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Breeding near heterospecifics as a defence against brood parasites: can redstarts  
lower probability of cuckoo parasitism using neighbours?

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## 18 Abstract

19 Breeding habitat choice based on the attraction to other species can provide valuable social  
20 information and protection benefits. In birds, species with overlapping resources can be a cue of  
21 good quality habitats; species with shared predators and/or brood parasites can increase joint  
22 vigilance or cooperative mobbing, while raptors may provide a protective umbrella against these  
23 threats. We tested whether the migratory common redstart (*Phoenicurus phoenicurus*) is  
24 attracted to breed near active nests of the great tit (*Parus major*), a keystone-information source  
25 for migrant passerine birds, or a top predator, the northern goshawk (*Accipiter gentilis*). This  
26 system is unique to test these questions because the redstart is a regular host for the common  
27 cuckoo (*Cuculus canorus*). Therefore, we also evaluated other possible benefits coming from the  
28 heterospecific attraction, especially in terms of reducing brood parasitism risk. We monitored  
29 redstart occupancy rates, onset of breeding, reproductive investment, and followed nest  
30 outcomes in terms of brood parasitism, nest predation risk and overall reproductive success.  
31 Redstarts avoided breeding near goshawks, but showed neither attraction nor avoidance to breed  
32 next to great tits. Both neighbours neither reduced brood parasitism risk nor affected overall  
33 nesting success in redstarts. Redstarts may not use heterospecific attraction for settlement  
34 decisions, as associations with other species can only exist when some benefits are gained. Thus,  
35 environmental cues may be more important than social information for redstarts when breeding  
36 habitat choice. Other front-line defence strategies may have a better impact reducing breeding  
37 negative interactions, such brood parasitism.

38

39    Keywords: nest-site choice, frontline defence, heterospecific attraction, reproductive success,  
40    interspecific interaction.

## Introduction

Breeding habitat selection is used to enhance reproductive outcomes. Occurring at the start of the breeding cycle, these decisions determine the breeding site and have a large influence on the likelihood of offspring survival (Rosenzweig 1981; Reynolds 1996; Mayor et al. 2009; Chalfoun and Schmidt 2012; Boualit et al. 2019). Adaptive location decisions should account for negative interactions (i.e., predation, brood parasitism and competition), food availability, and other environmental factors (Banks and Beebee 1987; Martin 1993, 1995; Morris 2003). Assessment is crucial as each location may present different risks and benefits (Goodenough et al. 2009; Lehtonen et al. 2013). However, because personal assessment is costly, the number of possible sites that can be sampled is limited (Doligez et al. 2003; Danchin et al. 2004; Nocera et al. 2009). One strategy to overcome this dilemma, given the spatial and temporal variation in habitat quality (Mayor et al. 2009), could be to use information about the habitat quality based on the choices of other individuals, coined social information use (Reed et al. 1999; Doligez et al. 2002; Danchin et al. 2004).

Social information can be gathered from conspecific or heterospecific individuals (Mönkkönen et al. 1990; Doligez et al. 2003; Parejo et al. 2005; Seppänen et al. 2007). Settlement decisions can be based on the presence or absence of others and their breeding success (Valone 2007; Forsman and Seppänen 2011; Pizzatto et al. 2016). Conspecifics should provide relevant information due to a shared niche (Danchin et al. 2004; Loukola et al. 2012), however, for the same reason, associating with or copying conspecifics will likely carry fitness costs associated with competition (Seppänen

et al. 2007; Szymkowiak et al. 2016). To decrease such costs, the use of heterospecifics with an overlapping yet not identical niche, should sometimes be preferred over conspecifics as sources of information (Parejo et al. 2005; Seppänen et al. 2007; Szymkowiak 2013). Moreover, direct information about the success of conspecifics is not available for the current breeding attempt, making heterospecifics that settle earlier an important information source (Seppänen et al. 2007; Tolvanen et al. 2020).

Attraction to, or associations with other species could also provide defence or protection benefits (Quinn and Ueta 2008, see also heterospecific flocks, Morse 1977; Colorado 2013). Heterospecific attraction should indirectly benefit the information user when the information source selects habitats with lower predator or brood parasite abundance. Direct benefits such as increased shared vigilance or cooperative mobbing can come from less-aggressive species (Hurd 1996; Forsman et al. 1998; Polak 2014), while more aggressive or top predator species may provide a protective umbrella effect for heterospecifics settling nearby (Quinn and Ueta 2008). For example, raptors (or “aggressive” species) may prey on potential nest predators thereby decreasing nest predation risk of neighbouring birds (e.g., Marti et al. 1993; Blanco and Tella 1997; Ueta 2007). However, trade-offs need to be considered, as raptors may prey on nestlings and adult birds (Ueta 1999; Quinn and Kokorev 2002; Mönkkönen et al. 2007). Even though the use of heterospecific as a source of information, and the attraction to predators/aggressive species for protective associations are well supported, very few empirical tests have considered them in the brood parasitism context (e.g., Grim 2008).

The evolutionary arms-race between host and brood parasites has produced several host strategies to combat brood parasitism (Davies 2000; Krüger 2007). Recently research has been focused on “front-line” strategies (i.e. defences before parasite egg-laying, Feeney et al. 2012), such as nest-site choice (Forsman and Martin 2009; Tolvanen et al. 2017a; Expósito-Granados et al. 2017). Previous studies have found that host nests far from neighbours appear more likely to suffer from brood parasitism (Brown and Lawes 2007; Feeney et al. 2012; Ma et al. 2018b; but see Kelly et al. 2019). Conspecific neighbours can be attracted by alarm calls, deterring cuckoos that avoid being mobbed by a larger group of individuals (Feeney et al. 2013; Ma et al. 2018a). Moreover, by breeding near to heterospecific host species, birds can reduce brood parasitism rates (Clark and Robertson 1979). However, whether non-host heterospecific neighbours can deter brood parasites and decrease the probability of hosts being parasitized remains to be explored.

In northern boreal forests, migrant songbirds that face short breeding seasons have been shown to select breeding locations near to both resident tits and breeding raptors to gain various breeding benefits (Mönkkönen and Forsman 2002; Thomson et al. 2003, 2006; Mönkkönen et al. 2007). The redstart (*Phoenicurus phoenicurus*) is part of this migrant forest bird community and is a regular, cavity-nesting host for the cuckoo (*Cuculus canorus*). In Finland, redstarts suffer relatively high brood parasitism rates (20% - 32%, Rutila et al., 2002; Samaš et al., 2016; Thomson et al., 2016), yet show weak egg-stage anti-parasitic strategies (Avilés et al. 2005; Grim et al. 2009; Thomson et al. 2016). Redstarts may evade cuckoo parasitism through front-line defences, for example, choosing nest-sites inaccessible for the cuckoo (Moreras et al. 2021). However, other

front-line strategies are largely unexplored (but see, Tolvanen et al. 2017a), as is also the case for many other brood parasite-host systems with apparent lack of host defence strategies.

Redstarts can be attracted to heterospecifics with overlapping niche and to protective raptors to get breeding benefits. Here we used the great tit (*Parus major*) and northern goshawk (*Accipiter gentilis*) as neighbours. Great tits can serve as an information source of good quality breeding habitats to redstarts. As residents and cavity nesters, great tits have more time to select where to breed (e.g., Forsman et al. 2002, Thomson et al. 2003), they also overlap in resources used during breeding (Van Balen et al. 1982; Valtonen et al. 2017; Baroni et al. 2020) and share both adult and nest predators (e.g., sparrowhawks, *Accipiter nisus*, woodpeckers, small rodents and mustelids) with redstarts (Nilsson 1984; Gotmark and Post 1996; Wesołowski 2002; personal observation). On the other hand, goshawks should act as a protective umbrella from predators and possibly cuckoos (Mönkkönen et al. 2007; Tornberg et al. 2015). It is known that occasionally goshawks prey upon cuckoos (Toyne 1998; Nielsen and Drachmann 1999; Rebollo et al. 2017, R. Tornberg, unpublished data), potentially making cuckoos avoid the proximity to active goshawk nests.

Great tits can plausibly inform redstarts about habitats with low cuckoo density. Great tits tend to avoid places where they have seen cuckoos (Davies and Welbergen 2008) and may also avoid breeding in habitats with high cuckoo density given that cuckoos have a hawk-like appearance resembling the sparrowhawk (Gluckman and Mundy 2013). Parids generally are attracted by others' alarm calls (Sieving et al. 2004), and may mob cuckoos, either due to their hawk-like appearance (Gluckman and Mundy 2013) or due to their potential history as a past host of the



cuckoo (Grim et al. 2014; Liang et al. 2016). Although great tits are currently believed to be a non-suitable host for the cuckoo, this might be biased information, because almost all knowledge about great tits comes from nest-box populations, where only nest-boxes with small entrances (ca. 3 cm in diameter) that prevent cuckoo parasitism are used (Grim et al. 2014; Moreras et al. 2021). Indeed, great tits can be naturally parasitized in cavities with larger entrances (e.g. 5 to 7 cm in diameter, woodpecker cavities, Grim et al. 2014). Moreover, some great tit populations still show high egg rejection rates (Liang et al. 2016), suggesting a co-evolutionary history with cuckoos.

