

Natureza & Conservação

Brazilian Journal of Nature Conservation



Supported by O Boticário Foundation for Nature Protection

Research Letters

The width and biotic integrity of riparian forests affect richness, abundance, and composition of bird communities

Camila Crispim de Oliveira Ramos^{a,*}, Luiz dos Anjos^b

^a Programa de Pós-graduação em Ecologia de Ambientes Aquáticos Continentais, Universidade Estadual de Maringá – UEM, Maringá, PR, Brazil ^b Departamento de Biologia Animal e Vegetal, Universidade Estadual de Londrina – UEL, Londrina, PR, Brazil

ARTICLE INFO

Article history: Received April 2013 Accepted April 2014

Keywords: Avian community Corridors Conservation Riparian habitats Transition zones

ABSTRACT

In riparian forests, width contributes most importantly to maximizing diversity. Therefore, corridors with different widths should differ in richness, abundance, and composition. We tested this hypothesis for the bird communities of two forests on the Upper Paraná River floodplain, Paraná, Brazil. Richness and abundance were higher in riparian forest with mean width of 50 m in each margin and lower anthropogenic disturbance. Species diversity increased 30%, with increase in total width from 40 m to 100 m on average. Bird species composition also differed, and groups with the strictest ecological requirements were better represented in the wider, better-preserved forest. This indicates that conservation of riparian forests has a positive effect on their bird communities. We suggest that these environments are prioritized for recuperation, and that a 50 m width on each side of a stream is necessary for riparian forests to effectively fulfill their function in the landscape. We also note that the recently discussed Brazilian Forest Code does not conform to this requirement.

© 2014 Associação Brasileira de Ciência Ecológica e Conservação. Published by Elsevier Editora Ltda.

Introduction

In fragmented landscapes, the survival of species depends on their ability to persist in fragments and/or move across the landscape (Lees & Peres 2008). Riparian forests are important corridors for many biological groups in fragmented landscapes, because they promote increased connectivity and hence species richness and flow of individuals (Lees & Peres 2008). These corridors increase genetic variability (Vieira & Carvalho 2008) and local biodiversity (Anjos *et al.* 2007), reduce climatic variations and their consequences (Marini *et al.* 2009), and allow forest organisms of adjacent biomes to disperse (Silva 1996). Riparian corridors are essential to maintain the diversity of plant and animal communities in many biomes, particularly in the Atlantic Forest (e.g., Metzger *et al.* 1997; Anjos *et al.* 2007).

^{*}Corresponding author at: Programa de Pós-Graduação em Ecologia de Ambientes Aquáticos Continentais, Universidade Estadual de Maringá, Av. Colombo, 5790, PEA, Bloco G90, 87020-900, Maringá, PR, Brazil.

E-mail address: mila_bru@yahoo.com.br (C.C.O. Ramos).

^{1679-0073/\$ -} see front matter. © Associação Brasileira de Ciência Ecológica e Conservação. Published by Elsevier Editora Ltda. DOI: 10.4322/natcon.2014.011

The available forest area along bodies of water is an important factor affecting the richness and species composition of a wide variety of organisms (Vieira & Carvalho 2008; Tubelis *et al.* 2004). The width of riparian corridors is the most important factor bene⊠ting biodiversity, and maximizing this width improves habitat quality by reducing edge effects (Metzger 2010). Other factors including the length, continuity, and degree of conservation of corridors (Lees & Peres 2008); and the surrounding matrix type, topography, and extent of the areas of riparian influence (Metzger *et al.* 1997) also influence the quality of riparian corridors.

The protection of these corridors, although present in the former Brazilian Forest Code (Federal Law No. 4.771 of September 15, 1965), was not effective. It is common to observe properties that contravene it, where the permanent preservation areas (PPA) along streams are fully or partially occupied (Sparovek et al. 2011). According to this law, riparian forests on the margins of streams up to ten meters wide should be 30 m in width on each side, but the new Forest Code provides for regularization of deforestation in areas of consolidated use, with reconstitution of the PPA, according to farm size. This reconstitution would be negligible, ranging from Øve meters on each side to a maximum of 15 m, and still allowing the use of these areas for activities of agroforestry, ecotourism, and rural tourism. Thus, the recovery of the functionality of this environment is compromised, since there will be no more incentive to restore them and their use and exploration will still be allowed (Sparovek et al. 2011).

