

# Revealing the unknown world of the endangered Lear's macaw using GPS-tracking data: identification of critical habitats for conservation

FERNANDA R. PASCHOTTO<sup>1</sup> [orcid.org/0000-0002-6651-8318](https://orcid.org/0000-0002-6651-8318), ERICA C. PACÍFICO<sup>2</sup> [orcid.org/0000-0003-3518-1887](https://orcid.org/0000-0003-3518-1887), THIAGO FILADELFO<sup>3</sup> [orcid.org/0000-0002-0523-9788](https://orcid.org/0000-0002-0523-9788), GABRIELA R. FAVORETTO<sup>4</sup> [orcid.org/0000-00026382-140X](https://orcid.org/0000-00026382-140X), JULIA E. F. OSHIMA<sup>5</sup> [orcid.org/0000-0003-1545-768X](https://orcid.org/0000-0003-1545-768X), PAULO CARDOSO<sup>6</sup> [orcid.org/0000-0001-5458-998X](https://orcid.org/0000-0001-5458-998X), FRANCISCO V. DÉNES<sup>7</sup> [orcid.org/0000-0002-9870-7682](https://orcid.org/0000-0002-9870-7682)

<sup>1</sup>[contact e-mail: fernandarpaschotto@gmail.com](mailto:fernandarpaschotto@gmail.com) | Movement Ecology Lab, Department of Ecology, Institute of Biosciences, University of São Paulo. Rua do Matão, travessa 14, nº 321, Cidade Universitária, 05508-090, São Paulo/SP, Brazil

<sup>2</sup>[ericapacifico81@gmail.com](mailto:ericapacifico81@gmail.com) | Department of Animal Biology, Institute of Biology, State University of Campinas. Rua Monteiro Lobato, nº 255, Cidade Universitária, 13083-862, Campinas/SP, Brazil

<sup>3</sup>[thiago\\_bioufba@yahoo.com.br](mailto:thiago_bioufba@yahoo.com.br) | Rua Ana C. B. Dias, Condomínio Jardim Atlântico, Casa E8, Miragem, 42709-500, Lauro de Freitas/BA, Brazil

<sup>4</sup>[gabifavoretto@yahoo.com.br](mailto:gabifavoretto@yahoo.com.br) | Av. Mario Ybarra de Almeida, nº 841, Centro, 14800-420, Araraquara/SP, Brazil

<sup>5</sup>[juliaoshima@yahoo.com.br](mailto:juliaoshima@yahoo.com.br) | Movement Ecology Lab, Department of Ecology, Institute of Biosciences, University of São Paulo. Rua do Matão, travessa 14, nº 321, Cidade Universitária, 05508-090, São Paulo/SP, Brazil

<sup>6</sup>[paulo.c@bioinsight.pt](mailto:paulo.c@bioinsight.pt) | Bioinsight, Lda. Rua Antero de Quental, nº 52 B, Urbanização Colinas do Cruzeiro, 2675-690, Odivelas, Portugal

<sup>7</sup>[francisco.denes@ib.usp.br](mailto:francisco.denes@ib.usp.br) | Movement Ecology Lab, Department of Ecology, Institute of Biosciences, University of São Paulo. Rua do Matão, travessa 14, nº 321, Cidade Universitária, 05508-090, São Paulo/SP, Brazil

**Word count: 6,998**

1 *Article*

2 **Revealing the unknown world of the endangered Lear's macaw**  
3 **using GPS-tracking data: identification of critical habitats for**  
4 **conservation**

5 FERNANDA R. PASCHOTTO\*<sup>1,2</sup> [orcid.org/0000-0002-6651-8318](https://orcid.org/0000-0002-6651-8318), ERICA C. PACÍFICO<sup>2,3</sup>  
6 [orcid.org/0000-0003-3518-1887](https://orcid.org/0000-0003-3518-1887), THIAGO FILADELFO<sup>2</sup> [orcid.org/0000-0002-0523-9788](https://orcid.org/0000-0002-0523-9788),  
7 GABRIELA R. FAVORETTO<sup>2</sup> [orcid.org/0000-00026382-140X](https://orcid.org/0000-00026382-140X), JULIA E. F. OSHIMA<sup>1</sup>  
8 [orcid.org/0000-0003-1545-768X](https://orcid.org/0000-0003-1545-768X), PAULO CARDOSO<sup>4</sup> [orcid.org/0000-0001-5458-998X](https://orcid.org/0000-0001-5458-998X),  
9 FRANCISCO V. DÉNES<sup>1,5</sup> [orcid.org/0000-0002-9870-7682](https://orcid.org/0000-0002-9870-7682)

10

11 \*Corresponding author, [fernandarpaschotto@gmail.com](mailto:fernandarpaschotto@gmail.com)

12 <sup>1</sup>Department of Ecology, University of São Paulo, São Paulo, Brazil

13 <sup>2</sup>Lear's Macaw Research and Conservation Group (GPCA), São Paulo, Brazil

14 <sup>3</sup>Department of Animal Biology, State University of Campinas, Campinas, Brazil

15 <sup>4</sup>BE Bioinsight | ECOA, Environmental Consultants, Portugal

16 <sup>5</sup>Department of Migration, Max Planck Institute of Animal Behavior, Konstanz, Germany

17

18 **Abstract** Understanding space use and home range is essential for the conservation  
19 planning of threatened species as it helps to assess the suitability, extent, and placement  
20 of conservation areas that are imperative for species survival and protection. The  
21 Endangered Lear's macaw (*Anodorhynchus leari*), a highly mobile frugivore, feeding  
22 specialist and endemic to Brazil's Caatinga dry forest, faces ongoing habitat degradation.  
23 In this study, we identified critical habitats by examining the spatial distribution of  
24 feeding, resting, and roosting sites and investigating home range size and its temporal  
25 variation. We GPS-tracked juvenile macaws and estimated fortnightly home ranges with  
26 autocorrelated kernel density estimators. We assessed if extrinsic factors, such as tagging

27 site, seasonality, rainfall and vegetation productivity (proxies for food availability)  
28 influenced home range size. Our findings reveal considerable variation in home ranges,  
29 with an average of 850.15 km<sup>2</sup> (1.24-8,549.48 km<sup>2</sup>). Home ranges expanded significantly  
30 during the dry season (mean 1,097.06 km<sup>2</sup>), representing a 2.14-fold increase from the  
31 wet season. We also found that site and season primarily drove home range size, while  
32 vegetation productivity and rainfall had limited influence. This suggests that macaw  
33 movements may respond to complex interactions between rainfall, landscape  
34 composition and configuration, and food availability rather than direct resource  
35 fluctuations. This is the first study to estimate home ranges for Lear's macaw, providing  
36 critical insights into its spatial ecology. Our findings underscore the importance of  
37 preserving key roosting and feeding areas and highlight the need for continuous  
38 monitoring to address threats posed by environmental changes and human activity.

39

40 **Keywords** Caatinga, conservation, GPS tracking, habitat use, home range, Lear's  
41 macaw, licuri palm, parrot

42 The supplementary material and code on GitHub for the analysis in R for this article are  
43 available at <https://doi.org/xxx>

44

## 45 **Introduction**

46 Understanding space use and home range is essential for wildlife conservation, providing  
47 insights into movement patterns, species' distribution, ecological needs, and spatial  
48 requirements (Adamek, 2011; Walton et al., 2017). Habitat loss is one of the main drivers  
49 of species extinction, especially for endemic and/or specialist species (Morato et al.,  
50 2018; Oliveira et al., 2021), and this knowledge can help identify suitable environments  
51 for threatened species and guide conservation actions, such as protecting feeding and  
52 breeding sites across their distribution ranges, many of which remain poorly understood,  
53 and providing quantitative information that may help delineate adequate protected areas,  
54 evaluating their distribution and extent as needed for the maintenance of these species  
55 (Schofield et al., 2010; Goldingay, 2015; Yeap et al., 2021).

56         The Lear's macaw (*Anodorhynchus leari*), an Endangered frugivore endemic to  
57 the Caatinga dry forest in northeastern Brazil, exemplifies the challenges of conserving  
58 species with highly specialized habitat requirements (BirdLife International, 2020). The  
59 macaw's survival is closely tied to the availability of licuri palm (*Syagrus coronata*)  
60 fruits, and it nests and roosts primarily on sandstone cliffs (Pacífico et al., 2014).  
61 However, recent observations indicate that some individuals also roost in trees, though  
62 the ecological significance of this behaviour is not fully understood (E.C.P. pers.  
63 comm., 2023).

64         Despite its restricted range, significant knowledge gaps persist regarding the  
65 species' movement patterns, seasonal variation in space use, and the drivers of home  
66 range size. Habitat degradation, compounded by desertification and anthropogenic  
67 activities, threatens key feeding and roosting sites (Santos Neto & Camandaroba, 2008;  
68 Barbosa & Tella, 2019). Tracking macaws' movements in their often-inaccessible natural  
69 habitat is challenging, limiting our understanding of their spatial ecology and our ability

70 to identify and protect critical habitat for the species (Le Souef et al., 2013; Yeap et al.,  
71 2017; Brightsmith et al., 2021).

72 To address these gaps, we investigated Lear's macaw space use by examining  
73 home range size, seasonal variation, and the spatial distribution of feeding, resting, and  
74 roosting sites. Specifically, we formulated the following research questions: (1) Where  
75 are the feeding, roosting and diurnal resting sites? (2) How is the habitat of these sites  
76 characterized? (3) How extensive are individual home ranges? (4) Do these ranges and  
77 movement patterns vary seasonally?

78 By combining Global Positioning System (GPS) tracking data with remote  
79 sensing and environmental variables, we aim to provide critical information to inform  
80 conservation efforts for this emblematic species of the Caatinga.

81

## 82 **Study area**

83 The study region encompasses the distribution of Lear's macaw in northern Bahia, Brazil,  
84 within the Caatinga Phytogeographical Domain (Fig. 1; Moro et al., 2016; BirdLife  
85 International, 2020). The eastern range includes the nine known roosting sites (sandstone  
86 cliffs and trees) in *Raso da Catarina* region, concentrating the main population of  
87 macaws, with the protected areas *Estação Biológica de Camudos* and *Estação Ecológica*  
88 *do Raso da Catarina*. The western range, known as *Boqueirão da Onça* and  
89 located 230 km west, includes the recently established protected areas *Área de Proteção*  
90 *Ambiental* and *Parque Nacional Boqueirão da Onça*. This range holds another currently  
91 disjunct population that declined sharply in the last decades. By 2019, it was functionally  
92 extinct with only two non-breeding adult macaws. Since that year, a long-term

93 reintroduction and monitoring program has been releasing rescued, confiscated and  
94 captive-bred macaws (GPCA, 2024).

95         The Caatinga is one of the largest tropical semi-arid areas in the world (Moro et  
96 al., 2016), where the seasonally tropical dry forests prevail as dominant vegetation,  
97 consisting of a mosaic of different physiognomies spanning a broad range of woody plant  
98 densities and shrubs. The climate is semiarid, and the average mean temperature is  
99 constant and high over the year, ranging 25-30°C. The region presents a wide  
100 spatiotemporal variation in annual and interannual rainfall, with most of the  
101 territory (68.8%) receiving 600-1,000 mm of annual precipitation, and 20% receiving  
102 less than 600 mm. However, this rainfall is concentrated over a few months of the year  
103 (Silva et al., 2017). Thus, the vegetation is under seasonal hydric deficiency, experiencing  
104 long drought periods. Regionally, vegetation dynamics vary widely across time and space  
105 and are strongly correlated with rainfall. Most plants usually lose their leaves in the dry  
106 period and grow and produce flowers and fruits during the short-wet season (Silva et al.,  
107 2017; Fernandes et al., 2020).

108         The combination of acute human disturbances (e.g., conversion of large areas of  
109 native vegetation into roads, energy infrastructure, or commercial agriculture), chronic  
110 disturbances (e.g., slow but continuous native vegetation overexploitation, slash-and-  
111 burn agriculture, and browsing by livestock), proliferation of exotic species (e.g.,  
112 Africanized honey bees; Pacífico et al., 2020) and climate change can lead to severe  
113 degradation of natural habitats and disruption of critical ecological services (Silva et al.,  
114 2017). Thus, macaws face the impacts of several anthropogenic activities in their habitat,  
115 and food availability is a concern for population growth and persistence (Barbosa & Tella,  
116 2019).

117

## 118 **Methods**

### 119 **Data collection**

#### 120 **Macaw capture, bio-logger deployment and GPS data collection**

121 We used bio-loggers (tags) to track macaw movements. Juvenile birds were fitted  
122 with 15 g solar-powered tags (e-obs, Germany) equipped with Ultra High Frequency  
123 (UHF), GPS, and tri-axial accelerometry sensors, and attached using a Teflon ribbon  
124 harness in backpack-configuration. The total weight of the tag plus harness was  
125 about 2.4% of the mean adult body mass (720 g), following established tag weight-limit  
126 guidelines (Yeap et al., 2021). After tag deployment, we carried out monthly surveys of  
127 known roosting and feeding sites to search and monitor the tagged macaws, allowing GPS  
128 data retrieval via UHF data transmission. Before data analysis, the movement data was  
129 uploaded to and stored in Movebank (Kays et al., 2022). See Supplementary Material 1  
130 for details on tag deployments and GPS data collection.

131

### 132 **Data analysis**

#### 133 **Identification of feeding, resting and roosting sites and spatial distribution of the** 134 **locations**

135 We identified the main feeding, resting and roosting sites using the “Roost and Foraging  
136 Site Extraction” MoveApps workflow (Kölzsch et al., 2023), by extracting mean  
137 coordinates of daily diurnal and nocturnal locations of high GPS fix density where  
138 macaws stayed in a defined radius for a defined minimum duration, not moving faster  
139 than 1 m/s (GPS ground speed). Before the identification of these sites, all GPS fixes  
140 were classified as day or night positions, delineated by sunrise + 30 min and by  
141 sunset - 60 min (as macaws tend to leave roosting sites about 30 min after sunrise, and  
142 stay at foraging sites until about 60 min before sunset; E.C.P. field observation, 2008).

143 Feeding sites were defined as all sites at daylight where a macaw stayed for at  
144 least 5 h within a radius of 1,000 m. The resting sites were defined as all sites during  
145 daytime where a macaw stayed for at least 3 h (minimum resting duration) within a radius  
146 of 50 m. Lastly, the roosting sites were defined as all sites during nighttime where a  
147 macaw stayed for at least 9 h within a radius of 500 m. Radii and time intervals selected  
148 to identify feeding, resting and roosting sites were established considering the species'  
149 daily movements between roosting areas and feeding sites, and observed behaviour;  
150 the 9-hour minimum interval at night was defined based on the known roosting  
151 behaviour, and to be long enough to fully include the tag battery-saving sleep  
152 mode (7pm – 4am). See Supplementary Material 2 for details on the MoveApps analysis  
153 protocol.

154 To spatially characterize the identified locations, we calculated the distances  
155 between the resting sites to the closest feeding sites, and the distances between the resting  
156 and feeding sites to the closest roosting sites using the “Distance Matrix” tool in QGIS  
157 (QGIS Development Team, 2023).

158

### 159 **Habitat characterization**

160 We characterized the environments visited by the macaws with variables representing  
161 anthropogenic features, land cover and topography. We derived rasters with the  
162 calculated Euclidean distances from both the medium-voltage power lines  
163 (ANEEL, 2024) and roads (DNIT, 2021) networks. We used land cover and land use  
164 rasters from MapBiomas to characterize the landscape composition with natural  
165 vegetation and anthropogenic activities (Franca Rocha et al., 2024; MapBiomas, 2024).  
166 With the same purpose, we used the Human Modification Degree index (Kennedy et al.,  
167 2019), a continuous metric varying from 0-1 that reflects the proportion of a landscape



168 that is modified based on modelling physical extents of 13 main anthropogenic stressors  
169 (with global coverage) and their estimated impacts. Finally, we also derived mean slope  
170 values (degrees) from Copernicus Global Digital Elevation Model (European Space  
171 Agency, 2024). The locations previously identified were then overlapped onto GIS  
172 remote sensing layers obtained. See Supplementary Table 1 for more detailed  
173 information.

174

### 175 **Movement data processing and generation of subsets prior to home range estimation**

176 To account for potential location errors before estimating individual home ranges, the  
177 user equivalent range error (UERE) was calculated using the `ctmm` package in R  
178 (Calabrese et al., 2016; Fleming et al., 2020). GPS points were collected while tags  
179 remained stationary for 10-14 hours in the study area to estimate the UERE values. The  
180 horizontal UERE of 0.437 m (95% CI: 0.377-0.497) was incorporated into the dataset.

181       Following this calibration, individual tracking data were divided into 15-day  
182 intervals (fortnightly subsets) using R (R Core Team, 2023), resulting in 211 subsets  
183 spanning May 2017 to May 2023 (Supplementary Table 2). These intervals allowed the  
184 assessment of temporal variation in range sizes. Using the fortnightly subsets, we  
185 generated 211 home range and core area estimates, as detailed in supporting materials  
186 (Supplementary Table 3).

187

### 188 **Home range and core area estimation, and environmental effects**

189 We estimated fortnightly individual home ranges and core areas (Supplementary Table 3)  
190 with GPS data using autocorrelated kernel density estimators (AKDE) with the `ctmm` R  
191 package (R Core Team, 2023). The home ranges were calculated using 95% AKDE,

192 corresponding to the individual's mobility potential; and the core areas,  
193 using 50% AKDE, corresponding to the areas the animal uses most frequently (Fleming  
194 et al., 2015; Calabrese et al., 2016). Calculating macaws' ranges every 15 days allowed  
195 us to assess changes in home range and core area size throughout the annual cycle and  
196 compare their variation size between seasons. See Supplementary Material 3 for more  
197 details on home range and core area estimation method.

198         Estimated ranges of birds with more than one year of movement data  
199 (Supplementary Table 4; IDs 5568, 5570, 6444, 9025) were evaluated concerning  
200 interannual and seasonal variation. The selection of predictor variables was based on their  
201 expected influence on the macaws' movement. Using a linear model (bbmle R package;  
202 Rencher & Schaalje, 2008), we explored if extrinsic factors were good predictors of home  
203 range and core area size. We evaluated the effect of site (i.e., roosting site where each  
204 macaw was born – or released – and tagged, and from where the individual performs daily  
205 movements to forage), season (“wet” versus “dry”), ordinal date, rainfall precipitation  
206 and vegetation productivity (both proxies of licuri palm fruit availability; see  
207 Supplementary Material 4 for details on environmental predictors and data processing,  
208 and Supplementary Table 5 for rainfall and Normalized Difference Vegetation Index  
209 (NDVI) data).

210

### 211 **Home range and core area size model**

212 We tested models that considered the influence of one variable (i.e., the variables were  
213 analysed independently), and models that considered the influence of two variables  
214 (pairwise), with and without interaction (Supplementary Tables 6 and 7 for home range  
215 and core area analysis, respectively). Pearson's correlation coefficients ( $r$ ) were calculated  
216 (Supplementary Table 8), and the predictor variables with  $|r| > 0.6$  were excluded from  
10

217 combined models. For the pairwise analysis, different variables selected to represent the  
218 seasonal variation (i.e., season, ordinal date, accumulated rainfall and NDVI data) were  
219 not evaluated together in the same model. Also, the variables “tagging site” and “year of  
220 monitoring” were not evaluated simultaneously in the same model due to correlation,  
221 because for a specific monitoring time frame, movement data was only available for  
222 individuals born in the same breeding site. In those cases, the variable “tagging site” was  
223 tested in the models.