In this study, we tested heterospecific attraction in redstarts, its impact on the breeding cycle (e.g., timing of breeding, nest predation rates, reproductive investment, and reproductive success) and its possible influence as a front-line defence against brood parasitism. If heterospecific attraction exists, we expect redstarts to preferentially nest in the neighbourhood of great tits, as well as, within goshawks territories. Moreover, for heterospecific attraction as a strategy should be adaptive, therefore, we expect benefits to redstarts in terms of earlier breeding time, larger clutch size, number of nestlings and lower predation rates in redstart having great tits and goshawks as neighbours. Finally, if the heterospecific attraction acts as an anti-parasite front-line defence, we expect redstarts breeding close to great tits to have lower parasitism rates, and the likelihood of cuckoo parasitism to increase as the distance from the goshawk nest increases.

## Materials and methods

### *Study site and basic protocol*

The study was conducted near Oulu, Finland (65°N, 25° 50' E), in scots pine (*Pinus sylvestris*) dominated forest patches suitable for redstarts, approximately 60 km<sup>2</sup> in area combined. We placed nest-boxes (approximately 400 permanent nest-boxes), which were monitored each breeding season by checking redstart occupancy every three to five days. We placed nest-boxes in pines approximately 1.5 m above the ground and 100–220 m apart since 2011. All nest-boxes had the same dimensions: 17.5 x 17.5 x 28 cm (width, depth and height), and an entrance hole diameter of 7 cm. We recorded the laying date of the first egg, events of brood parasitism and nest predation, and clutch size. Once the egg-laying was completed, nests were monitored once a week, and partly protected from nest predation by placing wire cages over the entrance of nest-boxes in early incubation (however, small predators can still enter the nest-box, Thomson et al. 2016). This was done to ensure that enough redstarts and cuckoos survive to make other concurrent studies possible. Therefore, for protected nests we only considered predation taking place during the *ca.* 7- 10 day laying and early incubation period (after which the nests were protected), underestimating the predation rates for the entire nesting period (*ca.* 35 days). Wire cages were only placed in nest-boxes with great tits as neighbours, but not in nest-boxes within goshawk territories (see below for details). For nests that were not protected (i.e., within goshawks territories), 29 predation events were documented in 416 redstart nests in 6 years. Nests that were predated before the fifth redstart egg was laid, were not considered for calculated parasitism rates, since it was impossible to determine if the nest was previously parasitized or not.

169 During the nestling period, we registered the number of nestlings. The number of fledglings was  
170 derived by subtracting from the nestling number any dead chicks found in the nest after fledging.  
171 A nest was defined as successful if at least one nestling fledged.

172

### 173 ***Nest-box experiment set-ups***

#### 174 *Great tit as neighbour*

175 Between 2014 and 2017, we selected 80 nest-boxes each year from the larger redstart nest-box  
176 population. These nest-boxes had 7 cm diameter entrance hole (hereafter “redstart box”), while  
177 nest-boxes with a 3 cm diameter entrance hole (hereafter “tit box”) were used for settling great  
178 tits. However, it is important to note that all species could freely choose any nest-box. We created  
179 pairs of redstart nest-boxes (40 pairs per year), where we randomly assigned one control and one  
180 neighbour nest-box. The ‘neighbour’ redstart boxes had one tit box placed approx. 25 m from the  
181 redstart box, and the ‘control’ redstart boxes did not have a tit box placed nearby (Figure 1). To  
182 control for habitat quality, the ‘neighbour’ and ‘control’ boxes of each pair were separated only  
183 by approx. 200 m, while the different neighbour-control pairs of redstart boxes were separated by  
184 at least 300 m. Between 2014 and 2015 the same 40 pairs of boxes were used, but treatments  
185 were swapped between years. Then we selected 40 different pairs of boxes and used them in 2016  
186 and 2017, where treatments were also swapped between years. We divided the study site into 10  
187 different patches to account for possible spatial differences in habitat. Of the 40 ‘neighbour’ boxes  
188 in each year, only those that at the time of redstart settlement had a great tit pair actively breeding  
189 in the tit box were considered in the analyses. “Control” boxes of the pairs where the “neighbour”

box was not considered were also excluded from the analysis. Finally, redstart boxes occupied by other species (great tits or pied flycatchers, *Ficedula hypoleuca*) were not considered. We excluded 54 pairs of nest-boxes.

#### *Goshawk as neighbour*

Between 2011 and 2018, we placed redstart boxes around the nest sites of 15 active and successfully breeding goshawk nests. A goshawk territory may include several nest sites, but only one is used each year. We used 9 different nest sites in 5 territories. As goshawks tend to breed in the same location over the years, some nest sites and territories were used for multiple years: three territories for two, five and six years and two nest sites for three and five years. Redstart boxes were placed at distances ranging from 20 to 2000 m from the active goshawk nest (distribution shown in Figure 2). The location of each nest box was fixed within the breeding season and between years. However, in some nest sites, the number of nest boxes varied as we added some redstart boxes in later years. Depending on the habitat suitability for redstart, we set up between 15 and 45 redstart boxes around each goshawk nest. The goshawk territories and nest sites have been followed as part of a long-term study on goshawk ecology (see Tornberg et al. 2009; Tolvanen et al. 2017b). Based on local goshawk ecology, all territories had different goshawk pairs, while within a territory the same pair was likely breeding for multiple years. In total, 7 out of the 15 goshawk breeding attempts were known or likely to involve unique goshawk pairs (at least one parent different). Redstart boxes were monitored as described above (see *Study site and basic protocol*). Note that both set-ups (i.e., great tit as neighbour and goshawk as neighbour) were in different areas and there was no overlap between them.

212

## 213 *Statistics*

214 All statistical analyses were conducted using R (version 3.6.2, R Development Core Team 2019).  
215 We built full models evaluating the heterospecific attraction of redstarts and its consequences  
216 with generalized linear mixed models (GLMMs) for great tit attraction and generalized additive  
217 mixed models (GAMMs) for goshawk attraction. Full models included all relevant explanatory  
218 variables (see below for details). Then we searched for a parsimonious final model sets by fitting  
219 all subset models within the full model, without excluding the fixed explanatory variable of main  
220 interest: the neighbour presence (in the great tit attraction) or distance to the goshawk nest (in  
221 the goshawk attraction).

222

223 Instead of a single polynomial over the whole domain (i.e. transformation), one can rather use  
224 piecewise polynomial curves (i.e. a spline curve) in regions defined by knot using GAMM models  
225 (Wood 2017). The GAMM models are composed of a sum of smooth functions of covariates in  
226 addition to the standard linear covariate effects to capture the non-linear relationships using a  
227 basis function. A penalized estimation of the spline is used to find the best-fitting line, while  
228 penalizing more complex smoothers, ensuring that the model fit is not too complex. If the data  
229 does not support a non-linear trend the spline curve will reduce to a linear trend. In our case the  
230 distance effect in goshawk attraction was modelled as a non-linear cubic regression spline. For the  
231 cubic regression the x-axis (i.e., distance) is divided into a certain number of knots. In knot  
232 intervals, a cubic polynomial is fitted (this is a model of the form:  $Y_i = \alpha + \beta_1 \times X_i +$   
233  $\beta_2 \times X_i^2 + \beta_3 \times X_i^3$ ), and the fitted values per segment are then glued together to form the

smoothing curve using first-order and second-order derivatives (Zuur et al. 2009; Perperoglou et al. 2019).

Firstly, we evaluated neighbour attraction for redstarts by testing differences in probability of redstart occupancy (yes/no) using a binomial GLMM/GAMM with logit link function. For the goshawk as neighbour analysis, as some nest boxes were added in later years for some nest sites, we added as a fixed effect a binary “new box” variable indicating whether the nest box was set up in the beginning of the season (new box = yes) or had already been available in earlier year(s) (new box = no). Then possible benefits gained from this attraction (i.e. lower brood parasitism and nest predation rates, earlier laying date, higher clutch size and fledgling number, and higher success) were also evaluated. We tested parasitism and predation rates using binomial GLMMs/GAMMs with logit link function, with the laying date included as a fixed effect in the full model to account for possible variation due to timing of breeding. However, for the great tit as neighbour, predation events were too few for a statistical analysis (9 events in total: 4 cases registered in restart nests with neighbours and 5 cases registered in nests without neighbour). We also compared laying date, clutch size and number of fledglings, using Poisson GLMMs/GAMMs with log link function. Finally, we compared the probability of success using binomial GLMMs/GAMMs with logit link function, with the laying date and probability of being parasitized as fixed effects. Full models of all response variables in the great tit as neighbours’ analysis included the interaction of great tit presence and the year the data was collected to account for possible weather and other conditions that vary over the years and might contribute to the nesting choice made by redstarts. For the goshawk as neighbour analysis, full models included year as a random-intercept effect.

