

# 1 **Rainfall is associated with divorce in the socially monogamous Seychelles warbler**

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11

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### 28 **Conflict of interest**

29 All authors have no competing interests.

30

### 31 **Author's contributions**

32 FJDS and HLD conceived the study question. AAB, FJDS, and HLD designed the hypotheses and  
33 methodology. AAB and FJDS performed the data selection. TB, JK, DSR, and HLD maintain the long-  
34 term dataset. DSR managed and undertook fieldwork over the period involved. AAB analyzed the data  
35 and wrote the manuscript with input from FJDS, HLD, DSR, and JK. All authors gave final approval  
36 for publication.

37

### 38 **Statement on inclusion**

39 The aid and input of local stakeholders from Nature Seychelles, who manage Cousin Island, in the  
40 conservation of, and research into, the Seychelles warbler is an important part of our work. As a result  
41 of this close interaction, underpinned by a memorandum of understanding, DSR is affiliated with  
42 Nature Seychelles, and we reciprocate by providing scientific support, intellectual input, and funding  
43 for facilities. Nature Seychelles is also included as an author affiliation in all Seychelles warbler  
44 publications.

45

### 46 **Data availability**

47 Data will be available on the University of Groningen dataverse.

48

49

50 **Abstract**

- 51 1. Divorce – terminating a pair bond while both members are alive – is a mating strategy observed  
52 in many socially monogamous species often linked to poor reproductive success. As  
53 environmental factors directly affect individual condition and reproductive performance, they  
54 can indirectly influence divorce. Given current climate change, understanding how  
55 environmental fluctuations affect partnership stability has important implications, including for  
56 conservation. Yet, the relationship between the environment and divorce remains largely  
57 unstudied.
- 58 2. We examined the influence of temporal environmental variability on the prevalence of within-  
59 and between-season divorce and the possible underlying mechanisms in a socially  
60 monogamous passerine.
- 61 3. Analyzing 16 years of data from a longitudinal dataset, we investigated the relationship  
62 between rainfall and divorce in the Seychelles warbler (*Acrocephalus sechellensis*). First, we  
63 performed climate window analyses to identify the temporal windows of rainfall that best  
64 predicted reproductive success and divorce. Then, we tested the effects of these temporal  
65 windows of rainfall on reproductive success and divorce and the influence of reproductive  
66 success on divorce while controlling for covariates.
- 67 4. Annual divorce rates varied from 1–16%. The probability of divorce was significantly  
68 associated with the quadratic effect of 7 months of total rainfall before and during the breeding  
69 season, with divorce increasing in years with low and high rainfall. Extreme rainfall events  
70 drove this quadratic relationship. Although the same temporal window of rainfall predicting  
71 divorce significantly influenced reproductive success, we found no significant correlation  
72 between reproductive success and divorce.
- 73 5. Our findings suggest that rainfall impacts divorce. Given that this effect is likely not directly  
74 mediated by reproductive success, we discuss the possible role of physiological stress. By

75 adding to the growing body of literature showing that environmental conditions influence the  
76 stability of socially monogamous partnerships, we provide novel insights that may also be  
77 important for conservation efforts in times of climate change.

78

79 **Keywords:** Climate window analysis, Divorce, Environmental conditions, Habitat-mediated  
80 hypothesis, Passerine, Rainfall, Seychelles warbler, Social monogamy.

81

## 82 **1 Introduction**

83 Social monogamy, the mating system where individuals have one social breeding partner at a time,  
84 occurs in over 90% of birds (Lack, 1968). In these systems, maintaining a pair bond across multiple  
85 breeding seasons can improve reproductive success by reducing the costs associated with mate  
86 searching and enhancing mate familiarity (Choudhury & Black, 1994; Sánchez-Macouzet *et al.*, 2014;  
87 Culina *et al.*, 2020). However, intra-sexual competition often constrains mate selection, resulting in  
88 suboptimal partnerships. Suboptimal partnerships may be corrected through divorce, whereby a pair  
89 bond is terminated while both partners are alive (Choudhury, 1995), which can either increase,  
90 decrease, or have no effect on future reproductive success for one or both partners (Culina *et al.*, 2015).

91

92 Divorce occurs in 92% of socially monogamous bird species (Jeschke & Kokko, 2008). With  
93 significant inter- and intra-species variation in divorce rates (Black, 1996), several hypotheses have  
94 been proposed to explain what causes divorce (Choudhury, 1995). For instance, divorce may correct  
95 for genetic or behavioral incompatibilities within partnerships (Wilson *et al.*, 2022) or enable  
96 individuals to choose a better-quality partner, such as one with a higher dominance status or one that  
97 occupies a better territory than their previous partner (Dhondt & Adriaensen, 1994; Otter & Ratcliffe,  
98 1996; Blondel *et al.*, 2000). Here, one or both pair-bonded individuals instigate divorce. Divorce can  
99 also be accidental, occurring due to temporal mismatches during migration (Gilsenan *et al.*, 2017) or

100 forced by the introduction of a third party (Jeschke *et al.*, 2007). Related to several of these hypotheses,  
101 previous reproductive success and divorce are often correlated, with reproductive failure being a strong  
102 predictor of partnership termination (Culina *et al.*, 2015). Notably, the effect of reproduction on  
103 divorce can vary depending on the stage of the breeding cycle, with failures at earlier breeding stages  
104 often being stronger predictors of divorce (Culina *et al.*, 2015).

105

106 As climate patterns create suboptimal environmental conditions that affect individual condition and  
107 reproductive performance, they can influence divorce. The ‘habitat-mediated’ hypothesis suggests  
108 divorce is more prevalent in unstable and lower-quality environments (Blondel *et al.*, 2000). This is  
109 because environmental factors can impact the decision-making process underpinning divorce by  
110 misinforming individuals about their partnership’s quality. For example, when partnerships perform  
111 poorly due to harsh environmental conditions, individuals within those partnerships may still attribute  
112 their poor performance to their chosen partner and not to the given circumstances (Ventura *et al.*,  
113 2021). Extreme weather can also increase physiological stress (Kitaysky *et al.*, 2010), an important  
114 factor influencing mate selection (Husak & Moore, 2008). Given the rapid timing of climate change,  
115 marked by more frequent extreme weather events, such as droughts and floods, and increased global  
116 temperatures (NOAA, 2022), which may limit possibilities for adaptation (Spooner *et al.*, 2018),  
117 understanding how climate patterns affect the stability of socially monogamous partnerships is critical.

118

119 The relationship between the ecological environment and divorce remains largely unstudied, with only  
120 a handful of publications (Blondel *et al.*, 2000; Heg *et al.*, 2003; Botero & Rubenstein, 2012; Ventura  
121 *et al.*, 2021; Lerch *et al.*, 2022). Existing studies are primarily cross-sectional, comparing the  
122 prevalence of divorce between species or populations of the same species. To our knowledge, Ventura  
123 *et al.* (2021) is the only longitudinal study to have analyzed the effects of climate-driven environmental  
124 conditions on divorce within the same population, discovering that, due to sea-surface temperatures

125 influencing food abundance and thus reproductive success, warmer sea-surface temperatures increased  
126 the probability of divorce in black-browed albatrosses (*Thalassarche melanophris*).

127

128 With climate change resulting in more frequent heavy rain and drought events (Marvel *et al.*, 2019),  
129 we aimed to investigate the relationship between rainfall and divorce by analyzing long-term  
130 longitudinal data from the socially monogamous Seychelles warbler (*Acrocephalus sechellensis*), a  
131 passerine endemic to the Seychelles archipelago. Extreme rainfall negatively impacts the warblers'  
132 reproductive output (Komdeur, 1996a; Borger *et al.*, 2023). As reproductive failures drive divorce in  
133 various bird species (Culina *et al.*, 2015), including the Seychelles warbler (Speelman *et al.*, 2024),  
134 we investigated whether 1) the temporal variability of rainfall affects the prevalence of divorce in the  
135 Seychelles warbler, 2) measures of reproductive success at four different stages of reproduction within  
136 the breeding season affect divorce, and 3) rainfall influences these four reproductive measures.

137

138 We predicted that extreme rainfall increases the prevalence of divorce (P1). As Seychelles warblers  
139 are insectivorous, low rainfall decreases food availability by impacting their prey's reproductive cycle  
140 (Komdeur, 1996a, Price, 1997). Conversely, high rainfall can affect the ability of birds to maintain  
141 optimal body temperatures and cause direct habitat and nest destruction (Kennedy, 1970; Wilson *et al.*,  
142 2004). Consequently, we predicted that low and high amounts of rainfall decrease reproductive  
143 success (P2). Specifically, due to decreased food availability, low rainfall impacts the ability of  
144 insectivorous birds to initiate breeding and produce a clutch (França *et al.*, 2020). Then, due to  
145 decreased food availability and increased metabolic demands in heavy rainfall conditions, low and  
146 high rainfall impact nestling and fledgling survival (Monadjem & Bamford, 2009; Heenen & Seymour,  
147 2012). The decreased reproductive success influenced by rainfall is predicted to increase the  
148 probability of divorce as reproductive success is used as a marker of a partnership's quality (P3).  
149 Overall, in line with the habitat-mediated hypothesis, we predicted that divorce would be more

150 prevalent following breeding seasons with poorer breeding conditions, with rainfall having a quadratic  
151 effect on divorce. Our findings may provide insights into how harsh environmental conditions affect  
152 reproduction and divorce in socially monogamous birds, which, in turn, can inform conservation  
153 efforts across multiple species in times of climate change, such as by informing population modeling.

154

## 155 **2 Materials and Methods**

### 156 **2.1 Study system**

157 Since 1985, mark-capture-recapture data have been collected on the Seychelles warblers on Cousin  
158 Island (4°19'53.5" S 55°39'43.2" E). From 1997, >96% of the population has been caught and given  
159 unique identifiers using colored bands and BTO-numbered metal rings (Richardson *et al.*, 2001). High  
160 annual resighting probabilities (98%) and no (<0.1%) inter-island dispersal enable accurate individual-  
161 level longitudinal measures of life-history traits (Komdeur *et al.*, 2004; Brouwer *et al.*, 2006).

162

163 The insectivorous Seychelles warbler forms long-term pair bonds and has a mean post-fledgling  
164 lifespan of 5.5 years and a maximum observed lifespan of 19 years (Hammers & Brouwer, 2017; Raj  
165 Pant *et al.*, 2020). Each of the ca. 110 territories on Cousin contains one dominant breeding pair.  
166 Dominant breeders are territorial, foraging most of their lives exclusively on their respective territories  
167 (Komdeur, 1991; Richardson *et al.*, 2007). Cooperative breeding can occur: around half of the  
168 territories including 1–5 sexually mature subordinates, some of which (20% of males and 42% of  
169 females) act as helpers, providing alloparental care to the dominant breeders' offspring (Richardson *et*  
170 *al.*, 2003; Hammers *et al.*, 2019). Due to resource competition, helpers are more present in higher-  
171 quality territories and can be maladaptive to breeders in lower-quality territories (Komdeur, 1998).

172

173 The main Seychelles warbler breeding season spans from June to October, and the minor breeding  
174 season from December to March (Komdeur & Daan, 2005). Our analyses focused on main breeding

175 seasons as data on breeding statuses are limited for minor breeding seasons and, although 30% of pairs  
176 breed (90% in main), breeding season type (main/minor) does not affect divorce in our study  
177 population (Speelman *et al.*, 2024). Most (87%) clutches contain a single egg but can consist of up to  
178 three (Richardson *et al.*, 2001). Additional eggs are often laid by co-breeding subordinates (Richardson  
179 *et al.*, 2003; Komdeur *et al.*, 2004). Insect abundance in a given month is predicted by rainfall two  
180 months prior (Komdeur, 1996a), likely cueing the onset of breeding to optimize food availability for  
181 nestlings. Although socially monogamous, there is a high rate of extra-pair paternity (EPP), with 44%  
182 of offspring sired by males other than the social partner (Richardson *et al.*, 2001; Hadfield *et al.*, 2006).  
183 Lastly, parents often provide up to 3 months of post-fledgling care to their offspring (Komdeur, 1991).

184

## 185 **2.2 Data collection**

186 We analyzed data from 1997 to 2015 as social pairs have been monitored intensively since 1997, and  
187 rainfall measurements were only available up to 2015. The years 1999 to 2001 were excluded due to  
188 limited fieldwork impacting the quality of partnership data required to classify divorces. During the  
189 main breeding seasons, all territories were monitored to determine the residency of ringed birds.  
190 Observations of foraging, singing, and non-aggressive and aggressive social interactions were used to  
191 assign territory boundaries and group membership (Bebbington *et al.*, 2017). The pair-bonded male  
192 and female in a territory, determined based on their courtship and nesting behaviors, were defined as  
193 the dominant birds (Richardson *et al.*, 2002). Breeding activity was monitored by following the  
194 dominant female for at least 15 mins every 1–2 weeks (Richardson *et al.*, 2007). We identified the  
195 number of helpers, which influences reproductive success (Hammers *et al.*, 2021), from nest watches  
196 of at least 60 minutes during the incubation and provisioning stages (van Boheemen *et al.*, 2019). In  
197 case of a failed breeding attempt before incubation or provisioning, subordinates were defined as non-  
198 helpers. The ages of unringed birds, which are usually caught before one year of age, were estimated  
199 using lay, hatch, or fledge dates and/or eye color (Komdeur, 1991). DNA was extracted from caught



200 individuals using brachial venipuncture blood samples (Richardson *et al.*, 2001). Up to 30  
201 microsatellite markers were genotyped to determine the relatedness between the dominant breeding  
202 pair and the parentage of the offspring (see supplementary material section ‘Pairwise relatedness’).  
203 Territory quality was measured using an index of insect availability, territory size, and foliage cover  
204 (see supplementary materials section ‘Territory quality’).

205

### 206 **2.2.1 Rainfall measurements**

207 As rainfall data were not available from Cousin, we obtained mean monthly rainfall measurements  
208 from a weather station on Praslin (4°18' 60.0" S 55°43'59.9" E), a neighboring island ca. 1.5 km  
209 northeast of Cousin (Seychelles Meteorological Authority, 2016). Mean monthly and annual rainfall  
210 varied greatly during the study period (monthly range: 0.8–716 mm; annual: 1349–3410 mm).

211

### 212 **2.3 Divorce classification**

213 Partnerships were classified as divorced when there was a change in the identity of dominant breeders  
214 across breeding seasons while both previously pair-bonded individuals were still alive. As breeding  
215 statuses were defined at the end of breeding seasons, divorce can occur within or between seasons.  
216 Temporary divorces, where pairs separate but reform after a breeding season, are rare but can occur  
217 (22 recorded cases: Speelman *et al.*, 2024). As we were solely interested in comparing the years when  
218 partnerships did or did not divorce, we excluded the years when partnerships terminated due to partner  
219 deaths or translocations undertaken for conservation (Richardson *et al.*, 2006; Wright *et al.*, 2014).  
220 Our dataset included 416 males and 392 females in 1321 partnerships, 84 (6.4%) of which divorced.