224         We used the AIC to rank models and selected the model with the lowest value  
225 (Akaike, 1987; Burnham & Anderson, 2002). Moreover, we calculated the adjusted  
226 coefficient of determination ( $R^2$ ) as a measure of how much of the observed variation was  
227 explained by the linear relationship with the explanatory variables included in the model  
228 (Su et al., 2012; bbmle and AICcmodavg R packages; R Core Team, 2023).

229

## 230 **Results**

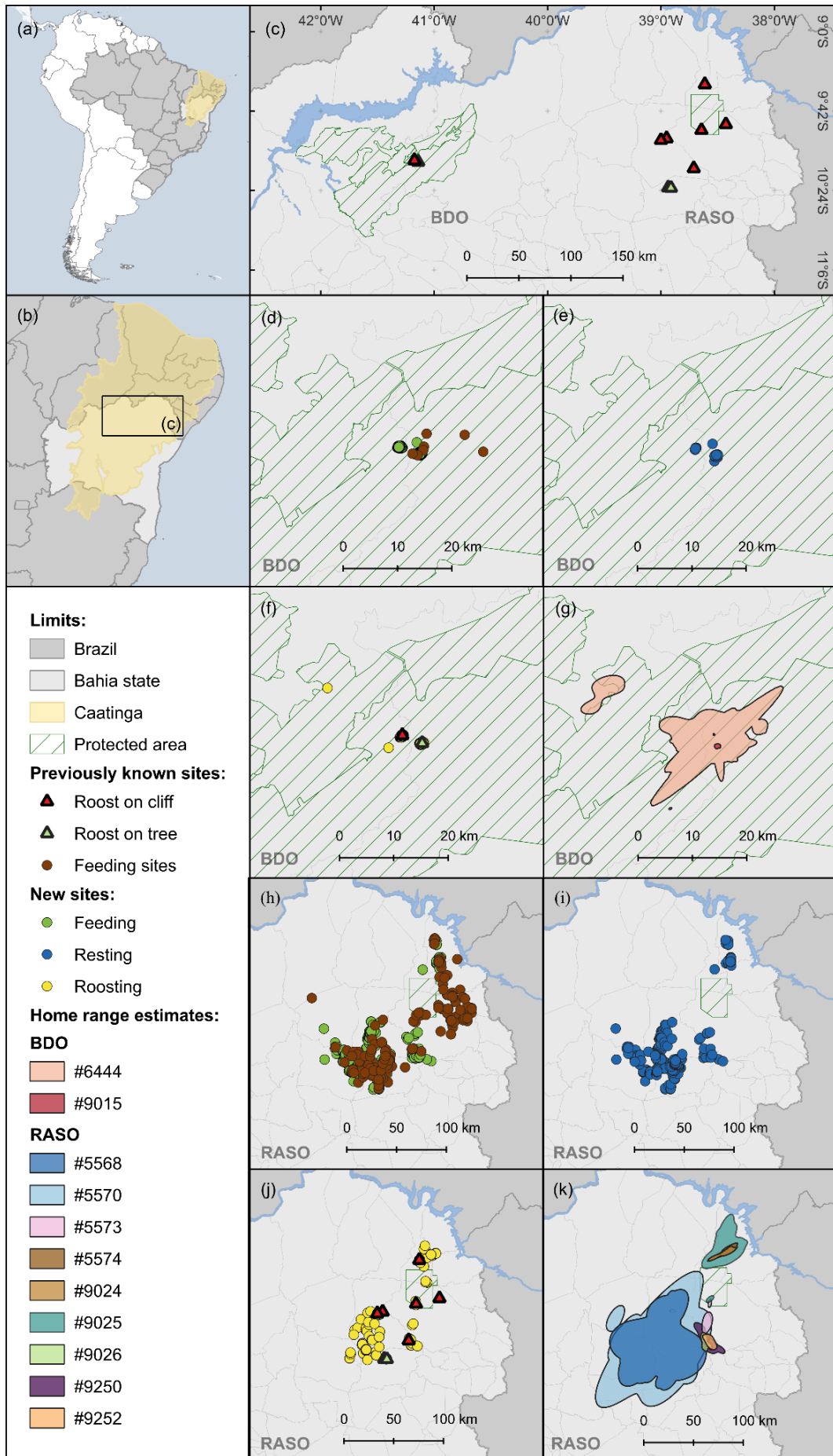
231 We collected GPS data from 2017 to 2024 (Supplementary Table 4). Although we  
232 retrieved movement data from 11 juvenile macaws tracked during this period, some  
233 individuals had limited tracking periods (< 45 days, or < 10 days of data; Supplementary  
234 Table 4). Data from these individuals represented 1% of the total GPS fixes of the  
235 telemetry dataset. While this data was included in the descriptive analysis and  
236 environmental characterizations, it was excluded from subsequent home range analyses.  
237 The tracking duration for individual macaws ranged from four to 1,346 days, and the  
238 average tracking period was 347 days (SD = 486.12). The final dataset consisted  
239 of 841,761 GPS fixes, spanning seven years (2017 – 2024; except the year 2020, for  
240 which no movement data was retrieved). The number of GPS fixes per individual ranged  
241 from 178 to 668,473 (Supplementary Table 4), with mean and median values of 76,524  
242 and 2,356 points, respectively.

243

## 244 **Identification of feeding, resting and roosting sites and spatial distribution of the** 245 **locations**

246 We identified 3,038 feeding, 1,228 resting and 2,923 roosting locations, distributed in  
247 *Boqueirão da Onça* (Fig. 1d,f) and *Raso da Catarina* regions (Fig. 1h,j).

248



250 FIG. 1 (a) Caatinga Domain in northeastern Brazil. (b) Study area in northern Bahia state,  
251 comprising the (c) *Boqueirão da Onça* (BDO; left) and *Raso da Catarina* (RASO;  
252 right). (d-f) Feeding, resting, roosting sites and (g) total home ranges of two macaws  
253 tagged in BDO; (h-j) Feeding, resting, roosting sites and (k) total home ranges of nine  
254 macaws tagged in RASO. Total home ranges were estimated by combining all the  
255 fortnightly individual ranges using the “Merge” tool in QGIS software  
256 (QGIS Development Team, 2023). Previous feeding areas database: GPCA data collected  
257 over the last 10 years, CEMAVE and Silva-Neto et al. (2012).

258

259 The resting sites were located very close to the feeding areas, with a median  
260 distance of only 19.8 meters to the nearest feeding site (Supplementary Table 9).  
261 Additionally, there were records of resting areas (2.5%) overlapping feeding locations  
262 (minimum distance = 0). The median distance to the nearest roosting site was 361.52 m;  
263 the feeding sites, in turn, were closer to the roosting sites (median distance = 201.13 m;  
264 Supplementary Table 9). Both feeding and resting sites were located as far as 36 km from  
265 the nearest roost.

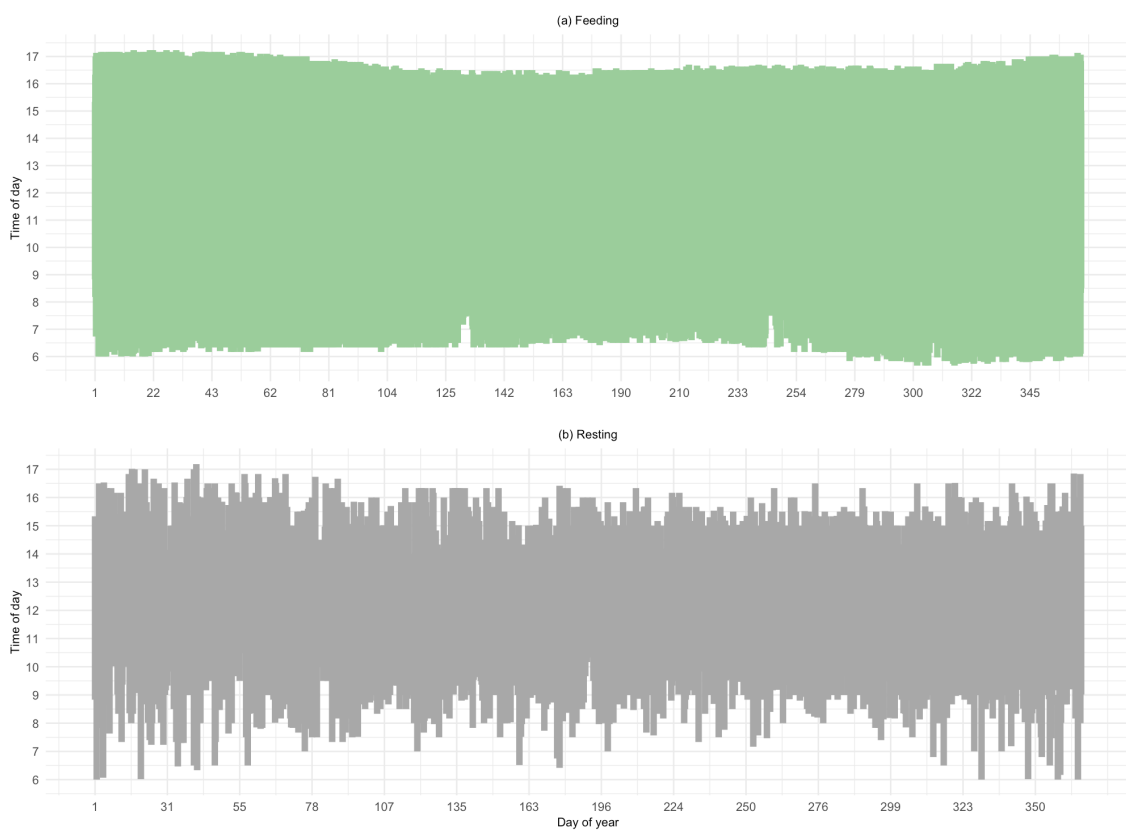
266

### 267 **Spatiotemporal analysis**

268 Macaws spent about 8 hours daily (median, ranging 5-11 h duration) in foraging sites,  
269 within a median radius of 431 meters (Supplementary Table 10). In comparison, the time  
270 spent in resting sites was almost two times shorter, with a median value of 4.2 h  
271 (ranging 3-10.5 h during the day), and a median radius of only 18.2 m. The night data, in  
272 turn, indicated that macaws roosted about 12 h per night (ranging 9-14 h; Supplementary  
273 Table 10), within a median radius of 160 m.

274 The feeding behaviour started early in the day, with macaws leaving roosts and  
275 initiating exploration of the foraging sites around 05:30-06:30 am (Fig. 2a). The start  
276 times of resting behaviour, in turn, were more distributed until mid-morning (Fig. 2b).  
277 Both the feeding and resting activities occurred throughout the day, lasting until late  
278 afternoon (Fig. 2a,b). Moreover, during the winter, macaws tended to arrive later and  
279 leave both the feeding and resting sites earlier (Fig. 2a,b).

280



281

282 FIG. 2 Daily activity patterns, throughout the year, of macaws arriving at feeding (a) and  
283 resting (b) sites and leaving these sites to return to their roosts. Data from 2017 to 2024.

284

### 285 **Habitat characterization**

286 Approximately 30% of feeding and resting sites were located close to medium-voltage  
287 power lines (zero up to 1,000 m distance), while only about 10% of roosting sites were

288 within the same range. 42% and 31% of feeding and resting sites, respectively, were also  
289 located within 10,000-12,000 m of electrical infrastructure. The feeding and resting sites  
290 also showed similar results regarding the central tendency of distance to power lines  
291 (Fig. 3a,e). For roosting sites, the median distance from power lines was about four times  
292 greater (Fig. 3i).

293 The feeding and resting sites also showed similar trends regarding the closeness  
294 to roads (not surprising, given that most power lines are built along roads), with more  
295 than 40% of the locations situated at distances up to 10 km (Fig. 3b,f). Almost 40% of  
296 roosting sites were also located within this range, although the median distance from  
297 roads was superior (Fig. 3j). On the other hand, about 30-40% of all the identified  
298 feeding, resting and roosting sites were located 30-35 km away from road surfaces.

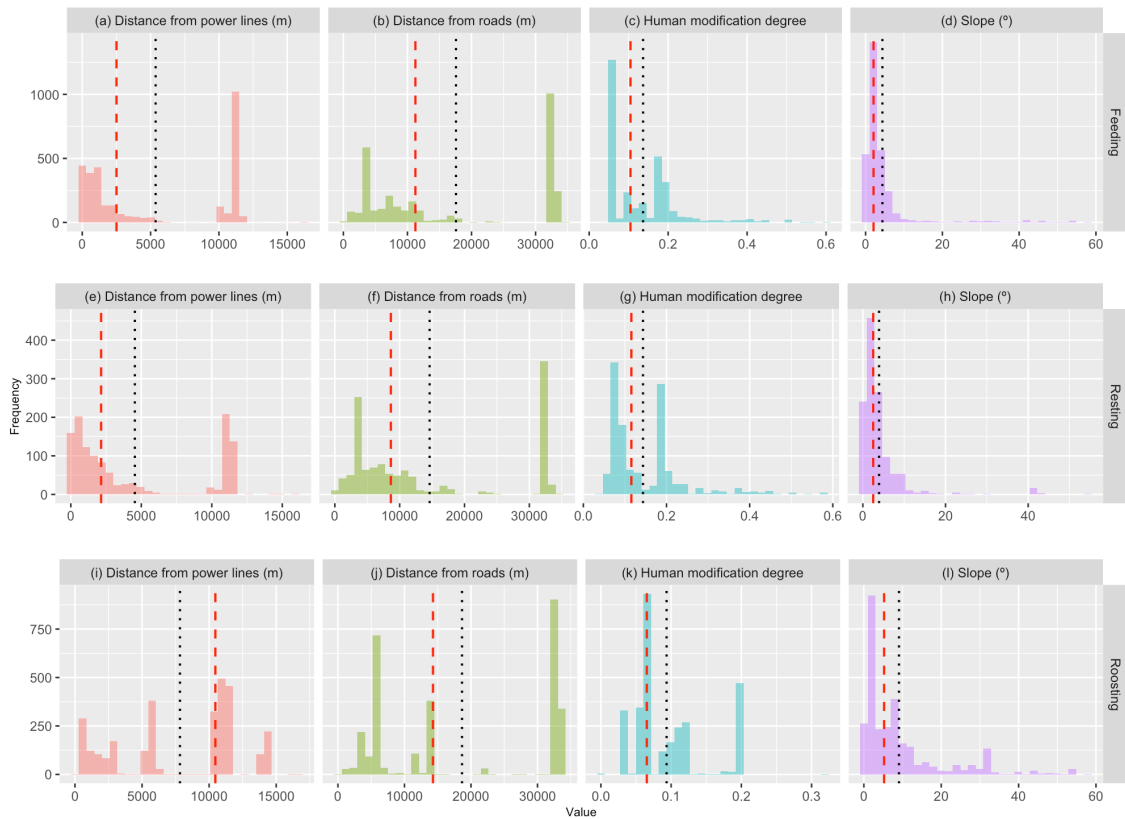
299 Strikingly, 100% of all lands identified had some indication of human activities  
300 ( $HMc > 0$ ). Once more, the feeding and resting sites presented similar results (Fig. 3c,g),  
301 with about half of their locations showing a moderate degree ( $0.10 < HMc \leq 0.40$ ) of  
302 human modification. Although the roosting sites showed a central tendency to be in lands  
303 with a lower degree of human modification ( $HMc \leq 0.1$ ; Fig. 3k), 35% of these sites were  
304 categorized as moderate modification ( $0.10 < HMc \leq 0.40$ ; the maximum degree value  
305 obtained was 0.31).

306 The median slope, although twice as high for roosting sites, showed low values  
307 (under 6 degrees) for all identified locations (Fig. 3d,h,l), indicating that macaws also  
308 used gentler slope surfaces as roosting areas. The feeding and resting sites had a very  
309 similar distribution of values, with a higher concentration of data at lower values  
310 (Fig. 3d,h). The slope of the terrain in roosting areas (Fig. 3l), on the other hand, showed  
311 greater variation, with a higher frequency of steeper slopes, compared to feeding and



312 resting sites. Moreover, all higher measurements (above 40 degrees) identified were  
 313 located neighbouring the previously mapped sandstone cliffs used as roosts, suggesting  
 314 that macaws use gentler surfaces during daylight activities.

315



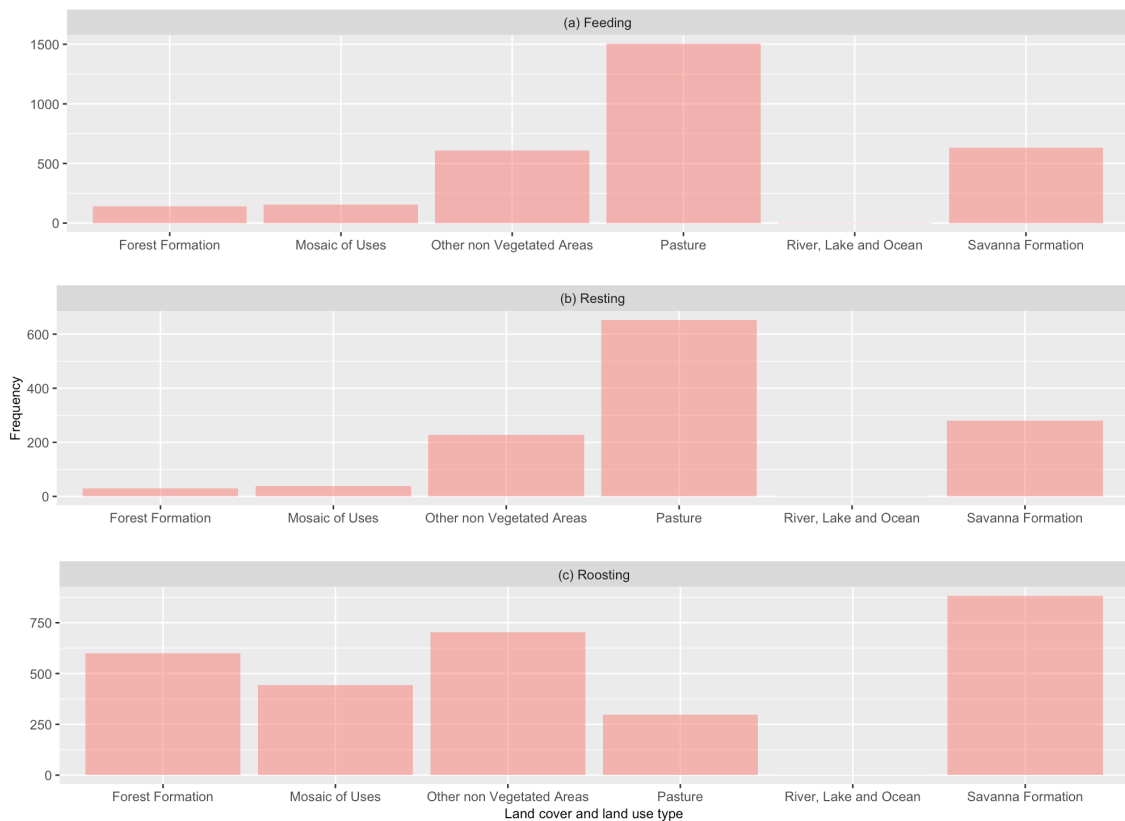
316

317 FIG. 3 Habitat characterization of feeding (a-d), resting (e-h) and roosting (i-l) locations  
 318 using the continuous variables selected to represent anthropogenic features and  
 319 topography. The median of each value is indicated by a red dashed line, the mean, by a  
 320 black dotted line. Note the variable scale of the y-axis.

