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1 Original research

2

3 **The impact of natural resource use on bird and reptile**  
4 **communities within multiple-use protected areas: evidence from**  
5 **sub-arid southern Madagascar**

6

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18

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25 **Abstract**

26 Multiple-use protected areas, in which sustainable levels of extractive livelihood activities are  
27 permitted, play an increasingly important role in the global protected area estate, and are  
28 expected to rise in prevalence. However, we know little about their effectiveness at  
29 conserving biodiversity. We surveyed bird and reptile communities in three areas across a  
30 forest disturbance gradient resulting from charcoal production and shifting cultivation within  
31 a multiple-use protected area in Madagascar's sub-arid spiny forest. We scored individual  
32 species using a Conservation Value Index (CVI; a simple metric based on rarity, threat and  
33 distinctiveness), and estimated the total conservation value of each treatment by calculating  
34 the sum of frequency-weighted CVI scores across all present species. Bird and reptile  
35 community responses to forest disturbance were idiosyncratic. Bird richness was greatest in  
36 the moderate-disturbance treatment, but the low-disturbance treatment had the superior  
37 conservation value due to higher frequencies of locally-endemic species. Reptile richness was  
38 the same in low- and moderate-disturbance treatments, but the conservation value of the latter  
39 was greater. The high-disturbance areas had lowest richness and conservation value for both  
40 groups. For birds, increasing disturbance levels were accompanied by community turnover  
41 from high-value to low-value species, a pattern highlighted by CVI that is masked by  
42 assessing species richness alone. Although some endemic species appear to be resilient to  
43 degradation, multiple-use protected areas in Madagascar may lose biodiversity since most  
44 endemic species are forest-dependent. Stricter protected area models may be more  
45 appropriate in areas where much of the high-value biodiversity is sensitive to habitat  
46 degradation.

47

48 **Keywords:** Conservation value; Degradation; Dry forest; Faunal communities; Sustainable  
49 Use

50

51

## 52 **1 Introduction**

53 The impacts of human activity now threaten most of the Earth's species and ecosystems  
54 (Ehrlich and Pringle 2008) and have precipitated the planet's sixth mass extinction (Barnosky  
55 et al. 2011). Our primary strategy to stem this biodiversity loss is the creation and  
56 management of protected areas, which cover over 15 % of the world's land area and  
57 constitute the largest planned land use in history (Juffe-Bignoli et al. 2014). All protected  
58 areas are spaces "recognised, dedicated and managed... to achieve the long-term conservation  
59 of nature with associated ecosystem services and cultural values" (Dudley 2008), but they  
60 vary greatly in management objective and approach. These differences form the basis for the  
61 World Conservation Union's (IUCN) protected area categorisation system (Dudley 2008;  
62 Dudley et al. 2010). For simplicity's sake the categories are often divided into 'strict'  
63 protected areas (generally categories I-IV), which seek to isolate nature from human  
64 processes that threaten it, and 'multiple-use' sites, which promote conservation through the  
65 sustainable extractive use of natural resources (category VI) or traditional land uses that  
66 sustain biodiversity (category V).

67

68 Recent decades have seen the number of multiple-use protected areas grow significantly in  
69 many parts of the world (Juffe-Bignoli et al. 2014). Although some strict sites have been  
70 downgraded (Mascia et al. 2014), this has been driven primarily by the predominance of  
71 multiple-use categories amongst new protected areas (Zimmerer et al. 2004). The trend can  
72 largely be attributed to: i) the lack of remaining 'wilderness' areas, with a low human  
73 footprint, suitable for the creation of strict categories (Leroux et al. 2010); and, ii) a paradigm  
74 shift in conservation, reflecting concern for the effects of exclusionary approaches on human  
75 wellbeing (Adams and Hutton 2007; Miller 2014), and the suggestion that sustainable use

76 may be a more effective long-term conservation strategy than strict protection (Rosser and  
77 Leader-Williams 2010). As a result, only 45 % of the world’s protected areas are assigned to  
78 categories I-IV (Jenkins and Joppa 2009), and category VI sites expanded from 14 to 32 % of  
79 the world’s protected area estate (by area) between 1990 and 2010 (Bertzky et al. 2012). This  
80 trend is expected to become even more pronounced in the future (McDonald and Boucher  
81 2011).

82

83 Signatories to the Convention of Biological Diversity are expected to increase the coverage  
84 of terrestrial protected areas to 17 % of their national territory by 2020 and ensure that they  
85 are “effectively managed” (CBD 2010, Aichi Target 11), a target that will require the most  
86 rapid expansion of protected areas in history (Venter et al. 2014). Thus, if new protected  
87 areas are expected to largely comprise multiple-use categories, it is important to know  
88 whether they are likely to be successful at achieving their objective – the long-term  
89 conservation of nature – in the face of authorised human impacts (Dudley et al. 2014; Watson  
90 et al. 2016). This is particularly apposite given longstanding debates over the contribution of  
91 multiple-use protected areas to conservation goals (Locke and Dearden 2005; Gaston et al.  
92 2008; Shafer 2015).

93

94 The effectiveness of protected areas depends on both their coverage (i.e. ensuring that  
95 maximum biodiversity is represented within them) and their success in buffering the  
96 biodiversity from the processes that threaten its viability (Gaston et al. 2008; Watson et al.  
97 2014). However, research tends to concentrate on the former (e.g. Montesino Pouzols et al.  
98 2014; Venter et al. 2014; Butchart et al. 2015; Polak et al. 2015; Visconti et al. 2015), with  
99 the result that we know little about the success of protected areas in maintaining their  
100 condition over time (Cabeza 2013; Geldmann et al. 2013; Beaudrot et al. 2016; Watson et al.

101 2016). This knowledge gap is particularly acute with regards to multiple-use categories.  
102 Global studies comparing across categories have found stricter protected areas to be more  
103 effective at slowing deforestation in some regions (Joppa and Pfaff 2011; Scharlemann et al.  
104 2010), whereas multiple-use sites demonstrate greater success in other countries (Ferraro et  
105 al. 2013; Nelson and Chomitz 2011). However, the use of remote sensed data within such  
106 analyses only allows us to quantify vegetation cover, therefore providing little insight into the  
107 ecological integrity of remaining natural vegetation and faunal communities beneath the  
108 canopy (Peres et al. 2006; Beaudrot et al. 2016). Less conspicuous changes to forest structure  
109 and composition (i.e. forest degradation) can stem from activities such as non-industrial  
110 selective logging, fuelwood collection, livestock grazing and the harvesting of non-timber  
111 forest products (NTFPs). Typically, these are precisely the types of activity that may be  
112 sanctioned within category V and VI protected areas (Dudley 2008). Indeed, conservationists  
113 still have a very limited understanding of species and community responses to habitat change,  
114 and our knowledge is largely derived from a small number of sites (Barlow et al. 2007; T.  
115 Gardner et al. 2009, 2010). Furthermore, few researchers have investigated the impacts of  
116 subsistence activities on biodiversity (Borghesio 2008; Brown et al. 2013).