221

### 222 **2.4 Statistical analyses**

223 All statistical analyses were performed in R 4.2.2 (R Core Team, 2022). Figures were created using  
224 *ggplot2* 3.4.1 (Wickham, 2016), and generalized linear mixed models (GLMMs) were run in *lme4* 1.1-  
225 31 (Bates *et al.*, 2015). The over or underdispersion of models and residual spatio-temporal

226 autocorrelations were checked (none were found) using *DHARMA* 0.4.6 (Hartig, 2022). Collinearity  
227 was determined using *car* 3.1-1 (Fox & Weisberg, 2019), and all variance inflation factors (VIF) were  
228 <3.0. Model predictions for visualization were produced using *AICcmodavg* 2.3-1 (Mazerolle, 2020)  
229 and *ggeffects* 1.1-5 (Lüdecke, 2018). To facilitate model convergence, all explanatory variables were  
230 mean-centered and divided by 1 standard deviation using the *scale* function in R. Unless stated  
231 otherwise, all estimates are given  $\pm$  SE and the term ‘significant’ refers to statistical significance.

232

#### 233 **2.4.1 Climate window analysis**

234 Following Bailey & van de Pol (2016) (see supplementary materials section ‘Climate window  
235 analysis’), we used *climwin* 1.2.3 to determine which temporal windows of rainfall best predicted  
236 divorce, reproductive success, and food availability. Previously, total rainfall from June to August was  
237 used to study the life-history effects of rain in the Seychelles warbler (Borger *et al.*, 2023). However,  
238 we performed an analysis of all possible temporal windows within 12 months before the end of the  
239 breeding season (28th of September), as we assumed that divorce is not an instantaneous decision but  
240 rather one that follows a long-term decision-making process influenced by multiple factors.

241

242 Firstly, we tested which temporal window of rainfall best predicted the probability of divorce (Y/N).  
243 Then, we tested which temporal windows of rainfall predicted measurements of reproductive success  
244 at four stages of reproduction: 1) The probability of attempting to breed - when a dominant breeding  
245 pair initiated nest building (Y/N); 2) The probability of producing a clutch - when the nest of a  
246 dominant breeding pair contained an egg (Y/N); 3) The probability of producing a fledgling - when an  
247 offspring fledged from the nest of a dominant breeding pair (Y/N); 4) The number of fledglings  
248 genetically related to the dominant female that survived until at least three months old (post-fledgling  
249 care period) - classified as a continuous response variable (from now on called: ‘genetic fledglings’).

250

251 As all measurements of reproductive success could include offspring resulting from EPP, we assumed  
252 that male social partners were unaware of cuckolding and cared for offspring sired by other males as  
253 if they were their own. Although a minority (11% of offspring; Sparks *et al.*, 2022), reproductive  
254 success measurements 1–3 could also include co-breeders' offspring. Therefore, we included the  
255 number of genetic fledglings in our analyses to exclude offspring assigned to co-breeding females.

256

257 Lastly, we tested which months of rainfall predicted territory quality (territory-level measure) and  
258 insect abundance (population-level measure; the mean number of insects found per unit leaf area across  
259 all monthly surveys). By comparing the temporal windows of rainfall predicting divorce against  
260 possible drivers of divorce, we aimed to examine potential patterns explaining the causality of divorce.

261

#### 262 **2.4.2 Population-level divorce rate**

263 We used a quasi-binomial generalized linear model (GLM) with a logit link function to model the  
264 annual population divorce rate as a function of rainfall and rainfall<sup>2</sup>. The measurement of rainfall  
265 included in the model was the total rainfall from the months that predicted divorce determined via the  
266 climate window analysis.

267

#### 268 **2.4.3 Partnership-level probability of divorce**

269 Using a binomial GLMM with a logit link function, we modeled the probability of divorce as a function  
270 of rainfall, rainfall<sup>2</sup>, the number of offspring (genetic fledglings), the number of helpers, partnership  
271 length (in years), pairwise relatedness, male age (in years), male age<sup>2</sup>, female age, and female age<sup>2</sup>  
272 (Komdeur, 1994; 1996a; 1996b; Richardson *et al.*, 2003; van Boheemen *et al.*, 2019; Hammers *et al.*,  
273 2019). All fixed effects were continuous variables. Next, we compared the effects of reproduction at  
274 four different stages - breeding attempted (Y/N), clutch produced (Y/N), fledgling produced (Y/N),  
275 and genetic fledglings produced (Y/N) - on divorce by including them in four separate models. We  
276 also tested our assumption that EPP did not affect divorce by including male and female EPP (Y/N) –

277 when the dominant male or female was assigned parentage of offspring and the opposite-sex parent  
278 assigned was not their social partner – in our model. Lastly, we included co-breeder presence (Y/N)  
279 in our model to separate helper and co-breeder effects on divorce. Population density effects were not  
280 tested as it does not affect divorce in our population (Speelman *et al.*, 2024).

281

### 282 **2.4.3 Partnership-level probability of reproductive success**

283 To explore the potential causal links between rainfall, reproductive success, and divorce, we examined  
284 whether rainfall during the months that best predicted divorce also influenced reproductive success.  
285 We used binomial GLMMs with logit link functions to model the probability of attempting to breed,  
286 producing a clutch, and producing a fledgling as functions of rainfall, rainfall<sup>2</sup>, the number of helpers,  
287 partnership length, pairwise relatedness, male age, male age<sup>2</sup>, female age, and female age<sup>2</sup>. Next, we  
288 used a Poisson GLMM with a log link function to model the effect of the same fixed effects on the  
289 number of genetic fledglings.

290

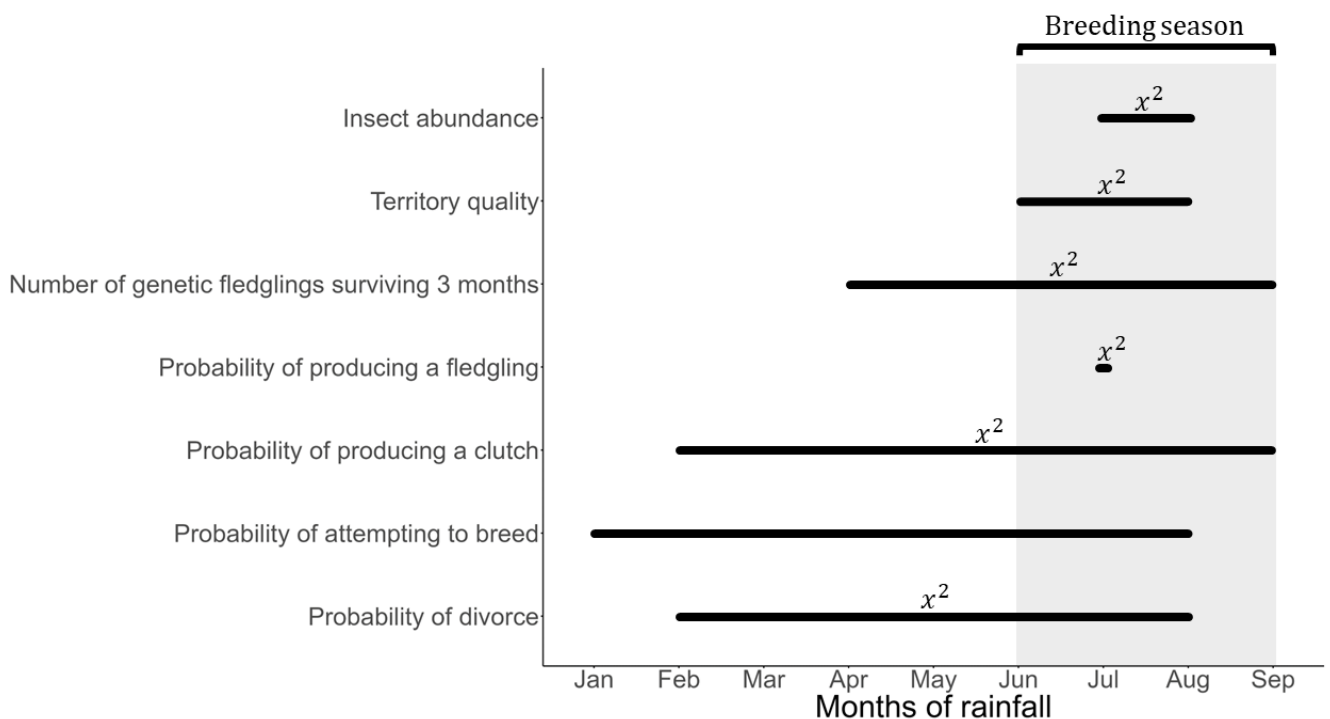
291 In all models, we tested whether partnership quality buffered the effects of rainfall by including  
292 interactions between partnership length and rainfall (all models) and reproductive success and rainfall  
293 (divorce models). Starting from a full model, we removed non-significant quadratic terms and  
294 interactions in order of least significance to interpret first-order effects. All models included the  
295 random effects: male ID, female ID, territory ID, and field period ID to control for birds sequentially  
296 performing worse or better than others and variable quality across territories and field periods (years).

297

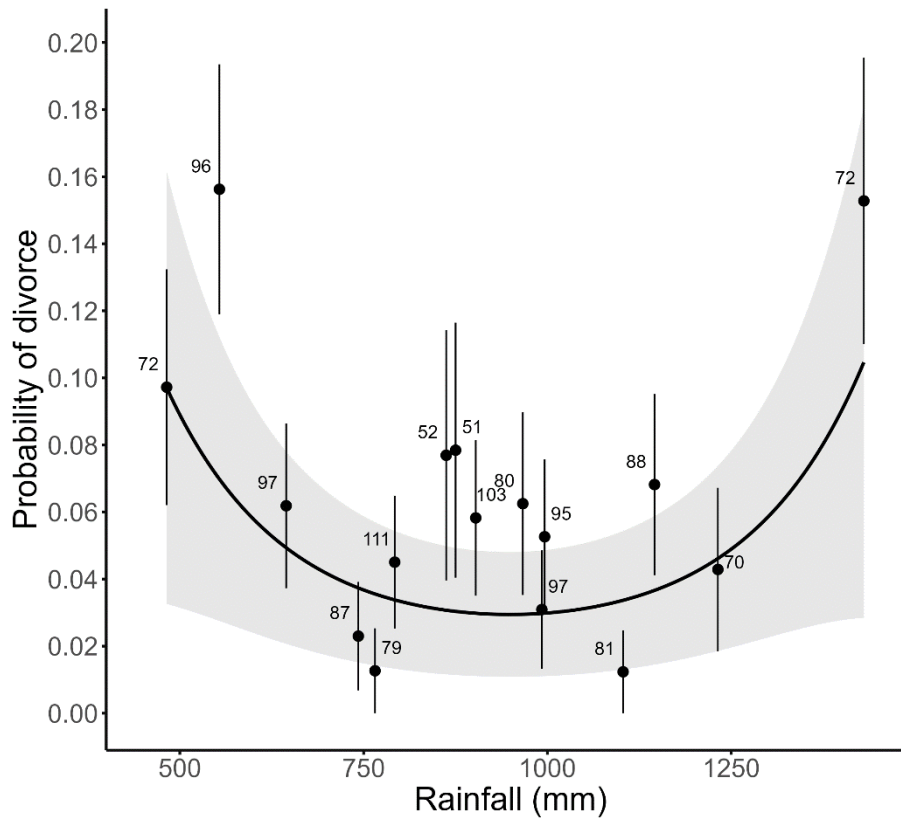
## 298 **3 Results**

### 299 **3.1 Effect of rainfall on divorce**

300 The mean annual divorce rate of Seychelles warbler partnerships during the study period was  $6.6 \pm$   
 301  $1.1\%$  and showed considerable inter-annual variability ( $1.2\text{--}15.6\%$ ; Figure S1a). Climate window  
 302 analyses revealed that the quadratic effect of total rainfall from February to August best predicted the  
 303 probability of divorce (Figure 1). At the population level, total rainfall from February to August had a  
 304 significant quadratic effect on the annual divorce rate (GLM, estimate =  $0.335 \pm 0.091$ ,  $p$ -value =  
 305  $0.003$ ), which increased in years with low and high rainfall (Figure S1b). Rainfall effects explained  
 306  $46.7\%$  of the annual divorce rate's variance ( $r^2 = 0.467$ ). At the partnership level, the quadratic effect  
 307 of total rainfall from February to August significantly affected the probability of divorce (Table 1;  
 308 Figure 2). Notably, the quadratic relationship between rainfall and the probability of divorce was driven  
 309 by extremely heavy rainfall in 1997, as excluding 1997 from the analysis revealed a significant  
 310 negative linear relationship between rainfall and divorce (Tables S1 & S2; Figure S2).



311 **Figure 1.** Temporal windows of rainfall that best predict seven response variables in the Seychelles  
 312 warbler on Cousin Island ( $n = 1321$  partnerships/15 years for insect abundance and territory quality)  
 313 as predicted by climate window analyses. Relationships between rainfall and the response variables  
 314 were quadratic if indicated by  $x^2$  and linear if not. The shaded area represents the main breeding season.



315

316 **Figure 2.** The effect of total rainfall from February to August on the probability of divorce in the  
 317 Seychelles warbler on Cousin Island ( $n = 1321$  partnerships) as predicted by a binomial generalized  
 318 linear mixed model. The solid line represents the predicted probability of divorce and the shading  
 319 indicates the 95% confidence intervals. Dots represent the mean observed divorce rate  $\pm$  SE, and labels  
 320 indicate the total number of partnerships observed in a given year.

321

### 322 **3.2 Effects of reproductive success and other partnership qualities on divorce**

323 Although we found a trend for reproductively successful partnerships to have lower divorce rates  
 324 (Figure S3; Tables S3-S6), we found no significant correlations between the probability of divorce and  
 325 measures of reproductive success (Figure S4; Tables S3-S6). However, the probability of divorce was  
 326 significantly negatively correlated with partnership length (Table 1), with shorter partnerships having  
 327 the highest probability of divorce. Notably, the correlation between rainfall and partnership length was  
 328 non-significant after excluding 1997 from the analyses (Table S1 & S2). The mean partnership length  
 329 in 1997 ( $0.8 \pm 0.1$ ) was considerably shorter than that of the full study period ( $2.2 \pm 0.07$ ). We also

330 found a significant interaction between partnership length and rainfall (Table S7), with heavy rainfall  
 331 increasing the probability of divorce in shorter but not longer-lasting partnerships (Figure S8).  
 332 However, this interaction was strongly influenced by outliers and subsequently removed from the final  
 333 model (see supplementary material section ‘Interaction between rainfall and partnership length’). EPP  
 334 and co-breeder presence were not associated with divorce (Table S9).

335

336 **Table 1.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 337 with rainfall, the length of the partnership, the number of offspring, the relatedness of the breeding  
 338 pair, the number of helpers, male age, and female age.  $n = 1321$  partnerships were analyzed using a  
 339 binomial generalized linear mixed model. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.480	0.330	-4.126 to -2.833	<b>&lt;0.001</b>
Rainfall	-0.143	0.109	-0.357 to 0.072	0.192
Rainfall <sup>2</sup>	0.323	0.085	0.157 to 0.490	<b>&lt;0.001</b>
Partnership length	-0.492	0.216	-0.915 to -0.070	<b>0.022</b>
Number of offspring	-0.108	0.141	-0.383 to 0.168	0.443
Pairwise relatedness	0.145	0.128	-0.106 to 0.395	0.257
Number of helpers	-0.207	0.156	-0.513 to 0.099	0.185
Male age	0.074	0.158	-0.236 to 0.383	0.641
Female age	0.195	0.144	-0.086 to 0.476	0.174
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	<0.001	416		
Female ID	0.251	392		
Field period ID	0.015	16		
Territory ID	0.358	158		

340

### 341 **3.3 Effect of rainfall on reproductive success**

#### 342 **3.3.1 Breeding attempted**

343 During the study period, 91% of partnerships attempted to breed. The probability of attempting to  
344 breed was best predicted by the linear increase in total rainfall from January to August (Figure 1). The  
345 probability of attempting to breed was also significantly positively correlated with the months of  
346 rainfall best predicting divorce (February to August; Table S13; Figure 3a).

