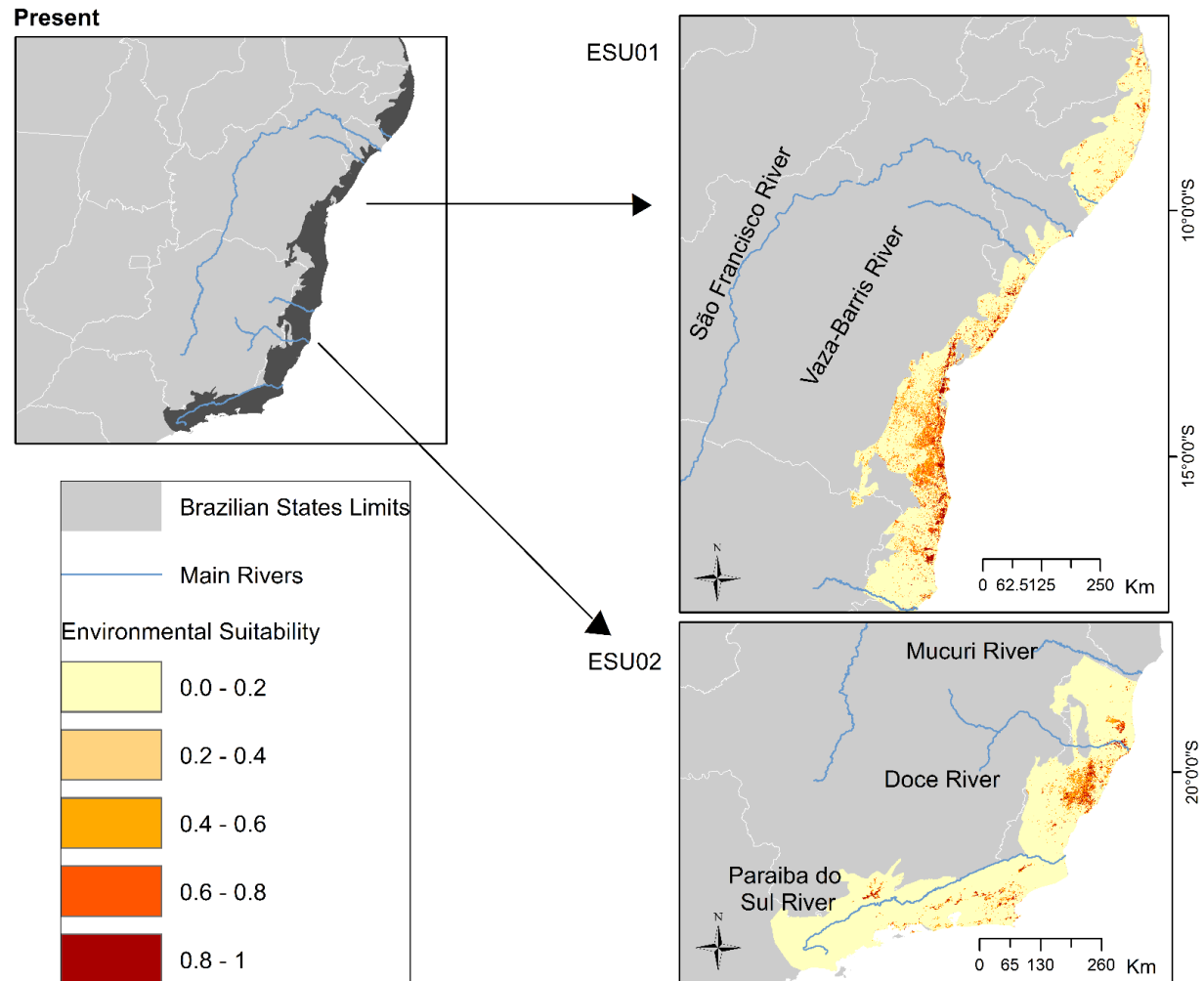
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4732 **Table S3** Maximum, minimum and mean values of the variables for both populations and in all scenarios.