321

322 More than half of feeding and resting sites were concentrated within areas directly  
 323 related to agricultural activity (Pasture and Mosaic of Uses; the latter characterized as  
 324 areas where it was not possible to distinguish between pasture and agriculture), with  
 325 about 50% located in non-natural Pasture lands (Fig. 4a,b). Native formations accounted

326 for ~25% of feeding and resting sites (Forest and Savanna). On the other hand, roosting  
 327 sites showed the opposite tendency, with the native vegetation cover classes accounting  
 328 for 50% of locations, while 25% of locations were within agricultural use lands (Fig. 4c).  
 329 It should be noted that about 20-25% of the feeding, resting and roosting sites were in  
 330 Other non-Vegetated Areas (Fig. 4a-c), characterized by non-permeable surfaces, such as  
 331 infrastructure, urban expansion or mining (these are clearly human-modified areas,  
 332 although these surfaces could not be mapped into their specific land use classes).  
 333



334  
 335 FIG. 4 Habitat characterization of (a) feeding, (b) resting and (c) roosting locations using  
 336 the land cover and land use variable. Note the variable scale of the y-axis.

337

338 **Home range and core area estimation**

339 All the movement models selected for the fortnightly subsets identified the residence  
340 behaviour of macaws (OU anisotropic, OUF and OUF anisotropic, and IID anisotropic  
341 (Calabrese et al., 2016); even for those individuals with less than 30 days of data;  
342 supporting information in Supplementary Table 2).

343 The size of fortnightly ranges of juvenile birds showed intra and interannual  
344 variation; also, the estimates varied across individuals (Fig. 5 and 6). The fortnightly  
345 analysis of overall range size indicated an average home range (95% kernel contour)  
346 of 850.2 km<sup>2</sup> (1.2-8,549.5 km<sup>2</sup>; SD = 1,292.3 km<sup>2</sup>), and an average core area (50% kernel  
347 contour) size of 198.6 km<sup>2</sup> (1.1-2,132 km<sup>2</sup>; SD = 322.7 km<sup>2</sup>); with greater variability in  
348 the size of core areas ( $CV_{\text{home range}} = 1.52$ ;  $CV_{\text{core area}} = 1.625$ ). Supporting information in  
349 Supplementary Table 3.

350 Macaws' dry season (June – December) ranges were highly variable,  
351 averaging 1,097.1 km<sup>2</sup> for the home ranges (1.7-8,549.5 km<sup>2</sup>; SD = 1,529.8 km<sup>2</sup>), while  
352 the core area size averaged 265.4 km<sup>2</sup> (1.7-2,132 km<sup>2</sup>; SD = 387.4 km<sup>2</sup>). The wet season  
353 (January – May) range analysis indicated the average of home ranges was 511.6786 km<sup>2</sup>  
354 (1.2-3,346.1 km<sup>2</sup>; SD = 753.1 km<sup>2</sup>), and the average of core areas, 107.1 km<sup>2</sup> (1.1-  
355 770 km<sup>2</sup>; SD = 165.9 km<sup>2</sup>).

356



357

358 FIG. 5 Home range size, in log scale, from 2017-2023, with the dry and wet seasons of  
 359 each year coloured in lighter orange and blue. Each point represents the fortnightly  
 360 estimate of individuals' home range. The Confidence Intervals are shown around each  
 361 estimate.

362



364

365 FIG. 6 Core area size, in log scale, from 2017-2023, with the dry and wet seasons of each  
 366 year coloured in lighter orange and blue. Each point represents the fortnightly estimate of  
 367 individuals' core area. The Confidence Intervals are shown around each estimate.

368

### 369 Home range and core area size model

370 The temporal variation in fortnightly home range size of juvenile Lear's macaws was best  
 371 explained by season and site (Supplementary Table 6), indicating that macaws tend to  
 372 have variable home range sizes depending on the site – larger in *Estação Biológica de*  
 373 *Canudos* roosting site, smaller in *Boqueirão da Onça* release area –, and larger home  
 374 ranges in dry season. However, the adjusted  $R^2$  value estimated for the selected model  
 375 ( $R^2 = 0.02256$ ) indicated that these predictor variables represented only ~2.25% of the  
 376 temporal variation observed in the sizes of the macaws' home ranges. Also, other equally  
 377 plausible models were selected ( $dAIC < 2$ ; Supplementary Table 6), including the null

378 model – therefore indicating only a slight influence of these variables on the home ranges  
379 size temporal variation.

380         Regarding the core areas, their temporal variation was linked to the accumulated  
381 rainfall over the last six months (Supplementary Table 7). However, the adjusted  $R^2$  value  
382 estimated for the top model ( $R^2 = 0.01699$ ) indicated that this predictor variable also  
383 represented only a small fraction ( $\sim 1.7\%$ ) of the temporal variation observed in the  
384 fortnightly sizes of the macaws' core areas; additionally, other equally plausible models  
385 were selected ( $dAIC < 2$ ; Supplementary Table 7).

386

## 387 **Discussion**

388 Our findings provide the first comprehensive estimates of Lear's macaw home ranges and  
389 core areas, offering novel insights into their spatial ecology and habitat use. Spatial  
390 distribution analysis of feeding, resting, and roosting sites demonstrated that macaws  
391 predominantly utilize areas outside protected zones, with roosting sites concentrated near  
392 sandstone cliffs; while feeding and resting sites are often in anthropogenically modified  
393 landscapes, scattered across individuals' ranges – providing empirical evidence for key  
394 habitats in need of restoration and conservation. We also found significant inter and intra-  
395 annual variation in home range sizes, with larger ranges during the dry season. We  
396 observed that, while tagging site and season influenced home range variation, the  
397 environmental productivity proxies – NDVI and rainfall – did not strongly explain the  
398 temporal fluctuations. These results underscore the complexity of factors influencing  
399 macaws' space use, suggesting a multifaceted interaction between rainfall, landscape  
400 composition and configuration and food availability (McIntyre & Wiens, 2000; Adamek,  
401 2011).

402 Deploying GPS tags on Lear's macaws presented logistical challenges but yielded  
403 valuable long-term movement data (Supplementary Table 4). Of the 38 tagged  
404 macaws, 44.7% experienced tag removal or destruction, highlighting the challenges  
405 associated with tracking of parrots in their natural habitats. This issue is common among  
406 parrot species that engage in preening and mutual preening behaviours (Le Souef et al.,  
407 2013; Groom et al., 2014). Therefore, understanding loss rates is crucial for planning  
408 future telemetry research, especially given the high cost of equipment. Additional factors  
409 may also contribute to data loss, such as tag malfunctions or the inability to locate tagged  
410 macaws during monitoring. Extended deployment periods, exceeding a year for some  
411 individuals, provided critical insights into seasonal and interannual movements. Ongoing  
412 refinement of harness attachment methods could further reduce logger loss and enhance  
413 data recovery rates in future studies (Brightsmith et al., 2021).

414 The spatial and temporal distribution of macaw activity hotspots revealed distinct  
415 patterns in the use of feeding, resting, and roosting sites. Roosting sites, primarily  
416 sandstone cliffs, remain critical despite new findings about tree roosts and served as stable  
417 nightly refuges while feeding and resting locations shifted more dynamically across the  
418 landscape. Interestingly, an amount of feeding and resting sites overlapped, suggesting  
419 that macaws utilize the same areas for multiple activities, thereby minimizing energy  
420 expenditure (Santos Neto & Camandaroba, 2008; Pacífico et al., 2014). These findings  
421 reflect the species' reliance on localized licuri palm patches, reinforcing the importance  
422 of conserving these fragmented but essential habitats.

423 Habitat characterization of feeding, resting, and roosting sites highlighted notable  
424 differences in landscape features. Feeding and resting sites were often situated closer to  
425 roads and power lines, with higher levels of human modification (HMc), compared to  
426 roosting sites. Roosting areas, on the other hand, were associated with steeper slopes and

427 less disturbed habitats, underscoring the critical role of sandstone cliffs. These  
428 distinctions emphasize the macaws' ability to navigate anthropogenic landscapes for  
429 foraging while depending on more pristine environments for roosting, reflecting an  
430 adaptive spatial strategy in response to habitat fragmentation (Tuomainen & Candolin,  
431 2011; Salinas-Melgoza et al., 2013; Renton et al., 2015).

432 Home range and core area estimates exhibited substantial variation across  
433 individuals. Also, the ranges' sizes were smaller for the reintroduced macaws  
434 (IDs 6444, 9015; Supplementary Table 3) than for the wild ones. The mean home range  
435 size of 850.15 km<sup>2</sup> aligns with estimates from other large macaws, although the observed  
436 variability (ranging 1.24-8,549.48 km<sup>2</sup>) highlights individual differences in space use  
437 (Brightsmith et al., 2021). This range disparity likely reflects differences in age,  
438 experience, and local resource distribution (Viana et al., 2018; Isted et al., 2023).  
439 Temporal variation in home range sizes, with larger ranges during dry season, supports  
440 the hypothesis that macaws expand their foraging areas in response to seasonal resource  
441 scarcity. This pattern mirrors findings on other tropical frugivores, where seasonal  
442 movements and dry-season range expansion is a common adaptive strategy (Salinas-  
443 Melgoza et al., 2013; Brightsmith et al., 2021). The variation in home range sizes over  
444 annual cycles has also been documented for other species and highlights the dynamic  
445 nature of home ranges (Chan et al., 2022; Isted et al., 2023; Broekman et al., 2024).  
446 Usually, animals tend to show smaller home ranges when habitats present higher  
447 productivity, leading to higher resource availability and so demanding smaller areas to  
448 find enough resources (Börger et al., 2008; Duncan et al., 2015). The spatiotemporal  
449 variation in resource availability is considered one of the key drivers of home range size  
450 variation (Broekman et al., 2024).



451           The licuri palm, a key resource for Lear's macaw, is influenced by rainfall patterns  
452 typical of the Caatinga dry forest (Pacífico et al., 2014). However, although the  
453 precipitation temporal scales we selected were based on its reproductive phenology, it is  
454 possible that its flowering and fructification are not immediately triggered by rainfall, as  
455 occurs with most Caatinga plants. During droughts, licuri palms may maintain vegetative  
456 growth, but fruit production can significantly decrease, further impacting macaw foraging  
457 behaviour (Griz & Machado, 2001; Alves et al., 2019; Noblick et al., 2020). Additionally,  
458 although NDVI is a well-established proxy for photosynthetic activity and green biomass  
459 (Pettorelli et al., 2005), its effectiveness as a proxy of licuri fruit availability may be  
460 limited, as palms can retain green foliage during different phenological stages, even  
461 during fruit-scarce periods (Alves et al., 2019). This discrepancy underscores the need for  
462 more tailored remote sensing techniques or ground-based phenological surveys to  
463 accurately capture the relationship between food availability and macaw space use.

464           The movement of macaws and variation in their home range sizes may be  
465 explained by multiple dynamic processes affecting their movement ecology (Isted et al.,  
466 2023). These factors can be both environmental and intrinsic, related to individuals'  
467 internal state (e.g., physiological restrictions) and/or individuals' traits (e.g., locomotion  
468 and/or navigation capacity; Nathan et al., 2008). Also, social interactions and other biotic  
469 factors may further shape macaw spatial use, highlighting the importance of incorporating  
470 diverse ecological drivers into future research.

471           Despite the observed variability in home range sizes, the consistent use of roosting  
472 and feeding areas emphasizes the residency pattern of Lear's macaws. This stability  
473 highlights the importance of restoring and protecting key areas that are critical for their  
474 survival, particularly licuri palm patches and sandstone cliffs, which serve as primary  
475 feeding and roosting sites. The reliance on fragmented landscapes and expansion of home

476 ranges during dry seasons underline the macaws' adaptive strategies in response to  
477 resource scarcity (Salinas-Melgoza et al., 2013), reinforcing the need for long-term  
478 conservation planning that addresses both the amount of suitable habitat and the seasonal  
479 and interannual habitat dynamics. Conservation efforts should focus on restoring and  
480 preserving native Caatinga vegetation and controlling land conversion and vegetation  
481 degradation to reduce anthropogenic disturbances, ensuring that critical resources like  
482 licuri fruits remain available throughout the year and support the long-term persistence  
483 of Lear's macaw populations (Pacífico et al., 2014; Barbosa & Tella, 2019).

484         Monitoring through telemetry studies continues to be an essential tool to identify  
485 and protect priority areas, guiding targeted actions to mitigate habitat degradation and  
486 support the persistence of macaws' populations in the long-term.

487

488 **Author contributions** Study design: FRP, ECP, FVD, JEFO; fieldwork: FRP, ECP,  
489 TF, GRF; data analysis, writing: FRP, ECP, FVD, JEFO, PC; reviewing: all authors.

490

491 **Acknowledgments** This study was funded by *Fundação de Amparo à Pesquisa do*  
492 *estado de São Paulo* (FAPESP), FVD Young Investigator Grant n° 2018/19389-9, FRP  
493 Masters Scholarship n° 2021/14510-7, JEFO Post-Doctoral Fellowship n° 2021/02132-8.  
494 Fieldwork activities and tags were also partially funded by *Volitalia* and *Enel Green*  
495 *Power* (2019-2023). We thank Dr. José L. Tella, Manuel de la Riva, Dr. Martin Wikelski,  
496 Dr. Kamran Safi, Dr. Andrea Koelzsch and Dr. Paulo Roberto Guimarães for scientific  
497 support; Dorivaldo Alves, Guilherme Feitosa, João Carlos Nogueira, Máximo Cardoso  
498 and Marcelo Oliveira for field work assistance; the National Center for Wild Bird  
499 Research and Conservation (CEMAVE), Chico Mendes Institute for Biodiversity

500 Conservation (ICMBio), *Fundação Biodiversitas* and Environment Authority for the  
501 Bahia state (INEMA) for logistic support. We also thank all fieldwork volunteers.

502

503 **Conflicts of interest** None.

504

505 **Ethical standards** This study abided by the Oryx guidelines on ethical standards. The  
506 macaws were captured and tagged under appropriate licenses granted by the Brazilian  
507 Ethics Committee (permit n° 381/2021) and ICMBio (permits n° 12763 and 59505).

508

509 **Data availability** The data that supported our findings is stored in Movebank Data  
510 Repository hosted by Max Planck Institute of Animal Behavior ([movebank.org](https://movebank.org), “(EBD)  
511 *Anodorhynchus leari* (Lear’s Macaw) Study”, study ID 322731302).

512

## 513 **References**

514 ADAMEK, K.A. (2011) *Temporal variation in space and resource use of macaws in the*  
515 *Southeastern Peruvian Amazon*. Masters’ Dissertation, Texas A&M University,  
516 Texas, USA.

517

518 AKAIKE, H. (1987) Factor analysis and AIC. *Psychometrika*, 52, 317–332.

519

520 ANEEL (2024) *Agência Nacional de Energia Elétrica. Sistema de Informação*  
521 *Geográfico Regulatório da Distribuição (SIG-R) e Base de Dados Geográfica da*  
522 *Distribuidora - BDGD*. [dadosabertos-](https://dadosabertos-aneel.opendata.arcgis.com/search?tags=distribuicao)  
523 [aneel.opendata.arcgis.com/search?tags=distribuicao](https://dadosabertos-aneel.opendata.arcgis.com/search?tags=distribuicao) [accessed November 2024].

524

525 ALVES, H., KUMAR, T.V.L., PAREDES, F., ELLIOTT, S. & AYUGA, J.G. (2019)  
526 Assessment of Caatinga response to drought using Meteosat-SEVIRI Normalized  
527 Difference Vegetation Index (2008 – 2016). *ISPRS Journal of Photogrammetry*  
528 *and Remote Sensing*, 148, 235–252.

529

530 BARBOSA, A.E.A. & TELLA, J.L. (2019) How much does it cost to save a species from  
531 extinction? Costs and rewards of conserving the Lear’s macaw. *Royal Society*  
532 *Open Science*, 6, 190190.

533  
534 BIRDLIFE INTERNATIONAL (2020) *Anodorhynchus leari* (amended version of 2019  
535 assessment). *The IUCN Red List of Threatened Species 2020*.  
536 [dx.doi.org/10.2305/IUCN.UK.2020-3.RLTS.T22685521A176030480.en](https://dx.doi.org/10.2305/IUCN.UK.2020-3.RLTS.T22685521A176030480.en) [accessed  
537 October 2021].  
538  
539 BÖRGER, L., DALZIEL, B.D. & FRYXELL, J.M. (2008) Are there general mechanisms of  
540 animal home range behaviour? A review and prospects for future research.  
541 *Ecology Letters*, 11, 637–650.  
542  
543 BRIGHTSMITH, D.J., BOYD, J.D., HOBSON, E.A. & RANDEL, C.J. (2021) Satellite  
544 telemetry reveals complex migratory movement patterns of two large macaw  
545 species in the western Amazon basin. *Avian Conservation and Ecology*, 16, art14.  
546  
547 BROEKMAN, M.J.E., HILBERS, J.P., HOEKS, S., HUIJBREGTS, M.A.J., SCHIPPER, A.M. &  
548 TUCKER, M.A. (2024) Environmental drivers of global variation in home range size  
549 of terrestrial and marine mammals. *Journal of Animal Ecology*, 93, 488–500.  
550  
551 BURNHAM, K.P. & ANDERSON, D.R. (2002) Model Selection and Multimodel Inference:  
552 A Practical Information-Theoretic Approach. In *Sociological Methods and*  
553 *Research* p. 2nd edition. Springer-Verlag, New York, USA.  
554  
555 CALABRESE, J.M., FLEMING, C.H. & GURARIE, E. (2016) ctm: an `<sc>r</sc>`  
556 package for analyzing animal relocation data as a continuous-time stochastic  
557 process. *Methods in Ecology and Evolution*, 7, 1124–1132.  
558  
559 CHAN, A.N., WITTEMYER, G., MCEVOY, J., WILLIAMS, A.C., COX, N., SOE, P., ET AL.  
560 (2022) Landscape characteristics influence ranging behavior of Asian elephants at  
561 the human-wildlands interface in Myanmar. *Movement Ecology*, 10, 6.  
562  
563 DNIT (2021) *Departamento Nacional de Infraestrutura de Transportes*.  
564 [servicos.dnit.gov.br/vgeo](https://servicos.dnit.gov.br/vgeo) [accessed October 2024].  
565  
566 DUNCAN, C., NILSEN, E.B., LINNELL, J.D.C. & PETTORELLI, N. (2015) Life-history  
567 attributes and resource dynamics determine intraspecific home-range sizes in  
568 Carnivora. *Remote Sensing in Ecology and Conservation*, 1, 39–50.  
569  
570 EUROPEAN SPACE AGENCY (2024) *Copernicus Global Digital Elevation Model*.  
571 Distributed by OpenTopography. <https://doi.org/10.5069/G9028PQB> [accessed  
572 November 2024].  
573  
574 FERNANDES, M.F., CARDOSO, D. & DE QUEIROZ, L.P. (2020) An updated plant checklist  
575 of the Brazilian Caatinga seasonally dry forests and woodlands reveals high  
576 species richness and endemism. *Journal of Arid Environments*, 174, 104079.  
577  
578 FLEMING, C.H., DRESCHER-LEHMAN, J., NOONAN, M.J., AKRE, T.S.B., BROWN, D.J.,  
579 COCHRANE, M.M., ET AL. (2020) A comprehensive framework for handling  
580 location error in animal tracking data. Cold Spring Harbor Laboratory. *bioRxiv*.  
581