347

#### 348 **3.3.2 Clutch produced**

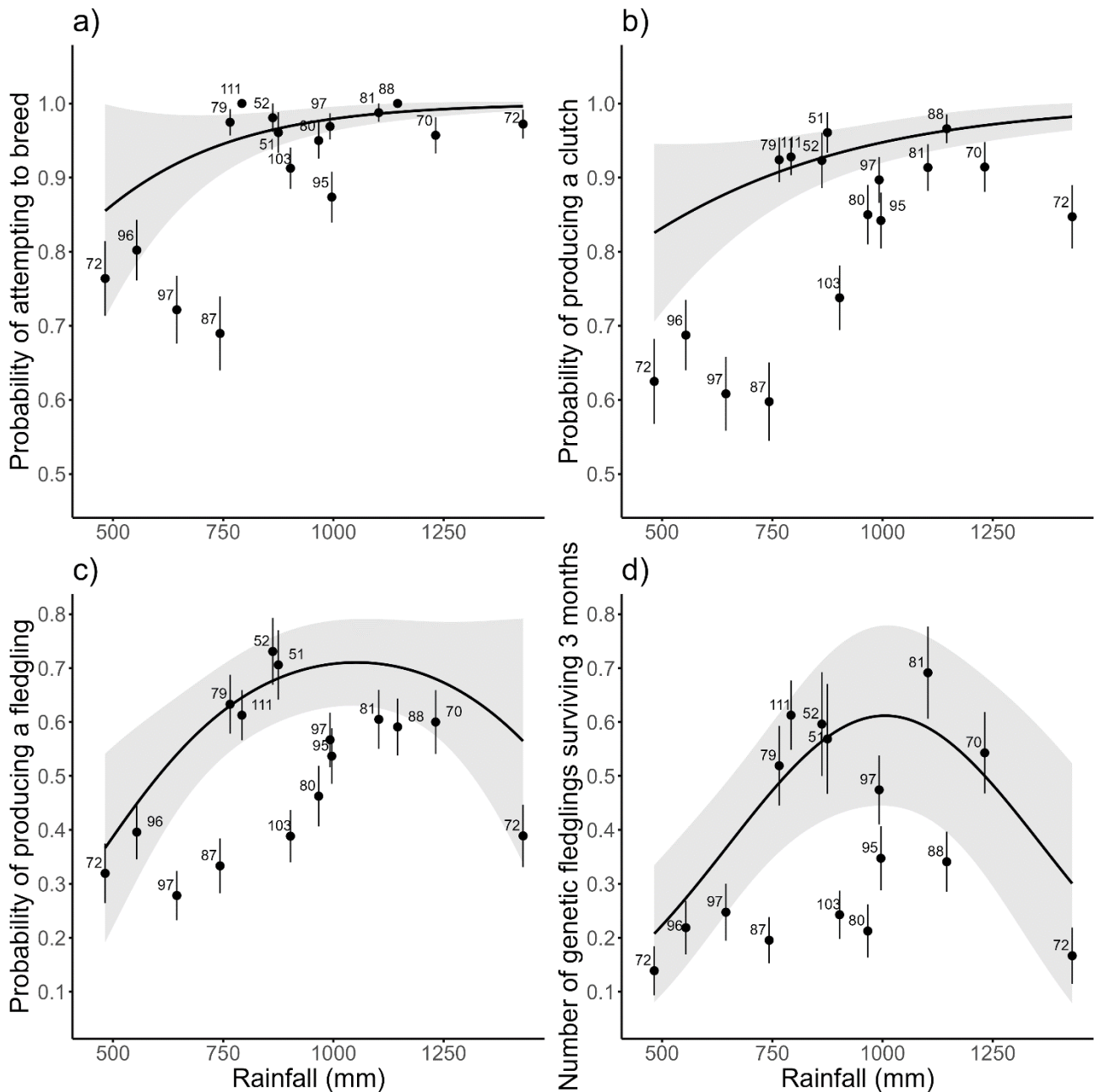
349 Overall, 83% of partnerships produced a clutch. The probability of producing a clutch was best  
350 predicted by the quadratic effect of rainfall from February to September (Figure 1), decreasing in years  
351 with low and high rainfall. Although the climate window analysis predicted a quadratic relationship,  
352 we found a significant positive linear correlation between total rainfall from February to August and  
353 the probability of producing a clutch (Table S13; Figure 3b). The quadratic effect of rainfall was  
354 marginal (Table S14).

355

#### 356 **3.3.3 Fledgling produced**

357 Overall, 50% of partnerships produced a fledgling. The probability of producing a fledgling was best  
358 predicted by the quadratic effect of rainfall in July, the peak of egg-laying (Figure 1), where both low  
359 and high amounts of rain decreased fledgling success. We also found that total rainfall from February  
360 to August had a significant quadratic effect on the probability of producing a fledgling (Table S13).  
361 Again, intermediate levels of rainfall were associated with the highest probabilities of fledgling success  
362 (Figure 3c).





363

364 **Figure 3.** The effect of total rainfall from February to August on the probability of Seychelles warbler  
 365 partnerships ( $n = 1321$ ): a) attempting to breed; b) producing a clutch; c) producing a fledgling; d) the  
 366 number of genetic fledglings surviving until three months old, as predicted by binomial (a, b, c) and  
 367 Poisson (d) generalized linear mixed models. The solid line represents the predicted probability of  
 368 divorce, and the shading indicates the 95% confidence intervals. Dots represent the mean observed  
 369 divorce rate  $\pm$  SE, and labels indicate the total number of partnerships observed in a given year.

370

### 371 **3.3.4 Genetic fledglings**

372 During the study period, 32% of partnerships produced a genetic fledgling (mean number of genetic  
373 fledglings surviving =  $0.40 \pm 0.02$ ). The number of genetic fledglings surviving was best predicted by  
374 the quadratic effect of rainfall from April to October (Figure 1). Here, low and high amounts of rain  
375 decreased genetic offspring survival post-fledgling care. Also, the number of genetic fledglings  
376 surviving was correlated with the quadratic effect of the total rainfall from February to August (Table  
377 S13). Again, the highest numbers of genetic fledglings surviving were associated with intermediate  
378 levels of rainfall (Figure 3d).

379

### 380 **3.4 Effect of rainfall on food availability**

381 A quadratic effect of total rainfall from June to August best predicted territory quality (Figure 1).  
382 Although quadratic, this relationship was skewed to high rainfall correlating strongly with high  
383 territory quality, while low and intermediate rainfall were associated with lower territory quality  
384 (Figure S6a). An increase in insect abundance was best predicted by the increase in total rainfall from  
385 July to August (Figure 1; Figure S6b).

386

## 387 **4 Discussion**

### 388 **4.1 Association between rainfall and divorce**

389 As predicted (P1), rainfall had a quadratic effect on divorce in the Seychelles warbler, where low and  
390 high amounts of rain significantly increased the population-level annual divorce rates and partnership-  
391 level divorce probabilities. Extremely heavy rainfall in 1997 (a super 'El Niño' event) drove the  
392 association between high rainfall and divorce; excluding 1997 from the analyses left a negative  
393 relationship between rainfall and divorce. However, we consider 1997 to be biologically valid, as it  
394 shows the effects of the heavy rainfall events predicted to become more prevalent because of future  
395 climate change (Pezza & Simmonds, 2008; Changnon, 2009; NOAA, 2022). Future investigations

396 incorporating more extreme rainfall years would allow us to estimate the robustness of the quadratic  
397 effect. The main Seychelles warbler breeding season spans from June to October, and total rainfall  
398 from February to August best predicted divorce. Thus, if divorce is a decision informed by the costs  
399 and benefits of staying with a partner, it is likely reinforced by various drivers linked to rainfall  
400 between February and August.

401

402 Compared to the high divorce rates of some migratory birds, including the congeneric great reed  
403 warbler (*Acrocephalus arundinaceus*) (85%: Bench & Hasselquist, 1991), the Seychelles warbler had  
404 a relatively low mean annual divorce rate (6.4%) similar to other birds with high site fidelity (3.7% in  
405 black-browed albatrosses: Ventura *et al.*, 2021), fitting with the prediction that birds with stable nesting  
406 sites are less likely to divorce (Choudhury, 1995). Nevertheless, inter-annual divorce rates varied  
407 considerably and were significantly associated with rainfall. As rainfall is associated with food  
408 abundance (discussed below), our study is one of few to provide empirical evidence supporting the  
409 habitat-mediated hypothesis of divorce (Ventura *et al.*, 2021).

410

#### 411 **4.2 Association between rainfall and reproductive success**

412 As predicted (P2), rainfall significantly influenced reproductive success. Borger *et al.* (2023)  
413 discovered that total rainfall from June to August had a quadratic effect on the number of genetic  
414 Seychelles warbler fledglings produced. We investigated this further by examining rain effects on all  
415 stages of reproduction. Total rainfall from January to August and February to September best predicted  
416 the probability of attempting to breed and producing a clutch, respectively. These reproductive  
417 measures were also significantly positively correlated with the temporal window of rainfall best  
418 predicting divorce. These large temporal windows support studies showing rain impacts birds'  
419 reproductive success by affecting the health of birds outside of the breeding season (Studds & Marra,  
420 2007). Rainfall can impact individual condition and reproductive success by influencing food

421 abundance (often insects), which explains why rainfall cues breeding for many birds (Lloyd, 1999;  
422 Cavalcanti *et al.*, 2016; França *et al.*, 2020).

423

424 On Cousin, the increase in total rainfall from June to August and July to August was associated with  
425 increased territory quality and population-wide insect abundance, respectively. As most insects lay  
426 their eggs in water, drought significantly limits their development (Price, 1997; Chen *et al.*, 2019),  
427 decreasing food availability for the warblers. As mean food availability at the end of the breeding  
428 season was best predicted by rainfall around the middle of the breeding season, our results support the  
429 two-month temporal window previously found by Komdeur (1996a). Thus, the Seychelles warbler  
430 likely uses rainfall to cue breeding to ensure adequate food availability for offspring. Consequently,  
431 by limiting the ability to invest in offspring, low rainfall decreases the probability of attempting to  
432 breed and producing a clutch.

433

434 The probability of producing a fledgling was predicted by rainfall in July, the month of peak egg-  
435 laying, which is consistent with studies that found that the probability of fledging in birds correlated  
436 with rainfall during the hatchling period (Monadjem & Bamford, 2009; Schöll & Hille, 2020). Next,  
437 the number of genetic fledglings produced was best predicted by rainfall from April to September.  
438 Here, rainfall can directly affect fledgling survival or do so indirectly by impacting parental care during  
439 the months of post-fledgling care. Total rainfall from February to August had a significant quadratic  
440 effect on both measures of fledgling success, where low and high amounts of rain decreased the  
441 probability of fledgling survival. Alongside the aforementioned effects of low rainfall on food  
442 availability, heavy rainfall, and the often accompanying strong winds, can be detrimental as they can  
443 destroy nests and make maintaining optimal body temperatures difficult for birds (Kennedy, 1970;  
444 Jones & Barnett, 1971; Wilson *et al.*, 2004). As nestlings often lack fully developed feathers, hindering  
445 their ability to maintain body temperature, it can be detrimental to their survival if they get wet

446 (Mertens, 1977; Newton, 1998). Similarly, heavy rainfall can increase the parental investment required  
447 to maintain optimal nest temperatures (Heenan & Seymour, 2012). If required parental investments  
448 increase during harsh weather conditions while their foraging ability is limited, they may face a trade-  
449 off between provisioning and their health (Radford *et al.*, 2001), impacting the survival of their  
450 offspring (Öberg *et al.*, 2015).

451

### 452 **4.3 Association between reproductive success and divorce**

453 The temporal window of rainfall that predicted divorce overlapped with the temporal windows of  
454 rainfall predicting measures of reproductive success. All measurements of reproductive success were  
455 also significantly correlated with total rainfall from the months that best predicted divorce, and there  
456 was a trend for higher mean divorce rates in partnerships with lower reproductive success. These  
457 results reveal a plausible pathway whereby rainfall influences reproductive success and low  
458 reproductive success is interpreted by individuals as a marker of poor partnership quality, resulting in  
459 divorce. Low reproductive success impacting divorce is in line with findings of previous studies  
460 (Culina *et al.*, 2015; Mercier *et al.*, 2021; Ventura *et al.*, 2021; Pelletier & Guillemette, 2022),  
461 including in our study population (Speelman *et al.*, 2024). However, when accounting for rainfall  
462 effects in our models, the direct effects of reproductive success on the probability of divorce were non-  
463 significant. Thus, reproductive success may not influence divorce in the Seychelles warbler as  
464 predicted (P3), and rainfall may influence divorce through alternative pathways.

465

466 Physiological stress may influence divorce in the Seychelles warbler. Harsh environmental conditions  
467 and food scarcity can increase the concentration of stress hormones in birds (Kitaysky *et al.*, 2010),  
468 which are positively associated with an individual's level of dissatisfaction with their social partner  
469 (Griffith *et al.*, 2011). Although the role of stress in divorce is currently unknown for the Seychelles  
470 warbler, research shows that lower territory quality correlates with higher levels of oxidative stress

471 because of increased foraging effort, especially during the early stages of reproduction (Komdeur  
472 1991; 1996b; van de Crommenacker *et al.*, 2011). Thus, rainfall and its effects on food availability and  
473 parental investments could increase physiological stress in the Seychelles warbler. Individuals may  
474 associate their heightened physiological stress with their choice of partner, causing individuals in  
475 resource-poor seasons to terminate partnerships regardless of reproductive output, signifying that  
476 stress could be the link between rainfall and divorce. Studies analyzing relationships between stress  
477 markers, such as glucocorticoids (Sapolsky *et al.*, 2000), rainfall (or other environmental effects), and  
478 divorce, are required to investigate this theory.

479

480 Divorce can be an adaptive strategy that improves reproductive success (Culina *et al.*, 2015). In times  
481 of climate change, behavioral plasticity may help animals minimize the negative consequences of  
482 coping with rapid environmental changes (Beever *et al.*, 2017). Our study introduces the possible  
483 consequences of climate change on partnership stability. However, further research into divorce  
484 consequences is required to determine whether rainfall-driven divorce is adaptive and can help the  
485 species overcome climatic challenges. An understanding of whether rainfall influences divorce in  
486 good- or bad-quality partnerships is currently lacking. If rainfall affects divorce by misinforming  
487 individuals about their partnership's quality, either through impacting stress or reproductive  
488 performance, divorce can occur in partnerships that may perform adequately in good conditions. Here,  
489 rainfall-driven divorce can be maladaptive, making climate change a concern to the future of this  
490 species. In the Seychelles warbler, no short or long-term reproductive costs of divorce have been  
491 detected (Speelman *et al.*, 2024). However, as this study did not test for divorce consequences in the  
492 context of environmental effects, studies disentangling divorce consequences in poor and high-quality  
493 years are required.