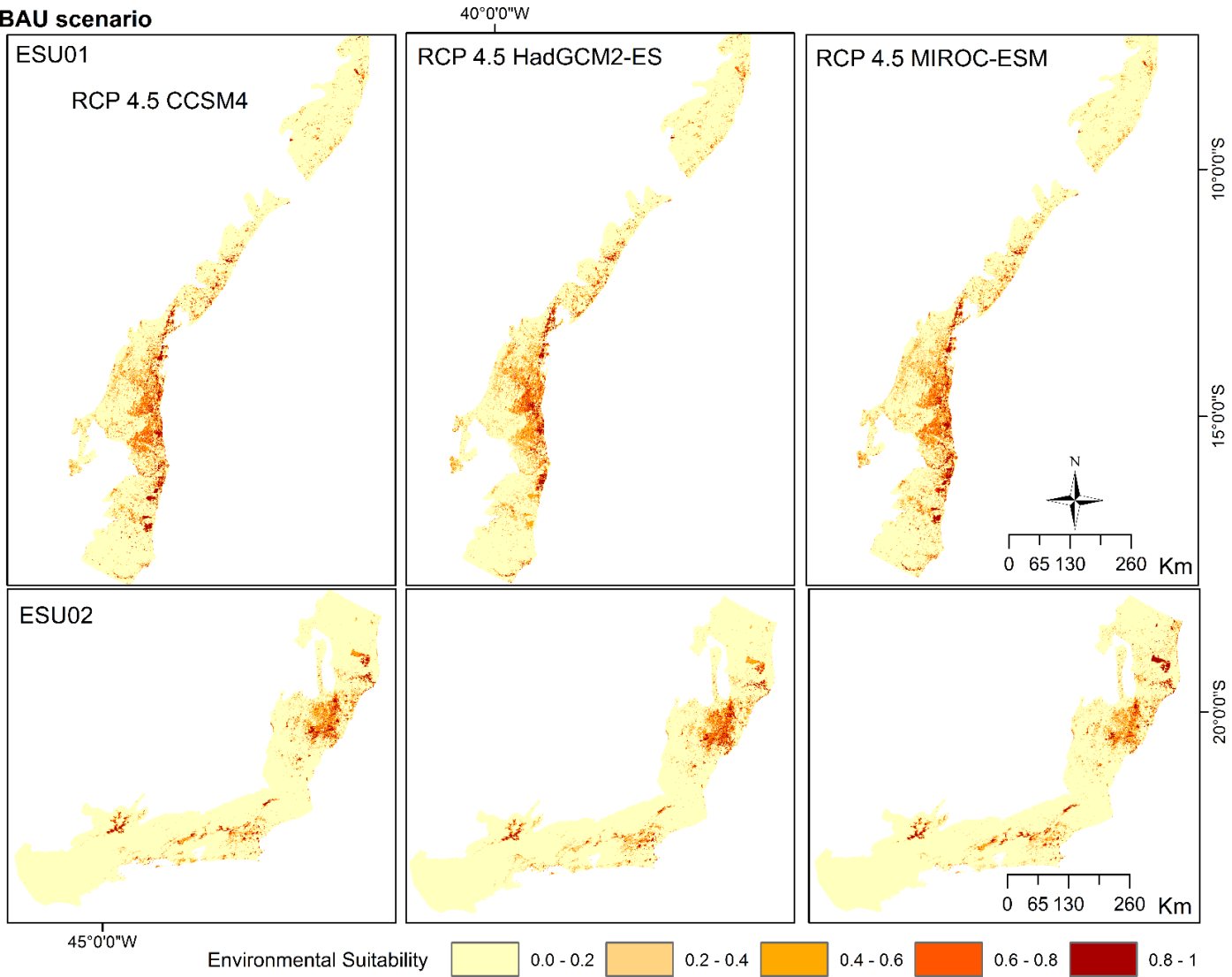
ESU01						
Scenario	Bio13 - Precipitation of the Wettest month (mm)			Bio15 - Precipitation Seasonality		
	Max	Min	Mean	Max	Min	Mean
Current Scenario	336	4	186.412	76	9	40.128
RCP45 CCSM4	275	84	162.629	78	14	42.200
RCP45 HadGCM2-ES	280	95	166.433	78	18	45.138
RCP45 MIROC-ESM	265	74	157.197	80	14	43.088
RCP85 CCSM4	249	76	156.710	78	11	42.600
RCP85 HadGCM2-ES	306	105	184.003	79	23	50.040
RCP85 MIROC-ESM	234	75	156.672	85	18	44.721
ESU02						
Scenario	Bio04 - Temperature Seasonality			Bio14 - Precipitation of the driest month (mm)		
	Max	Min	Mean	Max	Min	Mean
Current Scenario	2525	1444	1960.396	176	13	32.613
RCP45 CCSM4	2423	1427	1847.344	178	12	30.334
RCP45 HadGCM2-ES	2534	1515	2009.255	195	13	31.569
RCP45 MIROC-ESM	2354	1312	1786.010	159	10	27.033
RCP85 CCSM4	2380	1393	1908.077	159	12	28.866
RCP85 HadGCM2-ES	2602	1575	2088.178	197	12	29.145
RCP85 MIROC-ESM	2324	1343	1822.031	120	7	22.561