117

118 Madagascar is an example of a biodiversity-rich tropical developing country that is  
119 expanding its protected area system through the creation of new multiple-use sites. The island  
120 is a global conservation priority, boasting an unparalleled combination of species diversity  
121 and endemism (Brooks et al. 2006), with the majority of its endemic biota being forest  
122 dependent (Goodman and Benstead 2005). However, less than 16 % of the country retained  
123 forest cover by 2000 (Harper et al. 2007; McConnell and Kull 2014). Since 2003,  
124 Madagascar has been in the process of tripling the coverage of its protected area system, from  
125 1.7 to over 6 million ha, in response to lobbying from international conservation

126 organisations and funders (Corson 2014). Known as the ‘Durban Vision’ after the location of  
127 the fifth World Parks Congress at which it was launched, this ambitious programme has  
128 necessitated modifications to the country’s conception of protected areas and their  
129 governance. Previously, all protected areas were governed by the State, managed by the para-  
130 statal Madagascar National Parks, and comprised only strict categories (I, II and IV;  
131 Randrianandianina et al. 2003). Most of the new protected areas established as part of the  
132 Durban Vision are co-managed by non-governmental organisations (NGOs) and local  
133 communities, and are proposed or designated as categories V and VI (AGRECO 2012;  
134 Gardner 2011; Virah-Sawmy et al. 2014), with zoned areas where subsistence and low-level  
135 commercial natural resource use activities are permitted (e.g. Gardner et al. 2008; Virah-  
136 Sawmy et al. 2014; WWF 2010).

137

138 The goals of the expanded Madagascar Protected Area System (SAPM) are to conserve the  
139 country’s unique biodiversity and its cultural heritage, as well as promoting the sustainable  
140 use of natural resources for poverty alleviation and development (Commission SAPM 2006).  
141 The simultaneous achievement of these goals is particularly complex because most forms of  
142 traditional land and resource use in Madagascar have negative impacts on biodiversity  
143 (Gardner 2009, 2011; Irwin et al. 2010). Planning the management of new multiple-use  
144 protected areas requires an understanding of species and community responses to habitat  
145 degradation arising from permitted resource use, yet our knowledge of the influence this has  
146 on biodiversity is patchy for the country as a whole, and particularly for the globally-  
147 important spiny forest ecoregion (Irwin et al. 2010). Moreover, existing studies in  
148 Madagascar tend to mirror patterns in global research (Burivalova et al. 2014) by  
149 summarising assemblage-level change via species richness (e.g. Randriamiharisoa et al.  
150 2015; Scott et al. 2006). In other words, while studies may investigate the ecological or other

151 attributes of species remaining in degraded habitats, their results are usually reported in terms  
152 of species richness, but this measure has been criticised because it can mask community  
153 turnover from specialists to generalists (Barlow et al. 2007; Gardner et al. 2010). Here we  
154 investigate bird and reptile community responses to habitat change in a new protected area in  
155 the spiny forest ecoregion to ascertain the impacts of permitted and illegal livelihood  
156 activities (charcoal production and shifting cultivation respectively) on the conservation value  
157 of the vertebrate fauna. To overcome the issues associated with species richness as a metric,  
158 we use a novel Conservation Value Index (CVI) to examine the influence of habitat  
159 degradation on the two taxonomic assemblages.

160

## 161 **2 Methods**

### 162 **2.1 Study site**

163 Madagascar's spiny desert (or spiny forest), is a global priority ecoregion (Olson and  
164 Dinerstein 1998) and Endemic Bird Area (Stattersfield et al. 1998) with extremely high rates  
165 of local floral endemism (Phillipson 1996). Between 1990 and 2010 it suffered the fastest  
166 rates of deforestation of any ecoregion in the country (Harper et al. 2007; ONE et al. 2013)  
167 and, prior to 2003, it was the least represented ecoregion within the country's protected area  
168 network (Fenn 2003).

169

170 Ranobe PK32 is a new protected area that received temporary protected status within the  
171 Durban Vision framework in 2008, and is co-managed by local community associations, the  
172 regional Forest Service and the international NGO WWF (Virah-Sawmy et al. 2014). Lying  
173 north of the regional capital Toliara between the Fiherenana and Manombo rivers (Fig. 1), it  
174 is the richest landscape in the ecoregion in terms of its bird, reptile and lemur fauna (Gardner  
175 et al. 2009a,b; 2015a). However, the area is inhabited by approximately 90,000 people (WWF

176 2010), many of whom depend on natural resources from within and around the protected area  
177 for their subsistence and household income (Gardner and Davies 2014; Gardner et al. 2015b).  
178 Ranobe PK32 is thus proposed as a category VI protected area in which subsistence and low-  
179 level commercial livelihood activities (such as timber cutting, fuelwood collection and  
180 charcoal production, grazing and the harvesting of NTFPs) are permitted in sustainable use  
181 zones which cover 86.5 % of the protected area's 148,554 ha (Virah-Sawmy et al. 2014;  
182 WWF 2010).

183  
184 Charcoal is primarily produced in the western part of the protected area, due to the presence  
185 of the Route Nationale 9 (RN9) road that facilitates transportation. The industry is driven by  
186 the close proximity of Toliara, a city of approximately 200,000 people in which 98 % of  
187 households use wood or charcoal for cooking; demand from the city tripled between 2000  
188 and 2007, and is largely met by anarchic charcoal production along the RN9 (Gardner et al.  
189 2015b; Partage 2008). Since the region lacks fuelwood plantations, charcoal is produced  
190 entirely from natural forests (Bertrand et al. 2010). Charcoal producers select only hardwood  
191 trees (Randriamalala et al. 2016), thus causing forest degradation rather than outright  
192 deforestation (Casse et al. 2004).

193  
194 We conducted our study in the vicinity of Ranobe, a complex of three villages with a total  
195 population of approximately 2000 people (Gardner and Davies 2014), where the surrounding  
196 forests had been subjected to both charcoal production and shifting cultivation within recent  
197 years. We selected three areas within 3 km of the main village which, until recently, were  
198 part of a contiguous and relatively homogeneous forest block. Subsequently, the three areas  
199 have suffered varying levels of disturbance that are indicative of the habitat degradation  
200 gradient found across the whole landscape: i) a forest area showing minimal impacts of

201 human activity (low-disturbance, hereafter *Low*); ii) a forest area subject to intensive charcoal  
202 production (moderate-disturbance, *Mod*); and, iii) an area regenerating following shifting  
203 cultivation (high-disturbance, *High*). While *Low* and *Mod* retained a complex three-  
204 dimensional structure and can be termed forest, *High* was an open area dominated by shrubs,  
205 with only scattered trees (Fig. 1, Table 1). As there were no areas of forest near Ranobe that  
206 had not been subject to any human disturbance, it was not possible to include a control site  
207 representing intact habitat. Birds and reptiles were surveyed between January and March  
208 2010 in the rainy season, when both groups are most active (Glaw and Vences 2007; Safford  
209 and Hawkins 2013).

210

211 [Fig. 1]

212

213 [Table 1]

214

## 215 **2.2 Bird survey protocol**

216 We established 48 census stations within each area and used the point count method (Bibby  
217 et al. 1998) to estimate bird relative abundance. Access to the forest interior was hindered by  
218 the impenetrable nature of the vegetation at *Low* and *Mod*, so census stations were placed on  
219 a stratified random grid along existing ox-cart tracks. We positioned all stations at a  
220 perpendicular distance of 75 m from a track (following Jones et al. 1995) to minimise the  
221 influence of edge effects, and at least 150 m apart to minimise the risk of double counting.