In this context, we evaluated two riparian forests of different widths, with respect to the richness, composition, and abundance of their bird species. We tested the following hypotheses: the wider riparian forest (with less anthropogenic disturbance) will support greater species richness and abundance of individuals than the narrower forest (with greater anthropogenic disturbance); and species composition will differ between the forests, despite their spatial proximity (6 km apart). If these assumptions are correct, narrow riparian forests have limited importance for biodiversity conservation of forest birds in areas where loss of forest habitat is extreme and most remnants are restricted to the margins of water bodies.

Methods

Study areas

The study was conducted on the Upper Paraná River floodplain (UPR) in northwestern Paraná state, Brazil. This area is a transitional zone of the Atlantic Forest with the Cerrado (Mendonça *et al.* 2009). The riparian forests studied are west of the Paraná River, at an altitude of approximately 260 m, bordering the Caracu stream (22°45'55" S and 53°15'30" W, 4.5 ha), and the São Pedro River (22°44'58" S and 53°13'24" W, 11 ha) (Fig. 1). Both forests were intensively exploited and degraded by farming and urbanization, but in the last decade



Fig. 1 - Map of the Upper Paraná River floodplain, showing the riparian forests studied.

were fenced and allowed to regenerate naturally. The regional climate, according to the Köppen-Geiger system, is Cfa (humid subtropical mesothermal) with an annual mean temperature of 22°C (summer mean 26°C and winter 19°C) and mean annual rainfall of 1,500 mm. However, in some years, the climate may be Cwa (high-altitude tropical) with a tropical rainfall pattern and dry winters (Maack 2002). The vegetation is semideciduous seasonal forest (western boundary of the Atlantic Forest), which now covers only 1% of its original extent and occurs in a few small fragments and riparian remnants along the Paraná River and its tributaries (Campos & Souza 1997).

The Ørst riparian forest (RF1) averages 40 m wide (20 m on each side of the stream), but in some places narrows to 15 m in total. RF1 extends along the Caracu stream in the municipality of Porto Rico, between urban and rural areas. It is a discontinuous secondary forest, with a relatively open understory and some areas occupied by vines and lianas. The second riparian forest (RF2) averages 100 m wide (50 m on each side) in its narrower stretches, but the total width exceeds 100 m in several parts. This forest extends along the São Pedro River in the municipality of São Pedro do Paraná, in a rural area. It is a continuous secondary forest, with an open understory along one bank, and denser growth with tangles of vines and lianas on the opposite bank. RF2 has a better conservation level and lower anthropogenic disturbance.

Field method

We obtained the Øeld data along a transect 850 m in length through the two riparian forests, starting at the point where each stream enters the Paraná River. Data were gathered from September to November 2008, using the point count method (Blondel et al. 1981) as adapted by Anjos (2007) for studies of forest fragments, which was suficient to detect 75% of forest species diversity in these sites. Samples were taken monthly on four consecutive days, two in each forest. We allocated four points along predetermined trails in each forest, 200 m distant from each other and 100 m from the beginning of the transect. We began sampling at dawn and ended after four hours. Each point was sampled twice in the morning, in the sequence 1, 2, 3, 4, and then 4, 3, 2, 1. The next morning, the order was reversed. We remained for 15 min at each point, performing observations, with a 15-min interval between points. We included visual and auditory records of the species present within a radius limit, taking care not to record species outside the forest; each couple or group (for social species) was considered a contact. We recorded unrecognized vocalizations with a portable recorder and unidirectional microphone for later identiØcation at the Laboratory of Ornithology and Bioacoustics, Universidade Estadual de Londrina. From the quantitative survey data, the point abundance (IPA) for each species was calculated by dividing the number of contacts by the total number of points sampled in each area (Blondel et al. 1981).

Procedures for analysis

We included only species that are most dependent on forest habitats, according to Parker III *et al.* (1996); birds that live in the habitat matrix and are occasionally recorded in the forest edge were not included, e.g., *Crotophaga ani, Mimus saturninus*, and Furnarius rufus. We also excluded nocturnally active birds (families Strigidae, Caprimulgidae, and Nyctibiidae), and the families Accipitridae, Falconidae, and Throchilidae because of their high mobility and/or different habits.