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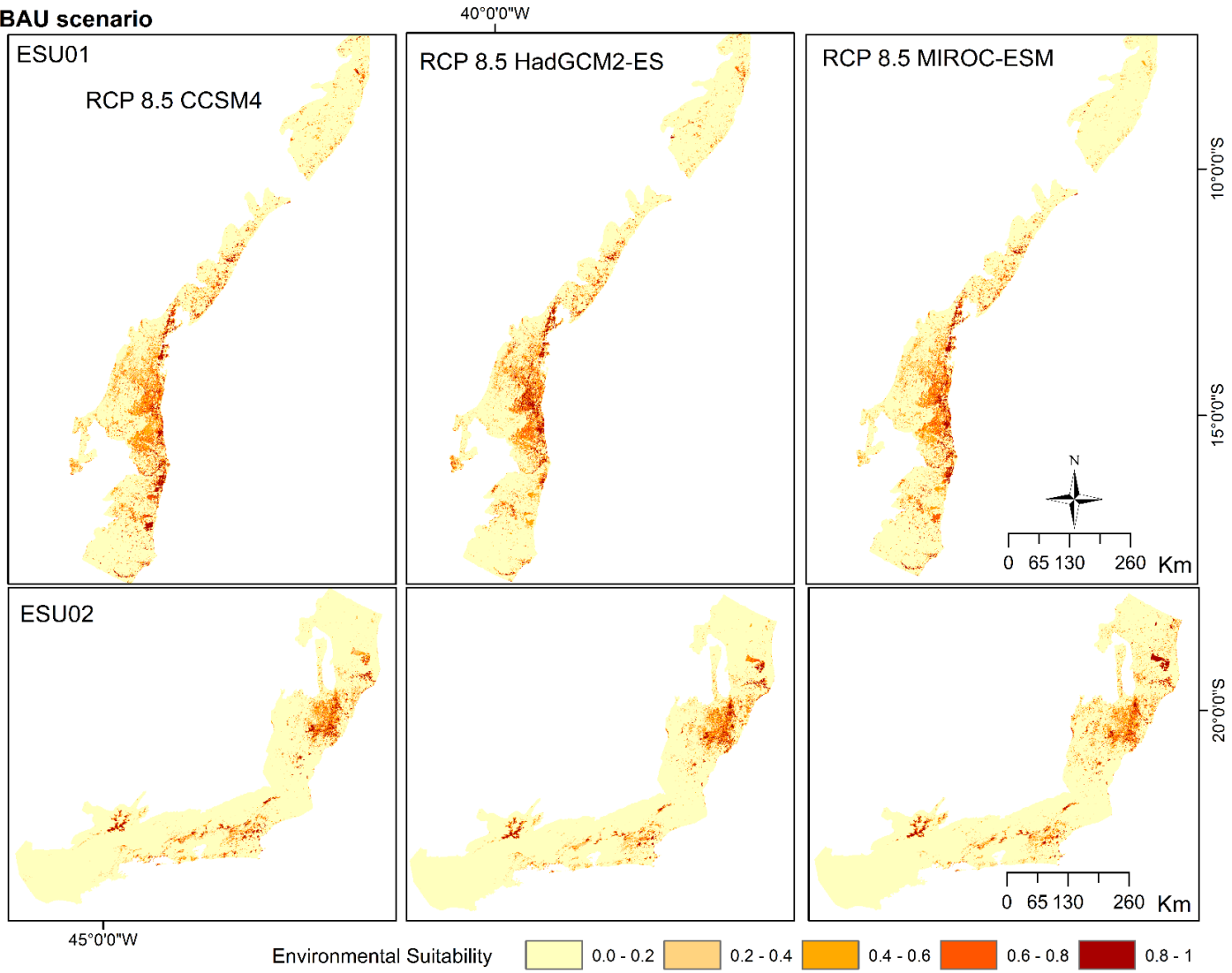
