Seasonal Variation in the Home Ranges of Black Curassow, *Crax alector*, in French Guiana

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Abstract

Cracidae is the most threatened avian family in the Neotropics, mainly because of habitat destruction, heavy hunting pressure and poaching. In French Guiana, Black Curassows are heavily hunted, although basic knowledge of the ecological and demographical traits of the species remains limited. Such a gap prevents any attempt to assess the impact of hunting and to help stakeholders to develop proposals ensuring hunting sustainability. The spatial relationship between animals and their habitat is important for conservation management, being related to population densities through complex patterns. Here, we report on a radio-tracking study of Black Curassows in tropical primary rainforest, in Nouragues National Reserve, French Guiana. The aims of the study were to estimate home range size and its variation across seasons, and to quantify movement patterns of the birds. We captured and fitted VHF tags to four adults, and tracked them for 10 to 21.5 months. Daily movements were recorded, and home ranges estimated using the Kernel Density method, for two consecutive wet seasons and one dry season. Using 95% and 50% Kernel densities, the average annual home range and core area were 96.3 ± 32.6 ha (SE) and 22.8 ± 2.8 ha respectively. Home ranges appeared spatially stable over the two years, and overlapped between neighbouring groups. During the dry season, Black Curassows did not migrate but tended to enlarge their home range, with greater daily movements and higher home range overlap. Although additional data are still needed, our results can help to improve the knowledge and management of this poorly studied species.

Keywords

spatial ecology, Cracids, tracking, Kernel method, Home range, movement pattern

Introduction

Cracids (curassows, guans, and chachalacas) comprise a group of large, forest-dwelling frugivorous birds and represent the most threatened avian family in the Neotropics (Brooks, 2006a; del Hoyo et al., 1994; Kattan et al., 2016; Strahl et al., 1997b), 50% of the species within this family being listed as “Vulnerable” or at higher risk of extinction (IUCN, 2018). Due to decreasing population trends, the Black Curassow *Crax alector* was uplisted in 2012 from “Least Concern” to “Vulnerable” in the IUCN red list (Birdlife International, 2016; A3c). It is considered to be one of the first species to disappear from areas subjected to heavy hunting pressure, having almost vanished from forests near the coast, around settled areas and villages, along large rivers, and wherever hunting pressure is high (Delacour & Amadon, 2004; Denis et al., 2016). In French Guiana, Niel et al. (2008) highlighted the risk of local overharvesting on this species, using the demographic invariant method (Niel & Lebreton, 2005), while Denis et al. (2016) further showed a negative correlation...
between hunting pressure and Black Curassow densities. Accordingly, in order to reduce the hunting pressure on this population, the trade of Black Curassows (mainly hunted for meat) was officially banned in 2007, and hunting was slightly constrained in 2012 by setting a bag limit of two Black Curassows harvested per hunter and hunting trip (with no limitation on the number of hunting trips). No stronger regulation could be set at that time given the absence of hunting licence and general hunting management laws in French Guiana. However, despite this critical situation, basic knowledge of the ecological and demographic traits of Black Curassows remain scarce.

Among those biological traits, spatial patterns as home range size and habitat use are of main interest. Home ranges reveal the relationships between animals and their habitat, and are closely related to population densities through complex relationships (Efford, 2014; Efford et al., 2015). Understanding ranging behaviour is a prerequisite for wildlife management and conservation, and basic spatial parameters are frequently needed to model abundance (Rowcliffe et al., 2008; Royle et al., 2016) and movement ecology. Moreover, knowledge of spatial patterns of habitat use is necessary in the assessment of hunting sustainability (Constantino, 2015; Levi et al., 2011; Shaffer et al., 2018). The previous sustainability assessment made on Black Curassow in French Guiana used home ranges size estimates extrapolated from other countries or cracid species, because the information was unavailable at that time (Niel et al., 2008).

To fill this knowledge gap, we therefore initiated a study to estimate home range size used by Black Curassows in 2010 in French Guiana (De Coster, 2011; Rocheteau, 2010). This local scale study was complementary to a larger scale approach undertaken over the whole territory, aiming at understanding the abundance and density of the species in relation to habitat type and hunting pressure (Denis et al., 2016; Richard-Hansen et al., 2015). The main goal of the present study was to increase our knowledge on Black Curassow behaviour and spatial ecology and for the first time to estimate home range size and its seasonal variation. The study was undertaken in the protected area of Nouragues National Reserve, where hunting and forest disturbance are currently absent, to get baseline data. The accurate evaluation of these parameters could highly improve the assessment of local hunting impact and help environmental managers and stakeholders to implement the best management decisions for hunting sustainability.