256 Additionally, to account for potential spatial variation in the responses and for using the same  
257 nest-boxes multiple times, for the great tit as neighbour analysis we tested random-intercept  
258 effects of the forest patch where the boxes were placed (10 patches) and nest-box ID, but we used  
259 only the one better supported (i.e. the model with the lower AICc). While for the goshawk as  
260 neighbour analysis, we tested random-intercept effects of goshawk territory ID and nest site ID,  
261 but we used only the one that was better supported (i.e. the model with the lower AICc).

262

263 For the model selection process, we ranked the models using the AICc criteria. We then followed  
264 Richards et al. (2011) and defined final model sets including the models within 6 AICc units, but  
265 excluding models that were more complex versions of a model with lower AICc (Table 1). If there  
266 were more than one model in the final model set, we focused on the best-ranking one but also  
267 note if the inferences based on the other model(s) in the final model set differ from the best one.  
268 We based our statistical inferences on the parameter mean estimates and their 95% confidence  
269 intervals. Collinearity between explanatory variables was acceptable in all models (variance  
270 inflation factors, VIF < 3 in all cases; Zuur et al. 2010). Finally, we graphically checked that the  
271 models did not violate other model assumptions using DHARMA package (Hartig 2018).

## Results

### *Great tit as neighbour*

Around half (51.2%) of the paired “neighbour-control” boxes were occupied by redstarts. Out of 160 pairs available during the four years of the study (2014-2017), 82 pairs of boxes were included in the analyses, and 77 redstart nests were recorded within these. Out of the 82 control boxes 38 were occupied by redstarts (46.3%) and out of the 82 neighbour boxes 39 were occupied by redstarts (47.6%, Fisher’s Exact test:  $p = 1.0$ ). For all the breeding parameters evaluated (i.e., occupancy, laying date, clutch size, number of fledglings, brood parasitism and success rates), the final model set included only one model (Table 1). Moreover, for most breeding parameters the final model included as random intercept effect the box ID, but for the occupation probability the forest patch ID was better supported than the box ID. Redstarts did not nest more frequently near to great tit nests (Table 2). However, we found differences in redstart occupancy rates between years (Table 2).

The mean laying date of the first egg was around the 30<sup>th</sup> of May (mean =  $30.1 \pm 6.6$  days,  $n=77$ ). Most nests produced at least one fledgling with a success rate of 76.1% ( $n=67$ ). We recorded 19 nests parasitized (27.1%), 12 cases in nests with neighbours (17.1%) and 7 cases in nests without neighbour (10.0%). Only 9 cases of predation were recorded during the study (11.7%), 4 cases in restart nests with neighbours (5.2%) and 5 cases in nests without neighbour (6.5%). This made it impossible to test differences in predation rates between treatments. None of the redstart



breeding parameters tested (laying date, parasitism risk, clutch size, number of fledglings or success rate) show any clear statistical trend whether great tits were neighbours or not (Table 2).

#### *Goshawk as neighbour*

Occupancy of redstarts varied between 13.6 % and 75% of the nest-boxes in the 5 different active goshawk territories. In total 175 out of 487 nest boxes were occupied during the 7 years we collected data (35.9%). Only one model was selected in the final model set for the occupancy (Table 1). This model included the distance effect, the variable describing if the nest-box was a new box or not, and the goshawk territory as a random intercept term (Table 1). Redstart occupancy increased linearly with the distance from the goshawk nest (Figure 3A). Moreover, nest-boxes placed in the beginning of the current season (i.e. new boxes), were more likely to be occupied by redstarts (Table 3).

Similar to great tit as neighbour data, the mean laying date of the first egg was around 30<sup>th</sup> of May (mean= 29.7 ± 8.7 days, n=174). Nest predation rate was 21.9% (n=160), while the brood parasitism rate was 22.9% (n=157) for redstarts with goshawks as neighbours. Therefore, more than half of redstart nests fledged at least one chick (success rate = 63.2%, n=155). For laying date, nest predation rate, brood parasitism rate and success rate, the final model set consisted of only one model including only the distance effect (Table 1). For clutch size and number of fledglings the final model set included two models, with the best model including the distance effect and laying date (Table 1). Moreover, for all breeding responses the best models included as random factor the nest ID (Table 1). For nest predation, the model included a non-linear smoother with

316 predation rate being lowest between approximately 800 m and 1400 m from the goshawk nest  
317 (Figure 3C), however the smoother was overall not statistically significant (Table 2). For parasitism  
318 rate there was no clear trend (Figure 3B). The laying date had a complex pattern, where females  
319 settling near to goshawk nest (200-800 m) tended to lay later than females settling far (> 800 m)  
320 from it (Figure 4A), showing a difference around 5 days between them (Figure 4A). The clutch size,  
321 the number of fledglings and nest success did not differ relative to the distance from the goshawk  
322 nest (Figure 4B, 4C, 3D).

## Discussion

We found no evidence of heterospecific attraction by redstarts to active nests of resident tits and predatory goshawks. Our results suggest that redstart nest site decisions are not obviously impacted by these species on a breeding territory scale, which is perhaps surprising given that resident tits and goshawks are regularly used in a similar context by other passerine species as cues of good quality habitat and/or safety (Forsman et al. 1998, 2002; Thomson et al. 2003; Mönkkönen et al. 2007; Quinn and Ueta 2008; Polak 2014). By tracking the consequences of redstart nest site decisions, our findings also showed that great tits and goshawks did not offer any protection from brood parasitism or nest predation risk. Other possible breeding benefits (i.e. earlier laying date, higher clutch size and fledgling number, and higher success) did not differ for redstart nests at different distances from these species, suggesting that redstarts do not use heterospecific attraction in breeding habitat decisions.

### *Great tit as neighbour*

Breeding redstarts did not associate with nesting great tits, even when social information from these residents has proven to attract other nesting migrants (e.g., pied flycatcher and chaffinch, *Fringilla coelebs*) breeding in the deciduous forests (Forsman et al. 2002, Thomson et al. 2003). Social information about the quality of the environment has previously been shown to be important to migrants with limited time to sample the area (Doligez et al. 2002, 2003; Mönkkönen and Forsman 2002; Danchin et al. 2004; Nocera et al. 2009). However, redstarts did not avoid breeding close to great tits either, suggesting low competitive or other costly interactions between

both species (but see, Ahola et al. 2007; Samplonius and Both 2019, to see competition between great tits and migrants). The breeding outcome of redstart nests supported this conclusion; redstarts showed no clear benefits nor costs of breeding next to great tits in terms of their reproductive investment and outcomes. Therefore, patches with or without great tits as neighbours may represent the same quality for redstarts even when our results show variation in occupancy rates between habitat patches.

Redstarts with great tit neighbours showed similar parasitism rates as those breeding alone. This suggests that great tits do not provide either indirect (i.e. habitats with lower cuckoo densities) nor direct defence (i.e cooperative vigilance and mobbing) against brood parasitism. However, as previously found in yellow warblers (*Setophaga petechia*), by breeding in habitats with lower brood parasite densities, birds may not ensure habitats with lower parasitism rates (Kelly et al. 2019). On the other hand, the direct defence of passerine heterospecific neighbours may vary depending on the risk (Forsman and Mönkkönen 2001). The hawk appearance of cuckoos may prevent great tits from responding to redstart alarm calls or from participating in cooperative nest defence, as they may not distinguish between cuckoos and hawks (Davies and Welbergen 2008; Welbergen and Davies 2011), considering cuckoos as a real mortality threat.

