

1 **Can hedgerow management mitigate the impacts of predation on songbird nest**
2 **survival?**

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20

21 **Abstract**

22 Nest predators can have significant impacts on songbird reproductive success. These
23 impacts may be amplified by habitat simplification and here we test whether
24 sympathetic management of farmland hedgerows can reduce nest depredation,
25 especially by corvids. We test whether songbirds select nest sites according to structural
26 features of hedgerows (including nest visibility and accessibility), and whether these
27 features influence nest predation risk. Songbirds selected nesting sites affording higher
28 vegetation cover above the nest, increased visibility on the nest-side of the hedgerow
29 and reduced visibility on the far side of the hedge. Nest survival was unrelated to corvid
30 abundance and only weakly related (at the egg stage) to corvid nest proximity. Nest
31 survival at the chick stage was higher where vegetation structure restricted access to
32 corvid-sized predators (averaging 0.78 vs. 0.53), and at nests close to potential vantage
33 points. Overall nest survival was sensitive to hedgerow structure (accessibility)
34 particularly at low exposure to corvid predation, while the overall impact of corvid
35 exposure was dependent on the relationship involving proximity to vantage points. Nest
36 survival over the chick stage was much higher (0.67) in stock-proof, trimmed and
37 mechanically cut hedgerows, (which tended to provide lower side visibility and
38 accessibility) than in recently laid, remnant or leggy hedgerows (0.18). Long-term
39 reductions in the management of British hedgerows may therefore be exposing nesting
40 songbirds to increased predation risk. We recommend regular rotational cutting of
41 hedgerows to maintain a dense woody structure and thereby reduce songbird nest
42 predation.

43

44 **Keywords:** nest predation, corvids, farmland birds, predator-habitat interactions,
45 farmland conservation

46

47 **1. Introduction**

48 Nest predation is the main cause of nestling mortality in birds (Ricklefs, 1969),
49 with losses to predators approaching 69% in some altricial species (Remes and Martin,
50 2002), sometimes leading to population sinks (Rogers et al., 1997). Species suffering
51 high levels of nest predation have evolved behavioural and life-history strategies to
52 minimise predation risk (Dunn et al., 2010; Eggers et al., 2005a; Martin, 1995) such as
53 shorter nestling periods and multiple broods each year (Martin, 1995). Parents tend to
54 reduce investment in a nest when predation risk is high, through reduced egg size
55 (Fontaine and Martin, 2006), clutch size (Julliard et al., 1997) and clutch mass (Fontaine
56 and Martin, 2006). High levels of activity around the nest may attract predators and
57 parents often reduce activity when the risk of nest predation is high (Conway and
58 Martin, 2000; Dunn et al., 2010; Eggers et al., 2005b).

59 Behavioural adjustment by adult birds to reduce nest predation risk (Dunn et al.,
60 2010; Eggers et al., 2005b) is dependent not only on predator activity, but also on the
61 cover around the nest and the availability of food for chicks (Eggers et al., 2008). In
62 areas where food abundance is low, high corvid abundance is associated with reduced
63 nestling growth in a farmland songbird (Dunn et al., 2010). Ecological factors affecting
64 the likelihood of nest predation include nest density (Cresswell, 1997; Schmidt and
65 Whelan, 1998), predator abundance and nest type (i.e. cavity vs. open-cup, Fontaine et
66 al. 2007). Nests that are more visible are more likely to be depredated at the egg stage
67 (Martin et al., 2000; Matessi and Bogliani, 1999). Predation rates tend to increase with
68 reduced vegetation cover, vegetation height, and nest height (e.g. Cresswell 1997), all
69 features that are likely to interact to influence nest detectability and accessibility
70 (Cresswell, 1997), although there is no evidence for nest size affecting predation risk
71 (Weidinger, 2004). Factors affecting nest predation risk may differ between predators:
72 corvids are more likely to depredate poorly concealed nests, whereas well concealed
73 nests are more likely to suffer depredation by rodents (Weidinger, 2002). There may be
74 a trade-off for nest survival between nest concealment and the ability of parent birds to
75 detect an approaching predator (Cresswell, 1997; Gotmark and Post, 1996; Weidinger,
76 2002).

77 Corvids are important nest predators, especially in farmland environments
78 (Andren, 1992; Luginbuhl et al., 2001), and their populations in the UK have increased
79 steadily since the 1960s, coincidental with the declines in many farmland songbirds
80 (Gregory and Marchant, 1995). Whilst no clear link has been found between declining
81 abundance of farmland songbirds and increasing abundance of corvids (Gooch et al.,
82 1991; Madden et al., 2015; Newson et al., 2010; Thomson et al., 1998), local examples