494

#### 495 **4.4 Non-environmental associations with divorce**

496 Older partnerships were less likely to divorce, fitting the prediction that divorce benefits are highest  
497 before individuals have gained the benefits associated with mate familiarity (Choudhury, 1995). While  
498 behavioral incompatibilities between individuals can also manifest early in partnerships and influence  
499 divorce by impacting reproductive success (Wilson *et al.*, 2022), we found no effect of partnership  
500 length on reproductive success. Consistent with studies showing that Seychelles warblers do not seem  
501 to avoid inbreeding (Eikenaar *et al.*, 2008), we also found no effect of pairwise relatedness on divorce,  
502 indicating that inbreeding avoidance or other genetic incompatibilities are unlikely drivers of divorce  
503 in our population (Hidalgo Aranzamendi *et al.*, 2016). Notably, the effect of partnership length on  
504 divorce was non-significant when excluding 1997 from the analysis. 1997 is when population  
505 monitoring intensified, and much more of the Seychelles warbler population became identity-tagged  
506 (>96% of the population; Richardson *et al.*, 2001). The mean partnership length in 1997 was  
507 considerably shorter than in other years. This may be because the limited nature of the data previous  
508 to this year meant that partnership lengths were underestimated that year, and consequently, removing  
509 it led to the loss of the significant interaction. Other earlier years (1999 to 2001) were already excluded  
510 from the analyses due to reduced partnership-level data – required to classify divorces - being collected  
511 in those years. Thus, biases in the data may drive the effect of partnership length on divorce.

512

## 513 **5 Conclusion**

514 We provide empirical evidence for an association between rainfall and divorce in a socially  
515 monogamous population, thereby contributing to a growing body of literature showing that harsh  
516 climates affect partnership stability. The prevalence of divorce in the Seychelles warbler was highest  
517 in years with low and high amounts of rainfall. We provide correlational evidence that this could result  
518 from rain impacting reproductive success, possibly by affecting food availability and parental trade-  
519 offs between investing in current versus future reproductive success. We also discuss alternative  
520 explanations involving the role of physiological stress, an important avenue for further research in this

521 and other species. Studies show that temperature influences divorce in birds, and now we find that  
522 rainfall does too. The climate can directly affect survival and indirectly influence population stability  
523 by restricting reproductive output. We do not yet understand whether rainfall-driven divorce in the  
524 Seychelles warbler is adaptive, maladaptive, or neutral. Therefore, studying the consequences of  
525 divorce in this species may highlight to what extent plasticity in breeding behavior can enable socially  
526 monogamous species to adapt to a rapidly changing world.

527

## 528 **References**

529 Bailey, L. D., & van de Pol, M. (2016). climwin: an R toolbox for climate window analysis. *PLOS*  
530 *ONE*, *11*(12), e0167980. <https://doi.org/10.1371/journal.pone.0167980>

531 Bates, D., Mächler, M., Bolker, B., & Walker, S. (2015). Fitting linear mixed-effects models using  
532 lme4. *Journal of Statistical Software*, *67*(1), 1–48. <https://doi.org/10.18637/jss.v067.i01>

533 Beever, E. A., Hall, L. E., Varner, J., Loosen, A. E., Dunham, J. B., Gahl, M. K., Smith, F. A., &  
534 Lawler, J. J. (2017). Behavioral flexibility as a mechanism for coping with climate change. *Frontiers*  
535 *in Ecology and the Environment*, *15*(6), 299–308. <https://doi.org/10.1002/fee.1502>

536 Bensch, S., & Hasselquist, D. (1991). Territory infidelity in the polygynous great reed warbler  
537 *Acrocephalus arundinaceus*: the effect of variation in territory attractiveness. *Journal of Animal*  
538 *Ecology*, *60*(3), 857–871. <https://doi.org/10.2307/5418>

539 Black, J. M. (1996). *Partnerships in Birds: The Study of Monogamy: The Study of Monogamy*. Oxford,  
540 UK: Oxford University Press.

541 Blondel, J., Perret, P., & Galan, M. J. (2000). High divorce rates in Corsican blue tits: how to choose  
542 a better option in a harsh environment. *Oikos*, *89*(3), 451–460. [https://doi.org/10.1034/j.1600-](https://doi.org/10.1034/j.1600-0706.2000.890304.x)  
543 [0706.2000.890304.x](https://doi.org/10.1034/j.1600-0706.2000.890304.x)



544 Borger, M. J., Richardson, D. S., Dugdale, H., Burke, T., & Komdeur, J. (2023). Testing the  
545 environmental buffering hypothesis of cooperative breeding in the Seychelles warbler. *Acta*  
546 *Ethologica*, 1–4. <https://doi.org/10.1007/s10211-022-00408-y>

547 Botero, C. A., & Rubenstein, D. R. (2012). Fluctuating environments, sexual selection and the  
548 evolution of flexible mate choice in birds. *PLOS ONE*, 7(2), e32311.  
549 <https://doi.org/10.1371/journal.pone.0032311>

550 Brouwer, L., Richardson, D. S., Eikenaar, C., & Komdeur, J. (2006). The role of group size and  
551 environmental factors on survival in a cooperatively breeding tropical passerine. *Journal of Animal*  
552 *Ecology*, 75(6), 1321–1329. <https://doi.org/10.1111/j.1365-2656.2006.01155.x>

553 Cavalcanti, L. M. P., Paiva, L. V. de, & França, L. F. (2016). Effects of rainfall on bird reproduction  
554 in a semi-arid Neotropical region. *Zoologia (Curitiba)*, 33, e20160018. [https://doi.org/10.1590/S1984-](https://doi.org/10.1590/S1984-4689zool-20160018)  
555 [4689zool-20160018](https://doi.org/10.1590/S1984-4689zool-20160018)

556 Changnon, S. A. (2009). Temporal changes in extremely damaging storms. *Physical Geography*,  
557 30(1), 17–26. <https://doi.org/10.2747/0272-3646.30.1.17>

558 Chen, C., Harvey, J. A., Biere, A., & Gols, R. (2019). Rain downpours affect survival and development  
559 of insect herbivores: the specter of climate change? *Ecology*, 100(11), e02819.  
560 <https://doi.org/10.1002/ecy.2819>

561 Choudhury, S. (1995). Divorce in birds: a review of the hypotheses. *Animal Behaviour*, 50(2), 413–  
562 429. <https://doi.org/10.1006/anbe.1995.0256>

563 Choudhury, S., & Black, J. M. (1994). Barnacle geese preferentially pair with familiar associates from  
564 early life. *Animal Behaviour*, 48(1), 81–88. <https://doi.org/10.1006/anbe.1994.1213>

565 Culina, A., Firth, J. A., & Hinde, C. A. (2020). Familiarity breeds success: pairs that meet earlier  
566 experience increased breeding performance in a wild bird population. *Proceedings of the Royal Society*  
567 *B: Biological Sciences*, 287(1941), 20201554. <https://doi.org/10.1098/rspb.2020.1554>

568 Culina, A., Radersma, R., & Sheldon, B. C. (2015). Trading up: the fitness consequences of divorce  
569 in monogamous birds. *Biological Reviews*, 90(4), 1015–1034. <https://doi.org/10.1111/brv.12143>

570 Dhondt, A. A., & Adriaensen, F. (1994). Causes and effects of divorce in the blue tit *Parus caeruleus*.  
571 *Journal of Animal Ecology*, 63(4), 979–987. <https://doi.org/10.2307/5274>

572 Eikenaar, C., Komdeur, J., & Richardson, D. S. (2008). Natal dispersal patterns are not associated with  
573 inbreeding avoidance in the Seychelles warbler. *Journal of Evolutionary Biology*, 21(4), 1106–1116.  
574 <https://doi.org/10.1111/j.1420-9101.2008.01528.x>

575 Fox, J., & Weisberg, S. (2019). *An R companion to applied regression*. Thousand Oaks CA: Sage.  
576 <https://socialsciences.mcmaster.ca/jfox/Books/Companion/>

577 França, L. F., Figueiredo-Paixão, V. H., Duarte-Silva, T. A., & Santos, K. B. dos. (2020). The effects  
578 of rainfall and arthropod abundance on breeding season of insectivorous birds, in a semi-arid  
579 neotropical environment. *Zoologia (Curitiba)*, 37, e37716.  
580 <https://doi.org/10.3897/zoologia.37.e37716>

581 Gilsenan, C., Valcu, M., & Kempenaers, B. (2017). Difference in arrival date at the breeding site  
582 between former pair members predicts divorce in blue tits. *Animal Behaviour*, 133, 57–72.  
583 <https://doi.org/10.1016/j.anbehav.2017.09.004>

584 Griffith, S. C., Pryke, S. R., & Buttemer, W. A. (2011). Constrained mate choice in social monogamy  
585 and the stress of having an unattractive partner. *Proceedings of the Royal Society B: Biological*  
586 *Sciences*, 278(1719), 2798–2805. <https://doi.org/10.1098/rspb.2010.2672>

587 Hadfield, J. D., Richardson, D. S., & Burke, T. (2006). Towards unbiased parentage assignment:  
588 combining genetic, behavioural and spatial data in a Bayesian framework. *Molecular Ecology*, *15*(12),  
589 3715–3730. <https://doi.org/10.1111/j.1365-294X.2006.03050.x>

590 Hammers, M., & Brouwer, L. (2017). Rescue behaviour in a social bird: removal of sticky ‘bird-  
591 catcher tree’ seeds by group members. *Behaviour*, *154*(4), 403–411.  
592 <https://doi.org/10.1163/1568539x-00003428>

593 Hammers, M., Kingma, S. A., Spurgin, L. G., Bebbington, K., Dugdale, H. L., Burke, T., Komdeur,  
594 J., & Richardson, D. S. (2019). Breeders that receive help age more slowly in a cooperatively breeding  
595 bird. *Nature Communications*, *10*(1), Article 1. <https://doi.org/10.1038/s41467-019-09229-3>

596 Hammers, M., Kingma, S. A., van Boheemen, L. A., Sparks, A. M., Burke, T., Dugdale, H. L., ... &  
597 Komdeur, J. (2021). Helpers compensate for age-related declines in parental care and offspring  
598 survival in a cooperatively breeding bird. *Evolution Letters*, *5*(2), 143-153.  
599 <https://doi.org/10.1002/evl3.213>

600 Hartig, F. (2022). DHARMA: residual diagnostics for hierarchical (multi-level / mixed) regression  
601 models. R package version 0.4.6. <http://florianhartig.github.io/DHARMA/>

602 Heenan, C. B., & Seymour, R. S. (2012). The effect of wind on the rate of heat loss from avian cup-  
603 shaped nests. *PLOS ONE*, *7*(2), e32252. <https://doi.org/10.1371/journal.pone.0032252>

604 Heg, D., Bruinzeel, L. W., & Ens, B. J. (2003). Fitness consequences of divorce in the oystercatcher,  
605 *Haematopus ostralegus*. *Animal Behaviour*, *66*(1), 175–184. <https://doi.org/10.1006/anbe.2003.2188>

606 Hidalgo Aranzamendi, N., Hall, M. L., Kingma, S. A., Sunnucks, P., & Peters, A. (2016). Incest  
607 avoidance, extrapair paternity, and territory quality drive divorce in a year-round territorial bird.  
608 *Behavioral Ecology*, *27*(6), 1808–1819. <https://doi.org/10.1093/beheco/arw101>

609 Husak, J. F., & Moore, I. T. (2008). Stress hormones and mate choice. *Trends in Ecology & Evolution*,  
610 23(10), 532–534. <https://doi.org/10.1016/j.tree.2008.06.007>

611 Jeschke, J. M., & Kokko, H. (2008). Mortality and other determinants of bird divorce rate. *Behavioral*  
612 *Ecology and Sociobiology*, 63(1), 1–9. <https://doi.org/10.1007/s00265-008-0646-9>

613 Jeschke, J. M., Wanless, S., Harris, M. P., & Kokko, H. (2007). How partnerships end in guillemots  
614 *Uria aalge*: chance events, adaptive change, or forced divorce? *Behavioral Ecology*, 18(2), 460–466.  
615 <https://doi.org/10.1093/beheco/arl109>

616 Jones, J. E., & Barnett, B. D. (1971). Body temperature of turkey poult exposed to simulated chilling  
617 rain. *Poultry Science*, 50(3), 972–974. <https://doi.org/10.3382/ps.0500972>

618 Kennedy, R. J. (1970). Direct effects of rain on birds: A review. *British Birds*. 63(10), 401-414.

619 Kitaysky, A. S., Piatt, J. F., Hatch, S. A., Kitaiskaia, E. V., Benowitz-Fredericks, Z. M., Shultz, M. T.,  
620 & Wingfield, J. C. (2010). Food availability and population processes: severity of nutritional stress  
621 during reproduction predicts survival of long-lived seabirds. *Functional Ecology*, 24(3), 625–637.  
622 <https://doi.org/10.1111/j.1365-2435.2009.01679.x>

623 Komdeur, J. (1991). Cooperative breeding in the Seychelles warbler, PhD thesis. Cambridge, UK:  
624 University of Cambridge.

625 Komdeur, J. (1994). Experimental evidence for helping and hindering by previous offspring in the  
626 cooperative-breeding Seychelles warbler *Acrocephalus sechellensis*. *Behavioral Ecology and*  
627 *Sociobiology*, 34(3), 175–186. <https://doi.org/10.1007/BF00167742>

628 Komdeur, J. (1996a). Seasonal timing of reproduction in a tropical bird, the Seychelles warbler: a field  
629 experiment using translocation. *Journal of Biological Rhythms*, 11(4), 333–346.  
630 <https://doi.org/10.1177/074873049601100407>

631 Komdeur, J. (1996b). Influence of age on reproductive performance in the Seychelles warbler.  
632 *Behavioral Ecology*, 7(4), 417–425. <https://doi.org/10.1093/beheco/7.4.417>

633 Komdeur, J. (1998). Long-term fitness benefits of egg sex modification by the Seychelles warbler.  
634 *Ecology Letters*, 1(1), 56–62. <https://doi.org/10.1046/j.1461-0248.1998.00009.x>

635 Komdeur, J., & Daan, S. (2005). Breeding in the monsoon: semi-annual reproduction in the Seychelles  
636 warbler (*Acrocephalus sechellensis*). *Journal of Ornithology*, 146(4), 305–313.  
637 <https://doi.org/10.1007/s10336-005-0008-6>

638 Komdeur, J., Richardson, D. S., & Burke, T. (2004). Experimental evidence that kin discrimination in  
639 the Seychelles warbler is based on association and not on genetic relatedness. *Proceedings of the Royal*  
640 *Society of London. Series B: Biological Sciences*, 271(1542), 963–969.  
641 <https://doi.org/10.1098/rspb.2003.2665>

642 Lack, D. L. (1968). *Ecological adaptations for breeding in birds*. London, UK: Methuen.

643 Lerch, B. A., Price, T. D., & Servedio, M. R. (2022). Better to divorce than be widowed: the role of  
644 mortality and environmental heterogeneity in the evolution of divorce. *The American Naturalist*,  
645 200(4), 518–531. <https://doi.org/10.1086/720622>

646 Lloyd, P. (1999). Rainfall as a breeding stimulus and clutch size determinant in South African arid-  
647 zone birds. *Ibis*, 141(4), 637–643. <https://doi.org/10.1111/j.1474-919X.1999.tb07371.x>

648 Lüdecke, D. (2018). ggeffects: tidy data frames of marginal effects from regression models. *Journal*  
649 *of Open Source Software*, 3(26), 772. <https://doi.org/10.21105/joss.00772>

650 Marvel, K., Cook, B. I., Bonfils, C. J. W., Durack, P. J., Smerdon, J. E., & Williams, A. P. (2019).  
651 Twentieth-century hydroclimate changes consistent with human influence. *Nature*, 569(7754).  
652 <https://doi.org/10.1038/s41586-019-1149-8>

653 Mazerolle, M. J. (2020). AICcmodavg: model selection and multimodel inference based on (Q)AIC(c).  
654 R package version 2.3-1. <https://cran.r-project.org/package=AICcmodavg>

655 Mercier, G., Yoccoz, N. G., & Descamps, S. (2021). Influence of reproductive output on divorce rates  
656 in polar seabirds. *Ecology and Evolution*, *11*(19), 12989–13000. <https://doi.org/10.1002/ece3.7775>

657 Mertens, J. A. L. (1977). Thermal conditions for successful breeding in great tits (*Parus major* L.).  
658 *Oecologia*, *28*(1), 1–29. <https://doi.org/10.1007/BF00346834>

659 Monadjem, A., & Bamford, A. J. (2009). Influence of rainfall on timing and success of reproduction  
660 in Marabou storks *Leptoptilos crumeniferus*. *Ibis*, *151*(2), 344–351. <https://doi.org/10.1111/j.1474->  
661 [919X.2009.00912.x](https://doi.org/10.1111/j.1474-919X.2009.00912.x)

662 Newton, I. (1998). *Population Limitation in Birds*. London, UK: Academic Press.

663 NOAA National Centers for Environmental Information (2022), monthly global climate report for  
664 december 2021, published online January 2022, retrieved on May 1, 2023 from  
665 <https://www.ncei.noaa.gov/access/monitoring/monthly-report/global/202200>.