To evaluate differences in species composition between the forests, the species were classi⊠ed in different groups based on data for habitat use, food habits, endemism, and distance of the UPR from the geographic distributional limit of the species. Regarding habitat use, we included strictly forest species, those that only occur in forest formations as de⊠ned by Parker III *et al.* (1996). Regarding food habits, we classi⊠ed the species as frugivores, insectivores, or omnivores according to Anjos & Schuchmann (1997), using the frugivore classi⊠cation of Mendonça *et al.* (2009). We considered endemic species to be those restricted to the Atlantic Forest or to central South America, according to Parker III *et al.* (1996).

Statistical procedures

We evaluated whether the sampling effort was suficient to obtain a representative number of bird species in each forest through accumulation curves and estimates of species richness. The abundance-based coverage estimator (ACE) was used, because it considers abundant species, i.e. those represented by more than ten individuals, besides considering singletons and doubletons (Lee & Chao 1994). Rarefaction curves of species for each area were constructed to evaluate the difference in richness between the areas. For both analyses, we used the software EstimateS 7.5.2 and Statistica 7.0. We used the G test with a correction factor to test for differences between the number of contacts (corrected for sampling effort, i.e., IPA x 100) of each species in both forests. When the expected number of contacts was less than Øve, we calculated the exact probabilities for the binomial test; the value of α was set at 0.01 for both tests.

We ordered the sampling points in RF1 and RF2 based on species composition and abundance. We used a correspondence analysis (CA) by assigning less weight to rare species, so they would not affect the ordination. As a criterion for retention and interpretation of the CA axes, we used only axes with eigenvalues greater than 0.20, as recommended in the literature (Manly 2008), and used PC-ORD 3.5 to perform this analysis. The scores from this analysis were submitted to a univariate analysis of variance (ANOVA) to investigate possible differences in these community structures. The community structure based on each particular group (habitat use, food habits, endemism, and distance from the distributional limit), was evaluated by the scores of species in parametric ANOVA, and by a nonparametric test (Kruskal-Wallis) when the conditions for the ANOVA were not satisized; both tests were performed with Statistica 7.0. The full summary of the research papers and software packages used in statistical methodology are provided in the supplementary material online.

Results

In the two riparian forests, we recorded a total of 74 species of forest birds: 49 in RF1 and 70 in RF2. The accumulation curve and estimation of species richness indicated that data from RF1

were closer to the asymptote, while in RF2, richness still tended to increase with sampling effort (Fig. S1A, supplementary material online). Rarefaction curves showed that richness (independently of abundance) in RF2 was higher than in RF1 (Fig. S1B, supplementary material online). Four species were unique to RF1 and 25 were unique to RF2. Among the 45 species common to both areas, four were more abundant in RF1, 14 more abundant in RF2, and 28 showed similar abundances (Table S1, supplementary material online).

In the correspondence analysis, the riparian forests formed two groups, separating the RF1 and RF2 sampling points (Fig. 2A). Only the Ørst CA axis was retained for interpretation (Table S2, supplementary material online). The sampling areas differed signi⊠cantly, as con⊠rmed by the analysis of variance from the scores generated for axis 1 ($F_{11, 84} = 16.47$, p < 0.001, Fig. 2B).

To evaluate how these communities were structured according to the species characteristics, we used the scores generated for the species, relative to axis 1, to construct the graphs. These graphs show how each group of species are organized in the space in a direct relationship with the ordination of the sample points. Regarding the use of habitat (Fig. 3A), edge species predominated in RF1, and forest species in RF2 (KW-H_{2, 74} = 7.26, p < 0.05, Fig. 3B).

Regarding feeding habits (Fig. 3C), insectivorous species predominated in RF2, and frugivores in RF1 (KW-H_{2, 74} = 7.76; p < 0.05, Fig. 3D).

Finally, endemic species (Fig. 3E) predominated in RF2 (KW-H $_{1.74}$ = 4.20, p < 0.05, Fig. 3F).

Discussion

The data from this study demonstrated that the difference in forest width, combined with different degrees of conservation of the riparian corridors, altered the richness, abundance, and species composition of forest birds. The ability of these riparian forests to support groups of species with stricter ecological requirements increased when both aspects of these particular forest formations (width and anthropogenic disturbance) were optimized. Forest bird species diversity increased 30%, with increase in total width from 40 m to 100 m on average. Studies of arboreal vegetation in the Atlantic Forest showed that corridors less than 100 m wide have limited effectiveness in maintaining diversity (Metzger *et al.* 1997). For birds in the Cerrado, Tubelis *et al.* (2004) postulated that the minimum width of native vegetation should be 120 m. According to the authors, this range allows the conservation both of birds associated with riparian forests, and those that are dependent on adjacent savannas.