**Methods**

**Study Site**

We conducted fieldwork totalling 336 days of tracking, from March 2010 to August 2013 (Table 1) in the Nouragues National Reserve and Nouragues Ecological Research Stations, French Guiana (4°05'15" N, 52°40'46" W). The Nouragues reserve is located ca.95 km inland and covers ca.2500 ha, and consists entirely of tropical primary rainforest of the Guiana Shield. The Research Station is located in the Nouragues creek watershed, a tributary of the Arataye River. Elevation varies between 100 m and 411 m asl, the height of the inselberg (granitic rocky mountain). The vegetation is dominated by *terra firme* forest, with canopy height between 30 and 50 m, with patches of palm-swamp, liana or bamboo forests (Bongers et al., 2001). Habitat and landscape type were classified as all-slope in the recent forest typology of *terra firme* French Guianan forests (Guitet et al., 2015), Caesalpiniaceae, Lecythidaceae, Sapotaceae and Burseraceae being among the most common tree families (Poncy et al., 2001). Two distinct study sites, ca.5 km apart, were used: the Pararé site was located along the Arataye river in the lower part of the area (25m − 170 m asl), and the Inselberg site was located at a higher elevation, with steeper slopes, and situated just beneath the ca.410 m inselberg.

In French Guiana, a dry season occurs from August to November, and a long rainy season from December to July, with a small decrease in rainfall around March. Mean monthly maximum temperatures rise from 29°C (January) to 34°C (October), while mean minimal ones stay around 22°C all year round. Average annual rainfall is 2990 mm, monthly rainfall being above 100 mm in September-October and up to 500 mm at the peak of rainy season (May; Bongers et al., 2001). Tree fruting peaks in March-May and is minimal in August-September (Norden et al., 2007, Sabatier & Puig, 1985). Among 48 trees and liana species, 54% showed an annual fruting pattern, 23% a masting fruting pattern (mainly animal-dispersed trees), and 23% an irregular fruting pattern (Norden et al., 2007).

**Capture**

Capture, tagging and tracking of Black Curassow were made in compliance with legal requirement (Décret n° 95-1299 portant création de la réserve naturelle des Nouragues (Guyane)). Capture was really challenging, and we needed various attempts using several methods, of which only two were successful. We first identified the tracks routinely used by family groups of curassows near the field research stations, and then we used two different types of traps: hoop net and leg laces. The hoop net was made of two converging 20 to 30 meters long walls of mist-net, deployed in a “V” shape on each side of the track. The last ten meters of the hoop “head” were covered by nets to prevent the bird to flush. When a bird entered the hoop net, we gently pushed it towards the
head of the hoop and caught it when it was trapped in the
net. Alternatively, we also positioned leg laces on the
birds’ tracks, the laces being attached to a 1.5 m long
rubber extensible spring to prevent the bird being injured.
As soon as the bird was captured, its head was covered
with a black sock to reduce stress. We weighed birds to the
nearest 10 g with a 5000-g Walter® spring balance.

Tracking
We tracked Black Curassows fitted with TW3 twin cells
Biotrack pip tags (weight 70g) using a Biotrack Sika
receiver and two types of handheld directional antennas
(3-element Yagi from Biotrack and RA-23K “H” antenna
from Telonics). Captured birds’ body mass ranged from
2.7 to 3.3 kg, thus the mass ratio between bird and trans-
mitter ranged from 2 to 2.6%. The transmitters were fas-
tened onto the birds’ back using a double loop harness
made with tubular Teflon ribbon (diameter 0.33 inch),
passing over each wing. We previously tested transmitter
design and mounting for several days on captive Black
Curassows from the Zoo of Cayenne. As also found in a
field study on captive and reintroduced neighbouring spe-
cies (Bernardo et al., 2011a), the backpack and antennas
were not conspicuous, being covered by feathers, and did
not modify the behavior of the birds held captive in the
zoo. Processing of the birds took on average 30 minutes.
The tracking period began one day after release and lasted
until the battery of the transmitter failed. Transmitters had
an expected battery life of 2 years and a theoretical
ground-to-ground detection range of 3 to 6 km.