- 582 FLEMING, C.H., FAGAN, W.F., MUELLER, T., OLSON, K.A., LEIMGRUBER, P. &  
583 CALABRESE, J.M. (2015) Rigorous home range estimation with movement data: a  
584 new autocorrelated kernel density estimator. *Ecology*, 96, 1182–1188.  
585
- 586 FRANCA ROCHA, W.J.S., VASCONCELOS, R.N., COSTA, D.P., DUVERGER, S.G., LOBÃO,  
587 J.S.B., SOUZA, D.T.M., ET AL. (2024) Towards Uncovering Three Decades of  
588 LULC in the Brazilian Drylands: Caatinga Biome Dynamics (1985–2019). *Land*,  
589 13, 1250.  
590
- 591 GOLDINGAY, R.L. (2015) A review of home-range studies on Australian terrestrial  
592 vertebrates: adequacy of studies, testing of hypotheses, and relevance to  
593 conservation and international studies. *Australian Journal of Zoology*, 63, 136.  
594
- 595 GPCA (2024) *Grupo de Pesquisa e Conservação da Arara-azul-de-lear (GPCA).*  
596 *Relatório anual 2023. São Paulo: 2024.*  
597 [acrobat.adobe.com/id/urn:aaid:sc:VA6C2:dd788c72-bed4-4191-98ca-](https://acrobat.adobe.com/id/urn:aaid:sc:VA6C2:dd788c72-bed4-4191-98ca-d5984143fcdf)  
598 [d5984143fcdf](https://acrobat.adobe.com/id/urn:aaid:sc:VA6C2:dd788c72-bed4-4191-98ca-d5984143fcdf) [accessed October 2024].  
599
- 600 GRIZ, L.M.S. & MACHADO, I.C.S. (2001) Fruiting phenology and seed dispersal  
601 syndromes in caatinga, a tropical dry forest in the northeast of Brazil. *Journal of*  
602 *Tropical Ecology*, 17, 303–321.  
603
- 604 GROOM, C., WARREN, K., LE SOUEF, A. & DAWSON, R. (2014) Attachment and  
605 performance of Argos satellite tracking devices fitted to black cockatoos  
606 (*Calyptorhynchus* spp.). *Wildlife Research*, 41, 571.  
607
- 608 ISTED, G.H., THOMAS, R.J., WARNER, K.S., STUBER, M.J., ELLSWORTH, E. & KATZNER,  
609 T.E. (2023) Ferruginous Hawk movements respond predictably to intra-annual  
610 variation but unexpectedly to anthropogenic habitats. *Ibis*, 165, 1156–1168.  
611
- 612 KAYS, R., DAVIDSON, S.C., BERGER, M., BOHRER, G., FIEDLER, W., FLACK, A., ET AL.  
613 (2022) The Movebank system for studying global animal movement and  
614 demography. *Methods in Ecology and Evolution*, 13, 419–431.  
615
- 616 KENNEDY, C.M., OAKLEAF, J.R., THEOBALD, D.M., BARUCH-MORDO, S. & KIESECKER,  
617 J. (2019) Managing the middle: A shift in conservation priorities based on the  
618 global human modification gradient. *Global Change Biology*, 25, 811–826.  
619
- 620 KÖLZSCH, A., LAMERIS, T.K., MÜSKENS, G.J.D.M., SCHREVEN, K.H.T., BUITENDIJK,  
621 N.H., KRUCKENBERG, H., ET AL. (2023) Wild goose chase: Geese flee high and far,  
622 and with aftereffects from New Year’s fireworks. *Conservation Letters*, 16.  
623
- 624 MAPBIOMAS (2024) *Coleção 8 da série anual de Mapas de Cobertura e Uso da Terra*  
625 *do Brasil.* [brasil.mapbiomas.org](https://brasil.mapbiomas.org) [accessed November 2024].  
626
- 627 MCINTYRE, N.E. & WIENS, J.A. (2000) A novel use of the lacunarity index to discern  
628 landscape function. *Landscape Ecology*, 15, 313–321.  
629

- 630 MORATO, R.G., CONNETTE, G.M., STABACH, J.A., DE PAULA, R.C., FERRAZ, K.M.P.M.,  
631 KANTEK, D.L.Z., ET AL. (2018) Resource selection in an apex predator and  
632 variation in response to local landscape characteristics. *Biological Conservation*,  
633 228, 233–240.  
634
- 635 MORO, M.F., NIC LUGHADHA, E., DE ARAÚJO, F.S. & MARTINS, F.R. (2016) A  
636 Phytogeographical Metaanalysis of the Semiarid Caatinga Domain in Brazil. *The*  
637 *Botanical Review*, 82, 91–148.  
638
- 639 NATHAN, R., GETZ, W.M., REVILLA, E., HOLYOAK, M., KADMON, R., SALTZ, D. &  
640 SMOUSE, P.E. (2008) A movement ecology paradigm for unifying organismal  
641 movement research. *Proceedings of the National Academy of Sciences*, 105,  
642 19052–19059.  
643
- 644 NOBLICK, L., WINTERGERST, S., NOBLICK, D. & LIMA, J.T. (2020) *Syagrus coronata*  
645 (Arecaceae) phenology and the impact of fire on survival and reproduction of the  
646 licuri palm. *SITIENIBUS série Ciências Biológicas*, 20, 1–9.  
647
- 648 OLIVEIRA, M. DA R., SZABO, J.K., JÚNIOR, A. DOS S., GUEDES, N.M.R., TOMAS, W.M.,  
649 CAMILO, A.R., ET AL. (2021) Lack of protected areas and future habitat loss  
650 threaten the Hyacinth Macaw (*Anodorhynchus hyacinthinus*) and its main food and  
651 nesting resources. *Ibis*, 163, 1217–1234.  
652
- 653 PACÍFICO, E.C., BARBOSA, E.A., FILADELFO, T., OLIVEIRA, K.G., SILVEIRA, L.F. &  
654 TELLA, J.L. (2014) Breeding to non-breeding population ratio and breeding  
655 performance of the globally Endangered Lear’s Macaw *Anodorhynchus leari* :  
656 conservation and monitoring implications. *Bird Conservation International*, 24,  
657 466–476.  
658
- 659 PACÍFICO, E.C., EFSTATHION, C.A., FILADELFO, T., HORSBURGH, R., CUNHA, R.A.,  
660 PASCHOTTO, F.R., ET AL. (2020) Experimental removal of invasive Africanized  
661 honey bees increased breeding population size of the endangered Lear’s macaw.  
662 *Pest Management Science*, 76, 4141–4149.  
663
- 664 PETTORELLI, N., VIK, J.O., MYSTERUD, A., GAILLARD, J.-M., TUCKER, C.J. &  
665 STENSETH, N.C. (2005) Using the satellite-derived NDVI to assess ecological  
666 responses to environmental change. *Trends in Ecology & Evolution*, 20, 503–510.  
667
- 668 QGIS DEVELOPMENT TEAM (2023) *QGIS Geographic Information System*. Open-Source  
669 Geospatial Foundation. [qgis.org](https://qgis.org)  
670
- 671 R CORE TEAM (2023) *R: A Language and Environment for Statistical Computing*. R  
672 Foundation for Statistical Computing, Vienna, Austria. [R-project.org/](https://r-project.org/)  
673
- 674 RENCHER, A.C. & SCHAALJE, G.B. (2008) *Linear Models in Statistics*. 2nd edition. John  
675 Wiley & Sons, New Jersey, USA.  
676

- 677 RENTON, K., SALINAS-MELGOZA, A., DE LABRA-HERNÁNDEZ, M.Á. & DE LA PARRA-  
678 MARTÍNEZ, S.M. (2015) Resource requirements of parrots: nest site selectivity and  
679 dietary plasticity of Psittaciformes. *Journal of Ornithology*, 156, 73–90.  
680
- 681 SALINAS-MELGOZA, A., SALINAS-MELGOZA, V. & WRIGHT, T.F. (2013) Behavioral  
682 plasticity of a threatened parrot in human-modified landscapes. *Biological*  
683 *Conservation*, 159, 303–312.  
684
- 685 SANTOS NETO, J.R. DOS & CAMANDAROBA, M. (2008) Mapeamento dos sítios de  
686 alimentação da arara-azul-de-lear *Anodorhynchus leari* (Bonaparte, 1856).  
687 *Ornithologia*, 3, 1–17.  
688
- 689 SCHOFIELD, G., HOBSON, V.J., LILLEY, M.K.S., KATSELIDIS, K.A., BISHOP, C.M.,  
690 BROWN, P. & HAYS, G.C. (2010) Inter-annual variability in the home range of  
691 breeding turtles: Implications for current and future conservation management.  
692 *Biological Conservation*, 143, 722–730.  
693
- 694 SILVA, J.M.C., LEAL, I.R. & TABARELLI, M. (2017) *Caatinga: The Largest Tropical*  
695 *Dry-Forest Region in South America*. Springer International Publishing, New  
696 York, USA.  
697
- 698 SILVA-NETO, G. F., SOUSA, A. E. B. A. & DOS SANTOS-NETO, J. R. (2012) Novas  
699 informações sobre a dieta da arara-azul-de-lear, *Anodorhynchus leari* Bonaparte,  
700 1856 (Aves, Psittacidae). *Ornithologia*, 5, 1–5.  
701
- 702 LE SOUEF, A.T., STOJANOVIC, D., BURBIDGE, A.H., VITALI, S.D., HEINSOHN, R.,  
703 DAWSON, R. & WARREN, K.S. (2013) Retention of transmitter attachments on  
704 black cockatoos (*Calyptorhynchus* spp.). *Pacific Conservation Biology*, 19, 55.  
705
- 706 SU, X., YAN, X. & TSAI, C. (2012) Linear regression. *WIREs Computational Statistics*,  
707 4, 275–294.  
708
- 709 TUOMAINEN, U. & CANDOLIN, U. (2011) Behavioural responses to human-induced  
710 environmental change. *Biological Reviews*, 86, 640–657.  
711
- 712 VIANA, D.S., GRANADOS, J.E., FANDOS, P., PÉREZ, J.M., CANO-MANUEL, F.J., BURÓN,  
713 D., ET AL. (2018) Linking seasonal home range size with habitat selection and  
714 movement in a mountain ungulate. *Movement Ecology*, 6, 1.  
715
- 716 WALTON, Z., SAMELIUS, G., ODDEN, M. & WILLEBRAND, T. (2017) Variation in home  
717 range size of red foxes *Vulpes vulpes* along a gradient of productivity and human  
718 landscape alteration. *PLOS ONE*, 12, e0175291.  
719
- 720 YEAP, L., SHEPHARD, J., BOUTEN, W., JACKSON, B., VAUGHAN-HIGGINS, R. & WARREN,  
721 K. (2017) Development of a tag-attachment method to enable capture of fine- and  
722 landscape-scale movement in black-cockatoos. *Australian Field Ornithology*, 34,  
723 49–55.  
724

725 YEAP, L., WARREN, K.S., BOUTEN, W., VAUGHAN-HIGGINS, R., JACKSON, B., RILEY, K.,  
726 ET AL. (2021) Application of tri-axial accelerometer data to the interpretation of  
727 movement and behaviour of threatened black cockatoos. *Wildlife Research*, 49,  
728 100–110.  
729  
730



# Revealing the unknown world of the endangered Lear's macaw using GPS-tracking data: identification of critical habitats for conservation

FERNANDA R. PASCHOTTO, ERICA C. PACÍFICO, THIAGO FILADELFO, GABRIELA R. FAVORETTO, JULIA E. F. OSHIMA, PAULO CARDOSO, FRANCISCO V. DÉNES

SUPPLEMENTARY MATERIAL 1 Details on tag deployment and GPS data collection.

In *Raso da Catarina* region, we deployed tags on 29 wild immature macaws (pre-fledging, approximately 3-months-old) captured inside their nests over seven breeding seasons between 2017 and 2023. Before the individual's capture, direct observations using telescopes and binoculars were undertaken to identify the nests occupied by breeding pairs (Pacífico et al., 2014). The cavity nests in the sandstone cliffs were accessed through rappel techniques, according to Pacífico et al. (2014), selecting the nestlings in the appropriate stage of development to receive the bio-loggers.

In *Boqueirão da Onça*, the release area, tags were deployed on seven captive-bred (two to five years old) and two rescued macaws (two and four years old) released between 2019 and 2022, after four months of training and adaptation in the release area. Tags were deployed during pre-release management conducted a few days prior to release, allowing behavioral observations of the tagged individuals both before and after the release.

Tags were programmed to record GPS locations at 10-min or 30-min intervals, between 4 am-7 pm (local time) every day year-round (frequency of GPS fixes depended on the available solar-powered battery charge).

SUPPLEMENTARY MATERIAL 2 MoveApps workflow analysis protocol performed for identification of feeding, resting and roosting sites.

To perform the identification of the feeding, resting and roosting sites, we accessed the “Roost and Foraging Site Extraction” workflow (Kölzsch, 2022) in the MoveApps (moveapps.org/workflows), a free platform for analysis of animal movement data. The Apps extracted, from the movement tracks, the mean coordinates of daily diurnal and/or nocturnal locations – depending on your settings – of high GPS fix density where the animals stayed in a certain radius area for a defined minimum duration, not moving faster than 1 m/s (GPS ground speed). The locations are given as .csv file output.

Within the main workflow “Roost and Foraging Site Extraction”, there are two available workflows: the ‘Foraging site extraction (day)’, with combined Apps to extract stationary locations during day; and the ‘Roost site extraction (night)’ to extract stationary locations during night. Be aware that there may be available more recent versions of the Apps within the workflows by the time you will use it. It is possible to upgrade the workflows and use the latest version.

We extracted both the feeding and resting sites running the analysis of ‘Foraging site extraction (day)’ workflow, changing a few parameters among them (detailed below); and extracted the roosting sites running the analysis of ‘Roost site extraction (night)’ workflow.

#### App ‘Movebank Location’ settings:

After adding the respective two workflows in our personal profile within the MoveApps platform, the next step (i.e., first App) was the selection of a data source, by configuring our Movebank login details to directly download the movement data stored in this platform – alternatively, it is also possible to upload movement-based files from a Cloud Storage or local systems.

We configured the Movebank location data source by providing the Movebank login, the study name, the animals for which we wanted to analyze the movement data, and the sensor types (GPS sensor). In this step, other settings were also requested. We chose to: not include outliers in this analysis, not use fast data reduction profile, and not use argument of minimization. Also, we selected a full data resolution, and the attribute of the defining track ID was a combination of animal and deployment. We repeated this procedure for both the ‘Foraging site extraction (day)’ and ‘Roost site extraction (night)’ workflows.

We saved the changes after configuring the parameters within each App.

#### App ‘Filter/Annotate by Season’ settings:

This App makes it possible to filter the data by season, comparing positions in different time intervals. In our study, we did not provide any time range to annotate positions to filter the data.

#### App ‘Daily Rest/Foraging Sites’ settings:

In both ‘Foraging site extraction (day)’ and ‘Roost site extraction (night)’ workflows, all GPS fixes were classified as day or night positions, delineated by sunrise +30 min (‘Sunrise adaptation time’) and by sunset -60 min (‘Sunset adaptation time’), as macaws tend to leave roosting sites about 30 min after sunrise, and stay at foraging sites until about 60 min before sunset; E.C.P. field observation, 2008). These settings depend on each species’ behavior.

For all the analysis, the ‘Maximum resting speed’ inserted was 1 m/s GPS ground speed (analysis default). Feeding sites were defined as all sites at daylight where a macaw stayed for at least 5 h (‘Minimum resting duration’) within a radius of 1,000 m (‘Maximum resting radius’). The resting sites were defined as all sites during daytime where a macaw stayed for at least 3 h (‘Minimum resting duration’) within a radius of 50 m (‘Maximum resting radius’). Lastly, the roosting sites were defined as all sites during nighttime where a macaw stayed for at least 9 h (‘Minimum resting duration’) within a radius of 500 m (‘Maximum resting radius’).

Radii and time intervals selected to identify feeding, resting and roosting sites were established considering the species’ movements and observed behavior.

#### Output:

The identified locations were given as .csv files output.

More detailed documentation of the Daily Rest/Foraging Sites MoveApps is available on the [authors’ GitHub repository](#).

SUPPLEMENTARY TABLE 1 Detailed information of the environmental variables ‘Distance from roads’, ‘Human modification’, ‘Distance from power lines’, ‘Land cover and land use’ and ‘Slope’ used for habitat characterization.

Variable	Type of information	Institution	Description	Unit	Year of original database	Original pixel resolution	Final pixel resolution	Primary or derived	Derived from	Source
Distance from roads	Anthropic	Departamento Nacional de Infraestrutura de Transportes (DNIT)	Raster with Euclidean distance calculated from roads shapefile	Meters	2021-2022	Vector	30m	Derived	Roads	Departamento Nacional de Infraestrutura de Transportes (DNIT) 2021. Visualizador de Informações Geográficas - Base oficial das rodovias federais e estaduais.
Human modification	Anthropic	The Nature Conservancy	Raster with values of human modification. It is a continuous 0-1 metric that reflects the proportion of a landscape modified based on modeling the physical extents of 13 anthropogenic stressors and their estimated impacts using spatially explicit global datasets with a median year of 2016	NA	2016	1km	30m	Primary	NA	Kennedy, C. M., Oakleaf, J. R., Theobald, D. M., Baruch-Mordo, S., & Kiesecker, J. (2019). Managing the middle: A shift in conservation priorities based on the global human modification gradient. <i>Global change biology</i> , 25(3), 811-826.
Distance from power lines	Anthropic	Agência Nacional de Energia Elétrica (ANEEL)	Raster with Euclidean distance calculated from medium voltage network	Meters	2020	Vector	30m	Derived	Medium voltage network	Agência Nacional de Energia Elétrica (ANEEL) 2024. Sistema de Informação Geográfico Regulatório da distribuição (SIG-R) e Base de Dados Geográfica da Distribuidora - BDGD
Land cover	Landscape	MapBiomas	Raster with values of land use and land cover classification	Land cover class	2016	30m	30m	Primary	NA	“MapBiomas - Coleção 8 da série anual de Mapas de Cobertura e uso da terra do Brasil, acessado em 2024 através do link: <a href="https://brasil.mapbiomas.org/">https://brasil.mapbiomas.org/</a> ”
Slope	Topography	European Space Agency	Raster with mean slope values derived from the Copernicus Digital Elevation Model	Degrees	2011-2015	30m	30m	Derived	Digital Elevation Model	European Space Agency (2021). Copernicus Global Digital Elevation Model. Distributed by OpenTopography. <a href="https://doi.org/10.5069/G9028PQB">https://doi.org/10.5069/G9028PQB</a> .

SUPPLEMENTARY TABLE 2 Summary of the fortnightly movement data series with identification of the Lear’s Macaws (“ID”) and the subsets, the beginning and end date of each subset (format year-month-day), the duration (in days) of the subsets, the number of points (i.e., geographic coordinates) registered for each subset, and the movement model selected.

\* Subsets that recorded less than 15 days of movement data.