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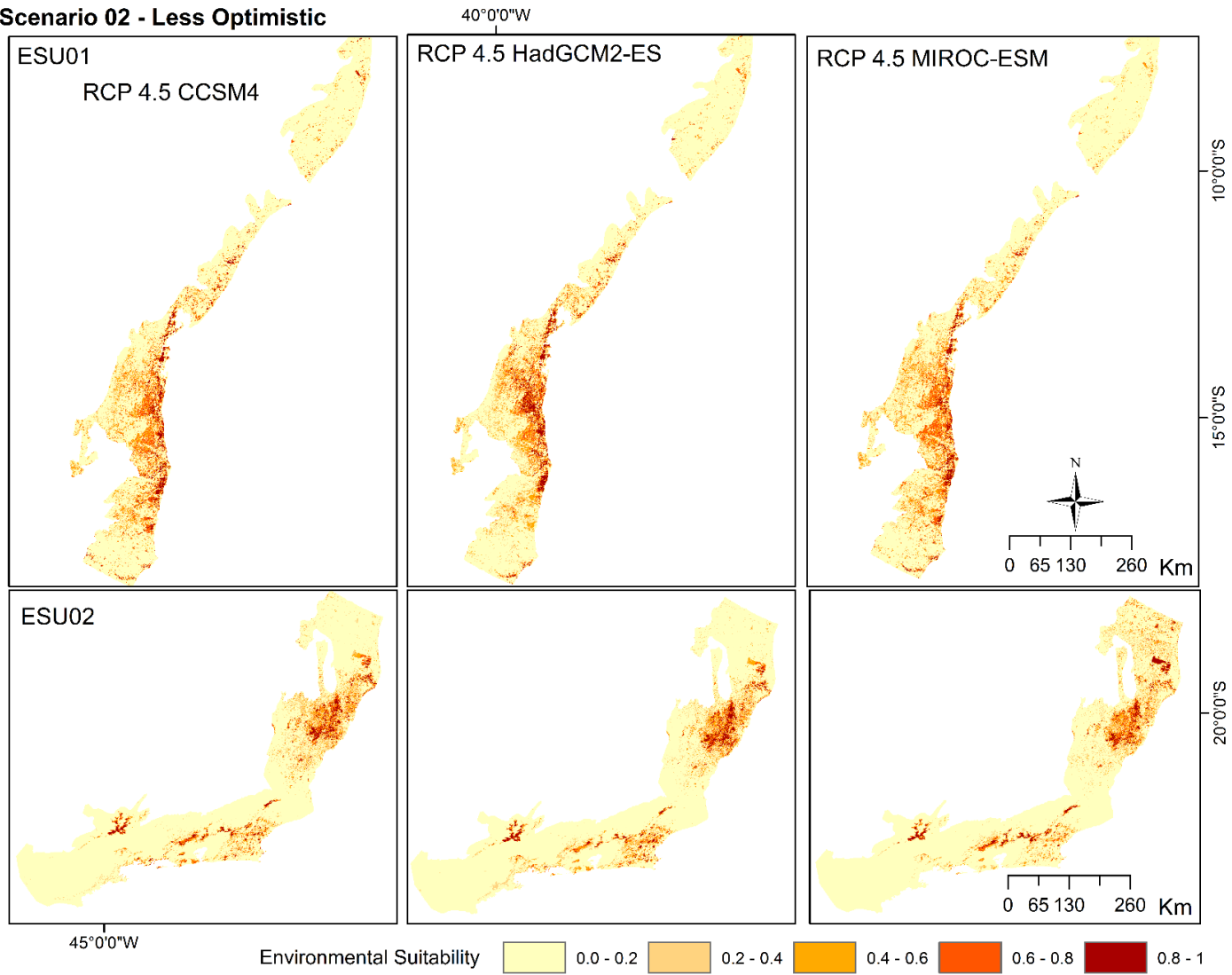
BAU scenario