We determined the sex of birds mainly by behavioural
observations: only males perform territorial vocaliza-
tions (booming) (Delacour & Amadon, 2004) and whistle
or clucking calls differ between sexes (fieldworker
observations during this study). Moreover, the colour
of eyes is dark for males and brownish for females
(Todd et al., 1992). As all birds within a family group
stay always very close to each other, the location of the
tagged bird generally corresponded to the location of the
whole group.

Locations were determined by moving in the direction
of a radio signal until we saw the bird. We then noted its
behaviour and, after the bird moved, we recorded the
location’s coordinates via a Garmin 60 CSX GPS unit.
The location error was estimated as 9.79 ± 0.29 m (SE).
When the thick vegetation prevented us to see the bird,
even at short distance (less than 50 m between the observ-
er and the bird), we estimated a most likely location. If
the bird moved in response to our approach, the location
of the bird prior to moving was recorded. We tracked
birds 3 to 4 times each day. The delay between successive
locations was at least 3 to 4 hours, in order to satisfy the
biological independence of locations, as curassows can
easily travel through their whole home range in less
than 3 hours. We chose the tracking hours randomly
every day for each bird, aiming to obtain, over the
whole study period, a roughly equivalent number of loca-
tions for every hour from dawn (06:00) to dusk (18:00).
We considered locations obtained at dawn and dusk as
representative of roosting sites.

Additionally, some days were dedicated to a continu-
ous tracking of birds. After some weeks, Male A and
Male B groups were very accustomed to human pres-
ence, and did not modify their behaviour as long as
the observer stayed at a minimum distance. We then
tracked animals by foot during the diurnal period, keep-
ing visual contact, as long as possible, while a GPS
recorded the track and a fix every minute. The GPS
human track was considered to match that of the
birds, and the behaviour was also noted every minute.
We recorded different behaviours (travelling, feeding,
resting, grooming and territorial behaviours), however
in the present study we only used travelling (hereafter
“movement”) data.

Home Range Estimation
We estimated home range size using both Minimum
Convex Polygon (MCP; White & Garrot, 1990) and
Kernel methods, with the href smoothing method
(Seaman & Powell, 1996). The MCP method encom-
passes all locations, and includes exploratory move-
ments, while the Kernel density estimator produces an
unbiased density estimate of presence probability direct-
ly from data and is not influenced by grid size or place-
ment (Silverman, 1986). The fixes recorded during the
continuous tracking days were not included in this anal-
ysis, because they were strongly autocorrelated. Data
were analysed using the software R (version 3.0.1; R
Core Team, 2014) and the packages ‘ade4’ (Chessel et
al., 2004), ‘adehabitat’ (Calenge, 2006) and ‘maps’
(Becker et al., 2007). We used the extension Hawth’s
Analysis Tools 3.27 on the GIS application ArcMap
9.2 to estimate MCP.

To standardize and compare the various estimates of
home-range size using the Kernel method within seasons
(wet, dry) or over the whole annual cycle (wet + dry
seasons), we first estimated 50% and 95% Kernel
home ranges using a smoothing parameter $h_{ref}$
(Seaman & Powell, 1996) for each individual separately.
Then we calculated a mean $h_{ref}$ within each time period
(wet season, dry season, annual cycle) and used this
value to run again the estimation of 50% and 95%
kernel home ranges.

Movement Patterns
We modelled the distance travelled by the birds (m) in
relation to time-period (hourly from 06:00 to 18:00 as a
Table 1. Tracking Periods for Each Tagged Black Curassow.

|                | 2010  | 2011  | 2012  | 2013  | fixes | tracking days |
|----------------|-------|-------|-------|-------|-------|---------------|
| Male A         |       |       |       |       | 555   | 252           |
| Male B + family|       |       |       |       | 345   | 252           |
| Female D (with A)   |       |       |       |       | 53    | 45            |
| Female D alone    |       |       |       |       | 42    | 32            |
| Female C         | 263   | 84    |       |       |       |               |

Black cells indicate months when tracking was performed.

categorical factor) and season (dry and wet), and included two-way interaction effects. For this purpose, we used a generalized linear mixed model (GLMM, function glmer (MASS package 7.3-45, (Venables & Ripley, 2002) with a negative binomial error structure to estimate distance travelled. We included individual birds (or female/male group), and the repetition of tracking hours as random effects. Finally, Wald tests were implemented to test fixed effects from season and time-period (and their interactions), discarding effects that do not differ significantly from zero (Bolker et al., 2009).