666 Öberg, M., Arlt, D., Pärt, T., Laugen, A. T., Eggers, S., & Low, M. (2015). Rainfall during parental  
667 care reduces reproductive and survival components of fitness in a passerine bird. *Ecology and*  
668 *Evolution*, *5*(2), 345–356. <https://doi.org/10.1002/ece3.1345>

669 Otter, K., & Ratcliffe, L. (1996). Female initiated divorce in a monogamous songbird: abandoning  
670 mates for males of higher quality. *Proceedings of the Royal Society of London. Series B: Biological*  
671 *Sciences*, *263*(1368), 351–355. <https://doi.org/10.1098/rspb.1996.0054>

672 Pelletier, D., & Guillemette, M. (2022). Times and partners are a-changin’: relationships between  
673 declining food abundance, breeding success, and divorce in a monogamous seabird species. *PeerJ*, *10*,  
674 e13073. <https://doi.org/10.7717/peerj.13073>

675 Pezza, A. B., & Simmonds, I. (2008). Large-scale factors in tropical and extratropical cyclone  
676 transition and extreme weather events. *Annals of the New York Academy of Sciences*, *1146*(1), 189–  
677 211. <https://doi.org/10.1196/annals.1446.005>

678 Price, P. W. (1997). *Insect Ecology*. New Jersey: John Wiley & Sons.

679 R Core Team (2022). R: A language and environment for statistical computing. R Foundation for  
680 Statistical Computing, Vienna, Austria. <https://www.R-project.org/>.

681 Radford, A. N., McCleery, R. H., Woodburn, R. J. W., & Morecroft, M. D. (2001). Activity patterns  
682 of parent great tits *Parus major* feeding their young during rainfall. *Bird Study*, *48*(2), 214–220.  
683 <https://doi.org/10.1080/00063650109461220>

684 Raj Pant, S., Hammers, M., Komdeur, J., Burke, T., Dugdale, H. L., & Richardson, D. S. (2020). Age-  
685 dependent changes in infidelity in Seychelles warblers. *Molecular Ecology*, *29*(19), 3731–3746.  
686 <https://doi.org/10.1111/mec.15563>

687 Richardson, D. S., Burke, T., & Komdeur, J. (2002). Direct benefits and the evolution of female-biased  
688 cooperative breeding in Seychelles warblers. *Evolution*, *56*(11), 2313–2321.  
689 <https://doi.org/10.1111/j.0014-3820.2002.tb00154.x>

690 Richardson, D. S., Burke, T., & Komdeur, J. (2007). Grandparent helpers: the adaptive significance of  
691 older, postdominant helpers in the Seychelles warbler. *Evolution; International Journal of Organic*  
692 *Evolution*, *61*(12), 2790–2800. <https://doi.org/10.1111/j.1558-5646.2007.00222.x>

693 Richardson, D. S., Jury, F. L., Blaakmeer, K., Komdeur, J., & Burke, T. (2001). Parentage assignment  
694 and extra-group paternity in a cooperative breeder: the Seychelles warbler (*Acrocephalus*  
695 *sechellensis*). *Molecular Ecology*, *10*(9), 2263–2273. [https://doi.org/10.1046/j.0962-](https://doi.org/10.1046/j.0962-1083.2001.01355.x)  
696 [1083.2001.01355.x](https://doi.org/10.1046/j.0962-1083.2001.01355.x)

697 Richardson, D. S., Komdeur, J., & Burke, T. (2003). Altruism and infidelity among warblers. *Nature*,  
698 422(6932). <https://doi.org/10.1038/422580a>

699 Sánchez-Macouzet, O., Rodríguez, C., & Drummond, H. (2014). Better stay together: pair bond  
700 duration increases individual fitness independent of age-related variation. *Proceedings of the Royal*  
701 *Society B: Biological Sciences*, 281(1786), 20132843. <https://doi.org/10.1098/rspb.2013.2843>

702 Sapolsky, R. M., Romero, L. M., & Munck, A. U. (2000). How do glucocorticoids influence stress  
703 responses? Integrating permissive, suppressive, stimulatory, and preparative actions. *Endocrine*  
704 *reviews*, 21(1), 55-89. <https://doi.org/10.1210/edrv.21.1.0389>

705 Schöll, E. M., & Hille, S. M. (2020). Heavy and persistent rainfall leads to brood reduction and nest  
706 failure in a passerine bird. *Journal of Avian Biology*, 51(7). <https://doi.org/10.1111/jav.02418>

707 Seychelles Meteorological Authority (2016) Monthly precipitation data for Praslin airstrip 1994-2016.  
708 Data available from: [info@meteo.gov.sc](mailto:info@meteo.gov.sc)

709 Sparks, A. M., Spurgin, L. G., van der Velde, M., Fairfield, E. A., Komdeur, J., Burke, T., Richardson,  
710 D. S., & Dugdale, H. L. (2022). Telomere heritability and parental age at conception effects in a wild  
711 avian population. *Molecular Ecology*, 31(23), 6324–6338. <https://doi.org/10.1111/mec.15804>

712 Speelman, F., Burke, T., Komdeur, J., Richardson, D., & Dugdale, H. (2024). Causes and  
713 consequences of divorce in a long-lived socially monogamous bird. *Authorea*.  
714 <https://doi.org/10.22541/au.170994368.84507116/v1>

715 Spooner, F. E. B., Pearson, R. G., & Freeman, R. (2018). Rapid warming is associated with population  
716 decline among terrestrial birds and mammals globally. *Global Change Biology*, 24(10), 4521–4531.  
717 <https://doi.org/10.1111/gcb.14361>



718 Studds, C. E., & Marra, P. P. (2007). Linking fluctuations in rainfall to nonbreeding season  
719 performance in a long-distance migratory bird, *Setophaga ruticilla*. *Climate Research*, 35(1–2), 115–  
720 122. <https://doi.org/10.3354/cr00718>

721 van Boheemen, L. A., Hammers, M., Kingma, S. A., Richardson, D. S., Burke, T., Komdeur, J., &  
722 Dugdale, H. L. (2019). Compensatory and additive helper effects in the cooperatively breeding  
723 Seychelles warbler (*Acrocephalus sechellensis*). *Ecology and Evolution*, 9(5), 2986–2995.  
724 <https://doi.org/10.1002/ece3.4982>

725 van de Crommenacker, J., Komdeur, J., Burke, T., & Richardson, D. S. (2011). Spatio-temporal  
726 variation in territory quality and oxidative status: a natural experiment in the Seychelles warbler  
727 (*Acrocephalus sechellensis*). *Journal of Animal Ecology*, 80(3), 668–680.  
728 <https://doi.org/10.1111/j.1365-2656.2010.01792.x>

729 Ventura, F., Granadeiro, J. P., Lukacs, P. M., Kuepfer, A., & Catry, P. (2021). Environmental  
730 variability directly affects the prevalence of divorce in monogamous albatrosses. *Proceedings of the*  
731 *Royal Society B: Biological Sciences*, 288(1963), 20212112. <https://doi.org/10.1098/rspb.2021.2112>

732 Wickham, H. (2016). Data Analysis. In H. Wickham (Ed.), *ggplot2: Elegant Graphics for Data*  
733 *Analysis* (pp. 189–201). New York: Springer. [https://doi.org/10.1007/978-3-319-24277-4\\_9](https://doi.org/10.1007/978-3-319-24277-4_9)

734 Wilson, G. R., Cooper, S. J., & Gessaman, J. A. (2004). The effects of temperature and artificial rain  
735 on the metabolism of American kestrels (*Falco sparverius*). *Comparative Biochemistry and*  
736 *Physiology Part A: Molecular & Integrative Physiology*, 139(3), 389–394.  
737 <https://doi.org/10.1016/j.cbpb.2004.10.009>

738 Wilson, K. M., Nguyen, M., & Burley, N. T. (2022). Divorce rate varies with fluidity of passerine  
739 social environment. *Animal Behaviour*, 183, 51–60. <https://doi.org/10.1016/j.anbehav.2021.10.018>

740 Wright, D. J., Shah, N. J., & Richardson, D. S. (2014). Translocation of the Seychelles warbler  
741 *Acrocephalus sechellensis* to establish a new population on Frégate Island, Seychelles. *Conserv Evid*,  
742 *11*, 20-24.

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758 Supplementary materials for the manuscript titled “Rainfall is associated with divorce  
759 in the socially monogamous Seychelles warbler”.

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## 762 **Materials & Methods**

### 763 **Pairwise relatedness**

764 Seychelles warbler DNA was extracted from brachial venipuncture blood samples using a Qiagen  
765 DNeasy Blood and Tissue Kit (2013 onwards) or modified ammonium acetate protocol (before 2013).  
766 DNA samples were used to determine sex using 1 to 3 markers and genotyping using a panel of 30  
767 microsatellite markers (Richardson *et al.*, 2004; Raj Pant *et al.*, 2020; Sparks *et al.*, 2022). Parentage  
768 was assigned using MasterBayes 2.52 (Hadfield *et al.*, 2006), which was used to build a genetic  
769 pedigree (Sparks *et al.*, 2022). We calculated pairwise relatedness between partners using the Queller  
770 and Goodnight estimation using the R-package *related* 0.8 (Queller & Goodnight, 1989; Pew *et al.*,  
771 2015). This estimation of pairwise relatedness also reflects pedigree relatedness in the Seychelles  
772 warbler (Brouwer *et al.*, 2007), and heterozygosity across the microsatellite panel reflects genome-  
773 wide heterozygosity (van de Crommenacker *et al.*, 2011).

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783 **Territory quality**

784 Territory quality in the main breeding seasons was measured using an index of insect availability,  
785 territory size, and foliage cover (Komdeur, 1992; van de Crommenacker *et al.*, 2011). This was done  
786 using the equation  $A \times \sum(Cx \times lx)$ , where  $A$  is territory size in ha,  $Cx$  is the percentage of foliage cover  
787 for tree species  $x$ , and  $lx$  is the per unit leaf area mean monthly insect density for tree species  $x$  in  $\text{dm}^2$ .  
788 Insect abundance was estimated by counting the number of insects on the underside of 50 leaves for  
789 ten dominant tree species, once a month at 14 different island locations. Estimates of insect counts for  
790 all territories were estimated based on their proximity to one of these locations. Foliage cover was  
791 estimated by scoring the presence or absence of ten dominant tree species at various heights during the  
792 middle of the breeding season (typically July). This was done at 20 different points in all territories  
793 and each tree species' total number of presence scores was its estimated foliage cover. In 2002, no  
794 territory quality data was collected resulting in 15 years of food abundance data.

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808 **Climate window analysis**

809 We used *climwin* 1.2.3 (Bailey & van de Pol, 2016) to determine which temporal windows of rainfall  
810 best predicted divorce, measurements of reproductive success, and measurements of territory quality.

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812 The *slidingwin* function determined the months of rainfall best predicting the variation in response  
813 variables using a sliding window technique. July is the peak of breeding, which is when the most eggs  
814 are laid, and warblers can then provide up to three months of post-fledgling care. As a result, we used  
815 the end of the main breeding season (28th of September) as the *slidingwin* reference date. Breeding  
816 statuses, which define our characterization of divorce, are finalized at the end of the breeding season,  
817 and our study is interested in investigating what happened in between the moment we know a  
818 partnership was last together and no longer together. As a result, we tested for all possible temporal  
819 windows (all combinations of months) from 12 months leading up to the end of the main breeding  
820 season (28th of September). Thus, from one end of the main breeding season to the end of the next.

821

822 Importantly, *climwin* is designed to avoid issues regarding multiple comparisons through a  
823 randomization scheme that ensures temporal windows are not found due to chance. Thus, after finding  
824 a temporal window, we tested whether the result was found due to chance (which was never the case)  
825 using the function *randwin*. We performed the *randwin* randomization procedure 1000 times and  
826 confirmed that observing such a negative value for the  $\Delta AICc$  of the best model was statistically  
827 significant ( $p\Delta AICc < 0.001$ ).

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829 For all response variables, both the linear and quadratic functions of rainfall were tested. AIC values  
830 of the models created were used to determine whether the linear or quadratic relationship best fit the  
831 data. A better fit for the more complicated model (quadratic) was defined as a  $\Delta AIC > 7$  (Burnham *et*  
832 *al.*, 2011).