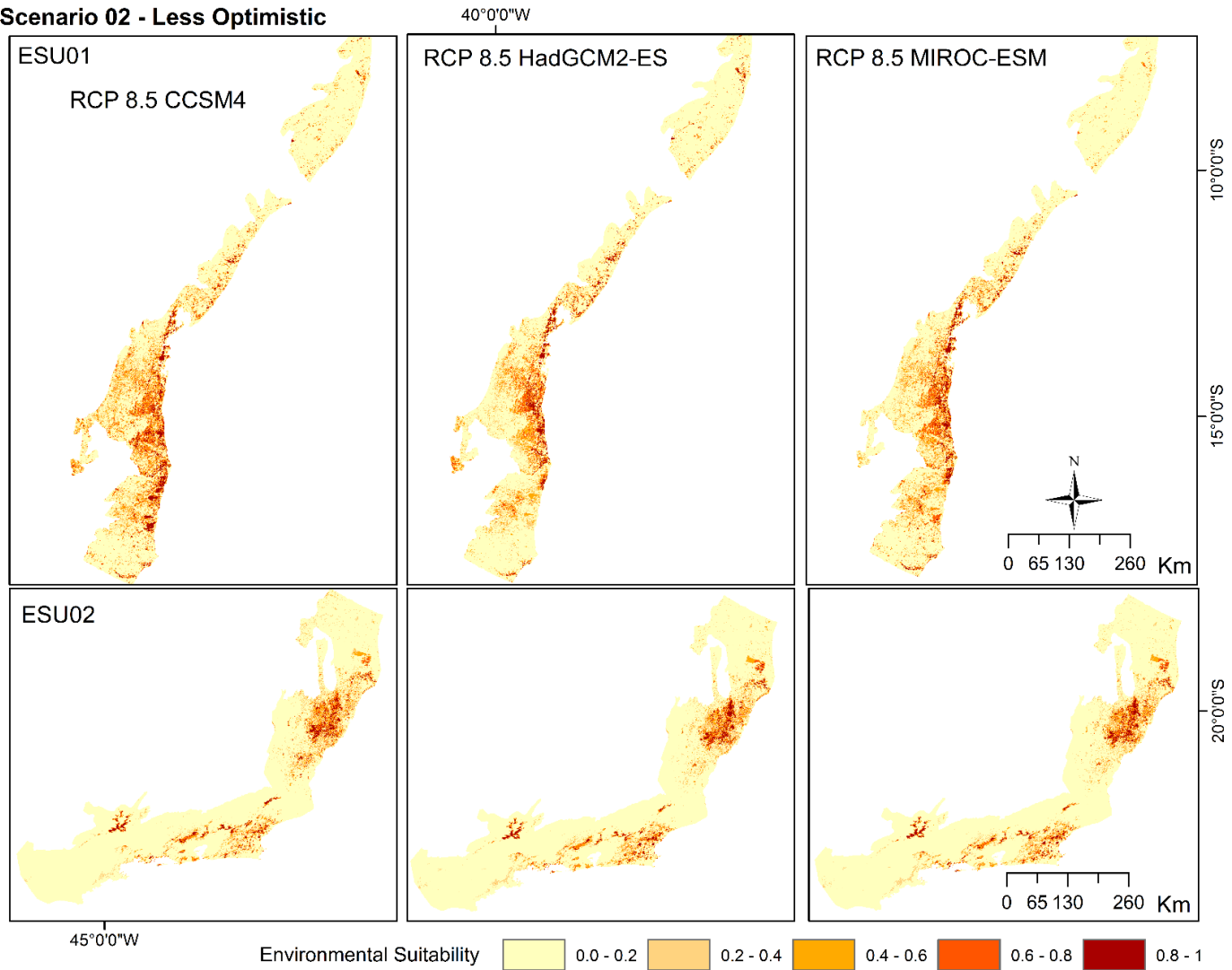
BAU scenario



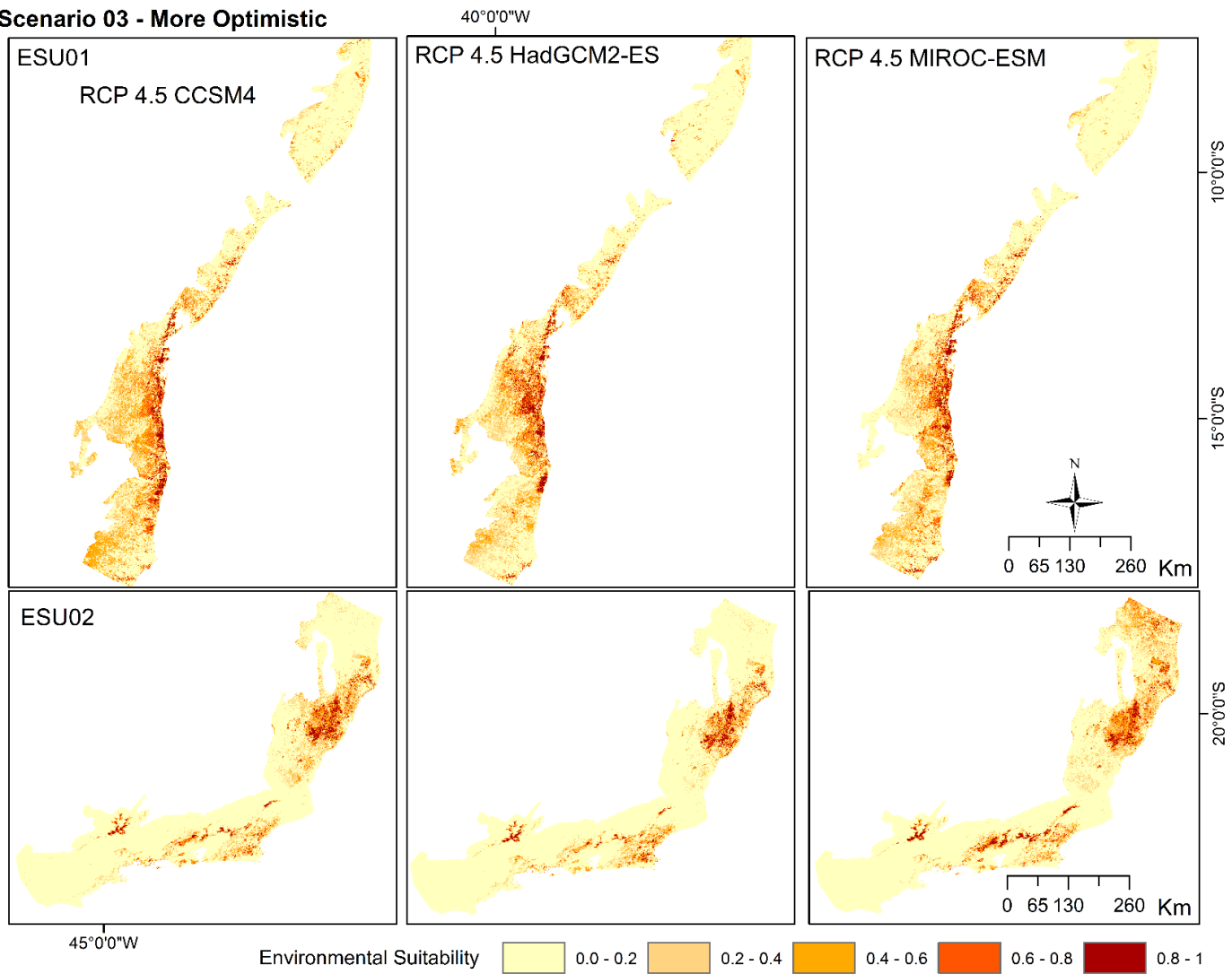
Scenario 02 - Less Optimistic

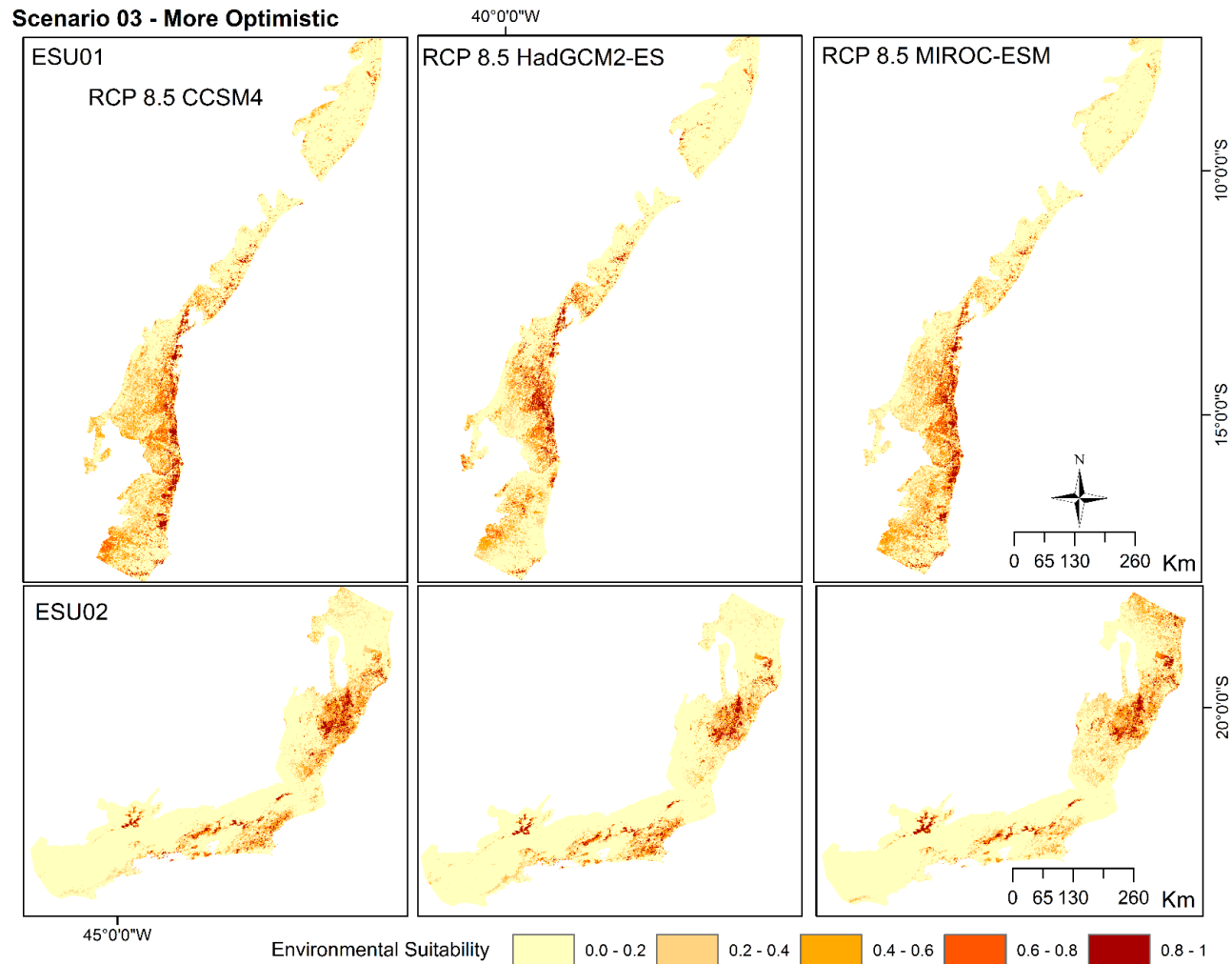


Scenario 02 - Less Optimistic



Scenario 03 - More Optimistic





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4743 **Fig S1** Current and future environmental suitability for both populations of *B.torquatus*. The environmental suitability is separated in different classes, and all
 4744 results from the different circulations models are represented here.

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Conclusões Gerais

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Nesta tese, buscamos identificar a influências de variáveis ambientais na ocupação da preguiça de coleira e na disposição das áreas mais adequadas para a espécie por meio de análises multi-espaciais e temporais. De modo geral, a ocupação da espécie em uma dada região irá depender de fatores múltiplos, como o ambiente local e a composição espacial dos elementos da paisagem. O futuro da espécie irá depender principalmente da dinâmica florestal e da climática. Além disso, a tese resultou em uma grande contribuição com a geração e o uso de dados abertos.