Results

Four Black Curassows, two males (A, B) and two females (C, D), were captured and tracked, three of them in the inselberg station, and one (C) in the Pararé station. The whole tracking period lasted a minimum of 10 months (female C) to a maximum of 21.5 months (male A) and the number of locations for each tagged bird ranged from 95 to 608 (see Table 1). Female D was initially paired with male A, but the pair split several times, and no home range could be estimated for the female D because she dispersed too far to be regularly located. Male A stayed in the same area and was observed paired with another female for the rest of the study.

The Male B group initially included an adult female and a fully feathered juvenile. From the 18 February 2011, we could see this group with two downy chicks; the juvenile was still there and, together with adults, was providing care to the chicks. One of the chicks disappeared in early May 2011. We did not see the juvenile after April 2012, presumably because it left the family group. At the end of the tracking study, the surviving chick was still with the adults and was in full-grown juvenile plumage.

Female C was initially paired with a male, and two chicks were observed in March 2013. However, we often observed female C joining another neighbouring group, without the initial male, which stayed in the same area.

Home Range

For groups A and B, we were able to estimate home ranges for two consecutive wet seasons and one dry season, and for female C, over one full annual cycle. On the basis of 95% and 50% Kernel estimates, the average annual home range and core area were 96.3 ± 32.6 ha (SE) and 22.8 ± 2.8 ha (Table 2; Figure 1A to C). Using MCP, this annual home range was 107.8 ± 32.6 ha (Table 2). Home ranges sizes of Male A and Female C were relatively similar but the home range and core area of Male B was much larger, twice the size when using MCP.

During the wet season, the average home range area was markedly lower than during the dry season (95% Kernel: 79 ± 4.4 ha versus 100.8 ± 18.6; MCP: 55.2 ± 4.3 ha versus 90.7 ± 36.9; Table 3; Figure 2A to C), particularly for Male B’s home range. Female C’s 95% Kernel home range was however slightly larger in wet season than in the dry season. Contrarily, the core area showed very little variation between seasons (wet season: 21.2 ± 1.1 ha, dry season 19.8 ± 2.5 ha; Table 3). The extended home range during the dry season was not associated with any dramatic changes in the area foraged by birds, since for groups A and B, the home range during the dry season (using 95% Kernel home range) overlapped respectively 70.5% and 80% of home ranges covered during the wet season.

The two family groups tracked in the Inselberg site showed an important overlap in their home ranges, reaching up to 63% of the group A annual home range (using 95% Kernel) and even 11.6% of the core area. This overlap was more pronounced during the dry season, reaching a maximum value of 60.1% of group A’s home range (30.9% for core area) versus 46.4% during the wet season (and no overlap for core area).

Movement Patterns

We recorded continuous tracking data on 35 diurnal sessions (18 during the wet season, 17 during the dry season), for a total of 232 hours (120 hours during the wet season, 112 hours during the dry season). The majority of the data was collected on males A and B.
(190 hours). The mean duration of the tracking sessions was 4h: 57 min ± 36 min (SE; min: 1h: 00, max: 12h: 00). All time-periods from 06:00 to 19:00 were sampled, and the mean hourly distance travelled by birds could be compared. The median number of tracking sessions per hour was 18 (range 5–25). The movement activity differed during the daytime (\( P < 0.001, \chi^2 = 707.2, \text{df} = 11 \)), and according to the season (\( P < 0.001, \chi^2 = 22.7, \text{df} = 1 \); interaction \( P < 0.001, \chi^2 = 247.4, \text{df} = 11 \)). On average, birds travelled 189 ± 32 m/hour (SE; overall 2455 m/day) in the wet season and 372 ± 57 m/hour (overall 4843 m/day) in the dry season. Animals travelled longer distances per minute during the dry season than during the wet season (Dry season 11.8 m/minute ± 0.21 (SE), Wet season 7.7 ± 0.19 (SE), difference: +4.1 m in dry season, \( P < 0.001 \)). We observed maximum movement earlier in the morning (6:00–7:00) and later in the evening (17:00–18:00) in the dry season than in the wet season (7:00–8:00 and 16:00–18:00). Minimum movement activity occurred at mid-day (11:00–12:00) for both seasons (Figure 3).