#### *Goshawk as neighbour*

Redstarts tended to breed far from active goshawk nests, even when goshawks represent little risk for small passerines (Solonen et al. 2019). Redstarts may fail to discriminate between different hawk species; a recent study suggested that given the fact that male goshawks closely resemble

female Eurasian sparrowhawks (*Accipiter nisus*, which preys on small birds) at community level small bird species avoid breeding near to goshawks (Burgas et al. 2021). We also found no support for the idea that goshawk may provide protection against nest predators (Mönkkönen et al. 2007). Given that nest predation risk was low overall in our study (18.6%, and completely absent in 2016), small birds may not benefit from associating with goshawks, but redstarts may encounter a negative outcome due to the attraction of mesopredators (Morosinotto et al. 2012). This is supported by a higher nest predation rate within goshawks territories than in habitats with great tits as neighbours (21.9% for goshawks as neighbours and 11.7% for great tits as neighbour). Moreover, redstart clutch size, number of fledglings and success rate did not change relative to the distance to the goshawk nest. Laying initiation date had a very non-linear pattern relative to the distance to goshawk nest, but it was not clearly earlier or later closer to goshawk nest. The lack of benefits of breeding near to goshawks may explain the lack of a positive association (Quinn and Kokorev 2002; Lima 2009). Nevertheless, it is important to highlight that the suitability of the habitat near to active goshawk nests was likely on average not as good for breeding redstarts as the habitat further away, potentially affecting the nest-site choice. All in all, the possible negative trade-offs of breeding near to goshawks support the avoidance pattern we found for redstarts, rather than the random occupancy pattern shown previously by pied flycatchers (Mönkkönen et al. 2007).

Redstarts breeding within goshawk territories did not experience a decrease in brood parasitism risk, despite that goshawks prey on cuckoos (Toyne 1998; Nielsen and Drachmann 1999; Rebollo et al. 2017, R. Tornberg, unpublished data). It is possible that redstarts breeding close to goshawks

would encounter goshawks more often and may become less attentive towards hawk-like birds such as cuckoos; similar to the mechanism proposed in the risk allocation hypothesis (see Lima and Bednekoff 1999; Thomson et al. 2011). This may facilitate brood parasitism events within goshawks territories and possibly counteracts any goshawk protection benefits, as cuckoos willing to seek host nests close within goshawk territories are not going to be perceived as a threat by redstarts. However, many host species discriminate cuckoos from hawks (Trnka and Prokop 2012; Trnka and Grim 2013; Li et al. 2015; Ma et al. 2018a), therefore, further research on redstart discrimination of cuckoos is needed.

#### *Brood parasitism and heterospecific neighbours*

Redstarts did not use heterospecific attraction to counteract brood parasitism. Neither aggressive neighbours (i.e., goshawks) nor other passerine species (i.e., great tits) seem to contribute to reduce brood parasitism risk. Other frontline defence strategies may be preferred for redstarts to reduce brood parasitism. There has been speculation that redstarts moved from ground-cavity to tree-cavity breeding due to cuckoo parasitism pressure (Avilés et al. 2005), and therefore, that redstarts could face lower selection pressure to evolve other defence strategies. Cavity-nesting in redstarts results in high rates of mislaid cuckoo eggs (around 70%, Samaš et al. 2016; Thomson et al. 2016), reducing successful brood parasitism risk. A recent study also found that cavity choice based on the entrance size is an adaptative strategy redstarts may use to prevent cuckoos to access the nest cup (Moreras et al. 2021). Other frontline strategies such as nest placement, breeding phenology and nest defence may be useful to reduce parasitism risk (see Feeney et al. 2012), yet, these have received little empirical research. Heterospecific attraction as a strategy to

411 reduce brood parasitism should be tested in other brood parasite-host systems, especially after  
412 considering that conspecific attraction can reduce brood parasitism risk (Brown and Lawes 2007;  
413 Feeney et al. 2012; Ma et al. 2018b; but see Kelly et al. 2019) and that communities can impact  
414 the life history of the host, and therefore, the development of adaptations.

415

416 In conclusion, redstarts do not use great tits or goshawks as part of a frontline defence of nest  
417 choice to prevent cuckoo brood parasitism. Redstarts seem to avoid breeding within goshawk  
418 territories, while they do not avoid nor are attracted to great tits. There were no apparent  
419 breeding benefits of having either as a heterospecific neighbour, and therefore, no need to be  
420 attracted to breed near heterospecifics. Other frontline strategies, such as nest attributes (i.e.,  
421 safer entrance size and cup location within the cavity for redstarts), and self-nest defence may  
422 constitute better trade-offs for redstarts. Further research on such frontline strategies is needed  
423 to better understand brood parasite-host co-evolution, especially in brood parasite-host systems  
424 where it has been thought that hosts possess no defence strategies.

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## 434 **Conflicts of interest/Competing interests**

435 The authors declare that they have no conflict of interest.

## 436 **Ethics approval**

437 All animal experiments were approved by the Finnish Center for Economics Development,  
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## References

- Ahola MP, Laaksonen T, Eeva T, Lehikoinen E (2007) Climate change can alter competitive relationships between resident and migratory birds. *J Anim Ecol* 76:1045–1052. <https://doi.org/10.1111/j.1365-2656.2007.01294.x>
- Avilés JM, Rutila J, Møller AP (2005) Should the redstart *Phoenicurus phoenicurus* accept or reject cuckoo *Cuculus canorus* eggs? *Behav Ecol Sociobiol* 58:608–617. <https://doi.org/10.1007/s00265-005-0941-7>
- Banks B, Beebee TJC (1987) Factors influencing breeding site choice by the pioneering amphibian *Bufo calamita*. *Ecography (Cop)* 10:14–21. <https://doi.org/10.1111/j.1600-0587.1987.tb00733.x>
- Baroni D, Korpimäki E, Selonen V, Laaksonen T (2020) Tree cavity abundance and beyond: Nesting and food storing sites of the pygmy owl in managed boreal forests. *For Ecol Manage* 460:. <https://doi.org/10.1016/j.foreco.2019.117818>
- Blanco G, Tella JL (1997) Protective association and breeding advantages of choughs nesting in lesser kestrel colonies. *Anim Behav* 54:335–342. <https://doi.org/10.1006/anbe.1996.0465>
- Boualit L, Pichenot J, Besnard A, et al (2019) Environmentally mediated reproductive success predicts breeding dispersal decisions in an early successional amphibian. *Anim Behav* 149:107–120. <https://doi.org/10.1016/j.anbehav.2019.01.008>
- Brown M, Lawes MJ (2007) Colony size and nest density predict the likelihood of parasitism in the colonial Southern Red Bishop *Euplectes orix* - Diderick Cuckoo *Chrysococcyx caprius* system. *Ibis (Lond 1859)* 149:321–327. <https://doi.org/10.1111/j.1474-919X.2006.00633.x>
- Burgas D, Ovaskainen O, Blanchet FG, Byholm P (2021) The Ghost of the Hawk: Top Predator Shaping Bird Communities in Space and Time. *Front Ecol Evol* 9:293. <https://doi.org/10.3389/fevo.2021.638039>
- Chalfoun AD, Schmidt KA (2012) Adaptive breeding-habitat selection: Is it for the birds? *Auk* 129:589–599. <https://doi.org/10.1525/auk.2012.129.4.589>
- Clark KL, Robertson RJ (1979) Spatial and temporal multi-species nesting aggregations in birds as anti-parasite and anti-predator defenses. *Behav Ecol Sociobiol* 5:359–371. <https://doi.org/10.1007/BF00292524>
- Colorado GJ (2013) Why animals come together, with the special case of mixed-species bird flocks. *Rev EIA* 10:49–66
- Danchin E, Giraldeau L-A, Valone TJ, Wagner RH (2004) Public Information: From Nosy Neighbors to Cultural Evolution. *Science (80- )* 305:487–491. <https://doi.org/10.1126/science.1098254>
- Davies NB (2000) Cuckoos, Cowbirds and Other Cheats. Poyser, London, UK
- Davies NB, Welbergen JA (2008) Cuckoo-hawk mimicry? An experimental test. *Proc R Soc B Biol Sci* 275:1817–1822. <https://doi.org/10.1098/rspb.2008.0331>
- Doligez B, Cadet C, Danchin E, Boulinier T (2003) When to use public information for breeding habitat selection? The role of environmental predictability and density dependence. *Anim Behav* 66:973–988. <https://doi.org/10.1006/anbe.2002.2270>

486 Doligez B, Danchin E, Clobert J (2002) Public information and breeding habitat selection in a wild bird  
487 population. *Science* (80- ) 297:1168–1170. <https://doi.org/10.1126/science.1072838>

488 Expósito-Granados M, Parejo D, Martínez JG, et al (2017) Host nest site choice depends on risk of cuckoo  
489 parasitism in magpie hosts. *Behav Ecol* 28:1492–1497. <https://doi.org/10.1093/beheco/arx113>

490 Feeney WE, Medina I, Somveille M, et al (2013) Brood parasitism and the evolution of cooperative  
491 breeding in birds. *Science* (80- ) 342:1506–1508. <https://doi.org/10.1126/science.1240039>

492 Feeney WE, Welbergen JA, Langmore NE (2012) The frontline of avian brood parasite-host coevolution.  
493 *Anim Behav* 84:3–12. <https://doi.org/10.1016/j.anbehav.2012.04.011>

494 Forsman JT, Martin TE (2009) Habitat selection for parasite-free space by hosts of parasitic cowbirds.  
495 *Oikos* 118:464–470. <https://doi.org/10.1111/j.1600-0706.2008.17000.x>