<b>ID</b>	<b>Subset</b>	<b>Beginning date</b>	<b>End date</b>	<b>Duration (days)</b>	<b>Number of points</b>	<b>Selected model</b>
5568	s1	2017-05-03	2017-05-17	15	1,339	OUF anisotropic
5568	s2	2017-05-18	2017-06-01	15	1,364	OUF anisotropic
5568	s3	2017-06-02	2017-06-16	15	1,364	OUF anisotropic
5568	s4	2017-06-17	2017-07-01	15	1,358	OUF anisotropic
5568	s5	2017-07-02	2017-07-16	15	1,365	OUF anisotropic
5568	s6	2017-07-17	2017-07-31	15	1,363	OUF anisotropic
5568	s7	2017-08-01	2017-08-15	15	1,357	OUF anisotropic
5568	s8	2017-08-16	2017-08-30	15	1,364	OUF anisotropic
5568	s9	2017-08-31	2017-09-14	15	1,354	OUF anisotropic
5568	s10	2017-09-15	2017-09-29	15	1,348	OUF anisotropic
5568	s11	2017-09-30	2017-10-14	15	1,358	OUF anisotropic
5568	s12	2017-10-15	2017-10-29	15	1,360	OUF anisotropic
5568	s13	2017-10-30	2017-11-13	15	1,360	OUF anisotropic
5568	s14	2017-11-14	2017-11-28	15	1,365	OUF anisotropic
5568	s15	2017-11-29	2017-12-13	15	1,358	OUF anisotropic
5568	s16	2017-12-14	2017-12-28	15	1,349	OUF anisotropic
5568	s17	2017-12-29	2018-01-12	15	1,353	OUF anisotropic
5568	s18	2018-01-13	2018-01-27	15	1,363	OUF anisotropic
5568	s19	2018-01-28	2018-02-11	15	1,362	OUF anisotropic
5568	s20	2018-02-12	2018-02-26	15	1,360	OUF anisotropic
5568	s21	2018-02-27	2018-03-13	15	1,362	OUF anisotropic
5568	s22	2018-03-14	2018-03-28	15	1,353	OUF anisotropic
5568	s23	2018-03-29	2018-04-12	15	1,349	OUF anisotropic
5568	s24	2018-04-13	2018-04-27	15	1,356	OUF anisotropic
5568	s25	2018-04-28	2018-05-12	15	1,365	OUF anisotropic
5568	s26	2018-05-13	2018-05-27	15	1,346	OUF anisotropic
5568	s27	2018-05-28	2018-06-11	15	1,359	OUF anisotropic
5568	s28	2018-06-12	2018-06-26	15	1,348	OUF anisotropic
5568	s29	2018-06-27	2018-07-11	15	1,364	OUF anisotropic
5568	s30	2018-07-12	2018-07-26	15	1,358	OUF anisotropic
5568	s31	2018-07-27	2018-08-10	15	1,365	OUF anisotropic
5568	s32	2018-08-11	2018-08-25	15	1,365	OUF anisotropic
5568	s33	2018-08-26	2018-09-09	15	1,365	OUF anisotropic
5568	s34	2018-09-10	2018-09-24	15	1,365	OUF anisotropic
5568	s35	2018-09-25	2018-10-09	15	1,362	OUF anisotropic
5568	s36	2018-10-10	2018-10-24	15	1,362	OUF anisotropic
5568	s37	2018-10-25	2018-11-08	15	1,362	OUF anisotropic
5568	s38	2018-11-09	2018-11-23	15	1,362	OUF anisotropic
5568	s39	2018-11-24	2018-12-08	15	1,365	OUF anisotropic
5568	s40	2018-12-09	2018-12-23	15	1,361	OUF anisotropic

<b>ID</b>	<b>Subset</b>	<b>Beginning date</b>	<b>End date</b>	<b>Duration (days)</b>	<b>Number of points</b>	<b>Selected model</b>
5568	s41	2018-12-24	2019-01-07	15	1,363	OUF anisotropic
5568	s42	2019-01-08	2019-01-22	15	1,362	OUF anisotropic
5568	s43	2019-01-23	2019-02-06	15	1,364	OUF anisotropic
5568	s44	2019-02-07	2019-02-21	15	1,365	OUF anisotropic
5568	s45	2019-02-22	2019-03-08	15	1,358	OUF anisotropic
5568	s46	2019-03-09	2019-03-23	15	1,362	OUF anisotropic
5568	s47	2019-03-24	2019-04-07	15	1,362	OUF anisotropic
5568	s48	2019-04-08	2019-04-22	15	1,365	OUF anisotropic
5568	s49	2019-04-23	2019-05-07	15	1,362	OUF anisotropic
5568	s50	2019-05-08	2019-05-22	15	1,363	OUF anisotropic
5568	s51	2019-05-23	2019-06-06	15	1,364	OUF anisotropic
5568	s52	2019-06-07	2019-06-21	15	1,363	OUF anisotropic
5568	s53	2019-06-22	2019-07-06	15	1,362	OUF anisotropic
5568	s54	2019-07-07	2019-07-21	15	1,362	OUF anisotropic
5570	s1	2017-05-03	2017-05-17	15	1,329	OUF anisotropic
5570	s2	2017-05-18	2017-06-01	15	1,344	OUF anisotropic
5570	s3	2017-06-02	2017-06-16	15	1,293	OUF anisotropic
5570	s4	2017-06-17	2017-07-01	15	1,105	OUF anisotropic
5570	s5	2017-07-02	2017-07-16	15	1,072	OUF anisotropic
5570	s6	2017-07-17	2017-07-31	15	1,136	OUF anisotropic
5570	s7	2017-08-01	2017-08-15	15	1,188	OUF anisotropic
5570	s8	2017-08-16	2017-08-30	15	1,163	OUF anisotropic
5570	s9	2017-08-31	2017-09-14	15	1,115	OUF anisotropic
5570	s10	2017-09-15	2017-09-29	15	1,106	OUF anisotropic
5570	s11	2017-09-30	2017-10-14	15	1,183	OUF anisotropic
5570	s12	2017-10-15	2017-10-29	15	1,246	OUF anisotropic
5570	s13	2017-10-30	2017-11-13	15	840	OUF anisotropic
5570	s14	2017-11-14	2017-11-28	15	1,085	OUF anisotropic
5570	s15	2017-11-29	2017-12-13	15	940	OUF anisotropic
5570	s16	2017-12-14	2017-12-28	15	811	OUF anisotropic
5570	s17	2017-12-29	2018-01-12	15	753	OUF anisotropic
5570	s18	2018-01-13	2018-01-27	15	1,144	OUF anisotropic
5570	s19	2018-01-28	2018-02-11	15	1,144	OUF anisotropic
5570	s20	2018-02-12	2018-02-26	15	949	OUF anisotropic
5570	s21	2018-02-27	2018-03-13	15	950	OUF anisotropic
5570	s22	2018-03-14	2018-03-28	15	1,028	OUF anisotropic
5570	s23	2018-03-29	2018-04-12	15	1,035	OUF anisotropic
5570	s24	2018-04-13	2018-04-27	15	1,026	OUF anisotropic
5570	s25	2018-04-28	2018-05-12	15	1,234	OUF anisotropic
5570	s26	2018-05-13	2018-05-27	15	940	OUF anisotropic
5570	s27	2018-05-28	2018-06-11	15	1,238	OUF anisotropic
5570	s28	2018-06-12	2018-06-26	15	1,126	OUF anisotropic
5570	s29	2018-06-27	2018-07-11	15	1,051	OUF anisotropic
5570	s30	2018-07-12	2018-07-26	15	1,222	OUF anisotropic
5570	s31	2018-07-27	2018-08-10	15	1,144	OUF anisotropic
5570	s32	2018-08-11	2018-08-25	15	1,105	OUF anisotropic
5570	s33	2018-08-26	2018-09-09	15	1,180	OUF anisotropic

<b>ID</b>	<b>Subset</b>	<b>Beginning date</b>	<b>End date</b>	<b>Duration (days)</b>	<b>Number of points</b>	<b>Selected model</b>
5570	s34	2018-09-10	2018-09-24	15	941	OUF anisotropic
5570	s35	2018-09-25	2018-10-09	15	1,015	OUF anisotropic
5570	s36	2018-10-10	2018-10-24	15	952	OUF anisotropic
5570	s37	2018-10-25	2018-11-08	15	1,228	OUF anisotropic
5570	s38	2018-11-09	2018-11-23	15	799	OUF anisotropic
5570	s39	2018-11-24	2018-12-08	15	929	OUF anisotropic
5570	s40	2018-12-09	2018-12-23	15	1,012	OUF anisotropic
5570	s41	2018-12-24	2019-01-07	15	806	OUF anisotropic
5570	s42	2019-01-08	2019-01-22	15	961	OUF anisotropic
5570	s43	2019-01-23	2019-02-06	15	929	OUF anisotropic
5570	s44	2019-02-07	2019-02-21	15	1,223	OUF anisotropic
5570	s45	2019-02-22	2019-03-08	15	871	OUF anisotropic
5570	s46	2019-03-09	2019-03-23	15	807	OUF anisotropic
5570	s47	2019-03-24	2019-04-07	15	972	OUF anisotropic
5570	s48	2019-04-08	2019-04-22	15	732	OUF anisotropic
5570	s49	2019-04-23	2019-05-07	15	599	OUF
5570	s50	2019-05-08	2019-05-22	15	407	OUF anisotropic
5570	s51	2019-05-23	2019-06-06	15	456	OUF anisotropic
5570	s52	2019-06-07	2019-06-21	15	556	OUF anisotropic
5570	s53	2019-06-22	2019-07-06	15	606	OUF anisotropic
5570	s54	2019-07-07	2019-07-21	15	426	OUF anisotropic
5570	s55	2019-07-22	2019-08-05	15	390	OUF anisotropic
5570	s56	2019-08-06	2019-08-20	15	687	OUF anisotropic
5570	s57	2019-08-21	2019-09-04	15	671	OUF anisotropic
5573	s1*	2018-06-26	2018-06-30	5	419	OUF anisotropic
5574	s1*	2018-07-07	2018-07-12	6	178	OU anisotropic
6444	s1	2021-02-09	2021-02-23	15	281	OU anisotropic
6444	s2	2021-02-24	2021-03-10	15	463	OU anisotropic
6444	s3	2021-03-11	2021-03-25	15	465	OU anisotropic
6444	s4	2021-03-26	2021-04-09	15	265	OU anisotropic
6444	s5	2021-04-10	2021-04-24	15	306	Ouf anisotropic
6444	s6	2021-04-25	2021-05-09	15	298	OU anisotropic
6444	s7	2021-05-10	2021-05-24	15	12	OU anisotropic
6444	s8	2021-05-25	2021-06-08	15	83	IID anisotropic
6444	s10	2021-06-24	2021-07-08	15	1,185	OUF anisotropic
6444	s11	2021-07-09	2021-07-23	15	1,220	OUF anisotropic
6444	s12	2021-07-24	2021-08-07	15	2,071	OUF anisotropic
6444	s13	2021-08-08	2021-08-22	15	4,570	OUF anisotropic
6444	s14	2021-08-23	2021-09-06	15	4,800	OUF anisotropic
6444	s15	2021-09-07	2021-09-21	15	4,711	OUF anisotropic
6444	s16	2021-09-22	2021-10-06	15	4,705	OUF anisotropic
6444	s17	2021-10-07	2021-10-21	15	4,573	OUF anisotropic
6444	s18	2021-10-22	2021-11-05	15	4,391	OUF anisotropic
6444	s19	2021-11-06	2021-11-20	15	4,741	OUF anisotropic
6444	s20	2021-11-21	2021-12-05	15	4,735	OUF anisotropic
6444	s21	2021-12-06	2021-12-20	15	4,525	OUF anisotropic
6444	s22	2021-12-21	2022-01-04	15	4,398	OUF anisotropic

<b>ID</b>	<b>Subset</b>	<b>Beginning date</b>	<b>End date</b>	<b>Duration (days)</b>	<b>Number of points</b>	<b>Selected model</b>
6444	s23	2022-01-05	2022-01-19	15	4,797	OUF anisotropic
6444	s24	2022-01-20	2022-02-03	15	4,415	OUF anisotropic
6444	s25	2022-02-04	2022-02-18	15	4,127	OUF anisotropic
6444	s26	2022-02-19	2022-03-05	15	2,680	OUF anisotropic
6444	s27	2022-03-06	2022-03-20	15	4,369	OUF anisotropic
6444	s28	2022-03-21	2022-04-04	15	4,590	OUF anisotropic
6444	s29	2022-04-05	2022-04-19	15	4,314	OUF anisotropic
6444	s30	2022-04-20	2022-05-04	15	4,571	OUF anisotropic
6444	s31	2022-05-05	2022-05-19	15	4,516	OUF anisotropic
6444	s32	2022-05-20	2022-06-03	15	4,883	OUF anisotropic
6444	s33	2022-06-04	2022-06-18	15	4,660	OUF anisotropic
6444	s34	2022-06-19	2022-07-03	15	4,629	OUF anisotropic
6444	s35	2022-07-04	2022-07-18	15	4,313	OUF anisotropic
6444	s36	2022-07-19	2022-08-02	15	7,427	OUF anisotropic
6444	s37	2022-08-03	2022-08-17	15	9,244	OUF anisotropic
6444	s38	2022-08-18	2022-09-01	15	10,528	OUF anisotropic
6444	s39	2022-09-02	2022-09-16	15	11,892	OUF anisotropic
6444	s40	2022-09-17	2022-10-01	15	12,455	OUF anisotropic
6444	s41	2022-10-02	2022-10-16	15	12,535	OUF anisotropic
6444	s42	2022-10-17	2022-10-31	15	12,002	OUF anisotropic
6444	s43	2022-11-01	2022-11-15	15	10,855	OUF anisotropic
6444	s44	2022-11-16	2022-11-30	15	12,673	OUF anisotropic
6444	s45	2022-12-01	2022-12-15	15	10,742	OUF
6444	s46	2022-12-16	2022-12-30	15	11,157	OUF anisotropic
6444	s47	2022-12-31	2023-01-14	15	11,442	OUF anisotropic
6444	s48	2023-01-15	2023-01-29	15	11,044	OUF anisotropic
6444	s49	2023-01-30	2023-02-13	15	11,130	OUF anisotropic
6444	s50	2023-02-14	2023-02-28	15	9,784	OUF anisotropic
9015	s1	2021-02-09	2021-02-23	15	805	OU anisotropic
9015	s2	2021-02-24	2021-03-10	15	1,365	OU anisotropic
9024	s1*	2021-07-09	2021-07-21	13	1,154	OUF anisotropic
9026	s1	2022-04-29	2022-05-13	15	1,338	OUF anisotropic
9250	s1	2022-04-30	2022-05-14	15	2,043	OUF anisotropic
9250	s2*	2022-05-15	2022-05-20	6	313	OUF anisotropic
9252	s1*	2022-06-27	2022-06-30	4	696	OUF anisotropic
9025	s1	2021-07-09	2021-07-23	15	1,330	OUF anisotropic
9025	s2	2021-07-24	2021-08-07	15	1,333	OUF anisotropic
9025	s3	2021-08-08	2021-08-22	15	1,295	OUF anisotropic
9025	s4	2021-08-23	2021-09-06	15	1,276	OUF anisotropic
9025	s5	2021-09-07	2021-09-21	15	1,292	OUF anisotropic
9025	s6	2021-09-22	2021-10-06	15	1,301	OUF anisotropic
9025	s7	2021-10-07	2021-10-21	15	1,317	OUF anisotropic
9025	s8	2021-10-22	2021-11-05	15	1,302	OUF anisotropic
9025	s9	2021-11-06	2021-11-20	15	1,261	OUF anisotropic
9025	s10	2021-11-21	2021-12-05	15	1,215	OUF anisotropic
9025	s11	2021-12-06	2021-12-20	15	1,219	OUF anisotropic
9025	s12	2021-12-21	2022-01-04	15	1,277	OUF anisotropic



<b>ID</b>	<b>Subset</b>	<b>Beginning date</b>	<b>End date</b>	<b>Duration (days)</b>	<b>Number of points</b>	<b>Selected model</b>
9025	s13	2022-01-05	2022-01-19	15	1,277	OUF anisotropic
9025	s14	2022-01-20	2022-02-03	15	1,257	OUF anisotropic
9025	s15	2022-02-04	2022-02-18	15	1,253	OUF anisotropic
9025	s16	2022-02-19	2022-03-05	15	1,185	OUF anisotropic
9025	s17	2022-03-06	2022-03-20	15	1,231	OUF anisotropic
9025	s18	2022-03-21	2022-04-04	15	1,204	OUF anisotropic
9025	s19	2022-04-05	2022-04-19	15	1,223	OUF anisotropic
9025	s20	2022-04-20	2022-05-04	15	1,229	OUF anisotropic
9025	s21	2022-05-05	2022-05-19	15	1,214	OUF anisotropic
9025	s22	2022-05-20	2022-06-03	15	1,126	OUF anisotropic
9025	s23	2022-06-04	2022-06-18	15	934	OUF anisotropic
9025	s24	2022-06-19	2022-07-03	15	666	OUF anisotropic
9025	s25	2022-07-04	2022-07-18	15	625	OUF anisotropic
9025	s26	2022-07-19	2022-08-02	15	458	OUF anisotropic
9025	s27	2022-08-03	2022-08-17	15	359	OUF anisotropic
9025	s28	2022-08-18	2022-09-01	15	383	OUF anisotropic
9025	s34	2022-11-16	2022-11-30	15	345	OUF anisotropic
9025	s35	2022-12-01	2022-12-15	15	389	OUF anisotropic
9025	s36	2022-12-16	2022-12-30	15	327	OUF anisotropic
9025	s37	2022-12-31	2023-01-14	15	352	OUF anisotropic
9025	s38	2023-01-15	2023-01-29	15	403	OUF anisotropic
9025	s39	2023-01-30	2023-02-13	15	297	OUF anisotropic
9025	s40	2023-02-14	2023-02-28	15	283	OUF anisotropic
9025	s41	2023-03-01	2023-03-15	15	200	OU anisotropic
9025	s42	2023-03-16	2023-03-30	15	100	OUF anisotropic
9025	s43	2023-03-31	2023-04-14	15	92	OUF anisotropic
9025	s44	2023-04-15	2023-04-29	15	77	OUF anisotropic
9025	s45	2023-04-30	2023-05-14	15	138	OUF anisotropic
9025	s46	2023-05-15	2023-05-29	15	234	OUF anisotropic

SUPPLEMENTARY TABLE 3 Estimated home ranges and core areas (“Estimated 95%” and “Estimated 50%”, corresponding to the estimates, in km<sup>2</sup>, of the 95% and 50% AKDE areas, respectively), for each fortnightly subset of macaw movement data (“Subset”), with identification of the individual (“ID”), the tagging site (*Estação Biológica de Canudos* (EBC), *Barreiras* and *B. do Chico* are breeding sites located in the *Raso da Catarina* region, and BDO is the release area in the *Boqueirão da Onça* region), the year of the respective fortnightly subset, the Confidence Intervals (CI) of each estimation (“Lower CI 95%” and “Upper CI 95%”, and “Lower CI 50%” and “Upper CI 50%”, corresponding to the CI lower and upper values, in km<sup>2</sup>, of the 95% and 50% AKDE estimated areas, respectively), and the scaled home ranges and core areas (“Scaled est. 95%” and “Scaled est. 50%”, corresponding to the scaled variables “Estimated 95%” and “Estimated 50%”, respectively).

\* Fortnightly estimates calculated using subsets with less than 15 days of movement data.