**Discussion**

This study presents the first estimation of *Crax alector* home range based on tracking data on more than one bird or pair, and over a full annual cycle or more. Literature on the ecology and spatial behaviour of Cracids relies upon a limited number of tracking data across their distribution range. To our knowledge, the
determination of annual home range size has been considered for a very limited number of cracid species (Crax alector (Bernal & Mejía, 1995; Erard & Sabatier, 1989); Crax daubentoni (Bertsch & Barreto, 2008a); Crax blumenbachii (reintroduced population; Bernardo et al., 2011b); Crax globulosa (Leite et al., 2018); M. salvini: (Parra et al., 2001; Santamaria & Franco, 2000; Yumoto, 1999). In many of those cases, it relies upon the monitoring of only one familial group. Moreover, the methodological tools used to estimate home range greatly differ between studies (MCP, Kernel, empirical estimation on the field...), which hampers eventual comparisons between studies. We summarize these different estimates in Table 4. Overall, our own estimate (95% kernel: 96 ha, MCP: 108 ha) tends to be smaller than results obtained by the other tracking studies available on Crax species (C. daubentoni 149 to 197 ha; C. blumenbachii 125 ha), and particularly C. globulosa in western amazon (468 ha or 804 ha with respectively 95% Kernel and 95% MCP). Bernal and Mejía (1995) in Leite (2017) estimated 185 ha for C. alector’s home range size in Bolivia. All these home ranges (our study included) were estimated in protected areas where no hunting occurs. Former values for C. alector home range sizes available in French Guiana were empirically estimated from the field without radio-tracking nor definite method (Erard & Sabatier, 1989) or from the literature for modelling purposes (Niel et al., 2008: 19.63 ha) and clearly underestimated the annual home range in comparison with our results. Our study also suggests that, at least for the two males, home ranges tend to remain stable from one year to the next. Detailed spatial data as daily movements also provides useful parameters for ecological functional approaches and modelling, as dispersal ability of juveniles (Bowman, 2003) or seed dispersal distance (Donoso et al., 2020), although Crax’ strong gizzard probably crushes larger seeds (Erard et al., 1991).

Temporal variation in habitat-use has been mentioned for various large Cracid species (Crax rubra griscomi (Martinez-Morales, 1999), Crax daubentoni: (Bertsch & Barreto, 2008b, Strahl et al., 1997a), Crax blumenbachii (Srbe-Araujo et al., 2012); Crax globulosa (Leite et al., 2018; Luna-Maira et al., 2013); Crax alector (Bernal & Mejia, 1995) and Mitu salvini (Parra et al., 2001; Santamaria & Franco, 2000; Yumoto, 1999). Parra et al. (2001) followed a pair of M. salvini over 6 months, including the wet and part of the dry season, and pointed

Figure 2. Location and Seasonal Home Ranges (95% and 50% href Kernel) for A and B and C Black Curassow Groups (Respectively A, B, C); Wet Season Home Ranges Are Figured With Dark Blue (95% Kernel) and Light Blue Lines (50% Kernel), and Dry Season Home Ranges Are Figured With Red (95% Kernel) and Orange Lines (50% Kernel), Grey Lines: Elevation.

Figure 3. Average Distance Travelled (M) per Minute in Each Hour of the Day for Black Curassow, by Season. Upper and lower 95% confidence intervals are figured by pink and blue areas.
out marked changes in habitat use over time. In contrast, Leite et al. (2018) noted for Crax globulosa the consistency in home range size across dry and wet seasons, the species locations being highly restricted to rivers and floodplains. A family group of Black Curassows followed for 7 months in Colombia had a smaller home range during the wet season (39 ha in September) than during the dry season (127 ha in December: Bernal & Mejía, 1995). The seasonal home ranges estimated in our study were similar to those estimated by Bernal and Mejía (1995) in both seasons, larger than those estimated for M. salvini during the dry season but apparently similar during the wet season. On the other hand, whatever the season, they were four to six times smaller than those estimated for C. globulosa (see Table 4). During the dry season, our tracked birds increased their home range but did not move significantly to other parts of the forest since home range overlap between the two seasons always exceeded 70%. This indicates that, as the dry season progressed, Black Curassows did not make migration-like movements towards other parts of the forest but rather extended the boundaries of their home range. For one female, however, the Kernel estimation of home range was not higher in the dry season, although the MCP estimation was.