496 Forsman JT, Mönkkönen M (2001) Responses by breeding birds to heterospecific song and mobbing call  
497 playbacks under varying predation risk. *Anim Behav* 62:1067–1073.  
498 <https://doi.org/10.1006/anbe.2001.1856>

499 Forsman JT, Mönkkönen M, Helle P, Inkeröinen J (1998) Heterospecific attraction and food resources in  
500 migrants' breeding patch selection in northern boreal forest. *Oecologia* 115:278–286.  
501 <https://doi.org/10.1007/s004420050517>

502 Forsman JT, Seppänen JT (2011) Learning what (not) to do: Testing rejection and copying of simulated  
503 heterospecific behavioural traits. *Anim Behav* 81:879–883.  
504 <https://doi.org/10.1016/j.anbehav.2011.01.029>

505 Forsman JT, Seppänen JT, Mönkkönen M (2002) Positive fitness consequences of interspecific  
506 interaction with a potential competitor. *Proc R Soc B Biol Sci* 269:1619–1623.  
507 <https://doi.org/10.1098/rspb.2002.2065>

508 Gluckman TL, Mundy NI (2013) Cuckoos in raptors' clothing: Barred plumage illuminates a fundamental  
509 principle of Batesian mimicry. *Anim Behav* 86:1165–1181.  
510 <https://doi.org/10.1016/j.anbehav.2013.09.020>

511 Goodenough AE, Elliot SL, Hart AG (2009) Are nest sites actively chosen? Testing a common assumption  
512 for three non-resource limited birds. *Acta Oecologica* 35:598–602.  
513 <https://doi.org/10.1016/j.actao.2009.05.003>

514 Gotmark F, Post P (1996) Prey selection by sparrowhawks, *Accipiter nisus*: relative predation risk for  
515 breeding passerine birds in relation to their size, ecology and behaviour. *Philos Trans R Soc London*  
516 *Ser B Biol Sci* 351:1559–1577. <https://doi.org/10.1098/RSTB.1996.0141>

517 Grim T (2008) Are Blackcaps (*Sylvia atricapilla*) defending their nests also calling for help from their  
518 neighbours? *J Ornithol* 149:169–180. <https://doi.org/10.1007/s10336-007-0257-7>

519 Grim T, Rutila J, Cassey P, Hauber ME (2009) The cost of virulence: an experimental study of egg eviction  
520 by brood parasitic chicks. *Behav Ecol* 20:1138–1146. <https://doi.org/10.1093/beheco/arp108>

521 Grim T, Samaš P, Procházka P, Rutila J (2014) Are tits really unsuitable hosts for the common Cuckoo?  
522 *Ornis Fenn* 91:166–177

523 Hartig F (2018) DHARMA: Residual Diagnostics for Hierarchical (Multi-Level / Mixed) Regression Models.

524 R package version 0.2.0. <https://CRAN.R-project.org/package=DHARMA>

525 Hurd CR (1996) Interspecific attraction to the mobbing calls of black-capped chickadees (*Parus*  
526 *atricapillus*). *Behav Ecol Sociobiol* 38:287–292. <https://doi.org/10.1007/s002650050244>

527 Kelly JK, Suckow NM, Ward MP (2019) Preferential settling at sites with higher conspecific density does  
528 not protect Yellow Warblers (*Setophaga petechia*) from brood parasitism. *Acta Oecologica* 96:24–  
529 28. <https://doi.org/10.1016/j.actao.2019.03.003>

530 Korpimäki E (1985) Diet of the Kestrel *Falco tinnunculus* in the breeding season. *Ornis Fenn* 62:130–135

531 Krüger O (2007) Cuckoos, cowbirds and hosts: adaptations, trade-offs and constraints. *Philos Trans R Soc*  
532 *B Biol Sci* 362:1873–1886. <https://doi.org/10.1098/rstb.2006.1849>

533 Lehtonen TK, Lindström K, Wong BBM (2013) Effect of egg predator on nest choice and nest  
534 construction in sand gobies. *Anim Behav* 86:867–871.  
535 <https://doi.org/10.1016/j.anbehav.2013.08.005>

536 Li D, Wei H, Zhang Z, et al (2015) Oriental reed warbler (*Acrocephalus orientalis*) nest defence behaviour  
537 towards brood parasites and nest predators. *Behaviour* 152:1601–1621.  
538 <https://doi.org/10.1163/1568539X-00003295>

539 Liang W, Møller AP, Stokke BG, et al (2016) Geographic variation in egg ejection rate by great tits across  
540 2 continents. *Behav Ecol* 27:1405–1412. <https://doi.org/10.1093/beheco/arw061>

541 Lima SL (2009) Predators and the breeding bird: Behavioral and reproductive flexibility under the risk of  
542 predation. *Biol. Rev.* 84:485–513

543 Lima SL, Bednekoff PA (1999) Temporal variation in danger drives antipredator behavior: The predation  
544 risk allocation hypothesis. *Am Nat* 153:649–659. <https://doi.org/10.1086/303202>

545 Loukola OJ, Seppänen JT, Forsman JT (2012) Intraspecific social information use in the selection of nest  
546 site characteristics. *Anim Behav* 83:629–633. <https://doi.org/10.1016/j.anbehav.2011.12.004>

547 Ma L, Yang C, Liang W (2018a) Hawk mimicry does not reduce attacks of cuckoos by highly aggressive  
548 hosts. *Avian Res* 9:35. <https://doi.org/10.1186/s40657-018-0127-4>

549 Ma L, Yang C, Liu J, et al (2018b) Costs of breeding far away from neighbors: Isolated host nests are more  
550 vulnerable to cuckoo parasitism. *Behav Processes* 157:327–332.  
551 <https://doi.org/10.1016/j.beproc.2018.07.017>

552 Marti CD, Korpimäki E, Jakšić FM (1993) Trophic Structure of Raptor Communities: A Three-Continent  
553 Comparison and Synthesis. In: *Current Ornithology*. Springer US, Boston, MA, pp 47–137

554 Martin TE (1995) Avian life history evolution in relation to nest sites, nest predation, and food. *Ecol*  
555 *Monogr* 65:101–127. <https://doi.org/10.2307/2937160>

556 Martin TE (1993) Nest Predation and Nest Sites. *Bioscience* 43:523–532.  
557 <https://doi.org/10.2307/1311947>

558 Mayor SJ, Schneider DC, Schaefer JA, Mahoney SP (2009) Habitat selection at multiple scales. *Ecoscience*  
559 16:238–247. <https://doi.org/10.2980/16-2-3238>

560 Mönkkönen M, Forsman JT (2002) Heterospecific attraction among forest birds: a review. *Ornithol Sci*

1:41–51. <https://doi.org/10.2326/osj.1.41>

Mönkkönen M, Härdling R, Forsman JT, Tuomi J (1999) Evolution of heterospecific attraction: Using other species as cues in habitat selection. *Evol Ecol* 13:91–104. <https://doi.org/10.1023/a:1006590215306>

Mönkkönen M, Helle P, Soppela K (1990) Numerical and behavioural responses of migrant passerines to experimental manipulation of resident tits (*Parus* spp.): heterospecific attraction in northern breeding bird communities? *Oecologia* 85:218–225. <https://doi.org/10.1007/BF00319404>

Mönkkönen M, Husby M, Tornberg R, et al (2007) Predation as a landscape effect: The trading off by prey species between predation risks and protection benefits. *J Anim Ecol* 76:619–629. <https://doi.org/10.1111/j.1365-2656.2007.01233.x>

Moreras A, Tolvanen J, Morosinotto C, et al (2021) Choice of nest attributes as a frontline defense against brood parasitism. *Behav Ecol* 32:1285–1295. <https://doi.org/10.1093/beheco/arab095>

Morosinotto C, Thomson RL, Hänninen M, Korpimäki E (2012) Higher nest predation risk in association with a top predator: Mesopredator attraction? *Oecologia* 170:507–515. <https://doi.org/10.1007/s00442-012-2320-1>

Morris DW (2003) Toward an ecological synthesis: A case for habitat selection. *Oecologia* 136:1–13

Morse DH (1977) Feeding Behavior and Predator Avoidance in Heterospecific Groups. *Bioscience* 27:332–339. <https://doi.org/10.2307/1297632>

Nielsen JT, Drachmann J (1999) Prey selection of Goshawks *Accipiter gentilis* during the breeding season in Vendsyssel, Denmark. *Dansk Orn Foren Tidsskr* 93:85–90

Nilsson SG (1984) The evolution of nest-site selection among hole-nesting birds: the importance of nest predation and competition. *Ornis Scand* 15:167–175. <https://doi.org/10.2307/3675958>