## 833 **Supplementary References**

834 Bailey, L. D., & van de Pol, M. (2016). climwin: an R toolbox for climate window analysis. *PLOS*  
835 *ONE*, *11*(12), e0167980. <https://doi.org/10.1371/journal.pone.0167980>

836

837 Brouwer, L., Richardson, D. S., Eikenaar, C., & Komdeur, J. (2006). The role of group size and  
838 environmental factors on survival in a cooperatively breeding tropical passerine. *Journal of Animal*  
839 *Ecology*, *75*(6), 1321–1329. <https://doi.org/10.1111/j.1365-2656.2006.01155.x>

840

841 Burnham, K. P., Anderson, D. R., & Huyvaert, K. P. (2011). AIC model selection and multimodel  
842 inference in behavioral ecology: some background, observations, and comparisons. *Behavioral*  
843 *Ecology and Sociobiology*, *65*, 23-35. <https://doi.org/10.1007/s00265-010-1029-6>

844

845 Hadfield, J. D., Richardson, D. S., & Burke, T. (2006). Towards unbiased parentage assignment:  
846 combining genetic, behavioural and spatial data in a Bayesian framework. *Molecular Ecology*, *15*(12),  
847 3715–3730. <https://doi.org/10.1111/j.1365-294X.2006.03050.x>

848

849 Komdeur, J. (1992). Importance of habitat saturation and territory quality for evolution of cooperative  
850 breeding in the Seychelles warbler. *Nature*, *358*(6386). <https://doi.org/10.1038/358493a0>

851

852 Pew, J., Muir, P. H., Wang, J., & Frasier, T. R. (2015). related: an R package for analysing pairwise  
853 relatedness from codominant molecular markers. *Molecular ecology resources*, *15*(3), 557-561.  
854 <https://doi.org/10.1111/1755-0998.12323>

855

856 Raj Pant, S., Hammers, M., Komdeur, J., Burke, T., Dugdale, H. L., & Richardson, D. S. (2020). Age-  
857 dependent changes in infidelity in Seychelles warblers. *Molecular Ecology*, 29(19), 3731–3746.  
858 <https://doi.org/10.1111/mec.15563>

859

860 Richardson, D. S., Komdeur, J., & Burke, T. (2004). Inbreeding in the Seychelles warbler:  
861 environment-dependent maternal effects. *Evolution; International Journal of Organic Evolution*,  
862 58(9), 2037–2048. <https://doi.org/10.1111/j.0014-3820.2004.tb00488.x>

863

864 Sparks, A. M., Spurgin, L. G., van der Velde, M., Fairfield, E. A., Komdeur, J., Burke, T., Richardson,  
865 D. S., & Dugdale, H. L. (2022). Telomere heritability and parental age at conception effects in a wild  
866 avian population. *Molecular Ecology*, 31(23), 6324–6338. <https://doi.org/10.1111/mec.15804>

867

868 van de Crommenacker, J., Komdeur, J., Burke, T., & Richardson, D. S. (2011). Spatio-temporal  
869 variation in territory quality and oxidative status: a natural experiment in the Seychelles warbler  
870 (*Acrocephalus sechellensis*). *Journal of Animal Ecology*, 80(3), 668–680.  
871 <https://doi.org/10.1111/j.1365-2656.2010.01792.x>

872

873 Queller, D. C., & Goodnight, K. F. (1989). Estimating relatedness using genetic markers. *Evolution*,  
874 43(2), 258–275. <https://doi.org/10.1111/j.1558-5646.1989.tb04226.x>

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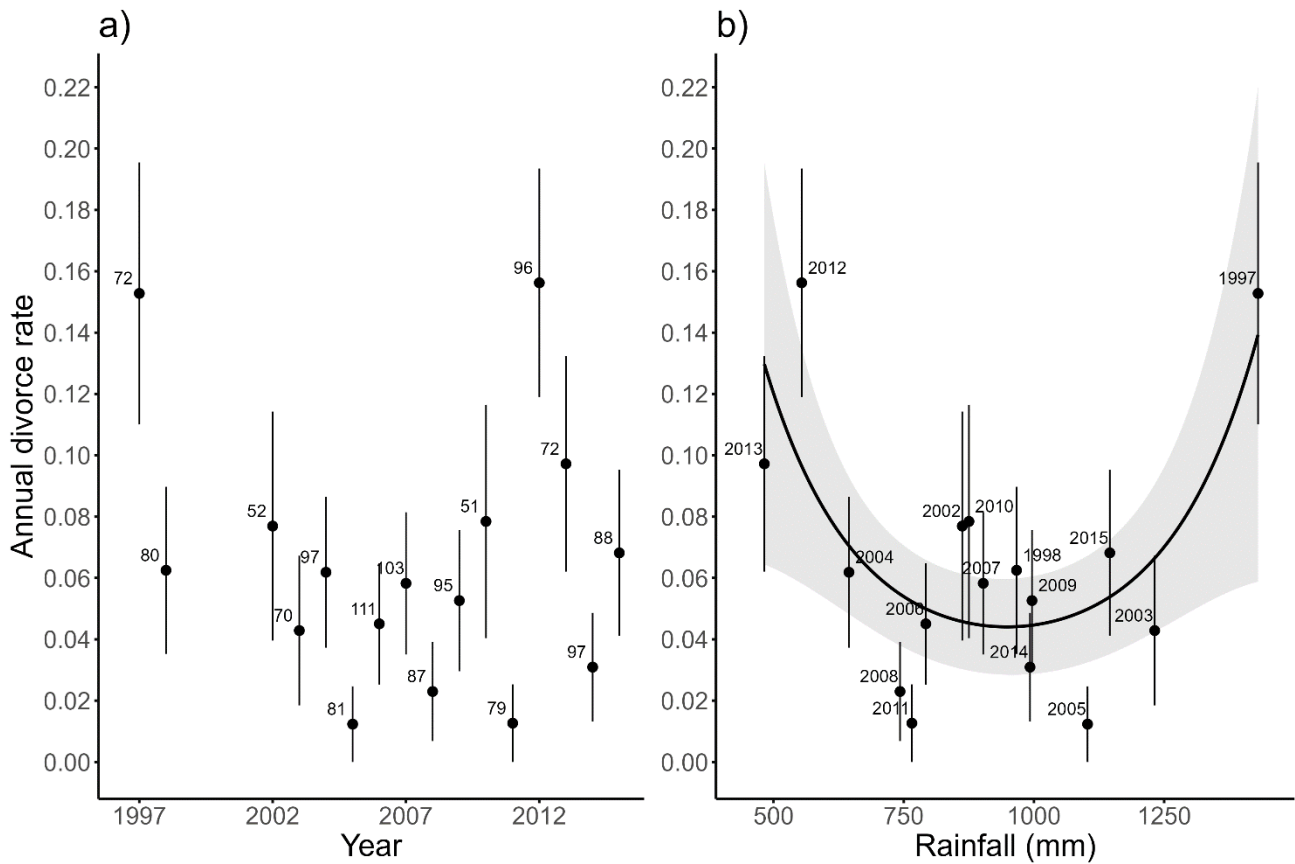
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881 **Results**



882

883 **Figure S1.** a) Variability in the annual divorce rate of the Seychelles warbler on Cousin Island ( $n =$   
 884 1321 partnerships) from 1997 to 2015. The years 1999, 2000, and 2001 were not included due to  
 885 limited fieldwork during those years. b) The effect of rainfall on the annual divorce rate as predicted  
 886 by a quasi-binomial generalized linear model. The solid line represents the predicted divorce rate, and  
 887 the grey shading indicates the 95% confidence intervals. Dots represent the mean observed annual  
 888 divorce rate  $\pm$  SE, and labels indicate the total number of partnerships in a given year (a) or the sample  
 889 year (b).

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894 **Table S1.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
895 with the total rainfall from February to August, the length of the partnership, the number of offspring,  
896 the relatedness of the breeding pair, the number of helpers, male age, and female age. A total of  $n =$   
897 1252 partnerships were analyzed using a binomial generalized linear mixed model. Significant  $p$ -  
898 values are in bold. Data from 1997 were removed from this analysis and the non-significant quadratic  
899 term of rainfall is included.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-4.012	0.694	-5.372 to -2.651	<b>&lt;0.001</b>
Rainfall	-0.233	0.172	-0.570 to 0.104	0.175
Rainfall <sup>2</sup>	0.285	0.164	-0.037 to 0.607	0.082
Partnership length	-0.319	0.251	-0.811 to 0.173	0.204
Number of offspring	-0.104	0.155	-0.408 to 0.200	0.502
Pairwise relatedness	0.142	0.161	-0.173 to 0.457	0.378
Number of helpers	-0.316	0.193	-0.696 to 0.063	0.102
Male age	-0.015	0.197	-0.402 to 0.372	0.939
Female age	0.184	0.180	-0.169 to 0.538	0.307
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	1.022	392		
Female ID	1.156	372		
Field period ID	0.000	15		
Territory ID	0.185	156		

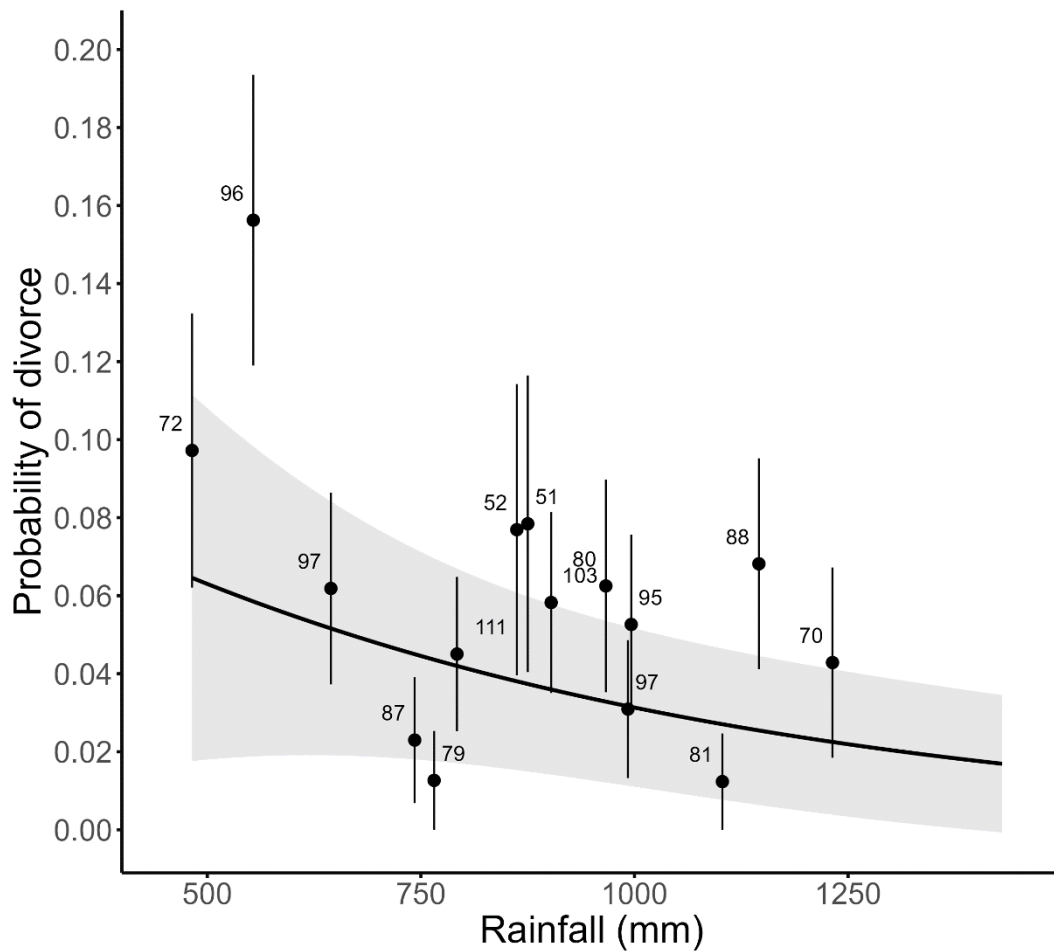
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903 **Table S2.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 904 with the total rainfall from February to August, the length of the partnership, the number of offspring,  
 905 the relatedness of the breeding pair, the number of helpers, male age, and female age. A total of  $n =$   
 906 1252 partnerships were analyzed using a binomial generalized linear mixed model. Significant  $p$ -  
 907 values are in bold. Data from 1997 were removed from this analysis and the non-significant quadratic  
 908 term of rainfall is excluded.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.280	0.320	-3.907 to -2.653	<b>&lt;0.001</b>
Rainfall	-0.345	0.169	-0.676 to -0.015	<b>0.040</b>
Partnership length	-0.415	0.221	-0.848 to 0.017	0.060
Number of offspring	-0.096	0.145	-0.381 to 0.189	0.509
Pairwise relatedness	0.105	0.134	-0.158 to 0.367	0.435
Number of helpers	-0.278	0.180	-0.631 to 0.075	0.123
Male age	-0.007	0.171	(0.343 to 0.328	0.966
Female age	0.171	0.153	-0.128 to 0.47	0.263
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.000	392		
Female ID	0.270	372		
Field period ID	0.076	15		
Territory ID	0.352	156		



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910 **Figure S2.** The effect of total rainfall from February to August on the probability of divorce in the  
 911 Seychelles warbler on Cousin Island ( $n = 1252$  partnerships) as predicted by a binomial generalized  
 912 linear mixed model (Table S2) where data from 1997 was removed from the analysis. The solid line  
 913 represents the predicted probability of divorce, and the grey shading indicates the 95% confidence  
 914 intervals. Dots represent the mean observed divorce rate  $\pm$  SE, and labels indicate the total number of  
 915 partnerships in a given year.