222

223 We surveyed each census station for 15 minutes (following a settling period of four minutes  
224 after arrival), during which we recorded all visual and auditory contacts within 50 m of the  
225 census station. To reduce time-of-day and weather-related effects, surveys were limited to

226 between 06.00 and 08.00 and were not conducted on rainy or windy days. The majority of  
227 bird contacts in spiny forest (> 85 % at *Low* and *Mod*) were auditory due to the dense  
228 vegetation, thus making it difficult to generate reliable distance estimates for bird contacts  
229 and, as such, we did not employ distance sampling methods. However, the non-visual nature  
230 of most contacts reduces the likelihood of a detectability bias arising from surveying in  
231 forests of varying degradation levels (Bibby and Buckland 1987). The auditory nature of  
232 most contacts also meant that we could not accurately count the number of individuals for  
233 social species, and thus we recorded the presence of groups not individuals. We did not  
234 include contacts with juvenile birds in our data analysis to reduce seasonality effects. Point  
235 count observations yielded both relative frequency (defined as the proportion of counts in  
236 which a given species was recorded) and relative abundance (mean number of contacts of a  
237 given species per count) data.

238

### 239 **2.3 Reptile survey protocol**

240 We calculated the relative abundance of reptiles based on capture in pitfall traps and area  
241 constrained refuge searches (transects), because observation and capture-based methods  
242 permit the sampling of different components of the reptile fauna (Raselimanana 2008). For  
243 pitfall trapping we followed a standard protocol widely used in Madagascar (D’Cruze et al.  
244 2007; Raselimanana 2008). The traps consisted of plastic buckets (270 mm deep, 290 mm  
245 internal diameter at top, 220 mm internal diameter at base) placed 10 m apart and buried in  
246 the ground with the rim level with the surface. Drainage holes were drilled in the bottom of  
247 each bucket and the handles were removed. Buckets were connected by a drift fence 500 mm  
248 high, passing directly over the centre of each bucket, constructed from a sheet of plastic  
249 supported by wooden stakes. The lower 50 mm of the fence was buried in the soil and  
250 covered with leaf litter to prevent animals passing underneath. Within each treatment we

251 established three trap lines (each of 10 or 11 buckets), placed randomly, but at least 150 m  
252 apart. Traps were constructed in the morning and left open for 13 nights, equating to 403 trap  
253 nights in total per area, and were checked at 07.00 and 16.00 each day. All captured animals  
254 were marked on the hind leg or ventral surface with nail polish, and released at the site of  
255 capture. Recaptured individuals were excluded from the data analysis.

256

257 We also established 38 transects along which we conducted active refuge searches. Each  
258 transect consisted of a 50 m rope erected adjacent to forest tracks based on a stratified  
259 random grid. Each transect was at least 150 m apart, ran perpendicular to a track and started  
260 10 m into the forest to reduce the influence of edge effects. We established each transect 24  
261 hours prior to surveying to minimise disturbance effects. During surveys, two observers  
262 moved slowly along each transect and searched for reptiles within 2 m of the central line,  
263 scanning the trunks and branches of trees, searching within tree holes, under bark, in the leaf  
264 litter and under/within dead branches. All reptiles initially observed within 2 m of the central  
265 line were recorded. Transects were walked from 08.00-10.00 (n = 22/site) and 15.00-17.00 (n  
266 = 16/site); we did not survey during periods of rain or thick cloud cover to minimise variation  
267 in weather-related detectability, which reduced the number of appropriate afternoon survey  
268 periods. Juveniles were omitted from the dataset to minimise any bias that might be  
269 associated with the effects of the breeding season. Transects and pitfalls generated density  
270 and capture rate data, respectively: we pooled the data and used total contacts for further  
271 analyses (not including rarefaction).

272

## 273 **2.4 Data analysis**

274 In order to compare species richness between treatments and estimate the completeness of  
275 our sampling, we generated individual-based observed species richness rarefaction curves

276 and associated 95 % confidence intervals using EstimateS v.9.0 (Colwell 2013). For reptiles,  
277 we combined the two datasets by assigning species to one or other method on the basis of  
278 substrate use, following a protocol adapted from Bicknell et al. (2015), whenever a species  
279 was recorded by both methods. Thus all arboreal species were assigned to transects and all  
280 terrestrial and fossorial lizards were assigned to pitfall traps. Remaining terrestrial species  
281 (snakes and a tortoise) were assigned to the method by which they were most frequently  
282 recorded. We used chi-squared contingency tables to test for homogeneity of observed  
283 species relative frequency (birds) or total contacts (reptiles) across treatments.

284

#### 285 *Conservation Value Index*

286 All species are not equal, and may differ in their value to conservationists on the basis of  
287 endemism, extinction risk (Mace et al. 2008), evolutionary distinctiveness (Tucker et al.  
288 2012; Hidasi-Neto et al. 2015), public appeal (Smith et al. 2012) or other attributes  
289 (Humphries et al. 1995; Joseph et al. 2009). This variation forms the basis of a range of  
290 protocols designed to elucidate the conservation value of species and, in turn, support the  
291 prioritisation of conservation actions or funding allocations (e.g. Huang et al. 2016; Isaac et  
292 al. 2007; Joseph et al. 2009). However, these protocols tend to be methodologically complex  
293 and require the collection of large datasets, diminishing their applicability in situations where  
294 non-academic conservation practitioners lack the training to apply them or where the  
295 necessary data are unavailable (Gardner et al. 2015a). In this study, we therefore used the  
296 novel Conservation Value Index (CVI) (adapted from Gardner et al. 2015a) to quantify the  
297 conservation value of individual species because it uses only readily available data and does  
298 not require the use of specialist software. As such, it can be easily applied in day-to-day  
299 decision-making by conservation practitioners. We assigned CVI scores to individual species

300 of bird and reptile before combining them to assess the impacts of natural resource use, and  
301 subsequent habitat degradation, on the conservation value of spiny forest habitats.

302

303 For the CVI we assigned scores to each individual species based on four attributes that reflect  
304 rarity, distinctiveness and threat. We use different combinations of attributes for the two  
305 taxonomic groups because the variation in conservation value within each group is driven by  
306 different factors. We scored rarity using geographical scale of endemism (*G*) and  
307 representation within SAPM (*R*), distinctiveness by taxonomic level of endemism (*E*), and  
308 threat on the basis of hunting and collection pressure (*C*) and degradation tolerance (*T*). We  
309 did not use *E* for reptiles because all species are endemic and there are no endemic families,  
310 so variation in the attribute is limited. Similarly, we did not use *C* for birds because most  
311 species in the Ranobe area are subject to comparable hunting pressure (Gardner and Davies  
312 2014).