Nocera JJ, Forbes GJ, Giraldeau LA (2009) Aggregations from using inadvertent social information: A form of ideal habitat selection. *Ecography (Cop)* 32:143–152. <https://doi.org/10.1111/j.1600-0587.2008.05614.x>

Parejo D, Danchin E, Avilés JM (2005) The heterospecific habitat copying hypothesis: can competitors indicate habitat quality? *Behav Ecol* 16:96–105. <https://doi.org/10.1093/BEHECO/ARH136>

Perperoglou A, Sauerbrei W, Abrahamowicz M, Schmid M (2019) A review of spline function procedures in R. *BMC Med Res Methodol* 19:1–16. <https://doi.org/10.1186/S12874-019-0666-3>

Pizzatto L, Stockwell M, Clulow S, et al (2016) Finding a place to live: conspecific attraction affects habitat selection in juvenile green and golden bell frogs. *Acta Ethol* 19:1–8. <https://doi.org/10.1007/s10211-015-0218-8>

Polak M (2014) Protective nesting association between the Barred Warbler *Sylvia nisoria* and the Red-backed Shrike *Lanius collurio*: an experiment using artificial and natural nests. *Ecol Res* 29:949–957. <https://doi.org/10.1007/s11284-014-1183-9>

Quinn JL, Kokorev Y (2002) Trading-off risks from predators and from aggressive hosts. *Behav Ecol Sociobiol* 51:455–460. <https://doi.org/10.1007/s00265-002-0466-2>

Quinn JL, Prop J, Kokorev Y, Black JM (2003) Predator protection or similar habitat selection in red-

599       breasted goose nesting associations: Extremes along a continuum. *Anim Behav* 65:297–307.  
600       <https://doi.org/10.1006/anbe.2003.2063>

601       Quinn JL, Ueta M (2008) Protective nesting associations in birds. *Ibis (Lond 1859)* 150:146–167.  
602       <https://doi.org/10.1111/j.1474-919X.2008.00823.x>

603       R Development Core Team (2019) A Language and Environment for Statistical Computing. R Found Stat  
604       Comput <https://www.R-project.org>

605       Rebollo S, García-Salgado G, Pérez-Camacho L, et al (2017) Prey preferences and recent changes in diet  
606       of a breeding population of the Northern Goshawk *Accipiter gentilis* in Southwestern Europe. *Bird*  
607       *Study* 64:464–475. <https://doi.org/10.1080/00063657.2017.1395807>

608       Reed JM, Boulonier T, Danchin E, Oring LW (1999) Informed Dispersal. *Curr Ornithol* 189–259.  
609       [https://doi.org/10.1007/978-1-4757-4901-4\\_5](https://doi.org/10.1007/978-1-4757-4901-4_5)

610       Reynolds JD (1996) Animal breeding systems. *Trends Ecol Evol* 11:68–72. [https://doi.org/10.1016/0169-](https://doi.org/10.1016/0169-5347(96)81045-7)  
611       5347(96)81045-7

612       Richards SA, Whittingham MJ, Stephens PA (2011) Model selection and model averaging in behavioural  
613       ecology: The utility of the IT-AIC framework. *Behav Ecol Sociobiol* 65:77–89.  
614       <https://doi.org/10.1007/s00265-010-1035-8>

615       Rosenzweig ML (1981) A Theory of Habitat Selection. *Ecology* 62:327–335.  
616       <https://doi.org/10.2307/1936707>

617       Rutila J, Latja R, Koskela K (2002) The common cuckoo *Cuculus canorus* and its cavity nesting host, the  
618       redstart *Phoenicurus phoenicurus*: A peculiar cuckoo-host system? *J Avian Biol* 33:414–419.  
619       <https://doi.org/10.1034/j.1600-048X.2002.02937.x>

620       Samaš P, Rutila J, Grim T (2016) The common redstart as a suitable model to study cuckoo-host  
621       coevolution in a unique ecological context. *BMC Evol Biol* 16:1–13.  
622       <https://doi.org/10.1186/s12862-016-0835-5>

623       Samplonius JM, Both C (2019) Climate Change May Affect Fatal Competition between Two Bird Species.  
624       *Curr Biol* 29:327–331.e2. <https://doi.org/10.1016/j.cub.2018.11.063>

625       Seppänen JT, Forsman JT, Monkkönen M, Thomson RL (2007) Social information use is a process across  
626       time, space, and ecology, reaching heterospecifics. *Ecology* 88:1622–1633.  
627       <https://doi.org/10.1890/06-1757.1>

628       Sieving KE, Contreras TA, Maute KL (2004) Heterospecific facilitation of forest-boundary crossing by  
629       mobbing understory birds in north-central Florida. *Auk* 121:738–751.  
630       <https://doi.org/10.2307/4090311>

631       Solonen T, Lokki H, Sulkava S (2019) Diet and brood size in rural and urban Northern Goshawks *Accipiter*  
632       *gentilis* in southern Finland. *Avian Biol Res* 12:3–9. <https://doi.org/10.1177/1758155919826754>

633       Szymkowiak J (2013) Facing Uncertainty: How Small Songbirds Acquire and Use Social Information in  
634       Habitat Selection Process? *Springer Sci Rev* 1:115–131. [https://doi.org/10.1007/s40362-013-0012-](https://doi.org/10.1007/s40362-013-0012-9)  
635       9

636       Szymkowiak J, Thomson RL, Kuczyński L (2016) Wood warblers copy settlement decisions of poor quality

637 conspecifics: support for the tradeoff between the benefit of social information use and  
 638 competition avoidance. *Oikos* 125:1561–1569. <https://doi.org/10.1111/oik.03052>

639 Thomson RL, Forsman JT, Mönkkönen M (2003) Positive interactions between migrant and resident  
 640 birds: Testing the heterospecific attraction hypothesis. *Oecologia* 134:431–438.  
 641 <https://doi.org/10.1007/s00442-002-1140-0>

642 Thomson RL, Forsman JT, Mönkkönen M (2011) Risk taking in natural predation risk gradients: Support  
 643 for risk allocation from breeding pied flycatchers. *Anim Behav* 82:1443–1447.  
 644 <https://doi.org/10.1016/j.anbehav.2011.09.029>

645 Thomson RL, Forsman JT, Sardà-Palomera F, Mönkkönen M (2006) Fear factor: Prey habitat selection  
 646 and its consequences in a predation risk landscape. *Ecography (Cop)* 29:507–514.  
 647 <https://doi.org/10.1111/j.0906-7590.2006.04568.x>

648 Thomson RL, Tolvanen J, Forsman JT (2016) Cuckoo parasitism in a cavity nesting host: Near absent egg-  
 649 rejection in a northern redstart population under heavy apparent (but low effective) brood  
 650 parasitism. *J Avian Biol* 47:363–370. <https://doi.org/10.1111/jav.00915>

651 Tolvanen J, Forsman JT, Thomson RL (2017a) Reducing cuckoo parasitism risk via informed habitat  
 652 choices. *Auk* 134:553–563. <https://doi.org/10.1642/auk-17-30.1>

653 Tolvanen J, Morosinotto C, Forsman JT, Thomson RL (2020) Information collected during the post-  
 654 breeding season guides future breeding decisions in a migratory bird. *Oecologia* 192:965–977.  
 655 <https://doi.org/10.1007/s00442-020-04629-5>

656 Tolvanen J, Pakanen VM, Valkama J, Tornberg R (2017b) Apparent survival, territory turnover and site  
 657 fidelity rates in Northern Goshawk *Accipiter gentilis* populations close to the northern range limit.  
 658 *Bird Study* 64:168–177. <https://doi.org/10.1080/00063657.2017.1309351>

659 Tornberg R, Mönkkönen M, Kivelä SM (2009) Landscape and season effects on the diet of the Goshawk.  
 660 *Ibis (Lond 1859)* 151:396–400. <https://doi.org/10.1111/j.1474-919X.2009.00910.x>

661 Tornberg R, Rytönen S, Välimäki P, et al (2015) Northern Goshawk (*Accipiter gentilis*) may improve  
 662 Black Grouse breeding success. *J Ornithol* 2015 1571 157:363–370.  
 663 <https://doi.org/10.1007/S10336-015-1292-4>

664 Toyne EP (1998) Breeding season diet of the Goshawk *Accipiter gentilis* in Wales. *Ibis (Lond 1859)*  
 665 140:569–579. <https://doi.org/10.1111/j.1474-919x.1998.tb04701.x>

666 Trnka A, Grim T (2013) Color plumage polymorphism and predator mimicry in brood parasites. *Front*  
 667 *Zool* 10:25. <https://doi.org/10.1186/1742-9994-10-25>

668 Trnka A, Prokop P (2012) The effectiveness of hawk mimicry in protecting cuckoos from aggressive  
 669 hosts. *Anim Behav* 83:263–268. <https://doi.org/10.1016/j.anbehav.2011.10.036>