As suggested in other studies on curassow species, seasonal differences in home range may reflect changes in habitat use in relation to variation in resource abundance throughout the year (Bertsch & Barreto, 2008b; Luna-Maira et al., 2013; Muñoz & Kattan, 2007; Parra et al., 2001; Santamaria & Franco, 2000; Srbek-Araujo et al., 2012). According to the study and species, movements may be reduced during fruit or water scarcity (Santamaria & Franco, 2000 for Mitu salvini and Leite et al., 2018 for Crax globulosa), or expanded (Parra et al., 2001, for Mitu salvini). More generally, movement ecology predicts that when resources become scarce, straighter and faster movements become more efficient than tortuous ones, as they result in high net displacement, thereby minimizing the chance of revisiting an already visited resource, increasing the chance of finding new resources, and giving time for stock to replenish (de Knegt et al., 2007; Viswanathan et al., 2008). This pattern would correspond to our results, showing an increase of travelled distances and home range size during the dry season, while the fructification peak is occurring during the wet season (February to May: Sabatier, 1985). Few tree species fruit during the dry season (Sabatier, 1985), and that may influence the ranging behaviour of frugivorous animals. For example, the Lauraceae Ocotea cf. cinerea, and the Moraceae Naucleopsis guianensis fruiting period occurred during the dry season only, and were heavily consumed at this time by the Black Curassows we tracked. These key resources were scattered, and located at the periphery of the home ranges in
our study area, probably contributing to the extension of the home range in dry season, (in the western part for group B, and in the southern part for animal C). During these months, birds were often observed walking straight for longer time towards these remote fruiting trees, and spending more time feeding on the same place, before going back to the core area for the night, while in the wet/fruiting season, feeding bouts were shorter and more regularly distributed during the day. Water availability may not greatly influence birds’ movement in our study area, as most streams flowed permanently, even during the dry season. Studies on Crax alector in French Guiana (Erard et al., 1991) and eastern Colombia (Jiménez et al., 2001) have shown little variation in diet composition among seasons. Consumption of seeds and seedlings increased during the rainy season in the Colombian study, while in French Guiana, the amount and diversity of fruits in the diet remained stable between seasons, but the specific composition was not studied (Erard et al., 1991).

Our results also revealed a high degree of home range overlap between family groups, also affecting the core area, and increasing during the dry season. Home range overlapping had already been mentioned in two Crax species (Delacour & Amadon, 2004) but not quantified, while home range of adult males in a reintroduced Red-billed Curassow population did not overlap and were often separated by distances of up to 3 km (Bernardo et al., 2011b).

We conducted our study in an area where hunting is absent and within a mountainous habitat: in French Guiana, mountainous-forested habitats have been shown to be favourable for this species and associated with the highest curassow densities measured (Denis et al., 2016). The locally high density of Black Curassows in our study area might therefore result in a high degree of overlapping in home ranges.

**Implications for Conservation**

This study allowed to improve our knowledge on the movement behavior of Black Curassows, which is crucial for the management and conservation of such key species (Allen & Singh, 2016). Overall, our estimates of both annual and seasonal home ranges were markedly larger than previous empirical estimates in French Guiana (Erard et al., 1991; Niel et al., 2008) but smaller than some estimates for the species from Colombia, at least for annual and wet season home ranges (Bernal & Mejia, 1995). The home range radius was included by Niel et al. (2008) as a parameter in the hunting sustainability index (SI), with a larger value increasing SI estimates by decreasing the Harvest rate (Harvest rate \( H = \text{Offtake/Area}; \) Niel & Lebreton, 2005; Slade et al., 1998). The impact of hunting is particularly sensitive to parameters such as the size of the hunted area, harvest rate and density, which need detailed ecological studies to be improved (Niel et al., 2008). Previous studies on hunting sustainability underlined the major role of source/sink processes (Richard-Hansen et al., 2019), in which the dispersal ability of species, related to their spatial behavior, has strong implications for population recovering. Moreover, the seasonal variation in home range area and overlap can potentially affect local densities throughout the year, and should be taken into account. Habitat also influences curassow densities (Denis et al., 2016), and resource abundance and distribution are likely to shape the birds movements and home range size. More data on resource needs, habitat use and spatial ecology in various areas will help to improve the management of this sensitive species in French Guiana.

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