313

314 Introduced species were removed from the dataset and scores assigned to indigenous taxa on  
315 a scale of 1-5 for each attribute (Table 2). For *G* we used different scoring systems for  
316 reptiles and birds because species distributions of the two taxonomic groups are best  
317 explained by different biogeographical models (Pearson and Raxworthy 2009). For birds we  
318 used distribution maps from Safford and Hawkins (2013) and followed Stattersfield et al.  
319 (1998) to classify microendemic species, whereas for reptiles we visually estimated range  
320 criteria using maps in Glaw and Vences (2007) and adopted 10,000 km<sup>2</sup> as the threshold for  
321 microendemic species (following Gardner et al. 2015a). *E* was assigned on the basis of  
322 taxonomy in Safford and Hawkins (2013), *R* scores were assigned on the basis of occurrence  
323 in 14 (birds) or 15 (reptiles) protected areas in the dry regions of Madagascar derived from  
324 the literature (Online resource 1), and values for *C* were based on occurrence in CITES

325 (Convention on International Trade in Endangered Species) appendices and the literature on  
326 reptile declines in Madagascar. *T* was attributed following the methods outlined in Gardner et  
327 al. (2015a) for reptiles, and were based on the literature (Safford and Hawkins 2013; Wilmé  
328 1996) for birds. Species for which no degradation tolerance data were available were scored  
329 as intolerant on the basis of the precautionary principle.

330

331 [Table 2]

332

333 The individual species CVI scores were calculated, producing a value in the range of 4-100,  
334 using the following formulae for reptiles and birds:

335 
$$CVI_{reptile} = (G + R) \times (C + T)$$

336 
$$CVI_{birds} = (G + E) \times (R + T)$$

337

338 The conservation value of a site can be considered a function of: i) the value of the species  
339 occurring there; and, ii) their abundance, because an area with a large population of a  
340 valuable species is more important than one with a small population. To understand the  
341 relative conservation value of each habitat treatment, we therefore wanted a metric that  
342 combined the CVI of each species with their relative abundance. However, simply weighting  
343 the CVI score by the relative frequency would heavily bias common species at the expense of  
344 rarer ones which are recorded only infrequently. We thus gave each species weightings  
345 standardised to the treatment where it was most frequent (e.g., a species with relative  
346 frequency of 0.36, 0.18 and 0.12 across each of the three treatments would be given  
347 weightings of 1, 0.5 and 0.33 respectively). In each treatment the CVI was then multiplied by  
348 the weighting to produce a frequency-weighted CVI score for each species, before these were  
349 summed to produce a conservation value score for each treatment.

350

## 351 **3 Results**

### 352 **3.1 Degradation impacts on birds**

353 We recorded 2385 bird contacts, comprising 53 species, in point counts across all treatments.  
354 Rarefaction curves approach an asymptote in all treatments, indicating that bird communities  
355 were sufficiently sampled (Fig. S1). Although observed richness was highest in the moderate-  
356 degradation treatment (*Low* – 36 spp.; *Mod* – 43 spp.; *High* – 37 spp.), rarefaction curves  
357 show no significant differences in richness since the 95 % confidence intervals overlap  
358 (Online resource 2). Total richness is estimated at 42.0 (*Low*), 46.8 (*Mod*) and 39.7 (*High*)  
359 species in the three treatments. Twenty-four species (45.3 %) were recorded in all treatments,  
360 one species (1.9 %) was restricted to *Low*, five species (9.4 %) were restricted to *Mod*, and  
361 seven (13.2 %) species were restricted to *High*: 17 species (32.1 %) were recorded only in  
362 forest habitats (*Low* and *Mod*).

363

364 Observed patterns of species relative frequency differed significantly for 22 species (41.5 %)  
365 across the three treatments. Three of these species (*Cuculus rochii*, *Hypsipetes*  
366 *madagascariensis* and *Dicrurus forficatus*) were observed more frequently in the low-  
367 degradation treatment, one species (*Ploceus sakalava*) in the moderate-degradation treatment,  
368 and six species (*Turnix nigricollis*, *Oena capensis*, *Agapornis canus*, *Cisticola cherina*,  
369 *Acridotheres tristis* and *Foudia madagascariensis*) in the high-degradation treatment. A  
370 further 12 species were recorded less frequently in the high-degradation treatment than in  
371 forest habitat (*Low* or *Mod*) (Online resource 3).

372

373 Patterns of species endemism varied across the degradation gradient (Fig. 2). While the  
374 proportion of endemic species was approximately equal in all treatments, the high-

375 degradation treatment contained a lower proportion of regionally-endemic birds (defined as  
376 restricted to Madagascar and the islands of the western Indian Ocean) and a higher proportion  
377 of non-endemic species. The vast majority (97.9 %) of contacts with introduced species  
378 (*Acridotheres tristis*) occurred in the high-degradation treatment.

379

380 [Fig. 2]

381

### 382 **3.2 Degradation impacts on reptiles**

383 We recorded 661 reptile contacts comprising 32 species, 27 of which were recorded at *Low*  
384 and *Mod*, and 15 species at *High*. Twenty-two species were observed during transects, and 27  
385 were captured in pitfall traps (Online resource 4). Twelve species (37.5 %) were recorded in  
386 all treatments, 17 species (53.1 %) were only recorded in forest habitats, and one species  
387 (*Lygodactylus tuberosus*) was recorded only in the high-disturbance site. Rarefaction curves  
388 indicate that *Low* and *Mod* had significantly higher species richness than *High*, as there is no  
389 overlap between confidence intervals (Online resource 5). Total richness is estimated at 30.5  
390 (*Low*), 34.2 (*Mod*) and 19.1 (*High*) species in the three treatments.

391

392 Observed patterns of reptile abundance, based on total contacts, were significantly  
393 heterogeneous for 11 species (34.4 %). Three species were recorded more frequently in the  
394 low-degradation treatment (*Chalarodon madagascariensis*, *Lygodactylus verticillatus* and  
395 *Oplurus cyclurus*), two species in the moderate-degradation treatment (*Madascincus* cf.  
396 *igneocaudatus* and *Tracheloptychus petersi*), and three species in the high-degradation  
397 treatment (*Lygodactylus tuberosus*, *Paroedura picta* and *Typhlops arenarius*). A further three  
398 species (*Geckolepis* c.f. *polypelis*, *Phelsuma mutabilis* and *Trachylepis elegans*) were  
399 recorded more frequently in the two forest areas than in the high-degradation treatment.

400

401 Forest disturbance affected distinct components of the reptile community differently,  
402 depending on their foraging substrate (Online resource 6). Terrestrial species decreased in  
403 frequency (capture rate and/or density) with increasing disturbance, while arboreal species  
404 demonstrated reduced frequency at *Mod* and reduced richness at *High* compared to the less  
405 degraded site. Fossorial and litter dwelling species reached peak frequency under conditions  
406 of moderate-intensity disturbance.

407

### 408 **3.3 Conservation value of species and sites**

409 The CVI allowed us to weight species on the basis of their conservation value. The six  
410 highest scoring bird species were locally-endemic forest specialists (Table 3), while the  
411 highest scoring reptile was the heavily harvested (and thus Critically Endangered) tortoise  
412 *Pyxis arachnoides* (Table 4). The relative conservation value of each treatment varied for the  
413 two taxonomic groups. Total bird conservation value was highest in *Low*, while total reptile  
414 conservation value was highest in *Mod*, although in both cases the differences between the  
415 two forest areas were small (Table 5). The high-degradation treatment had the lowest  
416 conservation value for both taxa.