The community composition found in RF1, where edge species predominated, mainly reflects the influence of the forest width on the community structure, although other factors, such as the intense human presence from the adjacent urban landscape and the resulting lower biotic integrity, likely also had an important role in the results obtained. The width of riparian forest affects habitat quality and regulates the area impacted by edge effects, i.e., the microclimatic changes and the increase in disturbances at the edges of these habitats (Metzger 2010). According to Metzger, these effects can vary in extent depending on the species and processes in question, and also according to the physical characteristics of the site, in particular the solar direction, latitude, and adjacent matrix type, which influence the amount of incident solar radiation. In general, stronger effects occur in the Ørst 100 m (Laurance et al. 2002), implying that corridors less than 200 m wide are essentially highly disturbed edge environments. Strictlyforest species would need corridors at least 200 m wide (Lees & Peres 2008).

In this study, examination of three parameters indicated that the better-preserved riparian forest harbors a set of species with stricter ecological requirements. The \boxtimes rst parameter, which takes into account habitat use, indicated that the presence of a higher proportion of strictly-forest species in RF2 indicates that it is a higher-quality habitat. Greater biotic integrity and a less-disturbed surrounding landscape allow the occurrence of more-specialized groups such as Dendrocolaptidae, which have stricter ecological requirements, according to Poletto *et al.* (2004).



Fig. 2 – Graphical representations of (A) ordination of samples per day for RF1 (RF11, RF12, RF13, RF14, RF15, RF16) and RF2 (RF21, RF22, RF23, RF24, RF25, RF26) and graphs of the variance analysis for the axis retained (Axis 1), with (B) scores of samples from both areas.



Fig. 3 – Graphical representation of scores generated for species grouped by preferential habitat (A), through the ranking of point samples from RF1 and RF2 and graphs of the variance analysis for the axis retained (Axis 1), with (B) scores of species grouped by habitat preference; graphical representation of scores generated for species grouped by feeding habit (C), through the ranking of point samples from RF1 and RF2 and graphs of the variance analysis for the axis retained (Axis 1), with (D) scores of species grouped by feeding habit; graphical representation of scores generated for species grouped by the presence of endemism (E), through the ranking of point samples from RF1 and RF2 and graphs of the variance analysis for the axis retained (Axis 1), with (D) scores of species grouped by feeding habit; graphical representation of scores generated for species grouped by the presence of endemism (E), through the ranking of point samples from RF1 and RF2 and graphs of the variance analysis for the axis retained (Axis 1), with (F) scores of species grouped by the presence of endemism.

The second parameter analyzed was the preferred feeding habit of the species, and we found that insectivores were more concentrated in RF2. Insectivores, especially those that forage in lower strata, are considered to be the most vulnerable species by many investigators (Ribon *et al.* 2003). Considering that this area also harbored more strictly-forest species and that many groups of insectivores have relatively limited mobility through the landscape (e.g., Lees & Peres 2009), these characteristics probably act to concentrate most species that are sensitive to fragmentation in this area. According to Uezu & Metzger (2011), to understand the sensitivity of species to habitat fragmentation, it is necessary to consider the multiple dimensions of features of the species.

We found a higher prevalence of endemic species - the third parameter analyzed - in RF2, which provides evidence that for the maintenance of species dynamics in a particular region, the quality of the corridors is a key aspect.

This study demonstrated that maintaining riparian forests intact can have a positive effect on bird conservation. The data presented here suggest that in the case of streams (where the requirement, according to the Forest Code, Law 4.771/65, is 30 m on each side), the PPAs should be expanded to a minimum of 50 m on each side of a stream, to aid in conserving species with stricter ecological requirements.

Acknowledgments

We thank Dilermando Pereira de Lima Júnior and Nadson Ressié Simões for revision of this manuscript. We are also grateful for the cooperation of the staff of Laboratório de Ornitologia e Bioacútica - UEL and Núpélia-UEM. Financial support was provided by CNPq (process no. 130294/2008-0), PELD - Site 6 and PEA/UEM.

Supplementary material

Supplementary material associated with this article can be found, in the online version, at www.naturezaeconservacao.com.br.