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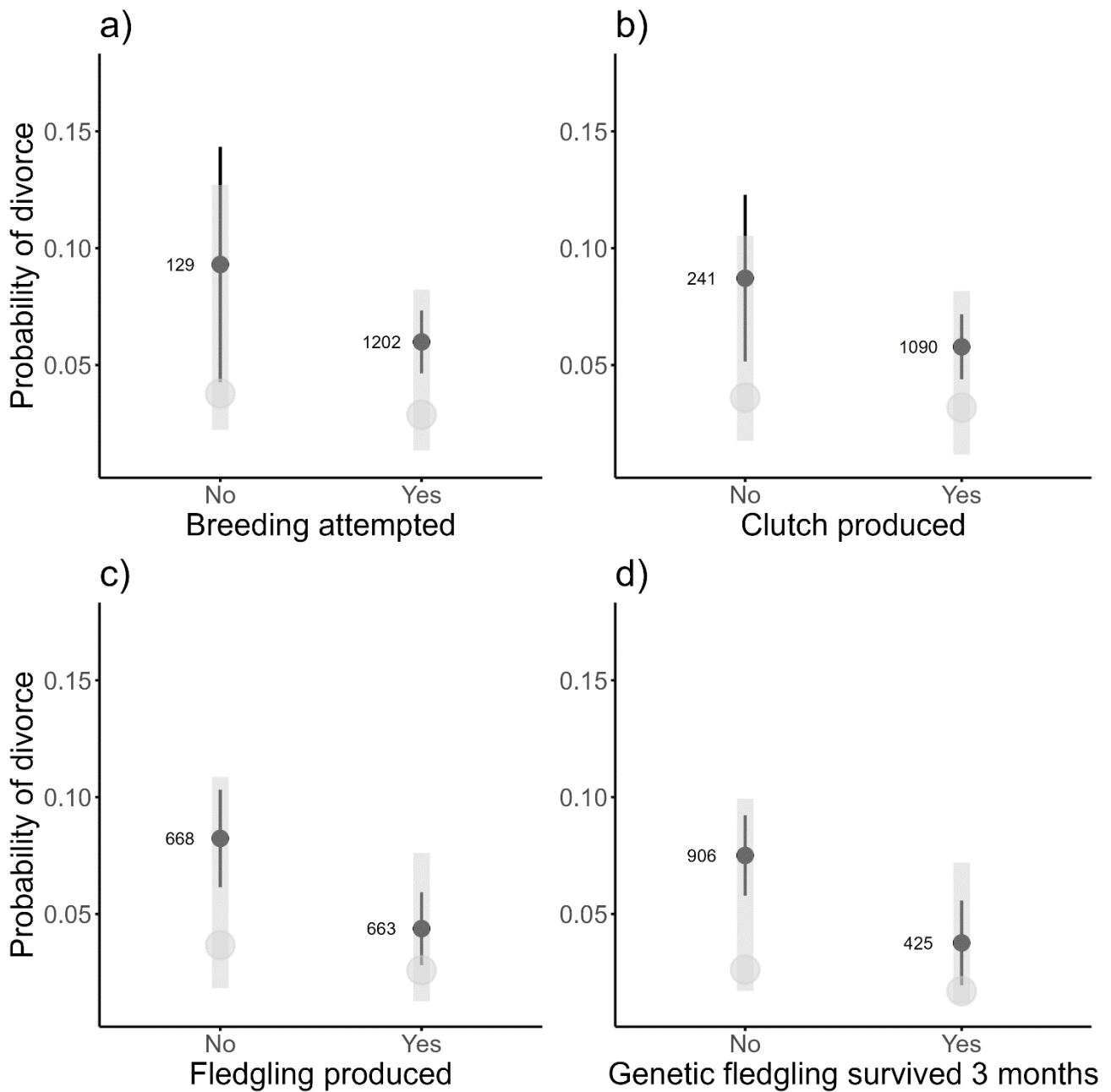
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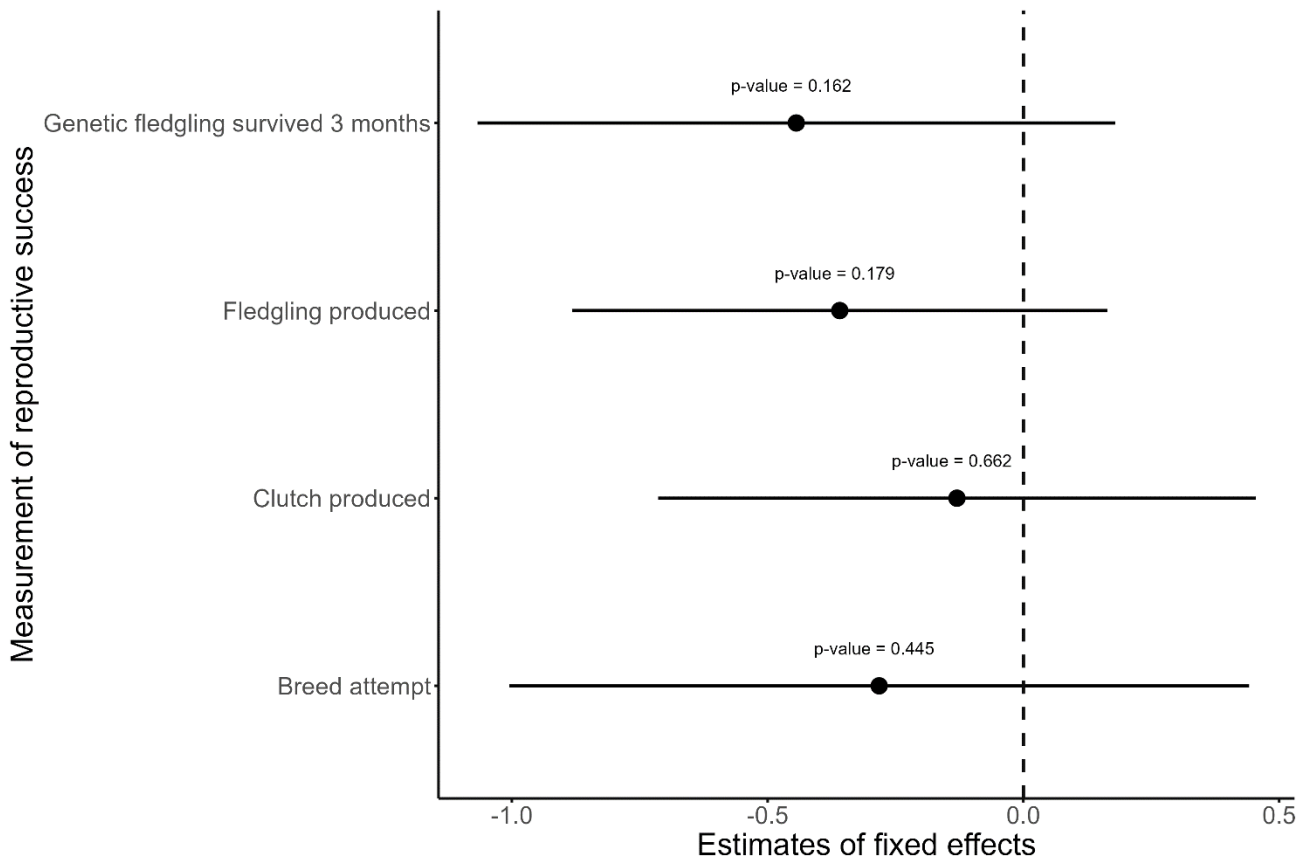
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923 **Figure S3.** The probability of divorce for Seychelles warbler partnerships ( $n = 1321$ ) that did or did  
 924 not: a) attempt to breed; b) produce a clutch; c) produce a fledgling; d) produce a fledgling genetically  
 925 related to the dominant female that survived till at the least three months old. The grey dots and shaded  
 926 area represent the probability of divorce  $\pm$  95% confidence intervals as predicted by binomial  
 927 generalized linear mixed model. The black dots indicate the mean observed divorce rate  $\pm$  95%  
 928 confidence intervals, and labels indicate the number of partnerships.



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930 **Figure S4.** The coefficient estimates (dots) and 95% confidence intervals (bars) of four measures of  
 931 reproductive success on the probability of divorce in the Seychelles warbler ( $n = 1325$  partnerships)  
 932 as predicted by binomial generalized linear mixed model. Each reproductive measure was  
 933 independently included in the model.  $p$ -values are indicated on the figure.

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937 The following tables (Tables S3 to S6) compare the effects of reproduction at four different stages -  
 938 breeding attempted (Y/N), clutch produced (Y/N), fledgling produced (Y/N), and genetic fledglings  
 939 (Y/N) - on divorce by including them one at a time in the partnership-level probability of divorce  
 940 model. The reproductive measures are italicized in the table legends and table contents for ease of  
 941 comparison between the four different model summary tables.

942 **Table S3.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 943 with rainfall, the length of the partnership, *breeding attempted*, the relatedness of the breeding pair,  
 944 the number of helpers, male age, and female age.  $n = 1321$  partnerships were analyzed using a binomial  
 945 generalized linear mixed model. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.239	0.469	-4.157 to -2.320	<b>&lt;0.001</b>
Rainfall	-0.130	0.115	-0.354 to 0.095	0.258
Rainfall <sup>2</sup>	0.329	0.086	0.160 to 0.497	<b>&lt;0.001</b>
Partnership length	-0.505	0.216	-0.929 to -0.081	<b>0.019</b>
<i>Breeding attempted</i>	-0.282	0.369	-1.006 to 0.442	0.445
Pairwise relatedness	0.151	0.128	-0.100 to 0.402	0.239
Number of helpers	-0.20	0.157	-0.508 to 0.108	0.203
Male age	0.075	0.158	-0.234 to 0.385	0.633
Female age	0.206	0.144	-0.076 to 0.488	0.152
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	<0.001	416		
Female ID	0.274	392		
Field period ID	0.020	16		
Territory ID	0.354	158		

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948 **Table S4.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 949 with rainfall, the length of the partnership, *clutch produced*, the relatedness of the breeding pair, the  
 950 number of helpers, male age, and female age.  $n = 1321$  partnerships were analyzed using a binomial  
 951 generalized linear mixed model. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.285	0.351	-3.973 to -2.598	<b>&lt;0.001</b>
Rainfall	-0.139	0.113	-0.361 to 0.082	0.218
Rainfall <sup>2</sup>	0.325	0.085	0.159 to 0.491	<b>&lt;0.001</b>
Partnership length	-0.521	0.208	-0.929 to -0.114	<b>0.012</b>
<i>Clutch produced</i>	-0.130	0.298	-0.714 to 0.453	0.662
Pairwise relatedness	0.146	0.124	-0.097 to 0.388	0.240
Number of helpers	-0.200	0.156	-0.505 to 0.106	0.201
Male age	0.079	0.154	-0.223 to 0.381	0.607
Female age	0.204	0.139	-0.068 to 0.475	0.142
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.000	416		
Female ID	0.000	392		
Field period ID	0.024	16		
Territory ID	0.365	158		

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953

954 **Table S5.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 955 with rainfall, the length of the partnership, *fledgling produced*, the relatedness of the breeding pair, the  
 956 number of helpers, male age, and female age.  $n = 1321$  partnerships were analyzed using a binomial  
 957 generalized linear mixed model. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.268	0.363	-3.98 to -2.557	<b>&lt;0.001</b>
Rainfall	-0.128	0.112	-0.348 to 0.092	0.253
Rainfall <sup>2</sup>	0.314	0.087	0.144 to 0.484	<b>&lt;0.001</b>
Partnership length	-0.488	0.214	-0.908 to -0.067	<b>0.023</b>
<i>Fledgling produced</i>	-0.359	0.267	-0.883 to 0.164	0.179
Pairwise relatedness	0.141	0.127	-0.107 to 0.390	0.265
Number of helpers	-0.153	0.161	-0.469 to 0.163	0.343
Male age	0.073	0.157	-0.234 to 0.381	0.640
Female age	0.198	0.142	-0.080 to 0.475	0.162
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.000	416		
Female ID	0.188	392		
Field period ID	0.024	16		
Territory ID	0.335	158		

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960 **Table S6.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 961 with rainfall, the length of the partnership, *genetic fledgling produced*, the relatedness of the breeding  
 962 pair, the number of helpers, male age, and female age.  $n = 1321$  partnerships were analyzed using a  
 963 binomial generalized linear mixed model. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.615	0.549	-4.690 to -2.539	<b>&lt;0.001</b>
Rainfall	-0.144	0.113	-0.366 to 0.078	0.203
Rainfall <sup>2</sup>	0.329	0.089	0.155 to 0.503	<b>&lt;0.001</b>
Partnership length	-0.441	0.232	-0.895 to 0.013	0.057
<i>Genetic fledgling produced</i>	-0.444	0.318	-1.067 to 0.179	0.162
Pairwise relatedness	0.155	0.142	-0.123 to 0.433	0.274
Number of helpers	-0.221	0.165	-0.545 to 0.103	0.181
Male age	0.077	0.172	-0.259 to 0.413	0.652
Female age	0.201	0.157	-0.107 to 0.509	0.202
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.532	416		
Female ID	0.636	392		
Field period ID	0.000	16		
Territory ID	0.260	158		

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967 **Interaction between rainfall and partnership length**

968 We found a significant interaction between partnership length and rainfall (Table S7), where the model  
969 predicted that heavy rainfall increased the probability of divorce in shorter-lasting but not longer-  
970 lasting partnerships (Figure S5). However, as the sample sizes of longer-lasting partnerships were  
971 small (Table S8), this relationship was strongly influenced by outliers. Outliers were defined as  
972 categories with less than 50 partnerships (low estimate) or less than 100 partnerships (high estimate).  
973 In both cases, removing outliers from the analysis removed the significant interaction (Table S10;  
974 Table S11). Also, running the model with breeding experience, defined as if a partnership has been  
975 together for at least one breeding season (Y/N), instead of partnership length removed the significant  
976 interaction (Table S12). As a result, the interaction was not included in the final model.

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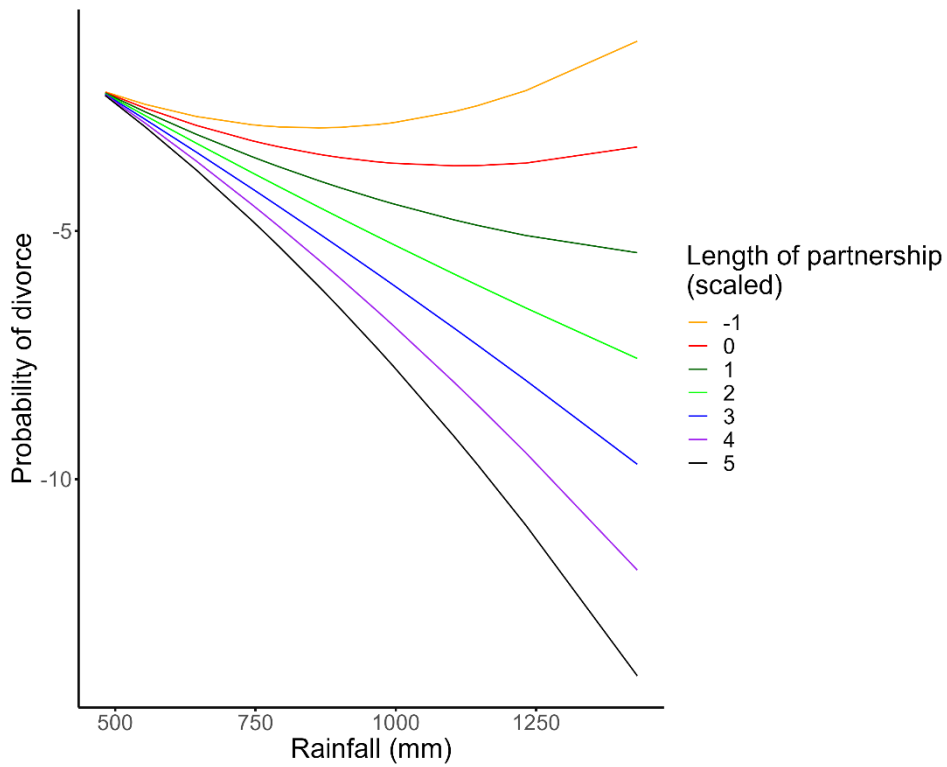
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991 **Table S7.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 992 with the total rainfall from February to August, length of partnership, the number of offspring, the  
 993 relatedness of the breeding pair, the number of helpers, male age, and female age.  $n = 1321$  partnerships  
 994 were analyzed using a binomial GLMM. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.840	0.542	-4.903 to -2.777	<b>&lt;0.001</b>
Rainfall	-0.344	0.164	-0.666 to -0.022	<b>0.037</b>
Rainfall <sup>2</sup>	0.240	0.117	0.011 to 0.469	<b>0.04</b>
Partnership length	-0.622	0.301	-1.212 to -0.032	<b>0.039</b>
Number of offspring	-0.089	0.147	-0.378 to 0.199	0.544
Pairwise relatedness	0.162	0.14	-0.112 to 0.435	0.247
Number of helpers	-0.197	0.164	-0.518 to 0.125	0.231
Male age	-0.072	0.200	-0.465 to 0.320	0.717
Male age <sup>2</sup>	0.190	0.112	-0.031 to 0.410	0.091
Female age	0.273	0.184	-0.088 to 0.634	0.138
Female age <sup>2</sup>	-0.121	0.118	-0.353 to 0.112	0.309
Rainfall * Partnership length	-0.484	0.217	-0.908 to -0.059	<b>0.026</b>
Rainfall <sup>2</sup> * Partnership length	-0.063	0.148	-0.353 to 0.227	0.672
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.411	416		
Female ID	0.555	392		
Field period ID	0.000	16		
Territory ID	0.314	158		

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997 **Figure S5.** The effect of total rainfall from February to August on the probability of divorce for  
 998 Seychelles warbler partnerships ( $n = 1321$ ) that have been together for different lengths of time on  
 999 Cousin Island as predicted by a binomial generalized linear mixed model.

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1001 **Table S8.** The number of available samples of Seychelles warbler partnerships that have been together  
 1002 for different lengths of time on Cousin Island.

Partnership length (scaled)	Number of partnerships
-1	534
0	496
1	183
2	73
3	30
4	10
5	5

1003 **Table S9.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 1004 with rainfall, the length of the partnership, the relatedness of the breeding pair, the number of helpers,  
 1005 male age, female age, male extra-pair-paternity (EPP; infidelity), female EPP, and co-breeder presence  
 1006 (Y/N).  $n = 1321$  partnerships were analyzed using a binomial generalized linear mixed model.  
 1007 Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.354	0.359	-4.058 to -2.650	<b>&lt;0.001</b>
Rainfall	-0.156	0.112	-0.376 to 0.064	0.164
Rainfall <sup>2</sup>	0.326	0.086	0.157 to 0.494	<b>&lt;0.001</b>
Partnership length	-0.488	0.215	-0.910 to -0.065	<b>0.024</b>
Number of offspring	0.031	0.217	-0.394 to 0.455	0.887
Pairwise relatedness	0.137	0.127	-0.112 to 0.385	0.282
Number of helpers	-0.211	0.159	-0.523 to 0.100	0.184
Male age	0.069	0.159	-0.243 to 0.380	0.666
Female age	0.193	0.143	-0.087 to 0.472	0.177
Male EPP	-0.016	0.171	-0.351 to 0.319	0.926
Female EPP	-0.324	0.286	-0.885 to 0.237	0.257
Co-breeder presence	0.413	0.333	-0.240 to 1.065	0.215
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.000	392		
Female ID	0.170	372		
Field period ID	0.019	16		
Territory ID	0.3650	156		

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1010 **Table S10.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 1011 with the total rainfall from February to August, length of partnership, the number of offspring, the  
 1012 relatedness of the breeding pair, the number of helpers, male age, and female age.  $n = 1296$  partnerships  
 1013 were analyzed using a binomial generalized linear mixed model. Significant  $p$ -values are in bold.  
 1014 Partnerships with a partnership length greater than 3 (scaled value) were removed from this analyses.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.468	0.378	-4.208 to -2.727	<b>&lt;0.001</b>
Rainfall	-0.295	0.159	-0.607 to 0.016	0.063
Rainfall <sup>2</sup>	0.256	0.120	0.021 to 0.490	<b>0.032</b>
Partnership length	-0.839	0.335	-1.495 to -0.182	<b>0.012</b>
Number of offspring	-0.083	0.144	-0.364 to 0.199	0.564
Pairwise relatedness	0.148	0.129	-0.105 to 0.400	0.252
Number of helpers	-0.201	0.165	-0.523 to 0.122	0.223
Male age	-0.011	0.191	-0.386 to 0.364	0.955
Male age <sup>2</sup>	0.133	0.115	-0.093 to 0.360	0.249
Female age	0.324	0.179	-0.026 to 0.674	0.070
Female age <sup>2</sup>	-0.291	0.150	-0.585 to 0.004	0.053
Rainfall * Partnership length	-0.359	0.213	-0.777 to 0.060	0.093
Rainfall <sup>2</sup> * Partnership length	-0.029	0.159	-0.340 to 0.283	0.857
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.000	416		
Female ID	0.296	392		
Field period ID	0.009	16		
Territory ID	0.289	158		

1015 **Table S11.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 1016 with the total rainfall from February to August, length of partnership, the number of offspring, the  
 1017 relatedness of the breeding pair, the number of helpers, male age, and female age.  $n = 1245$  partnerships  
 1018 were analyzed using a binomial generalized linear mixed model. Significant  $p$ -values are in bold.  
 1019 Partnerships with a partnership length greater than 2 (scaled value) were removed from this analyses.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.480	0.388	-4.241 to -2.720	<b>&lt;0.001</b>
Rainfall	-0.278	0.160	-0.592 to 0.037	0.083
Rainfall <sup>2</sup>	0.249	0.122	0.009 to 0.489	<b>0.042</b>
Partnership length	-0.778	0.353	-1.469 to -0.086	<b>0.027</b>
Number of offspring	-0.114	0.146	-0.401 to 0.173	0.436
Pairwise relatedness	0.128	0.130	-0.126 to 0.382	0.324
Number of helpers	-0.194	0.165	-0.518 to 0.129	0.240
Male age	-0.030	0.193	-0.408 to 0.349	0.878
Male age <sup>2</sup>	0.151	0.119	-0.082 to 0.385	0.204
Female age	0.321	0.178	-0.028 to 0.669	0.072
Female age <sup>2</sup>	-0.261	0.152	-0.559 to 0.037	0.086
Rainfall * Partnership length	-0.340	0.230	-0.791 to 0.111	0.140
Rainfall <sup>2</sup> * Partnership length	-0.035	0.173	-0.374 to 0.304	0.839
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	<0.001	416		
Female ID	0.312	392		
Field period ID	<0.001	16		
Territory ID	0.280	158		

1020 **Table S12.** Associations between the probability of divorce in the Seychelles warbler on Cousin Island  
 1021 with the total rainfall from February to August, breeding experience, the number of offspring, the  
 1022 relatedness of the breeding pair, the number of helpers, male age, and female age.  $n = 1321$  partnerships  
 1023 were analyzed using a binomial generalized linear mixed model. Significant  $p$ -values are in bold.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	-3.245	0.667	-4.551 to -1.938	<b>&lt;0.001</b>
Rainfall	0.014	0.158	-0.296 to 0.325	0.928
Rainfall <sup>2</sup>	0.245	0.116	0.016 to 0.473	<b>0.036</b>
Breeding experience	-0.983	0.429	-1.824 to -0.142	<b>0.022</b>
Number of offspring	-0.079	0.148	-0.368 to 0.210	0.594
Pairwise relatedness	0.160	0.140	-0.114 to 0.434	0.251
Number of helpers	-0.201	0.163	-0.520 to 0.118	0.217
Male age	-0.126	0.190	-0.499 to 0.247	0.508
Male age <sup>2</sup>	0.162	0.114	-0.062 to 0.387	0.156
Female age	0.225	0.176	-0.120 to 0.570	0.201
Female age <sup>2</sup>	-0.136	0.120	-0.371 to 0.099	0.256
Rainfall * Breeding experience	-0.333	0.233	-0.790 to 0.123	0.152
Rainfall <sup>2</sup> * Breeding experience	0.214	0.171	-0.121 to 0.548	0.210
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	0.419	416		
Female ID	0.557	392		
Field period ID	0.000	16		
Territory ID	0.321	158		

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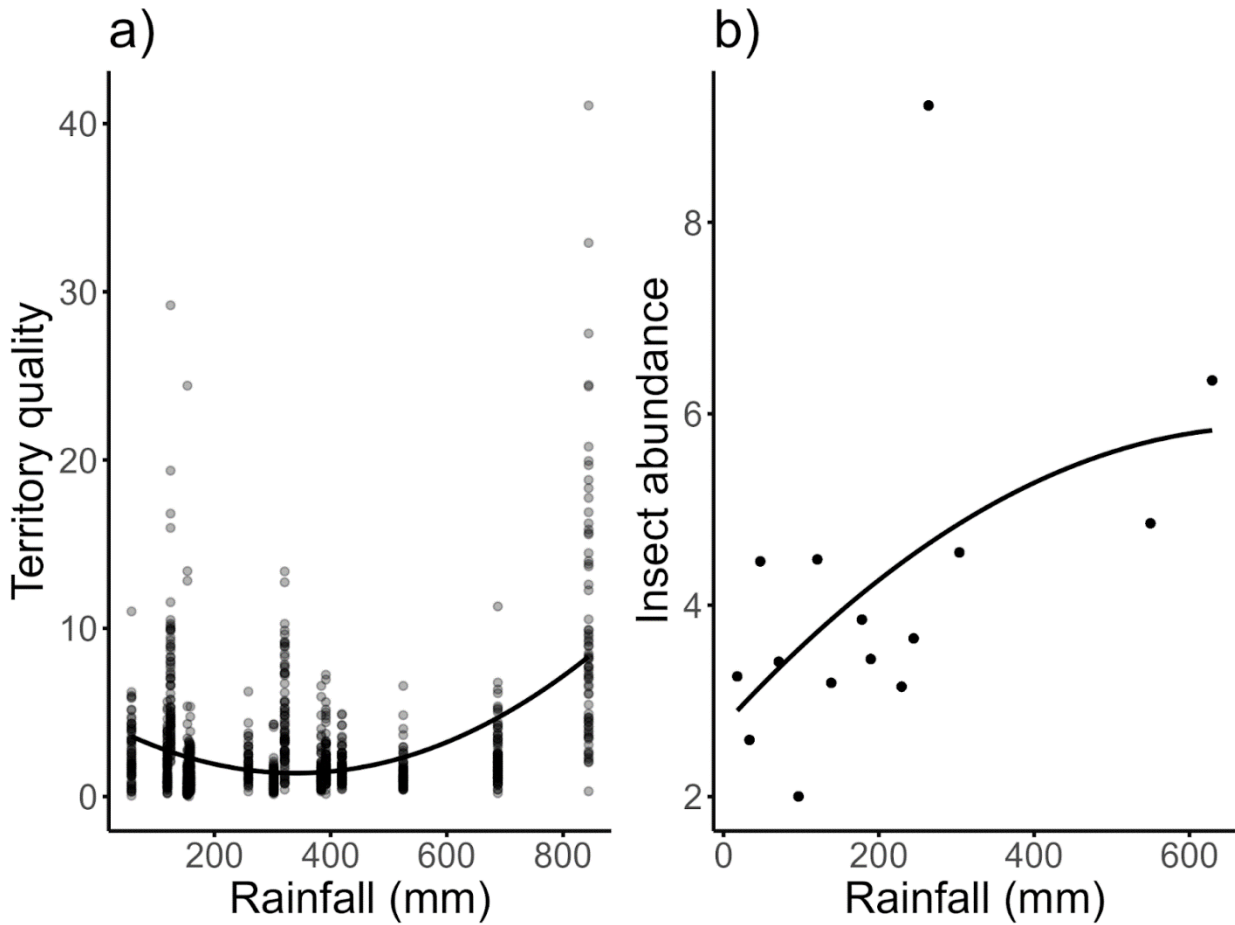
1026 **Table S13.** Associations between the probability of attempting to breed (model 1), producing a clutch  
 1027 (model 2), producing a fledgling (model 3), and the number of genetic fledglings surviving till at least  
 1028 three months old (model 4) in the Seychelles warbler on Cousin Island with rainfall, partnership length,  
 1029 pairwise relatedness, the number of helpers, male age, and female age.  $n = 1321$  partnerships were  
 1030 analyzed using binomial (models 1 to 3) and poisson (model 4) generalized linear mixed models.  
 1031 Significant terms are in bold.

	<b>Model 1</b>	<b>Model 2</b>	<b>Model 3</b>	<b>Model 4</b>
<b>Independent variables</b>	<b>Estimate (95% Confidence interval)</b>			
Intercept	<b>3.44***</b> (2.72 to 4.16)	<b>2.63***</b> (2.11 to 3.14)	<b>0.79***</b> (0.41 to 1.17)	<b>-0.54***</b> (-0.80 to -0.27)
Rainfall	<b>0.948**</b> (0.37 to 1.53)	<b>0.61**</b> (0.23 to 1.00)	<b>0.33**</b> (0.06 to 0.59)	<b>0.20*</b> (0.01 to 0.41)
Rainfall <sup>2</sup>	-	-	<b>-0.23*</b> (-0.44 to -0.05)	<b>-0.22**</b> (-0.38 to -0.06)
Partnership length	-0.16 (-0.50 to 0.19)	0.09 (-0.17 to 0.35)	0.14 (-0.07 to 0.35)	0.04 (-0.11 to 0.18)
Pairwise relatedness	0.00 (-0.24 to 0.24)	-0.03 (-0.21 to 0.14)	<b>-0.14*</b> (-0.28 to 0.00)	-0.08 (-0.17 to 0.01)
Number of helpers	-	<b>1.49***</b> (0.86 to 2.13)	<b>0.76***</b> (0.58 to 0.94)	<b>0.09*</b> (0.02 to 0.17)
Male age	<b>0.47*</b> (0.11 to 0.83)	0.10 (-0.12 to 0.33)	<b>0.22*</b> (0.01 to 0.43)	<b>0.21**</b> (0.06 to 0.35)
Male age <sup>2</sup>	<b>-0.23*</b> (-0.41 to -0.05)	-	<b>-0.19**</b> (-0.31 to -0.08)	<b>-0.16***</b> (-0.25 to -0.07)
Female age	0.26 (-0.03 to 0.55)	<b>0.37**</b> (0.14 to 0.61)	0.16 (-0.04 to 0.36)	0.03 (-0.12 to 0.17)
Female age <sup>2</sup>	-	<b>-0.30***</b> (-0.44 to -0.17)	<b>-0.28***</b> (-0.39 to -0.16)	<b>-0.21***</b> (-0.32 to -0.12)
<b>Random effects (Levels)</b>	<b>Variance</b>	<b>Variance</b>	<b>Variance</b>	<b>Variance</b>
Male ID (416)	0.00	0.00	0.02	0.00
Female ID (293)	0.34	0.00	0.22	0.00
Field period ID (16)	1.03	0.51	0.21	0.12
Territory ID (158)	0.43	0.35	0.07	0.00

1032 \*:  $p$ -value < 0.05; \*\*:  $p$ -value = 0.001; \*\*\*:  $p$ -value < 0.001

1033 **Table S14.** Associations between the probability of producing a clutch in the Seychelles warbler on  
 1034 Cousin Island with total rainfall from February to August, length of partnership, the relatedness of the  
 1035 breeding pair, the number of helpers, male age, and female age.  $n = 1321$  partnerships were analyzed  
 1036 using a binomial generalized linear mixed model. Significant  $p$ -values are in bold. The non-significant  
 1037 quadratic term of rainfall is included.

<b>Independent variables</b>	<b>Estimate</b>	<b>Standard error</b>	<b>95% Confidence interval</b>	<b><math>p</math>-value</b>
Intercept	2.899	0.294	2.322 to 3.476	<b>&lt;0.001</b>
Rainfall	0.682	0.182	0.325 to 1.038	<b>&lt;0.001</b>
Rainfall <sup>2</sup>	-0.264	0.136	-0.530 to 0.002	0.051
Partnership length	0.084	0.134	-0.178 to 0.347	0.528
Pairwise relatedness	-0.034	0.088	-0.206 to 0.138	0.702
Number of helpers	1.499	0.322	0.868 to 2.131	<b>&lt;0.001</b>
Male age	0.100	0.114	-0.123 to 0.324	0.378
Female age	0.373	0.121	0.135 to 0.611	0.002
Female age <sup>2</sup>	-0.303	0.069	-0.438 to -0.167	<b>&lt;0.001</b>
<b>Random effects</b>	<b>Variance</b>	<b>Levels</b>		
Male ID	<0.001	416		
Female ID	0.000	392		
Field Period ID	0.403	16		
Territory ID	0.355	158		



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1040 **Figure S6.** The effect of rainfall on: a) territory quality (scaled 1:10,000); b) insect abundance (the  
 1041 mean number of insects found per unit leaf area across all monthly surveys) on Cousin Island ( $n = 15$   
 1042 years), as predicted (solid line) by a *ClimWin* generated linear model. Territory quality was best  
 1043 predicted by rainfall from June to August, and insect abundance was best predicted by rainfall from  
 1044 July to August. Data are indicated as points where the shade of the points represents the sample size  
 1045 (darker represents more samples).

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