ID	Tagging site	Year	Subset	Estimated 95%	Lower CI 95%	Upper CI 95%	Estimated 50%	Lower CI 50%	Upper CI 50%	Scale est. 95%	Scaled est. 50%
5568	EBC	2017	s1	1,328.43	770.60	2,035.67	321.29	186.38	492.35	0.17	0.30
5568	EBC	2017	s2	1,312.42	897.17	1,805.40	342.39	234.06	471.00	0.15	0.39
5568	EBC	2017	s3	2,064.92	1,336.92	2,948.64	593.45	384.23	847.43	1.04	1.57
5568	EBC	2017	s4	673.26	457.95	929.41	157.39	107.05	217.26	-0.60	-0.47
5568	EBC	2017	s5	1,377.83	879.67	1,985.97	281.19	179.53	405.30	0.23	0.11
5568	EBC	2017	s6	960.46	631.39	1,357.46	169.41	111.36	239.43	-0.26	-0.41
5568	EBC	2017	s7	1,235.04	833.57	1,714.18	226.84	153.10	314.84	0.06	-0.14
5568	EBC	2017	s8	2,327.02	1,474.03	3,371.63	388.47	246.07	562.86	1.35	0.61
5568	EBC	2017	s9	2,172.70	1,360.77	3,171.53	613.00	383.93	894.81	1.17	1.66
5568	EBC	2017	s10	2,310.95	1,493.58	3,303.86	698.33	451.33	998.36	1.33	2.06
5568	EBC	2017	s11	838.61	578.89	1,145.70	164.84	113.79	225.20	-0.41	-0.43
5568	EBC	2017	s12	513.49	338.36	724.56	117.04	77.12	165.15	-0.79	-0.66
5568	EBC	2017	s13	834.95	568.37	1,151.98	162.81	110.83	224.63	-0.41	-0.44
5568	EBC	2017	s14	432.09	311.98	571.45	112.74	81.40	149.10	-0.89	-0.68
5568	EBC	2017	s15	714.43	482.91	990.57	151.69	102.54	210.33	-0.55	-0.49
5568	EBC	2017	s16	856.43	486.00	1,330.08	149.22	84.68	231.75	-0.39	-0.51
5568	EBC	2018	s17	615.68	416.33	853.41	123.55	83.54	171.25	-0.67	-0.63
5568	EBC	2018	s18	2,064.33	1,065.47	3,387.99	484.31	249.97	794.85	1.04	1.06
5568	EBC	2018	s19	1,598.39	1,029.02	2,291.14	311.70	200.67	446.79	0.49	0.25
5568	EBC	2018	s20	888.47	535.13	1,329.94	195.80	117.93	293.10	-0.35	-0.29
5568	EBC	2018	s21	114.91	77.14	160.09	17.37	11.66	24.19	-1.26	-1.12

<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
5568	EBC	2018	s22	1,178.40	699.83	1,779.61	165.79	98.46	250.38	-0.01	-0.43
5568	EBC	2018	s23	928.05	605.27	1,318.70	111.91	72.99	159.01	-0.30	-0.68
5568	EBC	2018	s24	1,749.35	927.45	2,827.70	342.22	181.44	553.18	0.67	0.39
5568	EBC	2018	s25	403.92	226.39	631.99	83.47	46.78	130.60	-0.92	-0.81
5568	EBC	2018	s26	1,716.65	962.48	2,685.43	363.45	203.77	568.56	0.63	0.49
5568	EBC	2018	s27	991.56	586.08	1,501.92	201.19	118.92	304.75	-0.23	-0.26
5568	EBC	2018	s28	1,749.45	1,200.53	2,400.09	383.69	263.30	526.39	0.67	0.59
5568	EBC	2018	s29	1,495.68	1,065.28	1,997.96	426.13	303.51	569.24	0.37	0.79
5568	EBC	2018	s30	1,342.91	927.71	1,833.67	323.28	223.33	441.42	0.19	0.31
5568	EBC	2018	s31	1,581.81	1,129.96	2,108.47	380.36	271.71	507.00	0.47	0.57
5568	EBC	2018	s32	4,763.80	2,443.15	7,846.32	1,116.04	572.37	1,838.20	4.22	4.00
5568	EBC	2018	s33	219.26	146.40	306.61	39.97	26.69	55.90	-1.14	-1.02
5568	EBC	2018	s34	748.20	466.30	1,095.68	139.43	86.90	204.18	-0.52	-0.55
5568	EBC	2018	s35	980.70	678.23	1,338.07	186.73	129.14	254.78	-0.24	-0.33
5568	EBC	2018	s36	611.51	461.57	782.23	127.12	95.95	162.61	-0.68	-0.61
5568	EBC	2018	s37	374.03	279.74	481.80	100.75	75.35	129.78	-0.96	-0.73
5568	EBC	2018	s38	581.25	430.90	753.75	143.19	106.15	185.69	-0.71	-0.53
5568	EBC	2018	s39	911.53	576.96	1,321.38	211.78	134.05	307.00	-0.32	-0.21
5568	EBC	2018	s40	270.39	207.41	341.60	45.90	35.21	57.99	-1.08	-0.99
5568	EBC	2018	s41	190.36	132.89	258.00	32.82	22.92	44.49	-1.17	-1.05
5568	EBC	2019	s42	241.47	159.65	339.94	45.97	30.39	64.72	-1.11	-0.99
5568	EBC	2019	s43	1,130.56	573.73	1,873.08	187.84	95.32	311.20	-0.06	-0.33
5568	EBC	2019	s44	1,356.97	736.17	2,164.46	206.34	111.94	329.12	0.20	-0.24
5568	EBC	2019	s45	2,350.45	1,395.74	3,549.91	424.69	252.19	641.41	1.37	0.78
5568	EBC	2019	s46	258.36	165.79	371.14	57.67	37.01	82.85	-1.09	-0.93
5568	EBC	2019	s47	1,663.29	922.00	2,619.66	278.70	154.49	438.94	0.56	0.10
5568	EBC	2019	s48	331.80	200.44	495.72	72.32	43.69	108.06	-1.01	-0.87
5568	EBC	2019	s49	721.86	436.33	1,078.09	157.16	95.00	234.71	-0.55	-0.47
5568	EBC	2019	s50	387.41	252.79	550.31	59.60	38.89	84.66	-0.94	-0.92
5568	EBC	2019	s51	488.40	293.97	731.37	90.57	54.51	135.63	-0.82	-0.78

<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
5568	EBC	2019	s52	2,379.73	1,469.18	3,506.31	518.55	320.14	764.04	1.41	1.22
5568	EBC	2019	s53	2,918.01	1,881.05	4,178.98	838.51	540.53	1,200.86	2.04	2.71
5568	EBC	2019	s54	1,730.42	1,105.75	2,492.71	272.01	173.82	391.84	0.64	0.07
5570	EBC	2017	s1	134.37	97.89	176.54	28.07	20.45	36.88	-0.85	-0.82
5570	EBC	2017	s2	255.99	191.57	329.60	52.78	39.50	67.95	-0.79	-0.77
5570	EBC	2017	s3	1,798.56	1,264.20	2,425.63	438.02	307.88	590.74	0.03	0.03
5570	EBC	2017	s4	631.31	453.55	838.00	137.02	98.44	181.88	-0.59	-0.60
5570	EBC	2017	s5	670.48	474.62	899.65	154.31	109.23	207.05	-0.57	-0.56
5570	EBC	2017	s6	1,407.21	1,026.55	1,846.98	310.97	226.85	408.16	-0.18	-0.24
5570	EBC	2017	s7	2,312.61	1,693.33	3,026.92	608.97	445.89	797.06	0.30	0.38
5570	EBC	2017	s8	2,306.96	1,635.56	3,092.01	589.41	417.88	789.99	0.30	0.34
5570	EBC	2017	s9	1,217.40	859.56	1,636.54	309.38	218.44	415.90	-0.28	-0.24
5570	EBC	2017	s10	1,138.71	812.32	1,519.36	228.99	163.35	305.54	-0.32	-0.41
5570	EBC	2017	s11	3,534.32	2,508.95	4,732.62	1,107.22	785.99	1,482.62	0.94	1.42
5570	EBC	2017	s12	1,676.79	1,202.45	2,228.77	504.43	361.74	670.49	-0.04	0.17
5570	EBC	2017	s13	1,536.62	1,111.61	2,029.35	303.87	219.83	401.31	-0.11	-0.25
5570	EBC	2017	s14	1,504.35	1,059.69	2,025.69	450.91	317.63	607.17	-0.13	0.06
5570	EBC	2017	s15	2,350.19	1,623.92	3,208.58	573.62	396.36	783.13	0.32	0.31
5570	EBC	2017	s16	529.87	398.69	679.42	135.85	102.22	174.18	-0.64	-0.60
5570	EBC	2018	s17	496.57	369.04	642.74	130.47	96.97	168.88	-0.66	-0.61
5570	EBC	2018	s18	2,618.85	1,418.99	4,180.26	594.68	322.22	949.24	0.46	0.35
5570	EBC	2018	s19	126.92	91.57	167.95	35.30	25.46	46.71	-0.86	-0.81
5570	EBC	2018	s20	39.99	30.50	50.75	8.80	6.71	11.17	-0.90	-0.86
5570	EBC	2018	s21	59.74	45.55	75.82	14.85	11.32	18.85	-0.89	-0.85
5570	EBC	2018	s22	40.63	31.93	50.35	7.87	6.18	9.75	-0.90	-0.87
5570	EBC	2018	s23	57.88	44.79	72.63	13.47	10.42	16.90	-0.89	-0.85
5570	EBC	2018	s24	81.67	62.70	103.11	22.62	17.37	28.56	-0.88	-0.83
5570	EBC	2018	s25	90.61	69.30	114.73	17.45	13.34	22.09	-0.88	-0.85
5570	EBC	2018	s26	2,615.37	1,195.75	4,581.04	654.59	299.28	1,146.56	0.46	0.48
5570	EBC	2018	s27	2,055.66	1,027.86	3,433.67	392.43	196.22	655.49	0.16	-0.07

<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
5570	EBC	2018	s28	1,654.93	1,181.90	2,206.33	345.92	247.05	461.18	-0.05	-0.16
5570	EBC	2018	s29	3,321.72	2,371.35	4,429.73	933.25	666.24	1,244.56	0.83	1.06
5570	EBC	2018	s30	1,005.53	687.44	1,383.15	263.93	180.44	363.05	-0.39	-0.33
5570	EBC	2018	s31	4,297.62	2,745.88	6,191.34	1,096.99	700.90	1,580.37	1.35	1.40
5570	EBC	2018	s32	3,586.92	2,598.26	4,732.49	776.98	562.82	1,025.13	0.97	0.73
5570	EBC	2018	s33	3,172.88	2,168.61	4,365.24	772.84	528.22	1,063.26	0.75	0.72
5570	EBC	2018	s34	2,018.25	1,480.08	2,638.57	418.18	306.68	546.72	0.14	-0.01
5570	EBC	2018	s35	2,681.82	1,962.44	3,511.80	683.10	499.86	894.51	0.49	0.54
5570	EBC	2018	s36	4,161.61	2,937.44	5,595.68	734.93	518.74	988.18	1.28	0.65
5570	EBC	2018	s37	4,490.48	2,454.93	7,130.45	965.99	528.10	1,533.90	1.45	1.13
5570	EBC	2018	s38	328.07	235.77	435.37	87.82	63.12	116.55	-0.75	-0.70
5570	EBC	2018	s39	246.68	176.60	328.30	67.69	48.46	90.08	-0.79	-0.74
5570	EBC	2018	s40	337.58	250.86	436.97	86.95	64.61	112.54	-0.75	-0.70
5570	EBC	2018	s41	157.30	115.84	205.01	33.87	24.94	44.14	-0.84	-0.81
5570	EBC	2019	s42	49.73	37.25	63.99	13.76	10.31	17.71	-0.90	-0.85
5570	EBC	2019	s43	1,032.62	583.79	1,607.35	148.90	84.18	231.78	-0.38	-0.57
5570	EBC	2019	s44	2,757.20	1,282.01	4,787.65	677.42	314.98	1,176.28	0.53	0.53
5570	EBC	2019	s45	1,957.72	552.99	4,234.12	371.16	104.84	802.75	0.11	-0.11
5570	EBC	2019	s46	831.11	130.77	2,165.59	187.04	29.43	487.36	-0.48	-0.49
5570	EBC	2019	s47	595.98	355.40	897.74	87.27	52.04	131.46	-0.61	-0.70
5570	EBC	2019	s48	114.66	86.87	146.26	24.96	18.91	31.84	-0.86	-0.83
5570	EBC	2019	s49	173.86	130.90	222.82	45.16	34.00	57.88	-0.83	-0.79
5570	EBC	2019	s50	258.39	162.48	376.19	58.18	36.58	84.70	-0.79	-0.76
5570	EBC	2019	s51	3,346.09	1,278.00	6,392.06	769.96	294.08	1,470.86	0.85	0.72
5570	EBC	2019	s52	8,549.49	3,684.12	15,427.60	2,007.25	864.96	3,622.09	3.60	3.29
5570	EBC	2019	s53	7,801.55	3,035.11	14,778.71	2,132.02	829.44	4,038.75	3.20	3.55
5570	EBC	2019	s54	111.69	83.13	144.41	20.55	15.29	26.57	-0.87	-0.84
5570	EBC	2019	s55	6,554.80	3,617.99	10,349.86	1,793.80	990.10	2,832.36	2.54	2.85
5570	EBC	2019	s56	1,164.52	845.41	1,533.93	320.68	232.81	422.41	-0.31	-0.22
5570	EBC	2019	s57	1,589.94	1,157.05	2,090.55	431.92	314.32	567.91	-0.08	0.02

<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
5573*	Barreiras	2018	s1	193.59	87.10	341.89	33.76	15.19	59.62	-	-
5574*	B. do Chico	2018	s1	156.57	96.88	230.36	43.13	26.69	63.45	-	-
6444	BDO	2021	s1	39.03	30.15	49.04	9.78	7.55	12.28	0.85	-0.17
6444	BDO	2021	s2	5.34	4.73	5.99	1.12	0.99	1.26	-0.64	-0.64
6444	BDO	2021	s3	6.43	5.65	7.26	1.31	1.15	1.48	-0.59	-0.63
6444	BDO	2021	s4	9.11	7.76	10.57	1.79	1.52	2.07	-0.47	-0.61
6444	BDO	2021	s5	7.22	6.34	8.16	1.13	0.99	1.28	-0.55	-0.64
6444	BDO	2021	s6	14.04	12.30	15.89	1.90	1.66	2.15	-0.25	-0.60
6444	BDO	2021	s7	1.37	0.10	4.23	30.34	2.26	93.85	-0.81	0.96
6444	BDO	2021	s8	45.15	35.91	55.43	6.10	4.85	7.49	1.11	-0.37
6444	BDO	2021	s10	1.70	1.49	1.93	17.50	15.33	19.82	-0.80	0.25
6444	BDO	2021	s11	2.98	2.46	3.56	54.95	45.26	65.57	-0.74	2.30
6444	BDO	2021	s12	4.44	3.89	5.02	61.70	54.08	69.82	-0.68	2.67
6444	BDO	2021	s13	17.47	15.39	19.68	1.67	1.47	1.88	-0.10	-0.61
6444	BDO	2021	s14	26.99	23.57	30.64	1.95	1.70	2.21	0.32	-0.60
6444	BDO	2021	s15	5.21	4.63	5.82	47.00	41.80	52.51	-0.64	1.87
6444	BDO	2021	s16	6.24	5.54	6.98	62.43	55.45	69.81	-0.60	2.71
6444	BDO	2021	s17	46.12	38.29	54.66	6.86	5.70	8.13	1.16	-0.33
6444	BDO	2021	s18	12.87	10.11	15.95	2.62	2.06	3.24	-0.30	-0.56
6444	BDO	2021	s19	9.62	7.72	11.73	1.77	1.42	2.16	-0.45	-0.61
6444	BDO	2021	s20	2.68	2.28	3.11	34.43	29.28	39.99	-0.75	1.18
6444	BDO	2021	s21	25.66	19.59	32.53	4.12	3.15	5.23	0.26	-0.48
6444	BDO	2021	s22	1.78	1.51	2.08	25.08	21.26	29.20	-0.79	0.67
6444	BDO	2022	s23	18.73	17.03	20.51	3.95	3.59	4.33	-0.05	-0.49
6444	BDO	2022	s24	1.25	1.03	1.48	20.35	16.86	24.17	-0.82	0.41
6444	BDO	2022	s25	2.84	2.27	3.48	43.57	34.84	53.26	-0.75	1.68
6444	BDO	2022	s26	27.58	24.27	31.10	4.21	3.70	4.75	0.34	-0.47
6444	BDO	2022	s27	2.62	2.14	3.15	37.87	30.88	45.56	-0.76	1.37
6444	BDO	2022	s28	14.75	13.39	16.18	3.19	2.90	3.50	-0.22	-0.53
6444	BDO	2022	s29	3.79	2.93	4.76	60.77	47.01	76.27	-0.70	2.62

<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
6444	BDO	2022	s30	21.53	17.88	25.52	4.09	3.40	4.85	0.08	-0.48
6444	BDO	2022	s31	12.89	10.61	15.40	2.72	2.23	3.24	-0.30	-0.56
6444	BDO	2022	s32	15.42	12.40	18.76	3.37	2.71	4.10	-0.19	-0.52
6444	BDO	2022	s33	14.07	11.81	16.52	2.43	2.04	2.85	-0.25	-0.57
6444	BDO	2022	s34	10.89	9.42	12.47	2.10	1.82	2.41	-0.39	-0.59
6444	BDO	2022	s35	16.05	13.65	18.63	3.18	2.70	3.69	-0.16	-0.53
6444	BDO	2022	s36	19.63	15.29	24.51	3.47	2.70	4.33	-0.01	-0.52
6444	BDO	2022	s37	23.00	18.68	27.76	4.59	3.73	5.54	0.14	-0.45
6444	BDO	2022	s38	24.03	20.69	27.61	4.03	3.47	4.63	0.19	-0.48
6444	BDO	2022	s39	35.70	27.59	44.83	7.56	5.84	9.49	0.70	-0.29
6444	BDO	2022	s40	147.28	112.74	186.36	13.52	10.35	17.11	5.60	0.04
6444	BDO	2022	s41	37.93	30.94	45.63	2.79	2.27	3.35	0.80	-0.55
6444	BDO	2022	s42	15.22	12.42	18.29	1.68	1.37	2.02	-0.20	-0.61
6444	BDO	2022	s43	34.99	29.58	40.85	4.69	3.97	5.48	0.67	-0.45
6444	BDO	2022	s44	46.20	37.70	55.55	4.04	3.30	4.86	1.16	-0.48
6444	BDO	2022	s45	18.41	14.78	22.43	2.55	2.05	3.11	-0.06	-0.57
6444	BDO	2022	s46	27.53	22.29	33.31	3.67	2.97	4.44	0.34	-0.50
6444	BDO	2023	s47	10.07	8.49	11.79	1.80	1.51	2.10	-0.43	-0.61
6444	BDO	2023	s48	21.59	20.43	22.78	1.33	1.26	1.40	0.08	-0.63
6444	BDO	2023	s49	42.16	37.17	47.45	6.03	5.31	6.79	0.98	-0.37
6444	BDO	2023	s50	12.31	9.62	15.33	1.87	1.46	2.32	-0.33	-0.60
9015	BDO	2021	s1	94.11	74.20	116.35	17.40	13.72	21.52	1.15	1.15
9015	BDO	2021	s2	9.43	8.74	10.16	1.57	1.45	1.69	-0.61	-0.57
9024*	B. do Chico	2021	s1	120.08	87.60	157.60	23.32	17.01	30.60	-	-
9026	Barreiras	2022	s1	87.67	65.77	112.66	24.23	18.18	31.14	-	-
9250	Barreiras	2022	s1	376.08	252.92	523.30	77.40	52.05	107.70	0.71	0.71
9250*	Barreiras	2022	s2	76.37	50.67	107.25	14.98	9.94	21.04	-0.71	-0.71
9252*	Barreiras	2022	s1	142.05	46.25	290.64	33.26	10.83	68.05	-	-
9025	B. do Chico	2021	s1	131.85	97.26	171.62	27.47	20.27	35.76	-0.54	-0.42
9025	B. do Chico	2021	s2	147.84	110.33	190.75	36.04	26.90	46.50	-0.48	-0.27