REFERENCES

Anjos L, 2007. A e⊠ciência do método de amostragem por pontos de escuta na avaliação da riqueza de aves. Ararajuba, 15(2):239-243.

Anjos L & Schuchmann K-L, 1997. Biogeographical af⊠nities of the avifauna of the Tibagi Basin, Paraná drainage system, southern Brasil. *Ecotropica*, 3(1):43-65.

Anjos L et al., 2007. The importance of riparian forest for maintenance of bird species richness in an Atlantic Forest remnant, southern Brazil. *Revista Brasileira de Zoologia*, 24(4):1078-1086.

Blondel D, Ferry C & Frochot B, 1981. Point counts with unlimited distance. Studies in Avian Biology, 6:414-420.

Campos JB & Souza MC, 1997. Vegetação. In Vazzoler AEAM; Agostinho AA & Hahn NS (eds.). Planície de inundação do alto rio Paraná: aspectos físicos, biológicos e socioeconômicos. Maringá: Eduem/Nupélia. p. 331-342.

Laurance WF et al., 2002. Ecosystem decay of Amazonian forest fragments: a 22-year investigation. *Conservation Biology*, 16: 605-618.

Lee S-M & Chao A, 1994. Estimating population size via sample coverage for closed capture-recapture models. *Biometrics*, 50(1): 88-97.

Lees AC & Peres CA, 2008. Conservation value of remnant riparian forest corridors of varying quality for Amazonian birds and mammals. *Conservation Biology*, 22: 439-449.

Lees AC & Peres CA, 2009. Gap-crossing movements predict species occupancy in Amazonian forest fragments. Oikos, 118: 280-290

Maack R, 2002. Geografia física do estado do Paraná. 3th ed. Curitiba: Imprensa O⊠cial do Paraná.

Manly BJF, 2008. Métodos estatísticos multivariados: uma introdução. 3th ed. Porto Alegre: Bookman.

Marini MA et al., 2009. Predicted climate-driven bird distribution changes and forecasted conservation conflicts in a neotropical savanna. *Conservation Biology*, 23: 1558-1567.

Mendonça LB, Lopes EV & Anjos L, 2009. On the possible extinction of bird species in the Upper Paraná River floodplain, Brazil. Brazilian Journal of Biology, 69(2 suppl):747-755.

Metzger JP, 2010. O Código Florestal tem base cientí 2 ca? Natureza & Conservação, 29:92-99.

Metzger JP, Bernacci LC & Goldenberg R, 1997. Pattern of tree species diversity in riparian forest fragments with different widths (SE Brazil). *Plant Ecology*, 133: 135-152.

Parker III TA, Stotz DF & Fitzpatrick JW, 1996. Ecological and distributional databases. In Stotz DF et al. (eds.). Neotropical birds. Ecology and Conservation Chicago: The University of Chicago Press. p.113-436.

Poletto F et al., 2004. Caracterização do microhabitat e vulnerabilidade de cinco espécies de arapaçus (aves: Dendrocolaptidae) em um fragmento florestal do norte do estado do Paraná, sul do Brasil. Revista Brasileira de Ornitologia, 12 (2):89-96.

Ribon R, Simon JE & de Mattos GT, 2003. Bird extinctions in Atlantic forest fragments of Viçosa region, southeastern Brazil. *Conservation Biology*, 17:1827-1839.

Silva JMC, 1996. Distribution of Amazonian and Atlantic birds in gallery forests of the Cerrado region, South America. Ornitologia neotropical, 7(1):1-18.

Sparovek G et al., 2011. A revisão do Código Florestal Brasileiro. Novos Estudos CEBRAP, 89:111-135.

Tubelis DP, Cowling A & Donnelly C, 2004. Landscape supplementation in adjacent savannas and its implications for the design of corridors for forest birds in the central Cerrado, Brazil. *Biological Conservation*, 118:353-364.

Uezu A & Metzger JP, 2011. Vanishing bird species in the Atlantic Forest: relative importance of landscape con⊠guration, forest structure and species characteristics. *Biodiversity and Conservation*, 20:3627-3643.

Vieira FA & Carvalho D, 2008. Genetic structure of an insectpollinated and bird-dispersed tropical tree in vegetation fragments and corridors: implications for conservation. Biodiversity Conservation, 17(10):2305-2321