670 Ueta M (2007) Effect of Japanese lesser sparrowhawks *Accipiter gularis* on the nest site selection of  
 671 azure-winged magpies *Cyanopica cyana* through their nest defending behavior. *J Avian Biol*  
 672 38:427–431. <https://doi.org/10.1111/j.2007.0908-8857.04172.x>

673 Ueta M (1999) Cost of Nest Defense in Azure-Winged Magpies. *J Avian Biol* 30:326.  
 674 <https://doi.org/10.2307/3677361>

675 Valone TJ (2007) From eavesdropping on performance to copying the behavior of others: A review of  
676 public information use. *Behav. Ecol. Sociobiol.* 62:1–14

677 Valtonen A, Latja R, Leinonen R, Pöysä H (2017) Arrival and onset of breeding of three passerine birds in  
678 eastern Finland tracks climatic variation and phenology of insects. *J Avian Biol* 48:785–795.  
679 <https://doi.org/10.1111/JAV.01128>

680 Van Balen JH, Booy CJH, Van Franeker JA, Osieck ER (1982) Studies on Hole-Nesting Birds in Natural Nest  
681 Sites. *Ardea* 70:1–24. <https://doi.org/10.5253/arde.v70.p1>

682 Welbergen JA, Davies NB (2011) A parasite in wolf's clothing: Hawk mimicry reduces mobbing of cuckoos  
683 by hosts. *Behav Ecol* 22:574–579. <https://doi.org/10.1093/beheco/arr008>

684 Wesołowski T (2002) Anti-predator adaptations in nesting Marsh Tits *Parus palustris*: The role of nest-  
685 site security. *Ibis (Lond 1859)* 144:593–601. <https://doi.org/10.1046/j.1474-919X.2002.00087.x>

686 Wood SN (2017) Generalized Additive Models. Chapman and Hall/CRC

687 Zuur AF, Ieno EN, Walker N, et al (2009) Mixed effects models and extensions in ecology with R.  
688 <https://doi.org/10.1007/978-0-387-87458-6>



689 **Table 1.** AICc statistics of the final set of models explaining variation in different breeding factors.  
690 Only models within 6 AICc units were included but excluding models that were more complex  
691 versions of a model with lower AICc.

Response variable	Model parameters	n	Df	AICc	dAICc	Akaike weight
<b><i>Great tit attraction</i></b>						
Occupancy	Tit box + Year	164	6	218.8	0.00	1.00
Laying date	Tit box	77	3	508.4	0.00	1.00
Success rate	Tit box	62	3	64.4	0.00	1.00
Parasitism rate	Tit box	70	3	87.0	0.00	1.00
Clutch size	Tit box	68	4	271.3	0.00	1.00
Number of fledglings	Tit box	46	3	192.5	0.00	1.00
<b><i>Goshawk attraction</i></b>						
Occupancy	Distance + Box seniority	418	6	534.2	0.00	1.00
Laying date	Distance	154	5	1124.1	0.00	1.00
Success rate	Distance	137	5	187.0	0.00	1.00
Predation rate	Distance	141	5	140.7	0.00	1.00
Parasitism rate	Distance	142	5	163.5	0.00	1.00
Clutch size	Distance + Laying date	131	6	519.3	0.00	0.58
	Distance	131	5	519.9	0.62	0.42
Number of fledglings	Distance + Laying date	137	6	796.1	0.00	0.75
	Distance	137	5	798.3	2.26	0.25

**Table 2.** Summary of the generalised linear mixed model of redstart breeding parameters depending on great tit attraction (present/absent). Parameter estimates for which the 95% CI excludes zero are in bold.

Fixed effects	Estimate	95% CI	Random effect	Variance
Occupancy				
Intercept	1.37	0.33, 2.41	Patch ID	0.51
Tit neighbour, yes	-0.09	-0.76, 0.58		
Year 2015	-0.99	-2.02, 0.02		
Year 2016	-1.92	-3.00, -0.85		
Year 2017	-1.97	-3.15, -0.79		
Laying date				
Intercept	3.41	3.34, 3.47	Nest ID	0.01
Tit neighbour, yes	0.03	-0.06, 0.11		
Success rate				
Intercept	1.24	0.03, 2.45	Nest ID	0.50
Tit neighbour, yes	0.13	-1.09, 1.34		
Parasitism rate				
Intercept	-1.29	-2.07, -0.50	Nest ID	<0.01
Tit neighbour, yes	0.59	-0.47, 1.66		
Clutch size				
Intercept	1.91	1.79, 2.03	Nest ID	0.00
Tit neighbour, yes	0.01	-0.17,0.19		
Number of fledglings				
Intercept	1.74	1.58, 1.91	Nest ID	0.00
Tit neighbour, yes	0.05	-0.19,0.29		

note: intercept is Tit neighbour "no" and Year 2014

**Table 3.** Summary of the generalised additive mixed models of redstart breeding parameters in the neighbourhood of active goshawk nest. Parameter estimates for which the 95% CI excludes zero are in bold. The distance as fixed effect has the smoother characteristic from generalised additive models. The estimated degrees of freedom (edf) reflect the degree of non-linearity of the smoother: an edf of 1 is equivalent to a linear relationship, an edf > 1 and ≤ 2 is a weakly non-linear relationship, and an edf > 2 indicates a highly non-linear relationship (Zuur et al. 2009).

Fixed effects	Estimate	95% CI	edf	p-value	Random effects	Variance
<b>Occupancy</b>						
Intercept	-1.19	-2.581, 0.036	-	-	Year	0.47
s(Distance)	-	-	<b>1</b>	<b>&lt;0.01</b>	Territory ID	0.97
New Box, yes	<b>0.87</b>	<b>0.278, 1.488</b>	-	-		
<b>Laying date</b>						
Intercept	<b>3.41</b>	<b>3.309, 3.501</b>	-	-	Year	<0.01
s(Distance)	-	-	<b>4.13</b>	<b>&lt;0.01</b>	Nest ID	0.01
<b>Success rate</b>						
Intercept	0.22	-0.836, 1.052	-	-	Year	0.00
s(Distance)	-	-	1	0.59	Nest ID	0.75
<b>Predation rate</b>						
Intercept	-0.99	-2.272, 0.391	-	-	Year	<0.01
s(Distance)	-	-	2.06	0.20	Nest ID	1.91
<b>Parasitism rate</b>						
Intercept	<b>-1.22</b>	<b>-1.996, -0.651</b>	-	-	Year	0.11
s(Distance)	-	-	1	0.40	Nest ID	0.00
<b>Clutch size</b>						
Intercept	<b>2.10</b>	<b>1.850, 2.356</b>	-	-	Year	0.00
s(Distance)	-	-	1	0.74	Nest ID	0.00
Laying date	-0.01	-0.014, 0.001	-	-		
<b>Number of fledglings</b>						
Intercept	<b>1.27</b>	<b>0.418, 1.987</b>	-	-	Year	<0.01
s(Distance)	-	-	1	0.09	Nest ID	0.69
Laying date	<b>-0.01</b>	<b>-0.026, -0.001</b>	-	-		

note: where applicable intercept is New box, no.

## 703 Figure legends

704 **Figure 1.** Redstart nest box pair design with one “control” and one “neighbour” box, set up  
705 approximately 200 m apart. The neighbour box had a great tit nest box placed approximately 25  
706 m away, while the control box had no great tit neighbour nearby. Redstart nest box pairs were  
707 only included in the dataset if the tit box contained an active great tit nest.

708

709 **Figure 2.** Frequency distribution of the number of nest-boxes available for redstarts from 20 m to  
710 2000 m from the active goshawk nest. The count bins are each 250 m.

711

712 **Figure 3.** The most supported generalized additive mixed model for redstart predicted relative  
713 probability of (a) occupancy, (b) parasitism and (c) predation rates and (d) the probability of  
714 success at different distance from an active goshawk nest. The dashed lines are the 95% CIs.

715

716 **Figure 4.** The most supported generalized additive mixed model for redstart predicted relative (a)  
717 laying date, (b) clutch size (c) number of fledglings at different distance from an active goshawk  
718 nest. The dashed lines are the 95% CIs.

Figure 1.

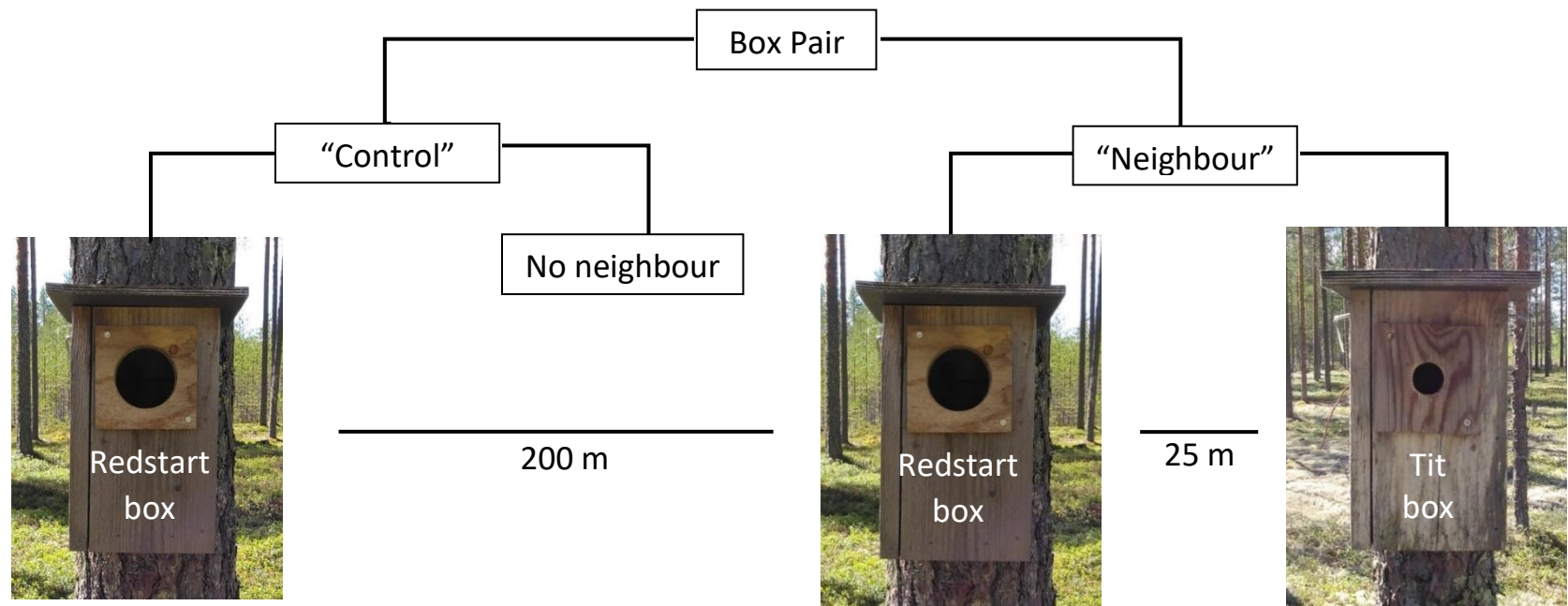


Figure 2.

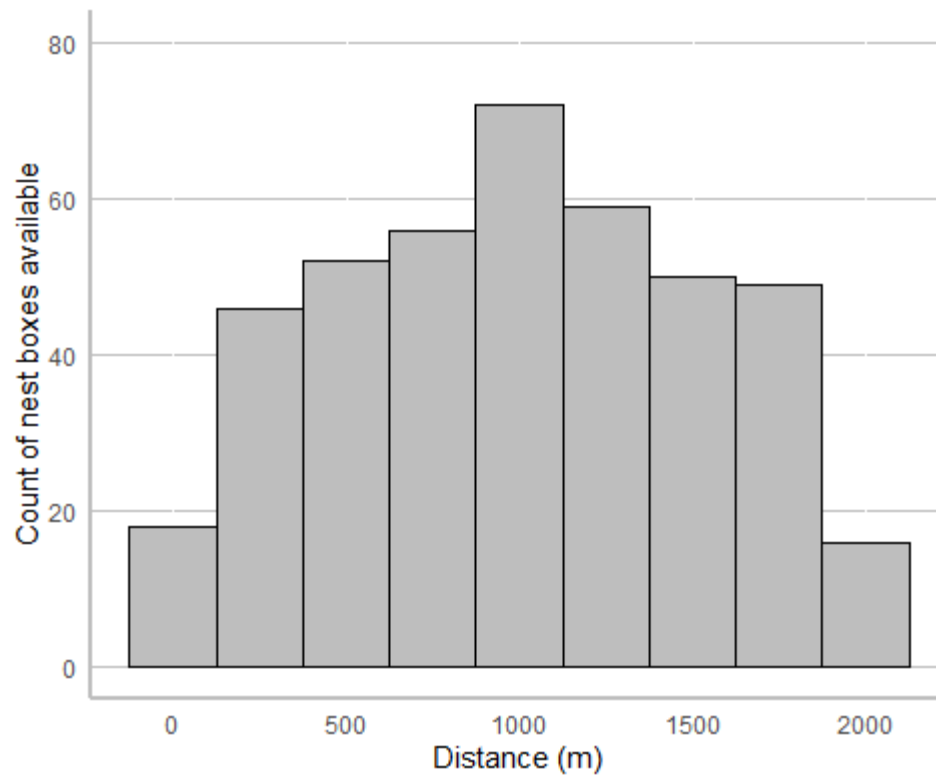
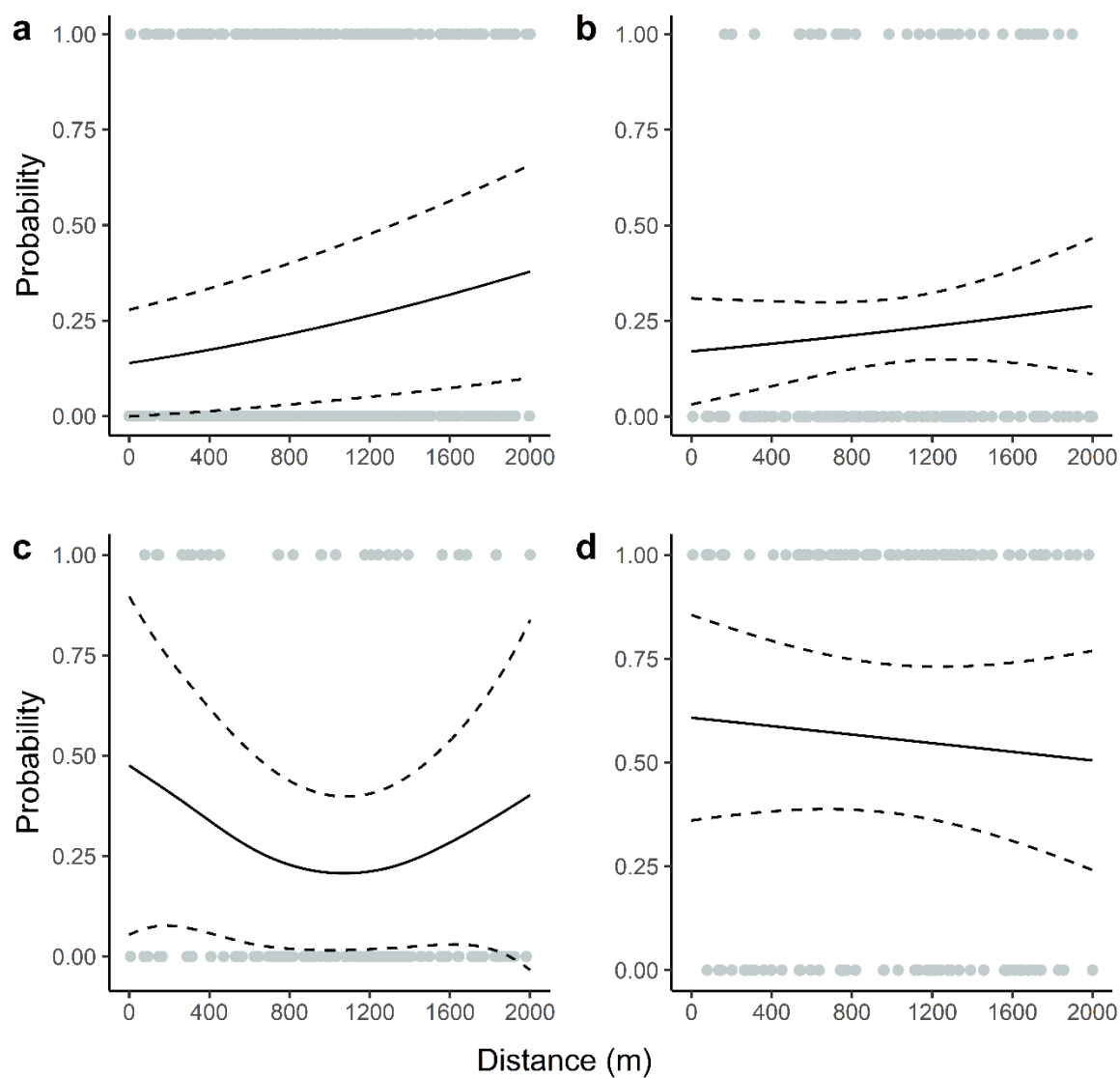


Figure 3.



Figure