<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
9025	B. do Chico	2021	s3	114.90	85.78	148.20	20.49	15.30	26.43	-0.61	-0.55
9025	B. do Chico	2021	s4	101.28	75.56	130.71	16.94	12.64	21.86	-0.67	-0.61
9025	B. do Chico	2021	s5	133.82	101.28	170.81	28.34	21.45	36.18	-0.54	-0.41
9025	B. do Chico	2021	s6	156.64	117.78	200.94	30.18	22.70	38.72	-0.44	-0.38
9025	B. do Chico	2021	s7	123.71	93.61	157.94	20.76	15.71	26.50	-0.58	-0.55
9025	B. do Chico	2021	s8	150.65	113.38	193.13	35.77	26.92	45.86	-0.47	-0.27
9025	B. do Chico	2021	s9	114.13	85.37	147.00	19.79	14.80	25.48	-0.62	-0.56
9025	B. do Chico	2021	s10	169.97	128.16	217.59	29.14	21.97	37.31	-0.39	-0.39
9025	B. do Chico	2021	s11	94.21	70.34	121.51	16.68	12.46	21.52	-0.70	-0.62
9025	B. do Chico	2021	s12	80.35	59.11	104.81	18.33	13.48	23.91	-0.75	-0.59
9025	B. do Chico	2022	s13	119.95	88.87	155.62	20.47	15.17	26.56	-0.59	-0.55
9025	B. do Chico	2022	s14	108.13	80.79	139.40	14.18	10.59	18.27	-0.64	-0.66
9025	B. do Chico	2022	s15	128.18	99.99	159.83	15.07	11.76	18.79	-0.56	-0.65
9025	B. do Chico	2022	s16	284.62	203.37	379.30	38.51	27.52	51.33	0.08	-0.22
9025	B. do Chico	2022	s17	223.58	163.03	293.54	26.51	19.33	34.81	-0.17	-0.44
9025	B. do Chico	2022	s18	464.88	335.30	615.29	79.44	57.30	105.14	0.82	0.52
9025	B. do Chico	2022	s19	291.77	204.56	394.21	49.22	34.51	66.50	0.11	-0.03
9025	B. do Chico	2022	s20	180.90	138.21	229.24	30.16	23.04	38.22	-0.34	-0.38
9025	B. do Chico	2022	s21	237.34	178.90	303.93	45.79	34.52	58.64	-0.11	-0.09
9025	B. do Chico	2022	s22	872.84	626.91	1,158.83	219.53	157.67	291.46	2.48	3.05
9025	B. do Chico	2022	s23	301.38	224.55	389.34	77.25	57.56	99.80	0.15	0.48
9025	B. do Chico	2022	s24	303.73	225.86	392.96	45.72	34.00	59.15	0.16	-0.09
9025	B. do Chico	2022	s25	397.24	290.75	520.10	59.02	43.20	77.27	0.54	0.15
9025	B. do Chico	2022	s26	835.29	570.13	1,150.29	213.00	145.39	293.33	2.33	2.93
9025	B. do Chico	2022	s27	525.45	384.15	688.53	92.27	67.46	120.90	1.06	0.75
9025	B. do Chico	2022	s28	558.66	393.57	752.21	89.03	62.72	119.88	1.20	0.69
9025	B. do Chico	2022	s34	606.44	336.86	953.96	161.22	89.56	253.61	1.39	2.00
9025	B. do Chico	2022	s35	67.04	51.57	84.50	15.44	11.88	19.46	-0.81	-0.64
9025	B. do Chico	2022	s36	114.84	87.71	145.58	19.94	15.23	25.28	-0.61	-0.56
9025	B. do Chico	2023	s37	134.91	103.18	170.81	22.63	17.31	28.65	-0.53	-0.51



<b>ID</b>	<b>Tagging site</b>	<b>Year</b>	<b>Subset</b>	<b>Estimated 95%</b>	<b>Lower CI 95%</b>	<b>Upper CI 95%</b>	<b>Estimated 50%</b>	<b>Lower CI 50%</b>	<b>Upper CI 50%</b>	<b>Scale est. 95%</b>	<b>Scaled est. 50%</b>
9025	B. do Chico	2023	s38	98.08	74.12	125.34	16.04	12.12	20.50	-0.68	-0.63
9025	B. do Chico	2023	s39	511.70	335.98	723.81	95.98	63.02	135.76	1.01	0.81
9025	B. do Chico	2023	s40	48.16	36.47	61.46	5.69	4.31	7.26	-0.89	-0.82
9025	B. do Chico	2023	s41	12.66	9.68	16.02	1.67	1.28	2.11	-1.03	-0.89
9025	B. do Chico	2023	s42	51.10	34.12	71.45	6.20	4.14	8.67	-0.87	-0.81
9025	B. do Chico	2023	s43	31.15	22.89	40.67	3.83	2.82	5.00	-0.95	-0.85
9025	B. do Chico	2023	s44	151.49	104.66	206.84	17.88	12.36	24.42	-0.46	-0.60
9025	B. do Chico	2023	s45	833.69	577.54	1,136.12	142.66	98.83	194.41	2.32	1.66
9025	B. do Chico	2023	s46	855.17	585.85	1,174.63	163.83	112.24	225.04	2.41	2.04

### SUPPLEMENTARY MATERIAL 3 Details on home range and core area estimation.

Using the ctm framework (Calabrese et al., 2016), we compared the fit of movement models (Independent Identically Distributed (IID), Ornstein-Uhlenbeck (OU) and Ornstein-Uhlenbeck Foraging (OUF)) to our data using the autocorrelation estimation method. Once the range residency was determined – required by the ctm package to calculate the AKDE ranges (Isted et al., 2023) –, the best model was selected (via Akaike Information Criterion – AIC) and applied to fit the function of the AKDE to estimate ranges' size (Calabrese et al., 2016). We calculated the home ranges using 95% AKDE (corresponding to the individual's mobility potential in the total area); and the core areas, using 50% AKDE (corresponding to the areas that the animal uses most frequently; Supplementary Table 3) (Fleming et al., 2015; Redpath et al., 2023).

Due to the large variation of home ranges' and core areas' estimates among individuals, instead of using the absolute values of home range and core areas estimates in the model analysis, we scaled them using “scale” function in base R software (R Core Team, 2023; the centralization argument used for the variables was the average of each individuals' estimates; Supplementary Table 4). The scaling procedure allowed fitting of models with data from multiple macaws simultaneously – comparatively observing the variation in fortnightly estimates of ranges over time to evaluate the influence of the predictor variables when adjusting the models.

### References

REDPATH, S.H.A., MARKS, N.J., MENZIES, F.D., O'HAGAN, M.J.H., WILSON, R.P., SMITH, S. ET AL. (2023) Impact of test, vaccinate and remove protocol on home ranges and nightly movements of badgers in a medium density population. *Scientific Reports*, 13, 2592.

SUPPLEMENTARY TABLE 4 Updated summary with identification of the tagged macaws for which we retrieved movement data, the tagging site (RASO = breeding sites in *Raso da Catarina* region, BDO = release area in *Boqueirão da Onça* region), the tagging date (year-month-day format), dates of the first and the last data download in the field, the total number of days (“Duration”) for which there is movement data recorded for each macaw, and the total number of points recorded for each individual.

<b>ID</b>	<b>Tagging site</b>	<b>Tagging date</b>	<b>1st download</b>	<b>Last download</b>	<b>Duration (days)</b>	<b>N° of points</b>
5568	RASO	2017-04-24	2017-05-03	2019-07-26	815	73,811
5570	RASO	2017-04-26	2017-05-03	2019-09-07	858	55,308
5573	RASO	2018-06-15	2018-06-26	2018-06-30	5	419
5574	RASO	2018-06-25	2018-07-07	2018-07-12	6	178
9015	BDO	2021-02-09	2021-02-09	2021-03-22	42	3,194
6444	BDO	2021-02-09	2021-02-09	2024-10-20	1,346	668,473
9024	RASO	2021-06-29	2021-07-09	2021-07-21	13	1,154
9025	RASO	2021-06-29	2021-07-09	2023-05-29	690	34,279
9026	RASO	2022-04-19	2022-04-29	2022-05-20	22	1,893
9250	RASO	2022-04-25	2022-04-30	2022-05-20	21	2,356
9252	RASO	2022-06-17	2022-06-27	2022-06-30	4	696

SUPPLEMENTARY MATERIAL 4 Details on environmental predictors and data processing, indicating how each variable was prepared for the home range and core area size temporal variation analysis.

We evaluated intra-annual and interannual variation in home range and core area size of tagged juvenile Lear's macaws using linear models (bbmle R package; Rencher & Schaalje, 2008; R Core Team, 2023). We investigated whether the predictor variables “tagging site” (communal roost site where the individual was born or released and tagged), “season”, “ordinal date”, “accumulated rainfall over the last six months”, “accumulated rainfall over the last twelve months”, “NDVI data relative to the real coordinates”, and “NDVI data relative to the random coordinates” influenced the temporal variation in size of the ranges estimated for the macaws.

For seasonal evaluation, considering that most of rainfall in Caatinga is concentrated in few consecutive months, and the rainy season is centered upon March, April and May, we considered the months from January to May as the “wet season”, and the “dry season”, from June to December (Tabarelli et al., 2003; da Silva, 2004; Silva et al., 2017). The “ordinal date” was also incorporated into the analysis as a predictor variable representing the seasonality throughout the year. In this analysis, the reference date for start counting the ordinal days (i.e., the ordinal date number 1) of a given subset of movement data always corresponded to January 1<sup>st</sup> of the year of that subset. Therefore, the count of consecutive days was restarted every new year – and the lower the count of the ordinal days, the closer the movement data subset was to the beginning of that year.

We also assessed if the licuri palm (*S. coronata*) fruit availability was related to temporal variation of the size of estimated ranges. The licuri palm fruit availability (therefore, the environmental productivity and resource availability) was inferred from climate remote sensing data (rainfall precipitation) and vegetation data (primary productivity index, more specifically, the Normalized Difference Vegetation Index – NDVI). The regional vegetation dynamics in semi-arid is strongly correlated with rainfall, and the NDVI, by measuring vegetation productivity/availability, is highly correlated to precipitation in the Caatinga – with the seasonal NDVI oscillation being related to the seasonal distribution of dry and wet periods, and also fluctuating according to the year rainfall (Pettorelli et al., 2005; Schucknecht et al., 2013; Barbosa et al., 2015; Silva et al., 2017).

Historical daily rainfall data series were extracted from meteorological stations within species' occurrence area from the Hidroweb platform (v3.2.7), managed by the Brazilian National Water and Basic Sanitation Agency (ANA). Aiming to investigate whether the movement of the macaws – and, therefore, the size variation of their ranges – would respond to changes in licuri palm fruit availability, the precipitation data was grouped into time series of accumulated rainfall (of both six and twelve months prior to start dates of each fortnightly movement data subset; supporting information in Supplementary Table 5). The temporal scales of rainfall were selected based on reproductive phenology of licuri palm and, therefore, on the estimated time for fruit development, ripening and availability. The *S. coronata* inflorescence requires five to ten months to fully develop. After this period, and after the time needed to mature the male and female flowers (ca. 40 days), the palm requires approximately another two months for fruits to ripen (Barbosa et al., 2021).

The NDVI was obtained from Movebank Environmental Data Automated Track Annotation System (Env-DATA) for the animals' tracking data (Dodge et al., 2013; Kays et al., 2022). We used NASA product MODIS Land Vegetation Indices MOD13Q1 V6.1 with spatial resolution/granularity = 250 m, and temporal resolution/granularity = 16-day period. To

incorporate the NDVI data into the linear models, the median of the NDVI values was calculated for each fortnightly range estimated (Supplementary Table 5).

Through the inverse distance weighted interpolation method, recommended in case of observation-driven variables, such as MODIS products (Dodge et al., 2013), Movebank annotated the tracking data by calculating estimated values of the variable for the location and time of each animal location – using values provided by the NDVI dataset for specific locations and times (Dodge et al., 2013; Kays et al., 2022). Considering that NDVI varies over time, each location was associated with the specific NDVI value corresponding to that location at the closest date (Viana et al., 2018).

Furthermore, aiming to understand whether the general productivity of the landscape would also influence on macaws' range sizes temporal variation, we included in the analysis, as another predictor variable, the median of NDVI values calculated for randomized geographic coordinates in the landscape – located within the bounding boxes of the estimated individual AKDE ranges, calculated in QGIS software (QGIS Development Team, 2023). The number of randomized geographic coordinates generated was the same as the real coordinates recorded for each individual, during its entire period of monitoring. The random NDVI data worked as a proxy for the availability of licuri in the environment. The random coordinates were also linked to the NDVI dataset by the Env-DATA (Dodge et al., 2013; Kays et al., 2022).

## References

- BARBOSA, C.M., MAIA, A.C.D., MARTEL, C., REGUEIRA, J.C.S., NAVARRO, D.M. DO A.F., RAGUSO, R.A., ET AL. (2021) Reproductive biology of *Syagrus coronata* (Arecaceae): sex-biased insect visitation and the unusual case of scent emission by peduncular bracts. *Plant Biology*, 23, 100–110.
- BARBOSA, H.A., LAKSHMI KUMAR, T.V., SILVA, L.R.M. (2015) Recent trends in vegetation dynamics in the South America and their relationship to rainfall. *Natural Hazards*, 77, 883–899.
- DODGE, S., BOHRER, G., WEINZIERL, R., DAVIDSON, S.C., KAYS, R., DOUGLAS, D., ET AL. (2013) The environmental-data automated track annotation (Env-DATA) system: linking animal tracks with environmental data. *Movement Ecology*, 1, 3.
- SCHUCKNECHT, A., ERASMI, S., NIEMEYER, I. & MATSCHULLAT, J. (2013) Assessing vegetation variability and trends in north-eastern Brazil using AVHRR and MODIS NDVI time series. *European Journal of Remote Sensing*, 46, 40–59.
- DA SILVA, V. DE P.R. (2004) On climate variability in Northeast of Brazil. *Journal of Arid Environments*, 58, 575–596.
- TABARELLI, M., VICENTE, A. & BARBOSA, D. C. A. (2003) Variation of seed dispersal spectrum of woody plants across a rainfall gradient in north-eastern Brazil. *Journal of Arid Environments*, 53, 197–210.

SUPPLEMENTARY TABLE 5 Rainfall data and Normalized Difference Vegetation Index (NDVI) data for each fortnightly subset of macaw movement data, with identification of the individual (“ID”), the fortnightly subset of macaw movement data (“Subset”), the month, season and ordinal date of the respective fortnightly subset, the identification of the rainfall station from which the rainfall data were extracted, the accumulated rainfall (in millimeters) for six (“Accum. rainfall 6”) and 12 months (“Accum. rainfall 12”), and the median of the NDVI data associated with the real coordinates (“Real NDVI median”) recorded for the macaws and the median of the NDVI data associated with the generated random coordinates (“Random NDVI median”).

<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
5568	s1	may	wet	122	Alto Redondo	114.4	193.4	0.37	0.41
5568	s2	may	wet	137	Alto Redondo	113.6	171.9	0.39	0.47
5568	s3	june	dry	152	Alto Redondo	149.3	201.6	0.42	0.49
5568	s4	june	dry	167	Alto Redondo	133.3	200.7	0.35	0.44
5568	s5	july	dry	182	Alto Redondo	144.9	203.5	0.47	0.50
5568	s6	july	dry	197	Alto Redondo	142.7	208.7	0.47	0.53
5568	s7	august	dry	212	Alto Redondo	148.1	211.9	0.40	0.45
5568	s8	august	dry	227	Alto Redondo	148.1	211.2	0.44	0.37
5568	s9	september	dry	242	Alto Redondo	148.1	208.9	0.42	0.40
5568	s10	september	dry	257	Alto Redondo	170.8	237.2	0.56	0.44
5568	s11	october	dry	272	Alto Redondo	121.7	238.7	0.52	0.63
5568	s12	october	dry	287	Alto Redondo	113.7	226.1	0.49	0.52
5568	s13	november	dry	302	Alto Redondo	111.7	226.1	0.38	0.37
5568	s14	november	dry	317	Alto Redondo	121.6	235.2	0.36	0.38
5568	s15	december	dry	332	Alto Redondo	85.9	235.2	0.33	0.35
5568	s16	december	dry	347	Alto Redondo	95	257.6	0.26	0.32
5568	s17	january	wet	362	Alto Redondo	83.4	228.3	0.30	0.36
5568	s18	january	wet	12	Alto Redondo	84.8	227.5	0.30	0.35
5568	s19	february	wet	27	Alto Redondo	85.7	233.8	0.26	0.33
5568	s20	february	wet	42	Alto Redondo	108.2	256.3	0.29	0.35
5568	s21	march	wet	57	Alto Redondo	179.2	327.3	0.36	0.45
5568	s22	march	wet	72	Alto Redondo	166.3	345.8	0.39	0.43
5568	s23	april	wet	87	Alto Redondo	165.6	339.5	0.37	0.42
5568	s24	april	wet	102	Alto Redondo	229.5	354.6	0.30	0.41

<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
5568	s25	may	wet	117	Alto Redondo	237.8	351.5	0.41	0.40
5568	s26	may	wet	132	Alto Redondo	254.6	63.8	0.36	0.47
5568	s27	june	dry	147	Alto Redondo	254.6	341.1	0.34	0.47
5568	s28	june	dry	162	Alto Redondo	244.4	349.6	0.34	0.45
5568	s29	july	dry	177	Alto Redondo	244.4	329.1	0.39	0.43
5568	s30	july	dry	192	Alto Redondo	235.5	321.2	0.42	0.39
5568	s31	august	dry	207	Alto Redondo	228.9	314.6	0.34	0.33
5568	s32	august	dry	222	Alto Redondo	206.4	314.6	0.31	0.33
5568	s33	september	dry	237	Alto Redondo	140.3	319.5	0.34	0.33
5568	s34	september	dry	252	Alto Redondo	121.8	301.5	0.30	0.31
5568	s35	october	dry	267	Alto Redondo	119.4	285	0.30	0.28
5568	s36	october	dry	282	Alto Redondo	52.1	281.6	0.28	0.27
5568	s37	november	dry	297	Alto Redondo	43.8	281.6	0.25	0.25
5568	s38	november	dry	312	Alto Redondo	17.1	281.6	0.25	0.24
5568	s39	december	dry	327	Alto Redondo	17.1	271.7	0.29	0.33
5568	s40	december	dry	342	Alto Redondo	148.8	415.6	0.41	0.48
5568	s41	december	dry	357	Alto Redondo	149	393.4	0.43	0.54
5568	s42	january	wet	7	Alto Redondo	149	384.5	0.36	0.47
5568	s43	january	wet	22	Alto Redondo	149	377.9	0.33	0.47
5568	s44	february	wet	37	Alto Redondo	154.2	360.6	0.34	0.49
5568	s45	march	wet	52	Alto Redondo	149.3	331.1	0.33	0.47
5568	s46	march	wet	67	Alto Redondo	159	282.8	0.31	0.40
5568	s47	march	wet	82	Alto Redondo	204.4	326.2	0.39	0.56
5568	s48	april	wet	97	Alto Redondo	332.6	384.7	0.45	0.60
5568	s49	april	wet	112	Alto Redondo	332.6	376.4	0.41	0.53
5568	s50	may	wet	127	Alto Redondo	359.3	376.4	0.41	0.49
5568	s51	may	wet	142	Alto Redondo	382.3	399.4	0.43	0.49
5568	s52	june	dry	157	Alto Redondo	241.9	390.7	0.50	0.58
5568	s53	june	dry	172	Alto Redondo	285.9	434.9	0.61	0.60
5568	s54	july	dry	187	Alto Redondo	287.5	436.5	0.40	0.60