417

418 [Table 3]

419

420 [Table 4]

421

422 [Table 5]

423

## 424 **4 Discussion**

425 We have generated some of the first data on the impacts of permitted livelihood activities  
426 within Madagascar's new generation of multiple-use protected areas. Our results show that  
427 charcoal production, an authorised activity within much of the Ranobe PK32 protected area,  
428 resulted in an overall reduction in the conservation value of habitats, although the responses  
429 of reptile and bird communities varied. However the impacts of charcoal production were less  
430 severe than the impacts of illegal shifting cultivation for both groups.

431

432 Although the impacts of habitat degradation on Madagascar's biodiversity have been well  
433 studied (reviewed in Gardner 2009; Irwin et al. 2010), the vast majority of research has been  
434 conducted in the country's humid and dry forests, ecosystems which greatly differ from the  
435 spiny forest in terms of biotic communities and abiotic conditions (Moat and Smith 2007;  
436 Goodman and Raheirilalao 2013). Within the spiny forest, degradation has been found to  
437 reduce species richness in both birds (Randriamiharisoa et al. 2015) and reptiles (Theisinger  
438 and Ratianarivo 2015). However, in our study, richness was maintained for reptiles and  
439 increased for birds at moderate degradation levels. Indeed the conservation value of reptiles  
440 was greatest at the moderate-disturbance site, perhaps reflecting an increase in microhabitat  
441 heterogeneity or structural complexity (MacArthur and MacArthur 1961; Tews et al. 2004).  
442 Bird communities were more responsive than reptiles to habitat degradation, undergoing  
443 extensive community turnover. This was reflected in the greater prevalence of birds adapted  
444 to open areas, and a decrease in the frequency of certain high-value, locally-endemic species  
445 such as *Monias benschi*, *Coua cursor* and *Newtonia archboldi*, with increasing degradation  
446 intensity.

447

448 Wilmé (1996) suggests that "the tolerance of [Madagascar's] endemic forest avifauna to  
449 forest degradation is proportional to its degree of taxonomic endemism". However, we

450 recorded seven members of endemic genera (*Coua cursor*, *Monias benschi*, *Neomixis*  
451 *striatigula*, *Newtonia brunneicauda*, *N. archboldi*, *Vanga curvirostris* and *Xenopirostris*  
452 *xenopirostris*) previously thought to occur only in undisturbed or slightly disturbed habitats,  
453 within a largely deforested habitat in our high-disturbance treatment. These findings lend  
454 some support to the hypothesis that faunal species of Madagascar's dry and spiny forests may  
455 be more tolerant of degradation than those same or congeneric species in the country's humid  
456 east and north (Gardner 2009). This may arise due to the more 'gentle' habitat modifications  
457 occurring in dry forests compared to rainforests (Irwin et al. 2010): for example, the  
458 increased light penetration in forest gaps is thought to make little difference to the understory  
459 in the spiny forest, because the sparse, deciduous nature of the canopy already allows  
460 illumination at ground level (Seddon and Tobias 2007). However, while tropical dry forests  
461 are thought to be more resilient than humid forests in terms of regeneration capacity (Lebrija-  
462 Trejos et al. 2008), little is known about the relative disturbance sensitivity of their respective  
463 faunas. Such research should be considered a priority since it has important repercussions for  
464 the implementation of multiple-use protected areas in different bioclimatic contexts.

465

466 The finding that moderate levels of degradation provoked an increase in richness of birds,  
467 and maintained richness in reptiles, is consistent with Connell's (1978) 'intermediate  
468 disturbance hypothesis', and reflects a pattern widely reported from other tropical  
469 environments, at least for some guilds (Burivalova et al. 2014; Child et al. 2009; Gray et al.  
470 2007; Martin and Blackburn 2010; Pons and Wendenberg 2005). However, all species are not  
471 equal, and the greater richness may often mask a turnover from range-restricted specialists to  
472 widespread generalists (Canaday 1997; Christian et al. 2009; Holbech 2005; Petit and Petit  
473 2003; Scott et al. 2006). The latter are of less importance to conservationists precisely  
474 because they adapt well to anthropogenic disturbance and thus do not require conservation

475 actions, such as protected areas, to maintain them (Harris and Pimm 2004; T. Gardner et al.  
476 2009). The use of species richness as a measure of conservation value has been widely  
477 criticised for this reason (Barlow et al. 2007; DeClercke et al. 2010; Fermon et al. 2005;  
478 Norris et al. 2010), but remains persistent (e.g., studies reviewed by Burivalova et al. 2014).  
479 Our use of the CVI provides further evidence of the inadequacies of richness in prioritising  
480 between sites or habitats, as the use of richness would indicate that forests degraded by  
481 charcoal production are more valuable for bird conservation in the spiny forest than less  
482 degraded habitats. Of course, the CVI does not represent a definitive quantification of  
483 conservation value, but is a useful heuristic tool to help conservationists prioritise action to  
484 where it is most needed (i.e. high-value species), and can be used without training, complex  
485 software or collecting new data.

486

487 Although the use of CVI provides novel insights into the impacts of habitat change on the  
488 conservation value of spiny forest bird and reptile assemblages, our results must be  
489 interpreted with caution. We carried out surveying during the rainy season when both groups  
490 are most active, and surveyed each site sequentially for logistical reasons. However, biases  
491 may have arisen due to changes in species detectability related to the advancing breeding  
492 season. In addition, the entry of new cohorts may have increased population size as surveying  
493 progressed. We minimised the latter problem by excluding all records of juveniles from the  
494 analysis, although it would have been preferable to repeat data collection over multiple years,  
495 or to survey each site simultaneously using multiple teams. Nonetheless, the latter approach  
496 has a number of drawbacks, including the extensive training needed to minimise the biases  
497 associated with potential differences in the bird detection abilities and/or identification skills  
498 of research assistants.

499

500 Although our observations appear to suggest that the majority of bird and reptile species in  
501 Ranobe are somewhat resilient to moderate or high levels of degradation, the presence of a  
502 species does not necessarily equate to its viability. It should not be assumed that local  
503 populations in disturbed areas will persist in the long-term because there are likely to be time  
504 lags associated with the impacts arising from perturbation, meaning that the degraded habitats  
505 at Ranobe may be carrying an ‘extinction debt’ (Kuussaari et al. 2009; Tilman et al. 1994).  
506 This is particularly true given that the habitat modifications that are the focus of this study are  
507 relatively recent (range: 3-15 years across the treatments). In addition, the persistence of  
508 some species within degraded habitats may be the result of source-sink dynamics, with  
509 populations maintained only by immigration from nearby areas of higher quality habitat  
510 (Hylander and Ehrlén 2013; Pulliam 1988; Tilman et al. 1994). The degraded habitats at  
511 Ranobe may therefore experience future local extinctions, even without further modification,  
512 and we may have over-estimated the value of these areas for bird and reptile diversity  
513 (Barlow et al. 2007; Sekercioglu et al. 2007). The scale of extinction debt can be influenced  
514 by habitat quantity, quality, or connectivity (Hylander and Ehrlén 2013). As such, when  
515 destructive activities such as charcoal production cannot be prevented within the ‘sustainable  
516 use zones’ of multiple-use protected areas, both the structural and functional connectivity  
517 between high-quality habitat patches should be maximised in order to maintain biodiversity  
518 and mitigate the negative impacts associated with resource exploitation.

519

520 The suggestion that Madagascar’s new generation of multiple-use protected areas may suffer  
521 the continued erosion of biodiversity as a result of the impacts of authorised livelihood  
522 activities has important ramifications for the objectives and management of multiple-use sites  
523 worldwide. In a multi-taxon assessment across a continuum of protection levels in East  
524 Africa, Gardner et al. (2007) found that multiple-use protected areas provide significant and

525 complementary conservation services to strictly-protected sites, maintaining species richness  
526 but conserving significantly different faunal communities to those occurring in national  
527 parks. Thus a spectrum of protected area categories may be appropriate to conserve the full  
528 complement of biodiversity in continental regions, if these possess a range of faunal  
529 assemblages adapted to a continuum of habitat types from dense forests to wooded savannahs  
530 and grasslands (Borghesio 2008; Gardner et al. 2007; Pons et al. 2003).

531

532 Madagascar, however, differs from continents in that the vast majority of the endemic biota is  
533 forest-dependent (Goodman and Benstead 2005), while non-forest areas typically contain  
534 floristically- and faunistically-impoverished assemblages characterised by non-endemic  
535 species of low conservation value (Irwin et al. 2010; Koechlin et al. 1974; Lowry II et al.  
536 1997). In this context, multiple-use sites essentially conserve the same communities as strict  
537 protected areas, but may do so less successfully than the latter. Thus, while multiple-use  
538 categories may be the only politically, ethically and logistically feasible option for many of  
539 Madagascar's new generation of protected areas, given the socioeconomic importance to  
540 rural communities of remaining forest resources (Gardner et al. 2013), it should not be  
541 assumed that they will be successful in maintaining the biodiversity they were established to  
542 conserve. Given that range-restricted habitat specialists are disproportionately likely to go  
543 extinct in modified habitats (Posa and Sodhi 2006; Scales and Marsden 2008), and are of  
544 greatest conservation interest worldwide, careful attention must be paid to the choice of  
545 protected area models in different contexts; in regions where the majority of priority species  
546 are disturbance-sensitive, strict protected areas may be a more appropriate model if they can  
547 be managed effectively.

548

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980 **Figure Legends**

981

982 **Fig. 1** Map of: A) Ranobe PK32 protected area (dotted line) showing location of five strict  
983 conservation zones (grey shading), wetlands and rivers (black shading/lines) and Ranobe  
984 village; and, B) location of three vegetation treatments used to survey bird and reptile  
985 communities across a gradient of degradation (forest cover, grey shading; wetlands, double  
986 line). Inset shows location of Ranobe PK32 within Madagascar (black square) and limits of  
987 spiny forest ecoregion following Goodman and Raherilalao (2013) (grey shading)

988

989 **Fig. 2** Endemism status of birds at Ranobe expressed as a percentage of contacts from 48  
990 point counts at three sites across a gradient of degradation. Black, Madagascar endemic; dark  
991 grey, regional endemic; light grey, indigenous non-endemic; white, introduced. Regional  
992 endemic species are defined as restricted to Madagascar and the western Indian Ocean islands  
993 (Comoros, Mascarene and Seychelles archipelagos)

994

995

996 **Table 1** Disturbance history and vegetation description of three habitat treatments used to  
997 investigate the impacts of degradation on birds and reptiles at Ranobe, southwest  
998 Madagascar.

999

<b>Treatment</b>	<b>Disturbance history</b>	<b>Habitat description</b>
Low disturbance ( <i>Low</i> )	Low level charcoal production since 2007	Relatively closed canopy dominated by <i>Didierea madagascariensis</i> and hardwood trees, with no understory shrub layer. Some charcoal production resulting in small openings, but canopy generally unbroken. Thick leaf litter layer.
Moderate disturbance ( <i>Mod</i> )	Intensive charcoal production since 1995	Broken canopy dominated by <i>Didierea madagascariensis</i> , with hardwood trees largely absent. Small openings are frequent and possess a dense shrub layer of regenerating stumps. Characterised by piles of dead branches and bark left over from charcoal production. Thin leaf litter layer.
High disturbance ( <i>High</i> )	Forest cleared for shifting cultivation in 2001, regenerating naturally since 2004/5	Dense shrub layer (height of 1-2m) of regenerating stumps dominated by <i>Cedrelopsis grevei</i> and <i>Fernandoa madagascariensis</i> , with no litter layer. Relict individual trees and small forest patches (< 1ha) occur within a mosaic pattern.

1000

1001

1002 **Table 2** Scoring criteria for Conservation Value Index (CVI) attributes, used to quantify the conservation value of individual bird and reptile  
 1003 species at Ranobe, southwest Madagascar. EBA = Endemic Bird Area (Stattersfield *et al.* 1998), PA = protected area.

1004

Taxonomic group	Score	Geographic scale of endemism (G)	Taxonomic level of endemism (E)	Representation in sample PAs (R)	Hunting/collection pressure (C)	Degradation tolerance (T)
Birds	1	Indigenous, non-endemic species	Indigenous, non-endemic species	Recorded in 12-14 PAs (n > 85%)	N/A	Tolerant of modified or artificial habitats
	2	Endemic to western Indian Ocean	Endemic species	Recorded in 8-11 PAs (55 > n < 85%)	N/A	N/A
	3	Widespread Madagascar endemic	Endemic genus	Recorded in 4-7 PAs (30 > n < 50%)	N/A	Tolerant of edge effects, medium-intensity degradation or secondary growth.
	4	Endemic to dry regions of Madagascar	Endemic subfamily	Recorded in 2-3 PAs (10 > n < 20%)	N/A	N/A
	5	EBA species	Endemic family	Recorded in only 1 PA (n < 10%)	N/A	Intolerant of low-intensity degradation
Reptiles	1	Indigenous, non-endemic species	N/A	Recorded in 12-15 PAs (n > 75%)	No known threat	Tolerant of modified or artificial habitats
	2	Widespread endemic, occurring in dry and humid regions	N/A	Recorded in 8-11 PAs (45 > n < 75%)	N/A	N/A
	3	Endemic to dry regions	N/A	Recorded in 4-7 PAs (20 > n < 45%)	Known threat (CITES Appendix I and II), but not known to cause local extirpations	Tolerant of edge effects, medium-intensity degradation or secondary growth.
	4	Endemic to one bioclimatic region <sup>a</sup>	N/A	Recorded in 2-3 PAs (10 > n < 20%)	N/A	N/A
	5	Local endemic, range size estimated as < 10,000 km <sup>2</sup>	N/A	Recorded in only 1 PA (n < 10%)	Threat known to have caused local extirpations or severe population declines	Intolerant of low-intensity degradation

1005 <sup>a</sup>Following Cornet 1974

1006 **Table 3** Bird species recorded at Ranobe showing attributes used in Conservation Value  
 1007 Index (CVI) and frequency-weighted CVI scores for three sites across a gradient of  
 1008 degradation: Low, Mod and High indicate low-, moderate- and high-degradation treatments.  
 1009 CVI attributes: *G* – geographic scale of endemism, *E* – taxonomic level of endemism, *R* –  
 1010 representation in sample protected areas, *T* – degradation tolerance. Asterisks indicate species  
 1011 endemic to the spiny forest Endemic Bird Area (Stattersfield *et al.* 1998).

Species	CVI attribute scores				CVI score	Frequency-weighted CVI		
	G	E	R	T		Low	Mod	High
* <i>Monias benschi</i>	5	5	4	5	<b>90</b>	90	22.5	22.5
* <i>Xenopirostris xenopirostris</i>	5	5	3	5	<b>80</b>	0	11.4	80
* <i>Coua cursor</i>	5	4	3	5	<b>72</b>	72	20.6	30.9
* <i>Uratelornis chimaera</i>	5	5	4	3	<b>70</b>	0	0	0
* <i>Thamnornis chloropetoides</i>	5	5	2	5	<b>70</b>	70	60.0	0
* <i>Newtonia archboldi</i>	5	5	2	5	<b>70</b>	70	47.6	22.4
<i>Coua ruficeps olivaceiceps</i>	4	4	2	5	<b>56</b>	40.0	56	0
<i>Calicalicus madagascariensis</i>	3	5	2	5	<b>56</b>	56	56	0
<i>Artamella viridis</i>	3	5	2	5	<b>56</b>	32.0	56	0
<i>Vanga curvirostris</i>	3	5	1	5	<b>48</b>	48	29.2	4.2
<i>Coua cristata</i>	3	4	1	5	<b>42</b>	42	36.6	25.7
<i>Falco zoniventris</i>	3	2	3	5	<b>40</b>	0	40	0
<i>Falcula palliata</i>	4	5	1	3	<b>36</b>	36	36	0
<i>Leptosomus discolor</i>	2	5	2	3	<b>35</b>	0	0	0
* <i>Nesillas lantzii</i>	5	2	2	3	<b>35</b>	35	0	11.7
<i>Newtonia brunneicauda</i>	3	5	1	3	<b>32</b>	32	30.7	14
<i>Leptopterus chabert</i>	3	5	1	3	<b>32</b>	19.2	16	32
<i>Aviceda madagascariensis</i>	3	2	3	3	<b>30</b>	0	30	0
<i>Neomixis striatigula</i>	3	3	2	3	<b>30</b>	21.5	30	10.8
<i>Cuculus rochii</i>	3	2	2	3	<b>25</b>	25	11.7	5
<i>Polyboroides radiatus</i>	3	2	1	3	<b>20</b>	20	10	0
<i>Buteo brachypterus</i>	3	2	1	3	<b>20</b>	0	20	0
<i>Mirafra hova</i>	3	2	3	1	<b>20</b>	0	0	20
<i>Copsychus albospecularis</i>	3	2	1	3	<b>20</b>	20	19.3	14.3
<i>Treron australis</i>	2	1	2	3	<b>15</b>	0	0	15
<i>Nectarinia notata</i>	2	1	2	3	<b>15</b>	0	15	3.75
<i>Ploceus sakalava</i>	4	1	2	1	<b>15</b>	0.7	15	5.0
<i>Accipiter francesiae</i>	2	1	1	3	<b>12</b>	0	12	0
<i>Turnix nigricollis</i>	2	1	1	3	<b>12</b>	2.1	0	12
<i>Nesoenas picturata</i>	2	1	1	3	<b>12</b>	9.7	12	2.9
<i>Coracopsis vasa</i>	2	1	1	3	<b>12</b>	12	12	0
<i>Coracopsis nigra</i>	2	1	1	3	<b>12</b>	5.0	12	0
<i>Phedina borbonica</i>	2	1	3	1	<b>12</b>	0	12	12
<i>Hirundo rustica</i>	1	1	5	1	<b>12</b>	0	0	0
<i>Hypsipetes madagascariensis</i>	2	1	1	3	<b>12</b>	12	2.6	6.8
<i>Terpsiphone mutata</i>	2	1	1	3	<b>12</b>	9.7	12	4.6
<i>Neomixis tenella</i>	3	3	1	1	<b>12</b>	12	11.7	9.3
<i>Cisticola cherina</i>	2	1	3	1	<b>12</b>	0	0	12
<i>Nectarinia souimanga</i>	2	1	1	3	<b>12</b>	11.5	12	8.8
<i>Dicrurus forficatus</i>	2	1	1	3	<b>12</b>	12	9	9.5
<i>Falco peregrinus</i>	1	1	4	1	<b>10</b>	0	0	10
<i>Agapornis canus</i>	3	2	1	1	<b>10</b>	3.3	3.3	10
<i>Tachymarptis melba</i>	1	1	4	1	<b>10</b>	0	10	0
<i>Eurystomus glaucurus</i>	1	1	2	3	<b>10</b>	0	10	0

<i>Upupa marginata</i>	3	2	1	1	<b>10</b>	10	6.4	8.6
<i>Falco newtoni</i>	2	1	2	1	<b>9</b>	4.1	3.3	9
<i>Caprimulgus madagascariensis</i>	2	1	2	1	<b>9</b>	9	0	0
<i>Falco concolor</i>	1	1	3	1	<b>8</b>	0	0	8
<i>Foudia madagascariensis</i>	3	1	1	1	<b>8</b>	0.2	0.8	8
<i>Milvus migrans</i>	1	1	2	1	<b>6</b>	0	0	0
<i>Oena capensis</i>	1	1	2	1	<b>6</b>	3.4	1.4	6
<i>Centropus toulou</i>	2	1	1	1	<b>6</b>	4.3	4.3	6
<i>Apus barbatus</i>	1	1	2	1	<b>6</b>	3	6	0
<i>Merops superciliosus</i>	1	1	2	1	<b>6</b>	3.7	3.3	6
<i>Corvus albus</i>	1	1	2	1	<b>6</b>	0	0	6
<i>Numida meleagris</i>	1	1	2	1	<b>6</b>	0	0	6
<b>Total conservation value of treatment</b>						<b>856.4</b>	<b>825.7</b>	<b>478.6</b>

1012

1013

1014 **Table 4** Reptile species recorded at Ranobe showing attributes used in Conservation Value  
 1015 Index (CVI) score and relative frequency-weighted CVI scores for three sites across a  
 1016 gradient of degradation: Low, Mod and High indicate low-, moderate- and high-degradation  
 1017 treatments. CVI attributes: *G* – geographic scale of endemism, *R* – representation in sample  
 1018 protected areas, *C* – collection/hunting threat, *T* – degradation tolerance. Locally-endemic  
 1019 species are indicated by an asterisk.

1020

Species	CVI attribute scores				CVI score	Frequency-weighted CVI		
	G	R	C	T		Low	Mod	High
<i>Pyxis arachnoides</i>	4	3	5	5	<b>70</b>	70	0	0
* <i>Voeltzkowia petiti</i>	5	4	1	5	<b>54</b>	14.7	54	0
* <i>Tracheloptychus petersi</i>	5	4	1	5	<b>54</b>	22.1	54	2.5
<i>Geckolepis polylepis</i>	4	4	1	5	<b>48</b>	48	32.8	0
<i>Paroedura androyensis</i>	4	3	1	5	<b>42</b>	14	42	0
* <i>Pygomeles braconnieri</i>	5	4	1	3	<b>36</b>	14.4	36	0
<i>Voeltzkowia rubrocaudata</i>	3	3	1	5	<b>36</b>	36	0	10.3
* <i>Zonosaurus quadrilineatus</i>	5	4	1	3	<b>36</b>	36	32	8
<i>Ithycyphus oursi</i>	3	3	1	5	<b>36</b>	0	36	0
* <i>Liophidium chabaudi</i>	5	4	1	3	<b>36</b>	36	28.8	21.6
<i>Madascincus igneocaudatus</i>	3	2	1	5	<b>30</b>	12	30	0
<i>Madagascarophis ocellatus</i>	4	3	1	3	<b>28</b>	28	0	0
<i>Blaesodactylus sakalava</i>	3	1	1	5	<b>24</b>	24	16	0
<i>Zonosaurus karsteni</i>	3	3	1	3	<b>24</b>	24	24	0
<i>Madagascarophis meridionalis</i>	3	3	1	3	<b>24</b>	0	24	0
<i>Trachylepis aureopunctata</i>	3	2	1	3	<b>20</b>	6.7	20	3.3
<i>Heteroliodon occipitalis</i>	3	2	1	3	<b>20</b>	10	20	0
<i>Leioheterodon geayi</i>	3	2	1	3	<b>20</b>	20	0	0
<i>Typhlops arenarius</i>	3	2	1	3	<b>20</b>	0	6.2	20
<i>Typhlops decorsei</i>	3	2	1	3	<b>20</b>	0	20	0
<i>Lygodactylus verticillatus</i>	4	4	1	1	<b>16</b>	16	4	0
<i>Phelsuma mutabilis</i>	3	1	3	1	<b>16</b>	16	10.3	2.3
<i>Amphiglossus ornaticeps</i>	2	2	1	3	<b>16</b>	9.6	16	0
<i>Oplurus cyclurus</i>	2	2	1	3	<b>16</b>	16	6.5	0.73
<i>Lygodactylus tuberosus</i>	4	3	1	1	<b>14</b>	0	0	14
<i>Paroedura picta</i>	3	2	1	1	<b>10</b>	3.8	1.9	10
<i>Furcifer verrucosus</i>	3	2	1	1	<b>10</b>	10	2.9	0
<i>Chalarodon madagascariensis</i>	2	2	1	1	<b>8</b>	8	4	3.0
<i>Trachylepis elegans</i>	2	1	1	1	<b>6</b>	4.9	6	2.9
<i>Dromicodryas bernieri</i>	2	1	1	1	<b>6</b>	6	6	6
<i>Mimophis mahfalensis</i>	2	1	1	1	<b>6</b>	4.5	6	5.3
<i>Hemidactylus mercatorius</i>	1	1	1	1	<b>4</b>	4	3.4	2.9
<b>Total conservation value of treatment</b>						<b>514.7</b>	<b>542.8</b>	<b>112.7</b>

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1024 **Table 5** Observed and estimated species richness and Conservation Value Index (CVI) score

1025 for birds and reptiles at three sites across a gradient of disturbance at Ranobe, southwest

1026 Madagascar.

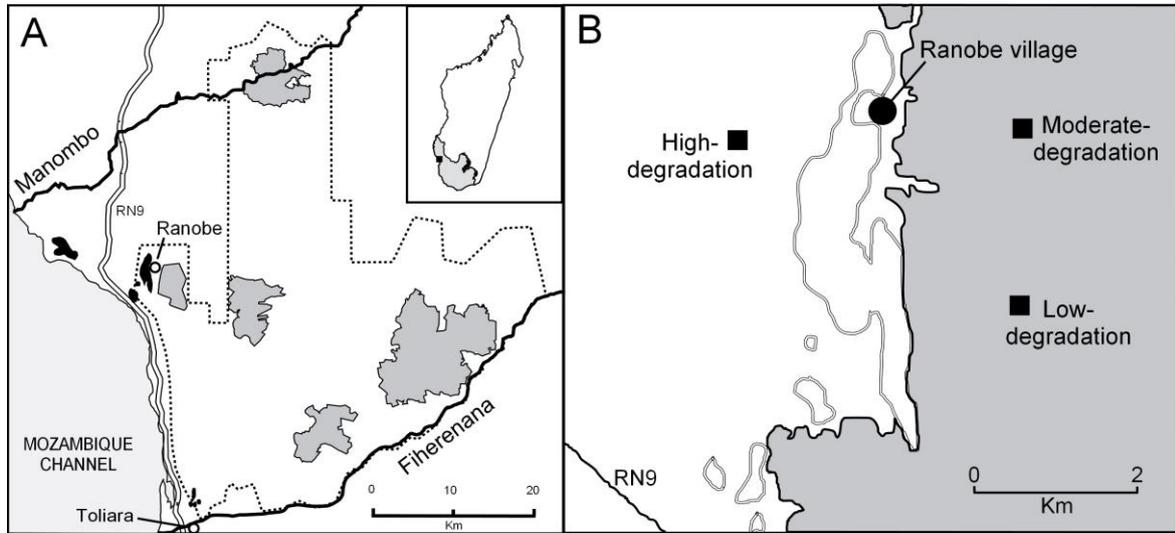
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	<b>Low disturbance</b>	<b>Moderate disturbance</b>	<b>High disturbance</b>
<b>Observed bird richness</b>	36	43	37
<b>Estimated bird richness</b>	42.0	46.8	39.7
<b>Bird CVI</b>	<b>856.4</b>	<b>825.7</b>	<b>478.6</b>
<b>Observed reptile richness</b>	27	27	15
<b>Estimated reptile richness</b>	30.5	34.2	19.1
<b>Reptile CVI</b>	<b>514.7</b>	<b>542.8</b>	<b>112.7</b>

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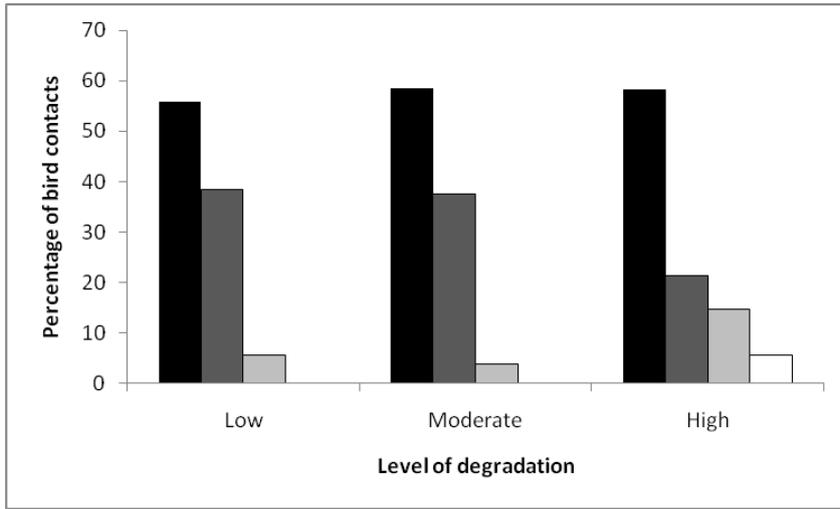
1030 **Fig. 1**



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1033 **Fig. 2**



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