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Embora a preguiça ocupe ambientes alterados – porém com boa cobertura floresta (> 35%), a sua ausência em áreas com menos de 20% de cobertura florestal acende um alerta em relação a sua conservação. Além disso, a presença de áreas abertas – mesmo em menor proporção – inviabiliza a presença da espécie em determinadas áreas. O alto requerimento energético para atravessar tais áreas e as ameaças intrínsecas nesses ambientes, como a predação oportunística, são as principais barreiras. Por outro lado, como já salientado em estudo anterior (SANTOS *et al.*, 2016), as preguiças preferem determinados tipos de ambientes florestais, como locais que possuam um dossel alto e conectado e que abranja árvores importantes para a sua dieta. As preguiças passam boa parte de sua vida nos dosséis e utiliza a conexão da copa entre as árvores para o seu deslocamento (MONTGOMERY; SUNQUIST, 1975). Assim, árvores altas são essenciais, por cumprirem função fundamental no comportamento termo regulatório desempenhado pelas preguiças, de subir no topo das árvores para se aquecer (MONTGOMERY; SUNQUIST, 1975; NAGY; MONTGOMERY, 1980). Assim, é possível concluir que existe uma série de fatores multi-escala responsáveis pela ocupação da espécie em determinado fragmento de mata.

4769 A necessidade de se obter mais dados sobre a preguiça-de-coleira nos levou a um
4770 importante passo nessa tese – o NEOTROPICAL XENARTHROS. A compilação e
4771 padronização das informações de ocorrência de espécies de xernatros, com mais de 30 mil
4772 registros de tatus, tamanduás e preguiças, produziu o maior banco de dados existentes para
4773 esse grupo ainda tão desconhecido. A análise desse banco de dados deixa clara a grande
4774 densidade de registros que se encontravam indisponíveis para o acesso do público em geral e
4775 o quanto ainda precisamos avançar nas pesquisas sobre o grupo. Todos os dados do NeoXen
4776 estão disponíveis no Github e a base de dados será alimentada à medida em que novos
4777 registros forem surgindo.

4778 Com os resultados do capítulo 01 e com a compilação de dados do capítulo 02 foi
4779 possível gerar modelos de distribuição de espécie voltados para a avaliação da adequabilidade
4780 do habitat para as preguiças. A presença de florestas, mais uma vez, se mostrou como a
4781 principal preditora da espécie, uma vez que a disponibilidade de boas áreas está relacionada
4782 com a presença de uma boa cobertura florestal. Adicionalmente, a alta cobertura de florestas
4783 também se mostra bastante importante para mitigar os efeitos das mudanças climáticas sobre
4784 a espécie – contribuindo na conservação de uma boa proporção de áreas adequadas.

4785 O estabelecimento de novas Unidades de conservação e a manutenção das já
4786 existentes é um importante passo para garantir a proteção das áreas naturais. Além disso, as
4787 Reservas Legais podem ser cruciais para resguardar áreas naturais, uma vez que, conforme
4788 mostrado nos capítulos 02 e 03, grande parte das áreas ocupadas e adequadas para a
4789 *B.torquatus* estão localizadas em propriedades particulares. Porém, para potencializar as suas
4790 funções, precisaria destituir as Áreas de Preservação Permanente do cálculo das RLs, uma
4791 vez que as duas categorias atuam em complementariedade na conservação e preservação de
4792 florestas. Da mesma forma, é necessário garantir o amparo legal das áreas potenciais para

4793 regeneração natural – ou as que já estão em processo de regeneração, a fim de evitar seu corte
4794 e a supressão

4795 Por fim, uma característica dessa tese é a sua ação conservacionista aplicada. Todo o
4796 escopo dessa tese está relacionado com diversas metas do Plano de Ação Nacional para a
4797 Conservação de Primatas e Preguiças (PPMA) - identificar áreas importantes, ampliar o
4798 conhecimento de ocorrência e ocupação e seus preditores ambientais - e os resultados gerados
4799 aqui irão integrar a avaliação da espécie. Além disso, todos os dados compilados no
4800 NEOTROPICAL XENARTHROS já estão sendo utilizados pelos órgãos ambientais. O
4801 Centro Nacional de Conservação dos Primatas Brasileiros (CPB/ICMBio), responsável por
4802 avaliar os xenartros, tem utilizado os dados de ocorrência do *datapaper* para desenvolver as
4803 fichas de avaliação das espécies, por meio de seu portal SALVE. O Instituto Nacional da
4804 Mata Atlântica (INMA) também utilizou os dados referentes aos xenartros no estado do
4805 Espírito Santo para elaborar o a lista estadual de espécie ameaçadas (INMA, 2019). Assim,
4806 essa tese vem cumprindo seu papel na conservação da biodiversidade, ao integrar academia,
4807 ações de organizações conservacionistas e órgãos públicos, federal e estaduais.

4808

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