<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
5570	s1	may	wet	122	Alto Redondo	114.4	193.4	0.33	0.41
5570	s2	may	wet	137	Alto Redondo	113.6	171.9	0.40	0.46
5570	s3	june	dry	152	Alto Redondo	149.3	201.6	0.46	0.48
5570	s4	june	dry	167	Alto Redondo	133.3	200.7	0.38	0.43
5570	s5	july	dry	182	Alto Redondo	144.9	203.5	0.50	0.48
5570	s6	july	dry	197	Alto Redondo	142.7	208.7	0.50	0.50
5570	s7	august	dry	212	Alto Redondo	148.1	211.9	0.41	0.46
5570	s8	august	dry	227	Alto Redondo	148.1	211.2	0.40	0.35
5570	s9	september	dry	242	Alto Redondo	148.1	208.9	0.42	0.35
5570	s10	september	dry	257	Alto Redondo	170.8	237.2	0.51	0.39
5570	s11	october	dry	272	Alto Redondo	121.7	238.7	0.59	0.47
5570	s12	october	dry	287	Alto Redondo	113.7	226.1	0.46	0.40
5570	s13	november	dry	302	Alto Redondo	111.7	226.1	0.38	0.36
5570	s14	november	dry	317	Alto Redondo	121.6	235.2	0.37	0.34
5570	s15	december	dry	332	Alto Redondo	85.9	235.2	0.34	0.32
5570	s16	december	dry	347	Alto Redondo	95	257.6	0.33	0.31
5570	s17	january	wet	362	Alto Redondo	83.4	228.3	0.40	0.34
5570	s18	january	wet	12	Alto Redondo	84.8	227.5	0.35	0.34
5570	s19	february	wet	27	Alto Redondo	85.7	233.8	0.29	0.31
5570	s20	february	wet	42	Alto Redondo	108.2	256.3	0.29	0.33
5570	s21	march	wet	57	Alto Redondo	179.2	327.3	0.33	0.34
5570	s22	march	wet	72	Alto Redondo	166.3	345.8	0.39	0.38
5570	s23	april	wet	87	Alto Redondo	165.6	339.5	0.45	0.42
5570	s24	april	wet	102	Alto Redondo	229.5	354.6	0.39	0.40
5570	s25	may	wet	117	Alto Redondo	237.8	351.5	0.42	0.43
5570	s26	may	wet	132	Alto Redondo	254.6	63.8	0.45	0.47
5570	s27	june	dry	147	Alto Redondo	254.6	341.1	0.44	0.47
5570	s28	june	dry	162	Alto Redondo	244.4	349.6	0.39	0.46
5570	s29	july	dry	177	Alto Redondo	244.4	329.1	0.44	0.45
5570	s30	july	dry	192	Alto Redondo	235.5	321.2	0.43	0.38



<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
5570	s31	august	dry	207	Alto Redondo	228.9	314.6	0.36	0.34
5570	s32	august	dry	222	Alto Redondo	206.4	314.6	0.33	0.31
5570	s33	september	dry	237	Alto Redondo	140.3	319.5	0.30	0.29
5570	s34	september	dry	252	Alto Redondo	121.8	301.5	0.29	0.28
5570	s35	october	dry	267	Alto Redondo	119.4	285	0.27	0.26
5570	s36	october	dry	282	Alto Redondo	52.1	281.6	0.26	0.26
5570	s37	november	dry	297	Alto Redondo	43.8	281.6	0.26	0.26
5570	s38	november	dry	312	Alto Redondo	17.1	281.6	0.26	0.23
5570	s39	december	dry	327	Alto Redondo	17.1	271.7	0.29	0.32
5570	s40	december	dry	342	Alto Redondo	148.8	415.6	0.40	0.47
5570	s41	december	dry	357	Alto Redondo	149	393.4	0.39	0.48
5570	s42	january	wet	7	Alto Redondo	149	384.5	0.33	0.42
5570	s43	january	wet	22	Alto Redondo	149	377.9	0.29	0.44
5570	s44	february	wet	37	Alto Redondo	154.2	360.6	0.34	0.45
5570	s45	march	wet	52	Alto Redondo	149.3	331.1	0.30	0.49
5570	s46	march	wet	67	Alto Redondo	159	282.8	0.29	0.46
5570	s47	march	wet	82	Alto Redondo	204.4	326.2	0.36	0.56
5570	s48	april	wet	97	Alto Redondo	332.6	384.7	0.48	0.53
5570	s49	april	wet	112	Alto Redondo	332.6	376.4	0.47	0.45
5570	s50	may	wet	127	Alto Redondo	359.3	376.4	0.39	0.44
5570	s51	may	wet	142	Alto Redondo	382.3	399.4	0.53	0.55
5570	s52	june	dry	157	Alto Redondo	241.9	390.7	0.58	0.57
5570	s53	june	dry	172	Alto Redondo	285.9	434.9	0.63	0.64
5570	s54	july	dry	187	Alto Redondo	287.5	436.5	0.77	0.77
5570	s55	july	dry	202	Alto Redondo	318.6	467.6	0.73	0.65
5570	s56	august	dry	217	Alto Redondo	313.4	467.6	0.63	0.50
5570	s57	august	dry	232	Alto Redondo	323.3	477.5	0.52	0.47
6444	s1	february	wet	39	Limoeiro	262.7	752.4	0.60	0.58
6444	s2	march	wet	54	Limoeiro	325.6	815.3	0.57	0.57
6444	s3	march	wet	69	Limoeiro	329	664.7	0.49	0.50

<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
6444	s4	april	wet	84	Limoeiro	335.1	599.9	0.53	0.53
6444	s5	april	wet	99	Limoeiro	344.5	535.9	0.50	0.50
6444	s6	may	wet	114	Limoeiro	400.8	405.9	0.49	0.51
6444	s7	may	wet	129	Limoeiro	287.8	408.4	0.50	0.61
6444	s8	june	dry	144	Limoeiro	225.8	408.4	0.52	0.53
6444	s10	july	dry	174	Limoeiro	225.8	403.3	0.46	0.53
6444	s11	july	dry	189	Limoeiro	204.8	403.3	0.40	0.49
6444	s12	july	dry	204	Limoeiro	175.8	403.3	0.37	0.47
6444	s13	august	dry	219	Limoeiro	140.6	403.3	0.33	0.51
6444	s14	august	dry	234	Limoeiro	77.7	403.3	0.31	0.46
6444	s15	september	dry	249	Limoeiro	83.3	412.3	0.30	0.45
6444	s16	september	dry	264	Limoeiro	77.2	412.3	0.28	0.40
6444	s17	october	dry	279	Limoeiro	67.8	412.3	0.29	0.40
6444	s18	october	dry	294	Limoeiro	11.5	412.3	0.37	0.42
6444	s19	november	dry	309	Limoeiro	91	378.8	0.53	0.58
6444	s20	november	dry	324	Limoeiro	145.4	385.2	0.62	0.67
6444	s21	december	dry	339	Limoeiro	209.4	435.2	0.52	0.59
6444	s22	december	dry	354	Limoeiro	257.4	483.2	0.60	0.70
6444	s23	january	wet	4	Limoeiro	377.2	582	0.65	0.65
6444	s24	january	wet	19	Limoeiro	399.2	575	0.64	0.73
6444	s25	february	wet	34	Limoeiro	421.2	569.8	0.52	0.72
6444	s26	february	wet	49	Limoeiro	429.2	516.9	0.51	0.67
6444	s27	march	wet	64	Limoeiro	448.7	535.4	0.55	0.70
6444	s28	march	wet	79	Limoeiro	489.7	573	0.54	0.56
6444	s29	april	wet	94	Limoeiro	489.7	557.5	0.59	0.66
6444	s30	april	wet	109	Limoeiro	526.2	594	0.67	0.66
6444	s31	may	wet	124	Limoeiro	444.2	535.2	0.57	0.63
6444	s32	may	wet	139	Limoeiro	409.2	554.6	0.50	0.60
6444	s33	june	dry	154	Limoeiro	345.2	554.6	0.48	0.59
6444	s34	june	dry	169	Limoeiro	297.2	554.6	0.55	0.59

<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
6444	s35	july	dry	184	Limoeiro	177.4	554.6	0.57	0.60
6444	s36	july	dry	199	Limoeiro	155.4	554.6	0.47	0.58
6444	s37	august	dry	214	Limoeiro	133.4	554.6	0.39	0.52
6444	s38	august	dry	229	Limoeiro	125.4	554.6	0.40	0.48
6444	s39	september	dry	244	Limoeiro	96.9	554.6	0.39	0.46
6444	s40	september	dry	259	Limoeiro	55.9	545.6	0.39	0.33
6444	s41	october	dry	274	Limoeiro	62.4	552.1	0.35	0.50
6444	s42	october	dry	289	Limoeiro	25.9	552.1	0.34	0.39
6444	s43	november	dry	304	Limoeiro	55.9	582.1	0.43	0.48
6444	s44	november	dry	319	Limoeiro	151.8	561	0.53	0.57
6444	s45	december	dry	334	Limoeiro	207.8	564	0.60	0.68
6444	s46	december	dry	349	Limoeiro	272.8	570	0.60	0.75
6444	s47	january	wet	364	Limoeiro	359.1	536.5	0.70	0.75
6444	s48	january	wet	14	Limoeiro	374.1	544	0.64	0.72
6444	s49	february	wet	29	Limoeiro	397.6	531	0.69	0.74
6444	s50	february	wet	44	Limoeiro	397.6	531	0.57	0.70
9025	s1	july	dry	189	Quixaba	204.8	350.6	0.48	0.49
9025	s2	july	dry	204	Quixaba	209.8	353.3	0.48	0.47
9025	s3	august	dry	219	Quixaba	194.6	362.5	0.48	0.46
9025	s4	august	dry	234	Quixaba	197.2	361.1	0.40	0.39
9025	s5	september	dry	249	Quixaba	194	358.9	0.33	0.33
9025	s6	september	dry	264	Quixaba	194	358.9	0.32	0.32
9025	s7	october	dry	279	Quixaba	170.1	364	0.32	0.32
9025	s8	october	dry	294	Quixaba	161.8	364	0.31	0.32
9025	s9	november	dry	309	Quixaba	182	294	0.32	0.34
9025	s10	november	dry	324	Quixaba	149.9	287.6	0.33	0.37
9025	s11	december	dry	339	Quixaba	139.1	297	0.41	0.46
9025	s12	december	dry	354	Quixaba	133.6	297.5	0.48	0.51
9025	s13	january	wet	4	Quixaba	138.8	343.6	0.51	0.54
9025	s14	january	wet	19	Quixaba	198.9	408.7	0.60	0.61

<b>ID</b>	<b>Subset</b>	<b>Month</b>	<b>Season</b>	<b>Ordinal date</b>	<b>Rainfall station</b>	<b>Accum. rainfall 6</b>	<b>Accum. rainfall 12</b>	<b>Real NDVI median</b>	<b>Random NDVI median</b>
9025	s15	february	wet	34	Quixaba	261.5	456.1	0.63	0.63
9025	s16	february	wet	49	Quixaba	302.8	500.7	0.63	0.62
9025	s17	march	wet	64	Quixaba	400.6	594.6	0.65	0.61
9025	s18	march	wet	79	Quixaba	400.6	594.6	0.53	0.55
9025	s19	april	wet	94	Quixaba	399.4	576.3	0.48	0.54
9025	s20	april	wet	109	Quixaba	405.6	574.2	0.49	0.56
9025	s21	may	wet	124	Quixaba	376.9	559.7	0.50	0.58
9025	s22	may	wet	139	Quixaba	379.1	530.1	0.51	0.59
9025	s23	june	dry	154	Quixaba	452.5	605.8	0.60	0.63
9025	s24	june	dry	169	Quixaba	485.3	623.2	0.56	0.63
9025	s25	july	dry	184	Quixaba	482.5	634.6	0.59	0.57
9025	s26	july	dry	199	Quixaba	457.5	657.5	0.58	0.56
9025	s27	august	dry	214	Quixaba	398.6	660.6	0.60	0.54
9025	s28	august	dry	229	Quixaba	384.3	687.8	0.44	0.49
9025	s34	november	dry	319	Quixaba	361.8	740.9	0.38	0.54
9025	s35	december	dry	334	Quixaba	320.7	782.1	0.59	0.65
9025	s36	december	dry	349	Quixaba	325.2	811	0.65	0.72
9025	s37	january	wet	364	Quixaba	374.5	857	0.61	0.72
9025	s38	january	wet	14	Quixaba	338.4	795.9	0.55	0.65
9025	s39	february	wet	29	Quixaba	319.5	718.1	0.48	0.59
9025	s40	february	wet	44	Quixaba	290.5	674.8	0.50	0.64
9025	s41	march	wet	59	Quixaba	289	604.7	0.45	0.58
9025	s42	march	wet	74	Quixaba	284.9	576.3	0.51	0.57
9025	s43	april	wet	89	Quixaba	437.5	725	0.55	0.63
9025	s44	april	wet	104	Quixaba	457.2	744.7	0.57	0.60
9025	s45	may	wet	119	Quixaba	457.2	727.7	0.61	0.60
9025	s46	may	wet	134	Quixaba	365.3	728.8	0.58	0.66

SUPPLEMENTARY TABLE 6 Candidate models set for the home ranges and the respective predictor variables showing performance of the top model relative to others in the model set. The models are ordered from the lowest to the highest delta AIC value (dAIC, i.e., calculated difference between AIC value of the observed model and the candidate model with the lowest AIC). The symbol ‘\*’ in the column “Model and predictor variables” indicates interaction between the two variables tested in respective model.

<b>Model and predictor variables (95% kernel)</b>	<b>AIC</b>	<b>dAIC</b>	<b>Adjusted R<sup>2</sup></b>
season + tagging site	568.7478	0.0	0.02256
season * tagging site	568.8816	0.1	0.03135
real coordinates median NDVI	569.6104	0.9	0.008693
accumulated rainfall 12 months	570.1034	1.4	0.006259
null	570.3730	1.6	
accumulated rainfall 6 months	571.4172	2.7	-0.0002577
random coordinates median NDVI	572.2937	3.5	-0.004629
tagging site	574.3730	5.6	-0.0101
ordinal date + tagging site	576.2354	7.5	-0.01453
ordinal date * tagging site	578.6583	9.9	-0.01693

SUPPLEMENTARY TABLE 7 Candidate models set for the core areas and the respective predictor variables showing performance of the top model relative to others in the model set. The models are ordered from the lowest to the highest delta AIC value (dAIC, i.e., calculated difference between the AIC value of the observed model and the candidate model with the lowest AIC). The symbol ‘\*’ in the column “Model and predictor variables” indicates interaction between the two variables tested in respective model.

<b>Model and predictor variables (50% kernel)</b>	<b>AIC</b>	<b>dAIC</b>	<b>Adjusted R<sup>2</sup></b>
accumulated rainfall 6 months	567.9202	0.0	0.01699
season * tagging site	568.9034	1.0	0.03125
real coordinates median NDVI	569.0634	1.1	0.01139
season + tagging site	569.1104	1.2	0.0208
null	570.3730	2.5	
random coordinates median NDVI	571.0147	3.1	0.001743
accumulated rainfall 12 months	572.1871	4.3	-0.004096
tagging site	574.3730	6.5	-0.0101
ordinal date + tagging site	576.2865	8.4	-0.01479
ordinal date * tagging site	580.1843	12.3	-0.02468

SUPPLEMENTARY TABLE 8 Correlation matrix showing the Pearson’s correlation coefficients (r) calculated for the numeric predictor variables: ordinal date of the respective fortnightly subset, median of the NDVI data associated with the real coordinates recorded for the macaws (“Real NDVI”), median of the NDVI data associated with the random coordinates (“Random NDVI”), accumulated rainfall (in millimeters) over the last six months (“Rainfall 6”), and accumulated rainfall (in millimeters) over the last twelve months (“Rainfall 12”).

	<b>Ordinal date</b>	<b>Real NDVI</b>	<b>Random NDVI</b>	<b>Rainfall 6</b>	<b>Rainfall 12</b>
<b>Ordinal date</b>	1.0	-0.2	-0.3	-0.4	-0.2
<b>Real NDVI</b>	-0.2	1.0	0.9	0.7	0.6
<b>Random NDVI</b>	-0.3	0.9	1.0	0.7	0.7
<b>Rainfall 6</b>	-0.4	0.7	0.7	1.0	0.7
<b>Rainfall 12</b>	-0.2	0.6	0.7	0.7	1.0

SUPPLEMENTARY TABLE 9 Summary of the values of mean, median, standard deviation (“SD”), minimum (“Min.”) and maximum (“Max.”) of the distances (in meters) calculated from each input location (“Feeding” or “Resting”) to the closest target site (“Roosting” or “Feeding”).

<b>Input locat.</b>	<b>Target locat.</b>	<b>Mean (m)</b>	<b>Median (m)</b>	<b>SD (m)</b>	<b>Min. (m)</b>	<b>Max. (m)</b>
Feeding	Roosting	2,330.72	201.13	4,135.24	0.10	35,824.69
Resting	Roosting	2,754.65	361.52	4,450.96	0.13	35,765.94
Resting	Feeding	75.45	19.81	281.24	0.00	7,608.64



SUPPLEMENTARY TABLE 10 Summary of the number of GPS fixes (positions), for the identified locations (“Feeding”, “Resting” and “Roosting”), where macaws stayed in a defined radius for a defined minimum duration (not moving faster than 1 m/s GPS ground speed), and the calculated values of minimum (“Min.”), maximum (“Max.”), mean and median of the time duration (i.e., time spent, in hours) inside the area of radius r (in meters).

		<b>Number of positions</b>	<b>Time duration (hours)</b>	<b>Radius (meters)</b>
<b>Feeding</b>	Min.	3	5	6
	Max.	121	11	999
	Mean	32	8	458
	Median	31	8	431
<b>Resting</b>	Min.	2	3	0.1
	Max.	121	10.5	49.9
	Mean	19.6	4.5	21
	Median	20	4.2	18.2
<b>Roosting</b>	Min.	2	9	1
	Max.	61	14	499
	Mean	11	12	178
	Median	10	12	160