83 have shown predation impacts through farming management. Organic farms harbour
84 more corvids, but fewer songbirds (Gabriel et al., 2010) and gamebird management
85 (corvid control and sympathetic habitat management) is associated with higher nest
86 survival and higher breeding densities of songbirds (Stoate and Szczur 2001, White et al.
87 2008, White et al. 2014). An extensive analysis of song thrush and blackbird nest record
88 cards found fine-scale spatial associations between corvid densities and nest survival
89 rates (Paradis et al., 2000). That corvids are responsible for high numbers of nest losses
90 is indisputable (Andren, 1992; Bradbury et al., 2000; Luginbuhl et al., 2001), and the
91 linear nature of hedgerows in farmland landscapes may increase the risk of nest
92 depredation (Chamberlain et al., 1995). Legal control of corvids is advocated and
93 practiced for game management, and specifically for songbird conservation, but the
94 control of one native species to benefit another is expensive and not universally
95 accepted as a management practice. Thus, reducing corvid nest predation through
96 habitat management would be desirable if possible, and, alongside measures to increase
97 food availability during summer and winter, providing productive nesting habitat might
98 help reverse population declines amongst some farmland bird species (Fuller et al.,
99 1995). To our knowledge, no previous study has investigated impacts of hedgerow
100 structure or management on nest predation risk in songbirds. The aims of the current
101 study were (1) to identify the structural features of farmland hedgerows that influence
102 nest site selection by songbirds, (2) to assess the relative importance of hedgerow
103 structure and corvid abundance / proximity in determining nest predation risk, and (3)
104 to identify aspects of hedgerow management that reduce nest predation risk.

105

106 **2. Methods**

107 **2.1 Study sites**

108 The fates of 399 songbird nests were monitored during April-July 2003 and
109 2004 across 11 farmland sites in total across two regions in eastern England (five in
110 Cambridgeshire, Bedfordshire and Hertfordshire, and six in Leicestershire and Rutland).
111 Not all sites were monitored in both years: 10 sites (5 in each region) were monitored in
112 2003 and seven sites (3 and 4 respectively) in 2004. The main species monitored were
113 Blackbird *Turdus merula* (n=140), Chaffinch *Fringilla coelebs* (83), Dunnock *Prunella*
114 *modularis* (17), Linnet *Carduelis cannabina* (103), Yellowhammer *Emberiza citrinella*
115 (28) and Song Thrush *Turdus philomelos* (8).

116

117 **2.2 Nest finding and monitoring**

118 Hedgerows were selected to cover the full range of hedgerow characteristics and
119 management types present across study sites. Studied hedgerows ranged from
120 intensively managed (usually less than 1 m high, with thin woody vegetation with gaps),
121 through managed (usually greater than 1 m high and cut or trimmed within the last 3 – 4
122 years) to unmanaged (usually >3 m high, not regularly cut or trimmed, often with trees
123 and tall shrubs). Hedgerows next to busy roads, gardens, woods or woodland strips, or
124 those planted within the last 5 years, were not selected for study.

125 Each hedgerow was cold-searched for nests at approximately weekly intervals
126 between early April and late July. For each nest, the species was noted and the nest was
127 inspected every 5–7 days until either the young fledged or the nest failed. Nest contents
128 and adult activity were recorded during each visit. Nest success was inferred by an
129 empty undamaged nest where the young were old enough to have fledged since the
130 previous visit. Nest failure was either known (nest contained cold eggs, egg fragments or
131 dead chicks) or was inferred from empty (often damaged) nests on a date prior to a
132 plausible fledging date. For analytical purposes the date of failure was assumed to be the
133 mid point between the last two visits. If there was evidence a nest had been pulled down
134 from below, we assumed predation by a mammal, although we acknowledge that
135 predator identification based on field signs is not always reliable (Pietz and Granfors,
136 2000). As we were specifically interested in corvid predation, nests for which
137 mammalian predation was presumed were excluded from analyses (n=11, 2.8 %), as
138 were any nests for which the outcome was uncertain (n=18, 4.5 %). A small number of
139 failures caused by starvation, abandonment, human interference, or egg infertility (eggs
140 not hatched or chicks found dead in the nest) were also excluded (n=24, 6.0 %).
141 Analyses were restricted to nests located within the woody vegetation of the hedge;

142 nests located on the ground or within field boundary vegetation were excluded. First egg
143 date (FED; a day-specific integer where 1st January = 1) was deduced from incomplete
144 clutches, hatch dates and estimated chick ages (e.g. Green, 2004). If FED could not be
145 determined to within 3 days then the nest was excluded from analyses.

146

147 **2.3 Nest site characteristics**

148 Data describing nest site characteristics were collected for 338 nests within ten
149 days of the nesting attempt ending and are defined in Table 1a (brief descriptions only
150 are given here). We recorded nest height above the ground, along with the shortest
151 horizontal depth and vertical depth between the nest and the hedgerow edge. We
152 measured nest dimensions to allow the calculation of nest volume, and identified
153 primary and surrounding plant species supporting the nest. Hedgerow height and width
154 at the nest were measured to calculate cross-sectional area at the nest site.

155 Nest concealment was assessed in three different ways: light penetration at the
156 nest (measured with a light meter), horizontal visibility (counts of white circles on a
157 black card positioned next to the nest) and vertical vegetation cover (assessed from a
158 digital image; see Table 1a for details). Nest accessibility (a binary variable) was
159 assessed by attempting to manoeuvre two different sized balls from the hedgerow edge
160 to the nest without breaking any woody vegetation (Table 1a). The smaller ball had a
161 circumference (30.5 cm) that was similar to the maximum body girth of a magpie *Pica*
162 *pica* (measured as 28 cm), and was intended to highlight potential accessibility to a
163 corvid. Accessibility with the larger ball (69cm) was intended to indicate easy access to a
164 foraging corvid.

165 Locations of carrion crow *Corvus corone* and magpie nests were recorded, and
166 the distance to the nearest corvid nest (corvid distance) subsequently calculated for
167 each songbird nest. We also recorded distance to the nearest wood or woodland strip
168 (wood distance), and distance to the nearest tree, pylon, telegraph pole or other vantage
169 point at least 5 m in height (vantage distance) as corvids are visually-oriented predators
170 known to utilise vantage points when searching for prey (Macdonald and Bolton, 2008).

171

172 **2.4 Corvid abundance**

173 The relative abundance of corvids (magpie, carrion crow, jackdaw *Corvus*
174 *monedula* and jay *Garrulus glandarius*) was assessed using a transect method (Stoate
175 and Szczur, 2001) on between 2 and 13 occasions (mean \pm SE: 5.08 ± 1.00 visits) at each
176 site during April-June of each year. Transects were spaced approximately 600 – 700 m
177 apart and followed field boundaries; mean transect length was 5.75 ± 1.15 km (± 1 SE).

178 Each transect was walked at a steady pace, and all corvids were recorded. The total
179 number of corvids per km within 100m of the observer, averaged over all transects, was
180 taken as an index of corvid abundance for each site in each year.

181

182 **2.5 Hedgerow characteristics and nest site selection**

183 Data describing hedgerow characteristics were collected for 391 nests and
184 described the section of hedgerow 30m either side of each nest. Firstly, the hedgerow
185 aspect was recorded, and hedgerow management and cutting style were categorised (as
186 in Table 1b). The percentage of gaps within the same stretch of hedgerow was
187 estimated, along with the number of trees. The occurrence and timing of a hedgerow cut
188 during the previous 5 years was determined during farmer interviews. The width of
189 vegetated margins on both sides of the hedgerow was measured.

190 In order to identify structural features of hedgerows that were selected or
191 avoided by nesting songbirds, we repeated the nest site measurements for primary
192 supporting and surrounding vegetation, horizontal visibility and vertical cover at six
193 locations spread at 10m intervals either side of the nest (if an interval fell within a
194 hedgerow gap, then a point 5 m either side of the gap was measured instead). The
195 measurements were collected at the same height in the hedgerow as the nest. These
196 data were collected for 333 nests.

197

198 **2.6 Statistical analysis**

199 *2.6.1 Nest site selection*

200 To determine the features of hedgerows selected by nesting birds, the key
201 features of nest sites thought to indicate aspects of nest visibility and accessibility
202 (vegetation cover, horizontal visibility from each side of the nest, primary species and
203 surrounding species; Table 1) were compared with the six adjacent non-nest locations
204 using conditional logistic regression stratified by nest identifier to allow for the non-
205 independence of nest and non-nest locations (Anteau et al., 2012).

206

207 *2.6.2 Likelihood of nest predation*

208 We used generalised linear mixed models (GLMMs) with binomial error
209 structures and logit link functions to determine whether nest site characteristics
210 influenced the likelihood of nest predation at the egg (n=202 nests) and chick (n=190)
211 stages. GLMMs were fitted using the 'glmer' function within the *lme4* package (Bates and
212 Maechler, 2009) in R v 2.10.1 for Mac (R Core Development Team, 2009). A hedgerow
213 identifier nested within farm was included as a random effect to control for the non-

214 independence of nests within the same hedgerow, or on the same farm, as well as to
215 control for spatial autocorrelation. The response variable was the daily whole nest
216 failure rate (DFR) in which nest outcome at the relevant nest stage (0 = successful, 1 =
217 depredated) was the binomial numerator and the number of exposure days during the
218 relevant nest stage declared as the binomial denominator (Aebischer, 1999; Hazler,
219 2004). Our aim was to identify predictors of nest survival associated with hedgerow
220 structure and corvid abundance / distance, and any interactions between the two. All
221 GLMMs initially included a set of fixed variables (irrespective of their statistical
222 significance) for factors that might have affected nest survival but were unrelated to
223 hedgerow structure or predator abundance (we call these 'base models'). These
224 included mean-centred FED (for egg stage survival) or hatch date (for chick stage
225 survival) as linear and quadratic terms to allow for non-linear temporal variation in
226 predation risk across the breeding season. They also included species, nest contents
227 (clutch or brood size for egg and chick stage models respectively) and year. We tested
228 each 'base variable' within the base model, and excluded those with $p > 0.10$ to avoid
229 overfitting, resulting in a 'final base model' which remained fixed for the rest of the
230 model selection.

231 We then followed a two-stage approach which aimed to identify predictors of
232 nest survival while balancing the likelihood of type I and type II errors (Pearce-Higgins
233 et al., 2009). First, each of the 15 hedgerow characteristics and corvid variables listed in
234 Table 1a (logarithm or arcsine transformed as necessary) was added to the final base
235 model one at a time. For those variables that were potentially influential on nest survival
236 ($p < 0.1$; Table 2) we checked for multicollinearity by examining correlations between
237 variable pairs (detailed in Appendix 1). This was done separately for nest site character
238 and hedgerow management variables. As potentially influential variables exhibited little
239 inter-correlation (all r values < 0.5 ; Appendix 1), all were retained in a second stage of
240 multivariate testing. This second stage involved the addition of all potentially influential
241 hedgerow/corvid variables to the final base model, followed by sequential backwards
242 deletion in which the least significant term (assessed using p value) was removed until
243 all remaining hedgerow/corvid terms were either formally significant ($p < 0.05$) or
244 potentially influential ($p < 0.1$). We report the latter to avoid type 2 errors but interpret
245 such relationships more cautiously. We finally tested two-way interactions between our
246 best measures of corvid exposure (corvid abundance and corvid distance) and our
247 measure of nest visibility most relevant to corvids flying overhead or walking along the
248 top of a hedgerow (vegetation cover), to assess whether more visible nests were more
249 likely to be depredated in areas of higher corvid exposure. Whilst stepwise model

250 construction has been criticised in the literature (Whittingham et al., 2006), it has since
251 been shown that stepwise approaches perform just as well as other methods (Murtaugh,
252 2009). Information theoretic methods were not employed as these require estimation
253 processes that approximate the likelihood rather than the model (Bolker et al., 2009).

254 To estimate the scale of effect at the whole nest level, we combined corvid and
255 nest access variables from our final models to predict overall nest survival rates for
256 accessible and inaccessible nests (small ball access) and high and low potential corvid
257 exposure (corvid distance and vantage distance). We predicted at both levels for binary
258 data, and at levels of the 10th and 90th percentiles from the raw data to provide whole
259 egg-stage, whole chick-stage and whole nest-stage survival proportions.

260

261 *2.6.3. Management associations with nest predation*

262 The GLMMs for egg (n=209) and chick stage (n=195) nest survival were
263 extended to test for any influence of our 8 hedgerow management variables (Table 1b).
264 Model selection proceeded as described above (2.5.2) with each management term
265 initially added in turn to the base model, followed by backwards deletion on the
266 significant one-at-a-time predictors. Because several of the management variables were
267 categorical and were likely to have co-varied, no interactions were considered.

268

269

270 3 Results

271

272 3.1 Nest site selection

273 Nest locations were characterised by higher vegetation cover compared to non-
274 nest sites, along with higher visibility from the side of the hedgerow closest to the nest
275 and lower visibility from far side of hedgerow (Table 3, Figure 1). Bramble was most
276 likely to be selected as the primary support for nests (rose the least), while ivy was the
277 preferred surrounding species (locations with rose or no surrounding species being
278 avoided; Table 3).

279

280 3.2 Effects of hedgerow structure and predators on nest failure rates

281 Mean DFR at the egg stage was 0.032, equivalent to 0.35 failure over a 13-day
282 incubation period. For nests reaching the chick stage, mean DFR was 0.035 equivalent to
283 0.38 failure over a 13-day chick-rearing period.

284 Egg stage DFRs declined significantly with increasing clutch size, and exhibited a
285 weak negative relationship with distance to the nearest corvid nest (Appendix 2; Table
286 4).

287 After allowing for a marked seasonal decline in chick stage failure rates, DFRs
288 were higher for nests that were accessible with a small ball (Figure 2a; DFRs of 0.047
289 and 0.019 for accessible and inaccessible nests respectively, equivalent to failure rates of
290 0.467 and 0.219 over a 13-day chick-rearing period), and for nests located further away
291 from vantage points (Figure 2b; Table 4).

292 The effect size of nest accessibility in terms of overall nest survival (averaged
293 between high and low corvid exposure) was 0.143, compared to a mean effect size of
294 0.031 for corvid exposure (averaged between accessible and inaccessible nests; Table
295 5). The sensitivity of overall nest survival to nest accessibility was particularly high
296 when corvid exposure was low (0.360 vs. 0.569, Table 5). When we excluded from these
297 calculations the (counterintuitive) positive relationship between chick stage nest failure
298 and vantage point distance, overall nest survival was similarly sensitive to corvid
299 exposure (mean effect 0.178) and nest accessibility (0.141; Table 5).

300

301 3.3 Effects of hedgerow management on nest failure rates

302 There was a statistically weak effect of time since last cut on egg stage DFR
303 (Table 6) with nests in recently cut hedgerows experiencing higher failure rates (nest
304 failure over the 13-day chick period was 0.693 in hedgerows cut during the preceding
305 year compared to 0.237 in hedgerows cut 4 years previously; Appendix 3). Chick stage

306 DFRs differed between hedgerow management (Table 6). In leggy, remnant and recently
307 laid hedgerows, the DFR averaged 0.125, equivalent to a nest failure rate of 0.824 over
308 the 13-day chick-rearing period. Conversely, in mechanically cut, trimmed but dense,
309 and stock-proof hedgerows, DFR averaged 0.030, equivalent to 0.327 nest failure across
310 the chick-rearing period (Figure 3). Hedgerow management categories associated with
311 this higher predation risk (leggy, remnant, recently laid) were characterised by
312 relatively high horizontal visibility and high small ball accessibility (Table 7).
313
314

315 4 Discussion

316 To our knowledge, this is the first study to consider the potential for hedgerow
317 management to ameliorate the impacts of avian nest predators on farmland songbirds.
318 In farmland environments, 50% of hedgerows have been removed since 1945 and the
319 diversity and quality of remaining hedgerows has declined (Robinson and Sutherland,
320 2002). This deterioration of linear hedgerow nesting habitat may have allowed songbird
321 nests to become more susceptible to nest predation in agricultural landscapes (Evans,
322 2004; Hinsley and Bellamy, 2000; Whittingham and Evans, 2004). We found songbirds
323 to select nest sites based on vegetation characteristics likely to provide concealment and
324 limit access to predators. We found evidence for both vegetation and corvid variables
325 influencing nest survival, suggesting that improving hedgerow structure can mitigate
326 corvid predation. Critically, we found that hedgerow management can influence nest
327 survival, with much lower nest failure rates in hedgerows that were managed to create a
328 dense structure (e.g. stock-proof or mechanically cut) compared to unmanaged (e.g.
329 leggy and remnant) hedgerows.

330

331 4.1 Nest site selection

332 Songbirds selected nest sites with high vegetation cover above the nest, which is
333 likely to afford a degree of protection from corvids flying overhead or foraging along the
334 top of hedgerows (Cresswell, 1997), as corvids tend to depredate more visible nests
335 (Matessi and Bogliani, 1999; Weidinger, 2002). Denser vegetation cover may also confer
336 protection from adverse weather. Songbirds also selected nest sites that conferred
337 relatively high visibility on the nest side of the hedge, and relatively low visibility on the
338 far side (Götmark et al., 1995). When faced with an approaching predator, incubating or
339 brooding parent birds tend to flush sooner when visibility from the nest is higher
340 (Burhans and Thompson, 2001), which may reduce the risk of attracting attention to the
341 nest, or disclosing the exact location of the nest by flushing late. Bramble was selected as
342 the primary nest support, possibly because its dense and thorny character may restrict
343 nest detection and access by predators. Ivy was selected for vegetation surrounding the
344 nest, probably as this evergreen species provides increased cover above the nest
345 especially early in the breeding season when well-hidden nest sites are less abundant
346 (e.g. White et al. 2008). Rose species were avoided for both primary and surrounding
347 vegetation, possibly because of their late leafing and relatively sparse leaf cover.

348

349 4.2 Effects of predator abundance / proximity and hedgerow structure on nesting 350 success

351 We found a weak negative effect of corvid nest distance on egg-stage nest failure
352 rates, which is likely to be a consequence of increased corvid activity close to corvid
353 nests. Conversely, we found a positive relationship between chick-stage failure rates and
354 distance to vantage point, which is surprising given the expectation that nests closer to
355 vantage points are more likely to be noticed by corvids (Macdonald and Bolton, 2008).
356 However, passerines are known to mediate predation risk through behavioural
357 modifications so may compensate for this increased exposure by reducing nest
358 visitation at times when corvids are present (Dunn et al., 2010; Eggers et al., 2005b). At
359 the chick stage, more accessible nests suffered higher failure rates, suggesting that
360 hedgerow structure can mitigate corvid predation (Evans, 2004). The relatively large
361 difference in nest survival between accessible and inaccessible nests translated into a
362 large effect of nest accessibility on overall nest survival especially when exposure to
363 potential corvid impacts was relatively low (raising average nest survival from 0.360 to
364 0.569: Table 5). This confirms that hedgerow vegetation structure confers considerable
365 protection to songbird nests against depredation, highlighting a role for hedgerow
366 management in songbird conservation. The overall impact of predator exposure on nest
367 survival depended on the inclusion of the (counterintuitive) positive relationship
368 between chick-stage nest failure and distance to vantage point (Table 5). Including this
369 relationship in predictions rendered overall nest survival relatively insensitive to corvid
370 exposure. However excluding this relationship (which may be artefactual) from
371 predictions, renders nest survival similarly sensitive to corvid exposure and hedgerow
372 vegetation structure (Table 5).

373

374 **4.3 Management influences on nest success and recommendations**

375 Although the effect of time since last cut on nest survival was statistically weak
376 the effect size was large, with nests more than twice as likely to survive in hedgerows
377 cut 4 years previously than during the past year. Nest survival rates were much higher
378 in stock-proof, trimmed or mechanically cut hedgerows (0.673) than in unmanaged or
379 recently laid hedges (0.176), probably because the more open vegetation structure in
380 unmanaged hedges limited the scope for nest concealment and protection from
381 predators (Cresswell, 1997; Matessi and Bogliani, 1999). Between 1984 and 2007 there
382 was a 24% reduction in the availability of 'managed' hedgerows in Great Britain (Carey
383 et al., 2008; Petit et al., 2003). Although hedgerow removal explains some of this loss up
384 until 1990, since then the loss of managed hedgerows is largely explained by a reduction
385 in hedgerow management activity and a transition to field boundaries dominated by
386 tree-lines and relict hedges especially in arable-dominated landscapes (Carey et al.,

387 2008). The loss of managed hedgerows from such landscapes, coupled with the increase
388 in corvid populations (Gregory and Marchant, 1995), may have increased the
389 vulnerability of hedgerow-nesting songbirds to nest predation (Evans, 2004).

390 Direct control of corvids can increase nest survival and breeding abundance of
391 some songbird species but is expensive and not universally accepted as a conservation
392 management strategy (White et al., 2014). Our study adds to the increasing literature
393 suggesting that negative impacts of corvids can be mitigated by improving habitat
394 quality (Dunn et al., 2010; Eggers et al., 2008; Evans, 2004). Associations with hedgerow
395 management were particularly clear-cut in determining chick survival with stock-proof
396 and trimmed hedgerows providing the highest nestling survival rates (Fig. 3). In
397 England, a new agri-environment scheme (Countryside Stewardship; Natural England,
398 2015) started in 2016 and promotes environmental management of hedgerows by
399 specifying minimum dimensions (2 m tall and 1.5 m wide) and cutting regimes (outside
400 the breeding season, no more than one year in three and leaving at least one-half of
401 hedgerows untrimmed each year). These cutting regimes can improve moth and
402 parasitoid diversity (Facey et al., 2014), and increase resources such as flowers and
403 berries (Staley et al., 2012), and our data suggest these guidelines should also benefit
404 nesting birds. Our data emphasise the importance of regular hedgerow trimming to
405 promote a dense woody structure and prevent succession to tree lines, and the
406 avoidance of overly frequent cutting (our data suggest a cut every 3-4 years might be
407 optimal to promote songbird nest survival; Appendix 3). Rotational hedge cutting
408 regimes within a farm (i.e. cutting 1/4 - 1/3 of hedgerows each year) is one way to
409 provide heterogeneity and ensure a continuous supply of other resources such flowers
410 and berries to meet other wildlife conservation objectives as well as improve passerine
411 nest survival.

412

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421

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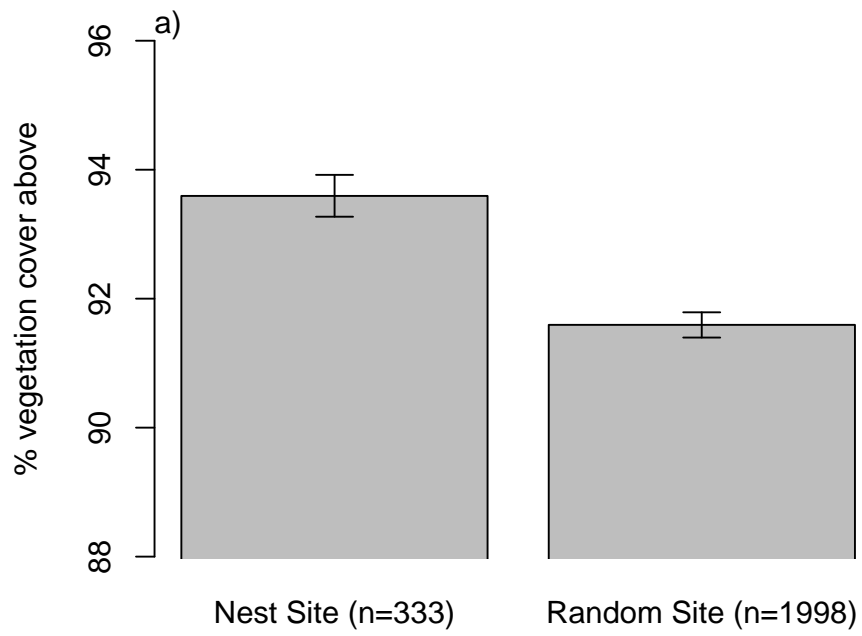
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567 Figure 1.

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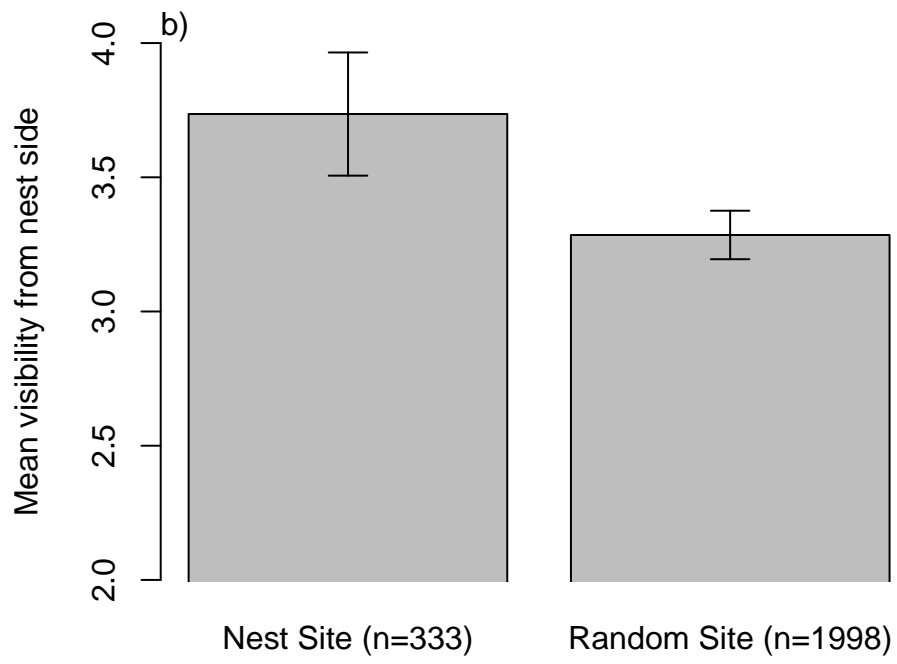
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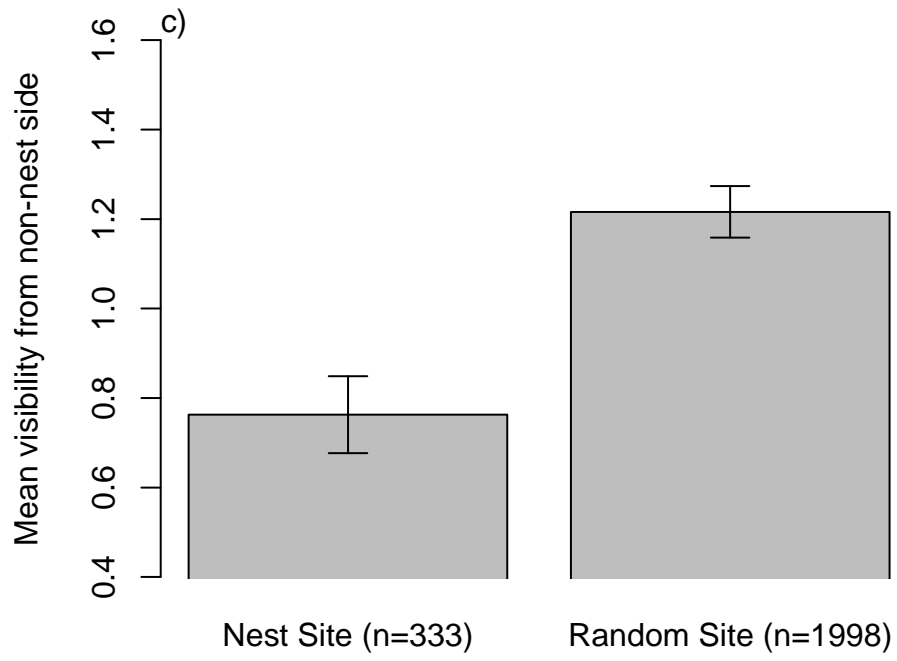
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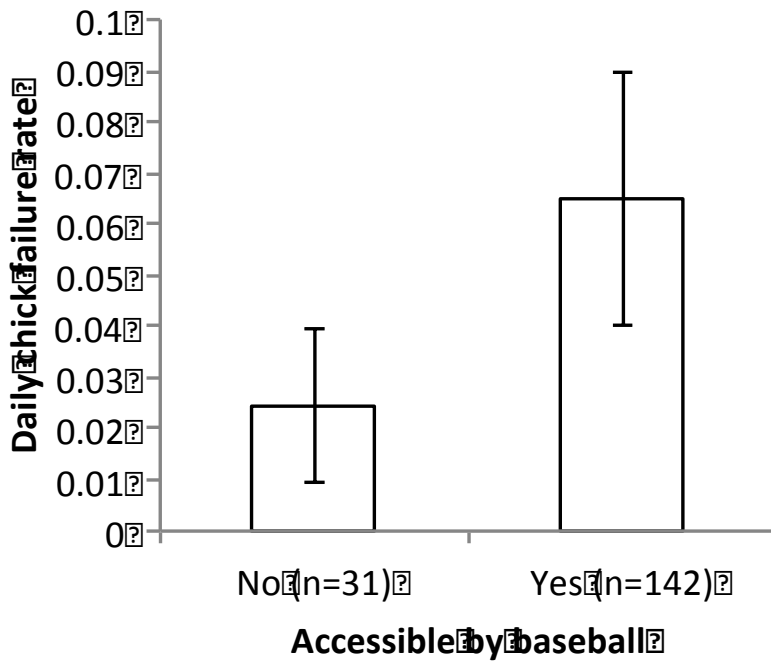


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581 Figure 2

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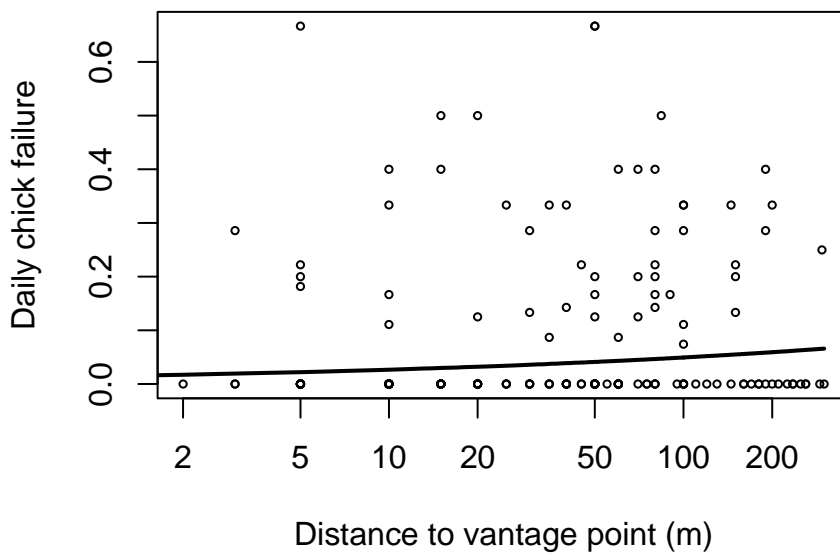
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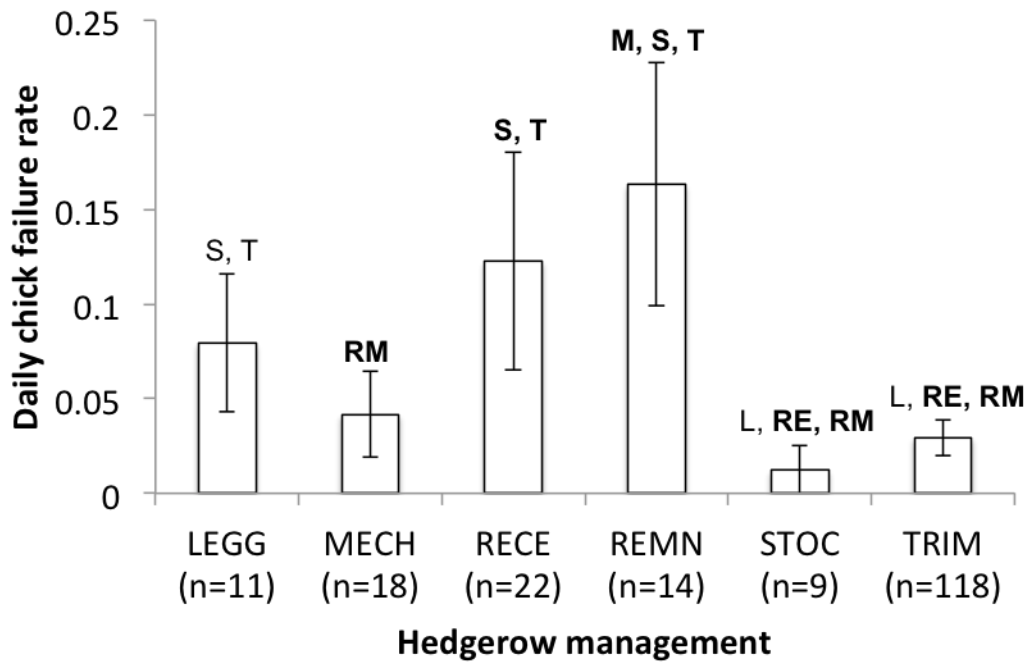
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590 Figure 3.

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599 **Figure and Table Legends**

600

601 Figure 1. Differences between nest sites and non-nest sites in 1) vegetation cover above
602 the nest, 2) visibility of the nest from the closest side of the hedgerow and 3) visibility
603 from the far side of the hedgerow. Bars show raw means \pm 1 SE.

604

605 Figure 2. Effect of a) nest accessibility (measured with a small ball) and b) distance from
606 the nearest vantage point, on chick stage failure rate. For a) Bars show predicted mean \pm
607 1 SE failure rates assuming mean hatch date. For b), points show raw data and the line is
608 predicted from the final model for accessible nests assuming mean hatch date.

609

610 Figure 3. Effect of hedgerow management on chick stage nest survival. Bars show
611 predicted means \pm 1 SE assuming mean hatch date and mean brood size, and numbers
612 indicate sample sizes. Categories of hedgerow management are: LEGG: Overgrown and
613 leggy; MECH: mechanically cut; RECE: recently laid; REMN: remnant; STOC: unclipped
614 and stock-proof; and TRIM: trimmed and dense. Letters above bars show where
615 significant differences lie (L = LEGG; M = MECH; Re = RECE, RM = REMN, S = STOC; T =
616 TRIM); letters in bold denote differences significant at $p < 0.05$, letters not in bold
617 denoted marginally significant ($P < 0.1$) differences.

618

619 Table 1. Descriptions of variables considered in analyses of a) nest and corvid
620 characteristics and b) management characteristics considered in nest survival models.

621

622 Table 2. Results of univariate tests for the initial base models to assess associations with
623 whole nest failure rates, followed by screening of explanatory variables against the final
624 base model for a) