



Deposited via The University of Sheffield.

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/id/eprint/119507/>

Version: Accepted Version

---

**Article:**

Kingma, S.A., Komdeur, J., Burke, T. et al. (2017) Differential dispersal costs and sex-biased dispersal distance in a cooperatively breeding bird. *Behavioral Ecology*, 28 (4). pp. 1113-1121. ISSN: 1045-2249

<https://doi.org/10.1093/beheco/arx075>

---

**Reuse**

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

**Takedown**

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing [eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk) including the URL of the record and the reason for the withdrawal request.

**Differential dispersal costs explain sex-biased dispersal distance in a cooperatively breeding bird**

Journal:	<i>Behavioral Ecology</i>
Manuscript ID	Draft
Manuscript Type:	Original article
Keywords:	cooperative breeding, delayed dispersal, habitat saturation, inbreeding, sex-biased dispersal

SCHOLARONE™  
Manuscripts

View Only

1  
2  
3 **1 Differential dispersal costs explain sex-biased dispersal distance**  
4  
5  
6 **2 in a cooperatively breeding bird**  
7  
8  
9  
10  
11

12 Short title: Sex-biased dispersal in Seychelles warblers  
13  
14  
15  
16  
17

18 **LAY SUMMARY**

19  
20 Why does the distance that young animals disperse between their natal- and breeding territory  
21 usually differ between males and females? We show that in cooperatively breeding  
22 Seychelles warblers, males are reluctant to disperse and disperse less far than females  
23 because for males dispersal is more costly due to attacks by conspecifics. We did not find  
24 evidence for regularly-invoked hypotheses for sex-biased dispersal (e.g. sex differences in  
25 territory defense and sex-biased dispersal as inbreeding avoidance mechanism).  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35

36 **ABSTRACT**

37  
38 In most bird species, dispersal distance from the natal territory to a breeding territory is  
39 greater for females than males. Sex-biased dispersal may have important implications for  
40 population dynamics and sexual reproduction because it can affect social recognition, genetic  
41 population structure, inbreeding and kin selection. Two main hypotheses have been proposed  
42 to explain sex-biased dispersal distance: 1) it serves as an inbreeding-avoidance mechanism  
43 or 2) it is linked to a sex difference in resource-holding potential. Additionally, in species  
44 where individuals delay dispersal and become subordinates in a natal territory, differences in  
45 benefits of philopatry (e.g. territory inheritance, opportunities for reproduction) may also  
46 affect sex-biased dispersal. We show that in the group-living Seychelles warbler,  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

1  
2  
3 25 *Acrocephalus sechellensis*, females disperse further to obtain a breeding position than do  
4  
5 26 males. However, female-biased dispersal distance is not explained by the above-mentioned  
6  
7 27 hypotheses: further dispersal does not lead to less-related partners, both sexes defend the  
8  
9 28 territory, both sexes can inherit a territory, and subordinate females are more likely to obtain  
10  
11 29 some reproduction than subordinate males. Instead, we provide evidence for a little-explored  
12  
13 30 hypothesis based on a sex difference in dispersal costs: extra-territorial forays to search for  
14  
15 31 limited vacancies are more costly, in terms of increased mortality, for males. In line with such  
16  
17 32 differential dispersal costs, males foray less far than females and often wait for local dispersal  
18  
19 33 opportunities, ultimately resulting in a shorter average dispersal distance than in females.  
20  
21 34 This explanation may help future studies in explaining sex-biased dispersal in social and  
22  
23 35 perhaps also non-social species.  
24  
25  
26  
27  
28  
29

30 **Key-words:** cooperative breeding, delayed dispersal, habitat saturation, inbreeding, sex-  
31  
32 biased dispersal  
33  
34  
35  
36  
37  
38  
39  
40

## 41 INTRODUCTION

42 In animals, the distance of dispersal from the natal territory or site to a place for independent  
43  
44 breeding is often sex-biased (Greenwood 1980; Pusey 1987; Clarke, Sæther and Roskaft  
45  
46 1997). Sex-biased dispersal can have important implications for the dynamics and the genetic  
47  
48 structure of populations (Aars and Ims 2000; Prugnolle and De Meeus 2002). Understanding  
49  
50 its causes and consequences is therefore important to understanding how processes like kin  
51  
52 cooperation and competition, resource defence and inbreeding avoidance can affect mating  
53  
54 systems and population dynamics.  
55  
56  
57  
58  
59  
60

1  
2  
3 49 Two main non-exclusive hypotheses have been invoked to explain sex-biased dispersal  
4  
5 50 (Greenwood 1980; Greenwood and Harvey 1982; Pusey 1987). First, the *inbreeding-*  
6  
7 51 *avoidance* hypothesis predicts that the risk of mating with closely related individuals is  
8  
9 52 reduced if dispersal distance is different between the sexes (Pusey and Wolf 1996; Perrin and  
10  
11 53 Mazalov 2000; Perrin and Goudet 2001). This mechanism may especially be important in  
12  
13 54 species with high levels of extra-pair mating, where dispersing females run the risk of mating  
14  
15 55 with their own fathers (who sired the female in an extra-pair fertilisation), whereas dispersing  
16  
17 56 males cannot mate with their mother, as their mothers are always from within the natal  
18  
19 57 territory. Second, the *resource-holding potential* hypothesis predicts that dispersal asymmetry  
20  
21 58 between the sexes is a consequence of bias in the degree of advantage gained from familiarity  
22  
23 59 with the area during intra-specific competition for resources towards the sex that defends  
24  
25 60 those resources (Greenwood 1980; Pusey 1987). Although both hypotheses have obtained  
26  
27 61 some degree of support (Johnson and Gaines 1990; Bowler and Benton 2005; Lawson  
28  
29 62 Handley and Perrin 2007), their respective importance remains unclear.  
30  
31  
32  
33

34 63 In species where opportunities for independent breeding are limited, such as family-  
35  
36 64 living and cooperatively breeding species, subordinate individuals either have to wait in a  
37  
38 65 resident territory for a breeding vacancy to arise nearby or to search for a vacancy in the  
39  
40 66 population (Cockburn 1998). Waiting in a resident territory may yield benefits to subordinate  
41  
42 67 individuals (Stacey and Ligon 1991; Koenig & Dickinson 2004) and, if such ‘benefits of  
43  
44 68 philopatry’ differ between males and females, this may lead to sex bias in motivation to  
45  
46 69 search for an independent breeding vacancy, resulting in differences in the ultimate dispersal  
47  
48 70 distance (Brown 1987; Cockburn 1998; Kingma et al. 2016a,b). As such, differences in the  
49  
50 71 ‘*reproductive benefits of philopatry*’ (i.e., the likelihood of obtaining a share in reproduction  
51  
52 72 in the resident territory, and/or inheriting the breeding position; e.g. Cockburn 1998; Kokko  
53  
54 73 and Ekman 2002; Richardson et al. 2002) might explain sex-biased dispersal. Sex differences  
55  
56  
57  
58  
59  
60

1  
2  
3 74 in the probability of territory inheritance may arise when the more competitive sex does not  
4  
5 75 accept a related individual as a partner and could expel either an inheriting offspring or the  
6  
7 76 remaining related breeder from the territory (e.g., Koenig and Stacey 1990; Nelson-Flower et  
8  
9 77 al. 2012). Alternatively, or additionally, individuals who delay dispersal and remain in a  
10  
11 78 resident territory may gain '*energetic benefits of philopatry*', such as access to food. It is not  
12  
13 79 immediately clear if and how such benefits differ between the sexes. However, the role that  
14  
15 80 any such benefits play in explaining delayed dispersal would also depend on the costs of  
16  
17 81 leaving, which may well be sex-specific (Perrin and Mazalov 2000; Gros et al. 2008).

20  
21 82 In species both with and without delayed dispersal, searching for an independent  
22  
23 83 breeding position involves extra-territorial forays through unfamiliar or unfavourable habitat  
24  
25 84 (Reed et al. 1999). In a number of species it has been shown that such forays are associated  
26  
27 85 with reductions in survival and body condition due to harassment by predators and  
28  
29 86 conspecifics, and such costs have been invoked as explanation for delayed dispersal (e.g.,  
30  
31 87 Yaber and Rabenold 2002; Griesser et al. 2006; Ridley, Raihani & Nelson-Flower 2008;  
32  
33 88 Ridley 2012; Kingma et al. 2016a). If such costs are different between both sexes, for  
34  
35 89 example because of differences in conspicuousness to predators or because attacks by  
36  
37 90 conspecifics may be more frequently directed at the sex that threatens the reproduction of the  
38  
39 91 resident individuals more, they may well explain sex-biased dispersal distance. Whether this  
40  
41 92 *dispersal-cost hypothesis* is supported is unclear, however, partly because extra-territorial  
42  
43 93 forays have received relatively little empirical attention (Reed et al. 1999) and because  
44  
45 94 studies of sex-biased dispersal intrinsically focus on proximate and ultimate factors  
46  
47 95 underlying dispersal, rather than the actual movement *per se* (Lawson Handley and Perrin  
48  
49 96 2007). Together, these hypotheses, in addition to the more conventional hypotheses of  
50  
51 97 inbreeding avoidance and sex bias in resource-holding potential, provide an interesting  
52  
53  
54  
55  
56  
57  
58  
59  
60

1  
2  
3 98 avenue to determine the importance of various social and ecological factors for the evolution  
4  
5 99 of sex-biased dispersal.  
6

7  
8 100 Here we tested all the above-mentioned hypotheses for female-biased dispersal distance  
9  
10 101 (see Table 1 and below) in the cooperatively breeding Seychelles warbler, *Acrocephalus*  
11  
12 102 *sechellensis*. This system is very suitable for testing these hypotheses for several reasons.  
13  
14 103 First, females on average disperse further from their natal territory than males (Eikenaar et al.  
15  
16 104 2008a). Second, distinguishing dispersal from mortality is generally difficult (Koenig et al.  
17  
18 105 1996), but the Seychelles warblers exist on a contained island with virtually no emigration  
19  
20 106 between islands, so individuals that have disappeared from the study-population almost  
21  
22 107 certainly died (Komdeur et al. 2004). Third, in this long-term study population nearly all  
23  
24 108 individuals are individually marked and followed throughout their life, so that their natal  
25  
26 109 territory, dispersal behavior, relatedness to other individuals, and dates of birth and death are  
27  
28 110 known. Fourth, habitat saturation inhibits independent breeding of subordinate individuals,  
29  
30 111 but individuals can improve the likelihood of obtaining an independent breeding position by  
31  
32 112 extra-territorial forays to find a position (Eikenaar et al. 2008a,b; Kingma et al. 2016a,b).  
33  
34 113 Although such behaviors may be difficult to assess in general (Reed et al. 1999), our detailed  
35  
36 114 monitoring allows us to make inferences about prospecting and floating.  
37  
38  
39

40  
41 115 We used a framework based on the above-mentioned hypotheses (see Table 1) to  
42  
43 116 develop and test predictions of how different proximate and ultimate factors may explain  
44  
45 117 female-biased dispersal distance in Seychelles warbler. Specifically, we assessed (1) whether  
46  
47 118 dispersal over greater distance leads to the acquisition of a less-related partner, and whether  
48  
49 119 this is especially the case for females who may mate with their extra-pair sire (*inbreeding-*  
50  
51 120 *avoidance*), (2) whether males and females differ in territory defence (*resource-holding-*  
52  
53 121 *potential*), (3) whether the probability of obtaining parentage as a subordinate and territory  
54  
55 122 inheritance rates differ between male and female subordinates (*reproductive-benefits-of-*  
56  
57  
58  
59  
60

1  
2  
3 123 *philopatry*), and (4) whether the costs of finding an independent breeding territory differ  
4  
5 124 between males and females in Seychelles warblers (*costly-dispersal*).  
6  
7  
8 125  
9  
10 126

## 127 **METHODS**

### 128 *Study system*

129 We studied a population of ca. 320 individually colour-ringed Seychelles warblers on Cousin  
130 Island, Seychelles (29 ha; 04°20'S, 55°40'E) during the main breeding seasons (June-  
131 September) from 2003 until 2014. Each of the ca. 110 territories are occupied year-round by  
132 a dominant breeding pair, of which approximately half are accompanied by 1 to 4  
133 independent subordinates. Dominant individuals rarely disperse and usually remain present in  
134 their territory until death (Hammers et al. 2015). Intruding conspecifics are physically  
135 attacked (Kingma et al. 2016a,b). Because of this, territory boundaries are easily determined  
136 based on border disputes between groups. Breeding vacancies are limited because all suitable  
137 habitat is occupied (Komdeur 1992) and Seychelles warblers are relatively long lived  
138 (average 5.5 years; Hammers et al. 2015). Individuals can improve their likelihood of finding  
139 a breeding vacancy by either temporarily (prospecting) or permanently (floating) leaving  
140 their territory to foray and search for vacancies across the island (Kingma et al. 2016b).

141 In each season we performed regular censuses (at least weekly per territory) to  
142 identify for each individual the resident territory (i.e., where birds were consistently observed  
143 foraging, performing reproductive tasks and/or involved in non-antagonistic interactions with  
144 other resident individuals) and breeding status (*dominant*: based on affiliative behavior  
145 between the pair members; *subordinate*: reproductively mature individuals but not involved  
146 in direct pair behaviors or initiation of breeding activities, or independent *juvenile*: 3-5  
147 months old). Birds were captured using mist nets and each bird was given a unique

1  
2  
3 148 combination of three colour rings and a numbered metal ring (if not already ringed). Body  
4  
5 149 mass ( $\pm 0.1$  g) and tarsus length ( $\pm 0.1$  mm) were measured, and a small blood sample was  
6  
7 150 taken to determine sex (following the protocol in Griffiths et al. 1998) and for genotyping  
8  
9 151 (see below).

10  
11  
12 152

### 13 153 *Inheritance, dispersal, prospecting and floating*

14  
15  
16 154 We determined whether each subordinate observed in a season was present by the beginning  
17  
18 155 of the next season as a subordinate on the same territory, obtained a breeding position, or had  
19  
20 156 died. For individuals that obtained a breeding position, we determined whether this was  
21  
22 157 achieved through inheritance of the natal territory or dispersal from that territory. We  
23  
24 158 determined dispersal distance as the minimum number of territories that an individual had to  
25  
26 159 cross between its natal territory and the territory where it obtained its breeding position  
27  
28 160 (following Eikenaar et al. 2008b). A small number of individuals ( $n = 3$  of 215 males and 5  
29  
30 161 of 218 females) settled as subordinates in a non-natal territory before obtaining a breeding  
31  
32 162 position elsewhere, but we assess here the dispersal distance between the original natal  
33  
34 163 territory and a breeding position only. The maximal possible distance that individuals could  
35  
36 164 disperse over the island ranged from 9 to 16 territories (median = 12), and this did not differ  
37  
38 165 between males (median: 12,  $n = 215$ ) and females (median: 12,  $n = 218$ ; generalized linear  
39  
40 166 mixed model with maximum possible dispersal distance as a response variable, sex as an  
41  
42 167 independent variable and 'natal-territory-identity' as a random variable:  $\beta = -0.004 \pm 0.028$ ,  $z$   
43  
44 168 =  $-0.13$ ,  $P = 0.90$ ). Some individuals were observed or caught while prospecting (defined as  
45  
46 169 individuals observed  $>2$  territories away from their resident territories and returned after  
47  
48 170 prospecting to that territory) or floating (individuals only observed on non-resident territories  
49  
50 171 multiple times throughout the season; see Kingma et al. 2016a,b for details). For each  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

1  
2  
3 172 prospector we determined the maximum number of territories it was seen away from its  
4  
5 173 resident territory, similarly as for determining dispersal distance.  
6  
7  
8 174

9  
10 175 *Statistical analyses*

11 176 For the statistical analyses, models were fitted in R 3.2.0. (R development core-team, 2016)  
12  
13 177 using the ‘lme4’ and ‘lmerTest’ packages (for linear mixed models and generalized linear  
14  
15 178 mixed models; Bates et al. 2015; Kuznetsova et al. 2016), unless stated otherwise. Non-  
16  
17 179 significant variables ( $P > 0.05$ ) were sequentially excluded from the model, starting with the  
18  
19 180 least significant variable, until the model only contained significant variables. Values for non-  
20  
21 181 significant variables were obtained by re-including them in turn in the final model to confirm  
22  
23 182 that the order of exclusion did not change the results. Mean values and model estimates ( $\beta$ )  
24  
25 183 are reported  $\pm$  standard error (SE).  
26  
27  
28  
29  
30 184

31  
32 185 *Sex-biased dispersal distance and prospecting behavior*

33  
34 186 We first confirmed findings from an earlier study (Eikenaar et al. 2008a), by testing whether  
35  
36 187 dispersal distances between natal and subsequent breeding territory were different for males  
37  
38 188 and females ( $n = 215$  and  $218$ , respectively) using a generalized linear mixed model with  
39  
40 189 Poisson error, including ‘natal-territory-identity’ as random variable.

41  
42  
43 190 We also tested whether the maximum distance that individuals prospected from the  
44  
45 191 natal territory was different between males and females, using a generalized linear model  
46  
47 192 with Poisson error.  
48

49  
50 193

51  
52 194 *Inbreeding avoidance*

53  
54 195 We tested whether dispersal distance and sex of the focal individual predicted the relatedness  
55  
56 196 between that bird and its new partner (response variable) using a general linear mixed model  
57  
58  
59  
60

1  
2  
3 197 with ‘natal-territory-identity’ as a random effect (excluding seven pairs for which relatedness  
4  
5 198 could not be determined due to an unsampled breeder). To test whether the effect of dispersal  
6  
7 199 distance differed between males and females, we added the interaction between sex and  
8  
9 200 distance. We excluded 41 individuals that inherited their natal territory from the analysis.  
10  
11 201 Relatedness of dispersing individuals to their (first) dominant partner (R) was calculated  
12  
13 202 using GenAlEx 6.5 (Peakall and Smouse 2012), using Queller and Goodnight (1989)  
14  
15 203 estimation. We used genotypes of all individuals involved in included breeding pairs, as  
16  
17 204 determined using 30 microsatellite markers (see for details: Richardson et al. 2000; Spurgin  
18  
19 205 et al. 2014).

206 To assess whether females are likely to end up in an incestuous relationship with their  
207 extra-group father, we used two approaches. First, we tested whether pair-relatedness was  
208 different between females that obtained a partner after short-distance dispersal (1 or 2  
209 territories from their natal territory) and those that dispersed further, using a linear mixed  
210 model with ‘natal-territory-identity’ as a random factor. Second, we tested whether pair-  
211 relatedness was different between short-distance dispersing males and females using a similar  
212 model. The distance of 1 or 2 territories for ‘short-distance dispersal that may lead to  
213 incestuous mating’ was chosen because two territories is the median distance between an  
214 extra-group offspring’s territory and its sire (62% of extra-group fathers lived within two  
215 territories distance; Richardson et al. 2001). Note however, that the results are similar if this  
216 distance would be chosen differently, because females who dispersed further did not pair with  
217 less related males (see Fig. 1b).

218

### 219 *Resource-holding potential*

220 We analysed two probable determinants of sex-differences in resource-holding potential.  
221 First, we assessed whether breeding males were larger and/or heavier than breeding females.

1  
2  
3 222 We compared tarsus length (averaged if an individual was measured more than once) of  
4  
5 223 males and females using a t-test. Subsequently, using all catches of each individual, we tested  
6  
7 224 in a linear mixed model whether body mass (as response variable) was different between the  
8  
9 225 sexes and added 'individual-identity', year and 'resident-territory-identity' as random factors  
10  
11 226 and time [morning (6:34-10:00), midday (10:00-14:00), afternoon (14:00-19:10)] of capture  
12  
13  
14 227 as an independent variable to account for temporal and spatial variation in body mass.  
15  
16 228 Second, using 121 opportunistically observed antagonistic interactions (observed during  
17  
18 229 weekly censuses in each territory) between (identified) resident individuals and intruders, we  
19  
20 230 determined whether male residents were more likely to be involved in antagonistic  
21  
22 231 interactions than female residents using a binomial test.  
23  
24  
25  
26

27 233 *Reproductive benefits of philopatry (parentage acquisition and territory inheritance)*

28  
29 234 Previously, it was shown that parentage success was substantially higher for female than male  
30  
31 235 subordinates (Richardson et al. 2002), so that we can exclude this factor as an explanation for  
32  
33 236 female-biased dispersal.  
34  
35

36 237 To assess whether the chances of territory inheritance are different for males and  
37  
38 238 females, we used a number of approaches. First, for 96 territory vacancies where a  
39  
40 239 subordinate was resident in a natal territory at the time a vacancy arose, we tested directly  
41  
42 240 whether inheritance occurred more for female vacancies than for male vacancies using a  $\chi^2$ -  
43  
44 241 test. Second, we used a generalized linear mixed model to test whether individuals were more  
45  
46 242 related to the opposite-sex breeder (response variable) if they inherited than if they dispersed,  
47  
48 243 and tested whether this effect was different between the sexes (included as independent  
49  
50 244 variables and their interaction), including 'natal-territory-identity' of the focal individual as a  
51  
52 245 random effect (excluding seven pairs for which relatedness could not be determined due to an  
53  
54 246 unsampled breeder). Third, if males can expel females, we would expect that males are more  
55  
56  
57  
58  
59  
60

1  
2  
3 247 likely than females to inherit the territory if the opposite-sex is related because males may  
4  
5 248 expel their mothers, but females not their fathers. Therefore, we tested whether the likelihood  
6  
7 249 that subordinates filled a vacancy in their territory (response variable) was dependent on the  
8  
9  
10 250 interaction between the sex of the subordinate and whether the opposite-sex breeder was  
11  
12 251 related or not (social parent or not; based on social pedigree data). We removed one  
13  
14 252 individual for which it was unknown whether the remaining breeder was related and 18 cases  
15  
16 253 where more than one same-sex subordinate was present, as the presence of a same-sex  
17  
18 254 subordinate reduces the chance that an individual would inherit the territory. Fourth, to  
19  
20 255 determine whether the probability of staying as breeder after inheriting a position with a  
21  
22 256 related partner was different between sexes, we determined whether females in incestuous  
23  
24  
25 257 pairs were more likely to divorce and leave than males.  
26  
27  
28 258

### 29 *Costly dispersal*

30  
31  
32 260 In order to make inferences about sex-biased costs of dispersal, we tested whether male and  
33  
34 261 female prospectors / floaters differed in the probability that before the next breeding season  
35  
36 262 they (1) died (Fisher exact test) and (2) obtained a breeding position ( $\chi^2$  test; including  
37  
38 263 individuals who died). As individuals only prospect or float before they are two years of age  
39  
40 264 (see Kingma et al. 2016b), we restricted the analyses to birds younger than two years. In each  
41  
42 265 season we determined whether individuals prospected or floated (see Kingma et al. 2016b for  
43  
44 266 details), and each individual was included in only one season, as only one female was  
45  
46  
47 267 observed prospecting for two years in a row (for which only the first season was included).

48  
49 268 On occasion, foraging birds were caught opportunistically together with resident  
50  
51 269 territory owners while they were involved in intraspecific chases (Kingma et al. 2016a).  
52  
53 270 Therefore, to make inferences about whether foraging males were more often attacked by  
54  
55  
56 271 territory owners than foraging females, we tested whether foraging (prospecting and floating  
57  
58  
59  
60

1  
2  
3 272 combined) males were more often caught with resident territory owners than foraging female  
4  
5 273 (using all catches). We fitted a GLMM with ‘bird-identity’ as a random factor to account for  
6  
7 274 multiple inclusions of individuals, whether or not an individual was caught with a resident  
8  
9 275 individual as a response variable, and whether individuals prospected or floated, and sex (and  
10  
11 276 their interaction), as independent variables.  
12  
13

14 277

### 17 278 *Ethical note*

18  
19 279 All protocols conformed to legal requirements for use of animals in research and were  
20  
21 280 approved by Seychelles Department of Environment and Seychelles Bureau of Standards  
22  
23 281 (permit: A0157).  
24  
25

26 282

## 28 283 **RESULTS**

### 30 284 **Sex-biased dispersal distance and prospecting behavior**

31  
32 285 To test for sex bias in dispersal distance and prospecting behavior we compared these two  
33  
34 286 measures between males and females. Females dispersed further from their natal territory  
35  
36 287 than males (Fig. 1a). Females (median = 5 territories distance; range = 3-12) were also  
37  
38 288 observed prospecting further than males (median = 3; range = 3-6;  $\beta = 0.460 \pm 0.156$ ,  $z =$   
39  
40 289 2.95,  $P = 0.003$ ). We have shown elsewhere that females prospect more often than males  
41  
42 290 (annually 19% of 175 females and 9% of 162 males; Kingma et al. 2016b).  
43  
44

45 291

### 48 292 **Inbreeding avoidance**

49  
50 293 Inbreeding avoidance is generally hypothesized to underlie sex-biased dispersal distance.  
51  
52 294 However, dispersal distance (excluding inheriting individuals) did not affect the relatedness  
53  
54 295 of the resulting breeding pair and this was similar for male and female dispersers (Table 2a;  
55  
56 296 Fig. 1b).  
57  
58  
59  
60

1  
2  
3 297 If females disperse further to avoid the risk of mating incestuously with an extra-  
4  
5 298 group father (nearby males), then we would expect to find that short-distance dispersal by  
6  
7 299 females would be more likely to result in incestuous pairs. However, short-distance  
8  
9 300 dispersing females were not more related to their new partner ( $R = 0.030 \pm 0.036$ ,  $n = 48$ )  
10  
11 301 than further-dispersing females ( $R = 0.004 \pm 0.018$ ,  $n = 143$ , Table 2c) or short-distance  
12  
13 302 dispersing males ( $R = 0.035 \pm 0.018$ ,  $n = 128$ ; Table 2d).  
14  
15  
16  
17

303

### 304 **Resource-holding potential**

305 Tarsus length and body mass were used as measures of competitive ability and resource-  
306 holding potential. On average, males ( $25.9 \pm 0.04$  mm,  $n = 309$ ) had a 6.3% longer tarsus  
307 length than females ( $24.4 \pm 0.03$  mm,  $n = 274$ ;  $t = -29.96$ ,  $P < 0.001$ ). Similarly, males (mean  
308 mass =  $16.5 \pm 0.03$  g,  $n = 784$  catches) were on average 9.6% heavier than females ( $15.0 \pm$   
309  $0.04$  g;  $n = 576$ ;  $\beta = -1.425 \pm 0.060$ ,  $t = -23.95$ ,  $P < 0.001$ ; correcting for time of capture  
310 (relative to morning): midday:  $\beta = 0.053 \pm 0.056$ ,  $t = 0.948$ ,  $P = 0.34$ , afternoon:  $\beta = 0.255 \pm$   
311  $0.055$ ,  $t = 4.65$ ,  $P < 0.001$ ).

312 Despite the asymmetry in size, however, both sexes were equally represented in  
313 agonistic interactions with intruders (62 of 121 attacks (51%) were performed by females and  
314 59 (49%) by males; binomial exact test:  $P = 0.86$ ). This included attacks by 19 subordinates  
315 (10 females, 9 males), but excluding these did not change the result ( $P = 0.92$ ).

316

### 317 **Reproductive benefits of philopatry (parentage acquisition and territory inheritance)**

318 If the reproductive benefits of philopatry (parentage acquisition and territory inheritance) are  
319 different between the sexes, then this might lead to sex-biased dispersal distance. As we  
320 could exclude differential parentage acquisition by subordinates as mechanism based on  
321 previous research (female subordinates are more likely to reproduce than male subordinates;

1  
2  
3 322 Richardson et al. 2002), we tested subsequently whether females were more likely to inherit  
4  
5 323 the territory than males. This was, however, not the case. First, the chances of territory  
6  
7 324 inheritance are equal for males and females: overall, 41 of the 433 breeding vacancies (9.5%)  
8  
9 325 were filled by inheritance (Fig. 1a), and if the respective-sex subordinate was present then  
10  
11 326 inheritance was equally likely for female (23 of 56 cases; 41.1%) and male vacancies (17 of  
12  
13 327 40 cases, 42.5%;  $\chi^2_1 < 0.01$ ,  $P = 1.00$ ). Second, as expected given that most subordinates are  
14  
15 328 retained offspring, inheriting individuals ( $R = 0.221 \pm 0.039$ ,  $n = 41$ ) were more related to  
16  
17 329 their partner than dispersing individuals ( $R = 0.020 \pm 0.011$ ,  $n = 385$ ) but this effect was not  
18  
19 330 different between males and females (Table 2b, Fig. 1b). Third, the likelihood of filling a  
20  
21 331 vacancy in the resident territory tended to be higher when subordinates were unrelated to the  
22  
23 332 opposite-sex breeder (12 of 23, 52.2%) than when the remaining breeder was a social parent  
24  
25 333 (16 of 54, 29.6%;  $\beta = -0.952 \pm 0.513$ ,  $t = -1.86$ ,  $P = 0.063$ ), but this effect did not depend on  
26  
27 334 the sex of the subordinate (interaction:  $\beta = 1.708 \pm 1.082$ ,  $z = 1.58$ ,  $P = 0.11$ ). Social fathers  
28  
29 335 were equally likely to accept their daughter as a partner (in 8 of 31 cases; 25.8%) as social  
30  
31 336 mothers were to accept their son (in 8 of 23 cases; 34.8%;  $\chi^2_1 = 0.171$ ,  $P = 0.68$ ). Fourth,  
32  
33 337 although the small sample of divorcing incestuous pairs did not permit adequate statistical  
34  
35 338 testing, only 1 of 8 father–daughter pairings and 3 of 8 mother–son pairings ended by  
36  
37 339 divorce, where in two cases the female left (daughter and mother) and in two cases the son  
38  
39 340 left.  
40  
41  
42  
43  
44  
45  
46

341

### 342 **Costly dispersal**

343 By assessing sex differences in mortality and intraspecific interactions of floaters and  
344 prospectors, we studied the potential sex-biased costs of dispersal. Male floaters were  
345 significantly more likely to die than female floaters before the beginning of the next season  
346 (Fig. 2), and male floaters tended to be less likely to obtain a breeding position before the

1  
2  
3 347 beginning of the next season (47% of 15 individuals) than female floaters (79% of 19)  
4  
5 348 (although this result was not significant;  $\chi^2_1 = 2.51$ ,  $P = 0.11$ ). Males and females that  
6  
7 349 engaged in temporary prospecting trips (before returning to their natal territory) were equally  
8  
9 350 likely to die (Fig. 2) and to obtain a breeding position before the beginning of the next season  
10  
11 351 (62% of 13 males and 54% of 39 females obtained a position;  $\chi^2_1 = 0.01$ ,  $P = 0.94$ ). Male  
12  
13 352 forayers (prospectors and floaters combined) were nearly twice as often caught with a  
14  
15 353 resident individual (29% of 24 catches) than female forayers (15% of 39 catches) but this  
16  
17 354 effect was not statistically significant, either alone ( $\beta = 0.804 \pm 0.658$ ,  $z = 1.22$ ,  $P = 0.22$ ) or  
18  
19 355 in interaction with whether individuals prospected or floated ( $\beta = -2.465 \pm 1.665$ ,  $z = -1.48$ ,  $P$   
20  
21 356  $= 0.14$ ). The likelihood of being caught with a resident individual, did not differ between  
22  
23 357 prospectors (20% of 35 individuals) and floaters (21% of 28;  $\beta = 0.206 \pm 0.744$ ,  $z = -0.28$ ,  $P$   
24  
25 358  $= 0.78$ ).

26  
27  
28  
29  
30 359

## 31 360 **DISCUSSION**

### 32 361 **Sex-biased dispersal distance in Seychelles warblers**

33  
34  
35 362 In line with many studies (see Greenwood 1980) and with previous work on Seychelles  
36  
37 363 warblers (Eikenaar et al. 2008a), we show that female subordinate Seychelles warblers  
38  
39 364 disperse further from their natal territory to obtain a breeding position than males. This effect  
40  
41 365 was not the result of females floating more often (i.e. permanently leaving a natal territory to  
42  
43 366 search for a vacancy, likely occurring because of eviction by breeders; Eikenaar et al. 2007,  
44  
45 367 Kingma et al. 2016b). However, female Seychelles warblers engage in temporary prospecting  
46  
47 368 trips more often than males (19% vs 9%; Kingma et al. 2016b) and they prospect over larger  
48  
49 369 distances than males (this study; see also Eikenaar et al. 2008a). This suggests that males  
50  
51 370 generally remain as subordinates within a territory and explains why males often shift to a  
52  
53 371 nearby vacancy when the opportunity arises (Fig. 1a), whereas females more often actively  
54  
55  
56  
57  
58  
59  
60

1  
2  
3 372 search for such vacancies, resulting in a breeding position further afield. There are several  
4  
5 373 potential explanations for sex-biased dispersal (Table 1), and below we discuss whether these  
6  
7 374 can explain sex-biased dispersal distance in Seychelles warblers.  
8  
9

10 375

11 376 **Benefits of philopatry, incest and inbreeding avoidance, and resource-holding potential**

12  
13  
14 377 We tested the predictions of several hypotheses for sex-biased dispersal (Table 1), but the  
15  
16 378 results indicate that we can reject most of these hypotheses to explain female-biased dispersal  
17  
18 379 distance in Seychelles warblers.

19  
20  
21 380 Sex-biased dispersal in species where subordinate individuals delay dispersal is  
22  
23 381 hypothesized to be based on differences in reproductive benefits-of-philopatry (i.e., in  
24  
25 382 reproduction as subordinate, territory inheritance or differences in competitive ability and  
26  
27 383 resource holding potential; Greenwood 1980; Zack and Rabenold 1989; Richardson et al.  
28  
29 384 2002). We can rule out sex differences in reproductive benefits gained by subordinates as an  
30  
31 385 explanation for female-biased dispersal distance in Seychelles warblers: subordinate females  
32  
33 386 are more likely to gain parentage than subordinate males (Richardson et al. 2001; Richardson  
34  
35 387 et al. 2002), and territory inheritance is rare (9.5% of positions) and achieved equally by  
36  
37 388 males and females. Furthermore, although incest avoidance inhibits territory inheritance to  
38  
39 389 some extent (whether or not the opposite-sex breeder was a social parent tended to predict ( $P$   
40  
41 = 0.06) whether subordinates inherited), incestuous pairs are formed in ca. 30% of the cases  
42  
43 390 and this was not more likely for female subordinates than for males. Moreover, although the  
44  
45 391 sample size was small, incestuous pairs between mothers and sons did not always end in  
46  
47 392 divorce (only in 3 of 8 cases) and, importantly, were not more likely to end in a female  
48  
49 393 leaving than a male. This suggests that males are not necessarily dominant over females and  
50  
51 394 that this cannot explain female-biased dispersal distance in this species. Similarly, although  
52  
53 395 males are larger and heavier, males did not appear to be more engaged in territory defence  
54  
55 396  
56  
57  
58  
59  
60

1  
2  
3 397 than females (as is, for example, the case in some migratory species where males arrive  
4  
5 398 earlier at the breeding ground to establish territories; Arlt and Pärt 2008), leaving the  
6  
7 399 'resource-holding potential hypothesis' unlikely as a direct explanation for our results. The  
8  
9  
10 400 latter is also confirmed by observations that, after the disappearance of a breeding male,  
11  
12 401 females are capable of holding the territory until she pairs with a new male (median duration  
13  
14 402 until the male vacancy was filled in an experimental removal of breeding males was 2 days;  
15  
16 403 Eikenaar et al. 2009). Thus, overall, these results suggest that differences in reproductive  
17  
18 404 benefits for philopatric individuals and resource-holding potential cannot explain sex-biased  
19  
20 405 dispersal distance in Seychelles warblers.

21  
22  
23 406 A previous study suggested that inbreeding avoidance may underlie sex-biased natal  
24  
25 407 dispersal in Seychelles warblers (Eikenaar et al. 2008). Based on considerable levels of extra-  
26  
27 408 pair paternity in this species (~40% of offspring; Richardson et al. 2001), Eikenaar et al.  
28  
29 409 (2008) hypothesized that females might generally disperse further than males because  
30  
31 410 females who pair with local individuals risk pairing incestuously with their extra-pair father.  
32  
33 411 Although females are indeed less likely to disperse to territories in the close vicinity (Fig. 1a),  
34  
35 412 our investigation of relatedness of breeding pairs in relation to dispersal distance does not  
36  
37 413 support this hypothesis. First, females pairing with a male within one or two territories of  
38  
39 414 their natal territory were not more related to that partner than either subordinate males pairing  
40  
41 415 with a female within a similar distance or females dispersing over a larger distances (Fig. 1b),  
42  
43 416 as would be expected if females had a high likelihood of pairing with an extra-pair father.  
44  
45 417 Second, although territory inheritance did frequently result in incestuous pairs (see above),  
46  
47 418 the distance that females and males dispersed did not predict the relatedness of the resulting  
48  
49 419 breeding pair, suggesting that inbreeding-avoidance does not underlie sex-biased dispersal  
50  
51 420 distance in this species.  
52  
53  
54  
55  
56  
57  
58  
59  
60

**422 Costly dispersal**

423 Our finding of lower mortality of female floaters compared to male floaters suggests that  
424 females may be more tolerated than males outside their resident territory. These differences  
425 in the costs of searching for an independent breeding territory may underlie the lower rate of  
426 prospecting by male subordinate Seychelles warblers than females, resulting in a shorter  
427 ultimate dispersal distance. The survival costs of prospecting did not differ between the  
428 sexes, but this may well be explained by prospecting individuals being able to return to their  
429 resident territory after an unsuccessful prospecting trip, thereby obtaining the benefits of  
430 philopatry like nepotistic benefits facilitating access to food (Kingma et al. 2016b).  
431 Nonetheless, several findings suggest that extra-territorial movement is more costly for male  
432 than for female Seychelles warblers. First, males experienced a higher mortality cost of  
433 floating than females (Fig. 2). Differential costs of floating are generally difficult to tease  
434 apart from variation in quality of individuals that leave (i.e. floaters may have been of poorer  
435 quality). However, that subordinate male Seychelles warblers search for vacancies (by means  
436 of prospecting) less often than females, is in line with males being reluctant to leave  
437 voluntarily because of relatively high costs. Second, Eikenaar et al. (2008a) found that males,  
438 but not females, in more densely-populated areas dispersed less far (independent of territory  
439 quality), which may suggest that dispersal by males in dense areas is more costly, perhaps  
440 because intruders are detected faster in denser areas resulting in an increasing number of  
441 attacks. Third, in line with reduced prospecting (and the resulting restriction to being able to  
442 only fill nearby vacancies), males were shown to obtain a breeding position on average at an  
443 older age than females (Eikenaar et al. 2009). While we can rule out predation (predation of  
444 adults is absent in this species), the exact mechanism for higher costs of dispersal for males is  
445 unknown so far. Although statistically not significant (perhaps due to limited sample size) our  
446 finding that foraging male subordinates were nearly twice as likely to be caught with an

1  
2  
3 447 individual resident in the intruded territory may indicate that males are attacked more often  
4  
5 448 than females. Perhaps males pose a greater threat to parentage of territory owners (e.g. due to  
6  
7 449 extra-pair mating) but females are equally involved in territory defence and such sex-biased  
8  
9 450 costs would only manifest during a brief period when individuals initiate breeding. Thus, the  
10  
11 451 underlying mechanism for sex-biased dispersal costs, and also to what extent these costs  
12  
13 452 apply in species where the habitat is not saturated, is an interesting topic for further  
14  
15 453 investigation. Ultimately, however, our results suggest that the fact that males often disperse  
16  
17 454 only one or two territories can be explained by the relatively high costs of extra-territorial  
18  
19 455 movement driving males to wait for local opportunities to disperse. Thus, asymmetry in costs  
20  
21 456 of dispersal may explain sex-biased dispersal distance in this, and possibly other species (see  
22  
23 457 also Perrin and Mazalov 2000; Gros et al. 2008; Pakanen et al. 2016).

24  
25  
26  
27 458

## 28 29 459 **Conclusions**

30  
31 460 We conclude that sex-biased dispersal distance in Seychelles warblers is unlikely to support  
32  
33 461 the currently often invoked hypotheses (see Table 1). Female-biased dispersal distance does  
34  
35 462 not seem to be explained by inbreeding-avoidance and differences in reproductive benefits of  
36  
37 463 philopatry and resource-holding potential. Instead, our data suggest that dispersal attempts  
38  
39 464 are more costly for males than for females. This may explain why selection favours reduced  
40  
41 465 male subordinate extra-territorial movement required to find a breeding vacancy, which in  
42  
43 466 turn likely leads to reduced dispersal distances and later acquisition of a breeding position  
44  
45 467 (Eikenaar et al. 2009). This additional mechanistic explanation may not only shed light on  
46  
47 468 sex-biased dispersal in family-living and cooperatively breeding species, but costs associated  
48  
49 469 with searching for an independent breeding position may potentially also play a role in  
50  
51 470 dispersal strategies in non-social species.

52  
53  
54  
55  
56 471  
57  
58  
59  
60

1  
2  
3 472 **REFERENCES**

- 4  
5 473 Aars J, Ims RA. 2000. Population dynamics and genetic consequences of spatial density-  
6  
7 474 dependent dispersal in patchy populations. *Am Nat.* 155:252–265.
- 8  
9  
10 475 Arlt D, Pärt T. 2008. Sex-biased dispersal: a result of a sex difference in breeding site  
11  
12 476 availability. *Am Nat.* 171:844-850.
- 13  
14 477 Bates D, Maechler M, Bolker B, Walker S. 2015. Fitting linear mixed-effects models using  
15  
16 478 lme4. *J Stat Softw.* 67:1-48.
- 17  
18 479 Bowler DE, Benton TG. 2005. Causes and consequences of animal dispersal strategies:  
19  
20 480 relating individual behaviour to spatial dynamics. *Biol Rev.* 80:205-225.
- 21  
22  
23 481 Brown JL. 1987. Helping and communal breeding in birds. Princeton: Princeton University  
24  
25 482 Press.
- 26  
27 483 Clarke AL, Sæther B-E, Roskaft E. 1997. Sex biases in avian dispersal: a reappraisal. *Oikos.*  
28  
29 484 79:429–438.
- 30  
31  
32 485 Cockburn A. 1998. Evolution of helping behaviour in cooperatively breeding birds. *Ann Rev*  
33  
34 486 *Ecol Syst.* 29:141–177.
- 35  
36 487 Eikenaar C, Richardson DS, Brouwer L, Komdeur J. 2008. Sex biased natal dispersal in a  
37  
38 488 closed, saturated population of Seychelles warblers *Acrocephalus sechellensis*. *J Avian*  
39  
40 489 *Biol.* 39:73-80.
- 41  
42  
43 490 Eikenaar C, Komdeur J, Richardson DS. 2008. Natal dispersal patterns are not associated  
44  
45 491 with inbreeding avoidance in the Seychelles Warbler. *J Evol Biol.* 21:1106-1116.
- 46  
47 492 Eikenaar C, Richardson DS, Brouwer L, Bristol R, Komdeur J. 2009. Experimental  
48  
49 493 evaluation of sex differences in territory acquisition in a cooperatively breeding bird.  
50  
51 494 *Behav Ecol.* 20:207-214.
- 52  
53  
54 495 Greenwood PJ, Harvey PH. 1982. The natal and breeding dispersal of birds. *Ann Rev Ecol*  
55  
56 496 *Syst.* 13:1-21.
- 57  
58  
59  
60

- 1  
2  
3 497 Greenwood PJ. 1980. Mating systems, philopatry and dispersal in birds and mammals. *Anim*  
4  
5 498 *Behav.* 28:1140-1162.  
6  
7 499 Griesser M, Nystrand M, Ekman J. 2006. Reduced mortality selects for family cohesion in a  
8  
9 500 social species. *Proc R Soc Lond B* 273:1881–1886.  
10  
11 501 Griffiths R, Double MC, Orr K, Dawson RJG. 1998. A DNA test to sex most birds. *Mol*  
12  
13 502 *Ecol.* 7:1071–1075.  
14  
15 503 Gros A, Hovestadt T, Poethke HJ. 2008. Evolution of sex-biased dispersal: The role of sex-  
16  
17 504 specific dispersal costs, demographic stochasticity, and inbreeding. *Ecol Model.*  
18  
19 505 219:226-233.  
20  
21 506 Hammers M, Kingma SA, Bebbington K, Van de Crommenacker J, Spurgin L, Richardson  
22  
23 507 DS, Komdeur J. 2015. Senescence in the wild: insights from a long-term study on  
24  
25 508 Seychelles warblers. *Exp Geront.* 71:69-79.  
26  
27 509 Johnson ML, Gaines MS. 1990. Evolution of dispersal: theoretical models and empirical tests  
28  
29 510 using birds and mammals. *Ann Rev Ecol Syst.* 21:449–480.  
30  
31 511 Kingma SA, Komdeur J, Hammers M, Richardson DS. 2016a. The cost of prospecting for  
32  
33 512 dispersal opportunities in a social bird. *Biol Lett.* 10.1098/rsbl.2016.0316.  
34  
35 513 Kingma SA, Bebbington K, Hammers M, Richardson DS, Komdeur J. 2016b. Delayed  
36  
37 514 dispersal and the costs and benefits of different routes to independent breeding in a  
38  
39 515 cooperatively breeding bird. *Evolution.* DOI: 10.1111/evo.13071.  
40  
41 516 Kuznetsova A, Brockhoff PB, Christensen RHB. 2016. lmerTest: Tests in linear mixed  
42  
43 517 effects models. R package version 2.0-30. [https://CRAN.R-](https://CRAN.R-project.org/package=lmerTest)  
44  
45 518 [project.org/package=lmerTest](https://CRAN.R-project.org/package=lmerTest)  
46  
47 519 Koenig WD, Dickinson J. 2004. Ecology and evolution of cooperative breeding birds.  
48  
49 520 Cambridge: Cambridge University Press.  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

- 1  
2  
3 521 Koenig WD, Van Vuren D, Hooge PN. 1996. Detectability, philopatry, and the distribution of  
4  
5 522 dispersal distances in vertebrates. *Trends Ecol Evol.* 11:514-517.  
6  
7 523 Koenig WD, Stacey PB. 1990. Acorn woodpeckers: group-living and food storage under  
8  
9 524 contrasting ecological conditions. In: Stacey PB, Koenig WD, editors. *Cooperative*  
10  
11 525 *breeding in birds: long-term studies of ecology and behaviour.* Cambridge: Cambridge  
12  
13 526 University Press. p. 413-453.  
14  
15  
16 527 Kokko H, Ekman J. 2002. Delayed dispersal as a route to breeding: territorial inheritance,  
17  
18 528 safe havens, and ecological constraints. *Am Nat.* 160:468-484.  
19  
20  
21 529 Komdeur J. 1992. Importance of habitat saturation and territory quality for the evolution of  
22  
23 530 cooperative breeding in the Seychelles warbler. *Nature.* 358:493-495.  
24  
25 531 Komdeur J, Piersma T, Kraaijeveld K, Kraaijeveld-Smit F, Richardson DS. 2004. Why  
26  
27 532 Seychelles warblers fail to recolonize nearby islands: unwilling or unable to fly there?  
28  
29 533 *Ibis.* 146:298-302.  
30  
31  
32 534 Lawson Handley LJ, Perrin N. 2007. Advances in our understanding of mammalian sex-  
33  
34 535 biased dispersal. *Mol Ecol.* 16:1559-1578.  
35  
36 536 Nelson-Flower MJ, Hockey PAR, O’Ryan C, Ridley AR. 2012. Inbreeding avoidance  
37  
38 537 mechanisms: dispersal dynamics in cooperatively breeding southern pied babblers. *J*  
39  
40 538 *Anim Ecol.* 81:876-883.  
41  
42  
43 539 Pakanen V-M, Koivula K, O’Rell M, Rytönen S, Lahti K. 2016. Sex-specific mortality costs  
44  
45 540 of dispersal during the post-settlement stage promote male philopatry in a resident  
46  
47 541 passerine. *Behav Ecol Sociobiol.* doi:10.1007/s00265-016-2178-z  
48  
49  
50 542 Peakall R, Smouse PE. 2012. GenAIEx 6.5: genetic analysis in Excel. Population genetic  
51  
52 543 software for teaching and research - an update. *Bioinformatics.* 28:2537-2539.  
53  
54  
55  
56  
57  
58  
59  
60

- 1  
2  
3 544 Perrin N, Goudet J. 2001. Inbreeding, kinship, and the evolution of natal dispersal. In:  
4  
5 545 Clobert J, Danchin E, Dhondt AA, Nichols JD, editors. Oxford: Oxford University  
6  
7 546 Press. p. 123-142.  
8  
9  
10 547 Perrin N, Mazalov V. 2000. Local competition, inbreeding, and the evolution of sex-biased  
11  
12 548 dispersal. *Am Nat.* 155:116–127.  
13  
14 549 Prugnolle F, De Meeus T. 2002. Inferring sex-biased dispersal from population genetic tools:  
15  
16 550 a review. *Heredity.* 88:161–165.  
17  
18 551 Pusey AE. 1987. Sex-biased dispersal and inbreeding avoidance in birds and mammals.  
19  
20 552 *Trends Ecol Evol.* 2:295-299.  
21  
22  
23 553 Pusey A, Wolf M. 1996. Inbreeding avoidance in animals. *Trends Ecol Evol.* 11:201-206.  
24  
25 554 Queller DC, Goodnight KF. 1989. Estimating relatedness using genetic markers. *Evolution.*  
26  
27 555 43:258-275.  
28  
29  
30 556 R core-team. 2016. *R: a language and environment for statistical computing*. R foundation  
31  
32 557 for Statistical Computing, Vienna, Austria. <http://www.R-project.org>.  
33  
34 558 Reed JM, Boulinier T, Danchin E, Oring LW. 1999. Informed dispersal: prospecting by birds for  
35  
36 559 breeding sites. In: Nolan JV, Ketterson ED, Thompson CF, editors. *Current ornithology*. New  
37  
38 560 York: Kluwer Academic/Plenum Publishers. p. 189-259.  
39  
40 561 Richardson DS, Jury F, Dawson D, Salgueiro P, Komdeur J, Burke T. 2000. Fifty Seychelles  
41  
42 562 warbler (*Acrocephalus sechellensis*) microsatellite loci polymorphic in Sylviidae  
43  
44 563 species and their cross-species amplification in other passerine birds. *Mol Ecol.*  
45  
46 564 9:2226–2231.  
47  
48  
49 565 Richardson DS, Jury FL, Blaakmeer K, Komdeur J, Burke T. 2001. Parentage assignment  
50  
51 566 and extra-group paternity in a cooperative breeder: the Seychelles warbler  
52  
53 567 (*Acrocephalus sechellensis*). *Mol Ecol.* 10:2263–2273.  
54  
55  
56 568 Richardson DS, Burke T, Komdeur J. 2002. Direct benefits explain the evolution of female  
57  
58 569 biased cooperative breeding in the Seychelles warblers. *Evolution.* 56:2313–2321.  
59  
60

- 1  
2  
3 570 Ridley AR. 2012. Invading together: the benefits of coalition dispersal in a cooperative bird.  
4  
5 571 Behav Ecol Sociobiol. 66:77–83.  
6  
7 572 Ridley AR, Raihani NJ, Nelson-Flower MJ. 2008. The cost of being alone: the fate of floaters  
8  
9 573 in a population of cooperatively breeding pied babblers *Turdoides bicolor*. J Avian  
10  
11 574 Biol. 39:389-392.  
12  
13 575 Spurgin LG, Wright DJ, Van der Velde M, Collar NJ, Komdeur J, Burke T, Richardson DS.  
14  
15 576 2014. Museum DNA reveals the demographic history of the endangered Seychelles  
16  
17 577 warbler. Ecol Appl. 7:1134-1143.  
18  
19 578 Yaber MC, Rabenold KN. 2002. Effects of sociality on short-distance, female-biased  
20  
21 579 dispersal in tropical wrens. J Anim Ecol. 71:1042-1055.  
22  
23 580 Zack S, Rabenold KN. 1989. Assessment, age and proximity in dispersal contests among  
24  
25 581 cooperative wrens - field experiments. Anim Behav. 38:235–247.  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

582 **FIGURE LEGENDS**

583 **Figure 1.** (a) Dispersal distance (minimum number of territories traversed between an  
584 individual's natal territory and the territory where they obtained a breeding position) of  
585 female (black bars) and male (grey bars) Seychelles warblers, and (b) the mean ( $\pm$ SE)  
586 relatedness of the ultimate breeding pairs in relation to the focal individual's dispersal  
587 distance. Numbers reflect number of individuals. Females dispersed on average further than  
588 males ( $\beta = 0.604 \pm 0.061$ ,  $z = 9.95$ ,  $P < 0.001$ ). However, dispersal distance did not predict  
589 relatedness to the obtained partner for both females and males, other than that inheriting  
590 individuals (dispersal distance 0) obtained a more related partner than dispersing individuals.

591

592 **Figure 2.** The likelihood that prospecting and floating Seychelles warbler females (black  
593 bars) and males (grey bars) died before the subsequent season. Whereas there was no  
594 significant difference between prospecting males and females (Fisher exact test:  $P = 1.00$ ),  
595 male floaters were more likely to die than female floaters (Fisher exact test:  $P = 0.03$ ).

596

## 597 TABLES

598 **Table 1.** An overview of the concepts of the main hypotheses of sex-biased dispersal in  
 599 cooperatively breeding birds. A set of predictions was developed to test whether these  
 600 hypotheses explain female-biased natal dispersal distance in Seychelles warblers. Whether  
 601 these predictions are met is stated in the final column.  
 602

Hypothesis	Concept	Prediction in Seychelles warblers (female-biased dispersal)	Prediction met?
Inbreeding avoidance	Dispersal leads to less related partner	Further dispersal leads to a less related partner	No (Fig. 1b)
		Females who disperse to nearby territories have a higher likelihood engage in an incestuous relationship (with extra-group father) than females who disperse further	No, locally dispersing females did not obtain a more related partner than far-distance dispersing females (Fig. 1b, Table 2c)  No, relatedness between short-distance dispersers and their obtained partner was not different between males and females (Fig. 1b, Table 2d)
Resource-holding potential	Territory establishment and defence biased to one sex	Males are larger and heavier than females	Yes, males are 6% larger and 10% heavier
		Females defend less than males	No, females defend equally
Reproductive benefits of philopatry	Sex-biased dispersal driven by sex differences in ability to reproduce as subordinate	Subordinate males reproduce more than subordinate females	No, females reproduce more <sup>1</sup>
		Sex-biased dispersal driven by differences in chance of territory inheritance	Males are more likely to inherit the territory than females
	Inheritance improves chance of incest differently between males and females	No, although inheritance improves the chance of incestuous pairing, this was not different between males and females (Table 1b)	
	Mothers accept sons as partner more than fathers accept daughters <sup>2</sup>	No, likelihood is equal (35% vs 26%)	
Costly dispersal	Costs of dispersal or floating are sex-specific	Males expel females after they inherit	No, only 4 of 16 incestuous pairs ended in divorce: in 2 of these the female left, and in 2 cases the male left
		Sex-bias in reproductive threat of floaters for territory owners	Yes, extra-pair paternity occurs, but no egg-dumping, so that males are more of a threat <sup>1</sup>
		Males are attacked more in foreign territories than females	Maybe: male floaters are attacked twice as much but not significant (limited statistical power)
		Floating males are more likely to die than floating females	Yes, male floaters are 3.5 times as likely to die than female floaters (Fig. 2)

603 <sup>1</sup> 44% of subordinate females lay an egg, but only 1 of 55 young was sired by a subordinate male (Richardson et  
 604 al. 2001). <sup>2</sup> In species with extra-group mating, the risk of engaging in an incestuous pair after territory  
 605 inheritance is larger for males than for females, predicting male-biased dispersal. However, since dispersal  
 606 distance was female-biased in Seychelles warblers, we did not include that hypothesis here.

607

608 **Table 2.** The effect of sex and/or dispersal distance on relatedness of Seychelles warbler  
 609 pairs, when the focal subordinate (a) dispersed, (b) inherited or not, (c) dispersed long vs  
 610 short distance (females only) and (d) dispersed a short distance.

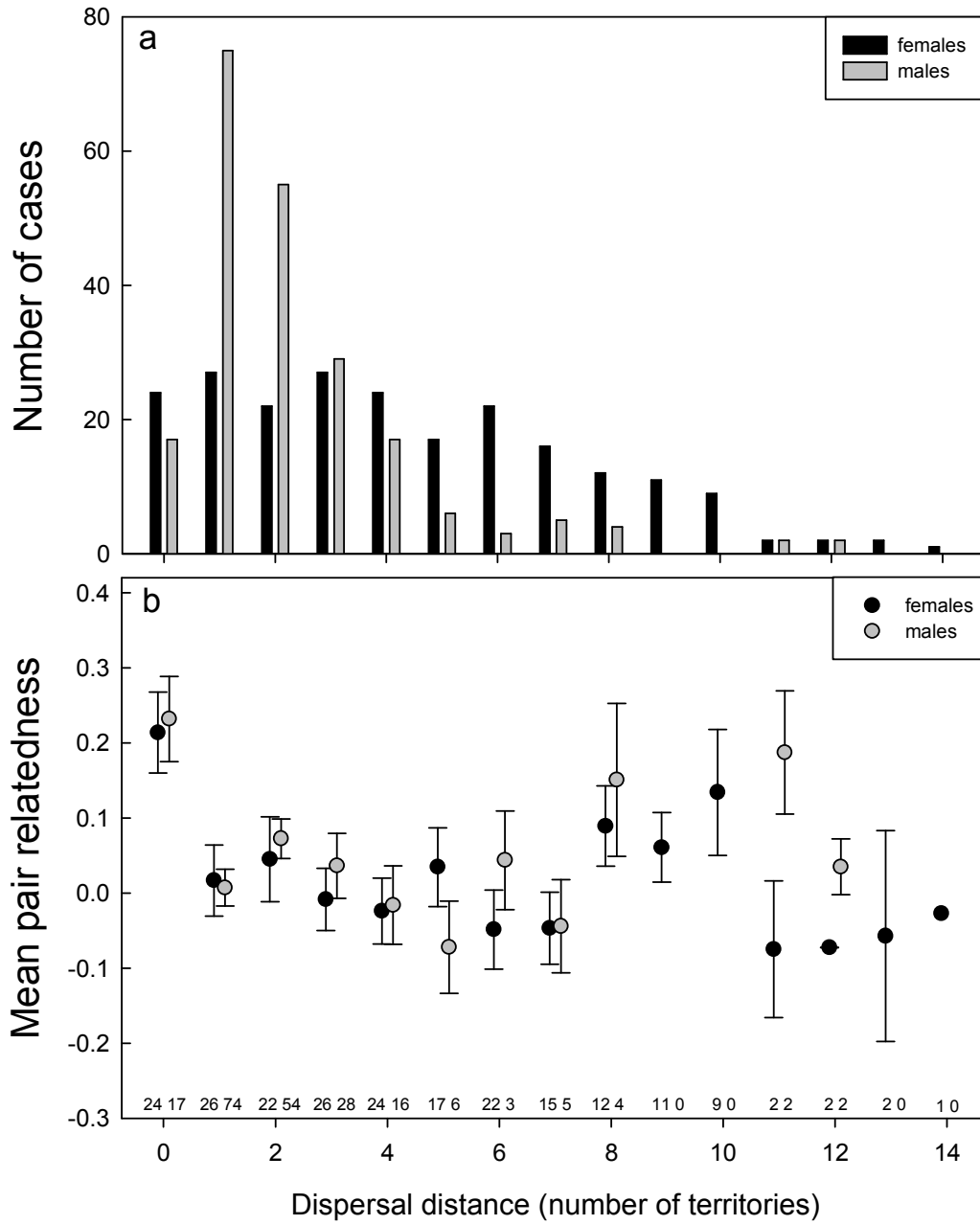
		$\beta$	SE	<i>t</i>	<i>P</i>
<b>a) Relatedness (non-inheriting pairs)</b>					
(385 individuals, 118 territories)	<i>Intercept</i>	0.020	0.011	1.82	0.07
	Dispersal distance	0.0004	0.004	0.10	0.92
	Sex	0.020	0.022	0.90	0.37
	Dispersal distance * sex	0.001	0.009	0.17	0.87
<b>b) Relatedness (inheritance vs. dispersal)</b>					
(426 individuals, 120 territories)	<i>Intercept</i>	0.020	0.011	1.81	0.07
	Inherited	0.201	0.036	5.67	< <b>0.001</b>
	Sex	0.019	0.021	0.91	0.36
	Inherited * sex	-0.001	0.072	-0.01	0.99
<b>c) Relatedness (females only)</b>					
(191 individuals, 86 territories)	<i>Intercept</i>	0.002	0.019	0.11	0.91
	Dispersal distance (short vs long)	0.027	0.037	0.72	0.47
<b>d) Relatedness (only short distance)</b>					
(176 individuals, 92 territories)	<i>Intercept</i>	0.033	0.016	2.02	<b>0.045</b>
	Sex	0.005	0.037	0.13	0.89

611

612

613 FIGURES

614 Figure 1.

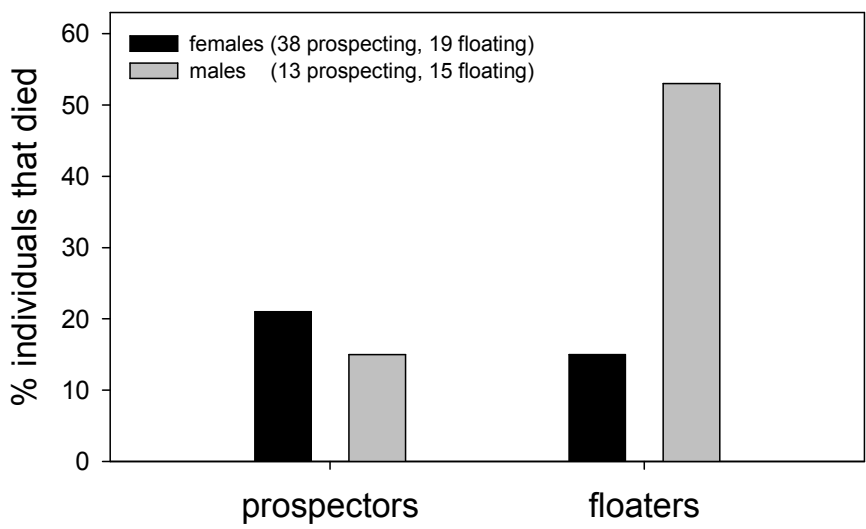


615

616

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

617 **Figure 2.**



618

Review Only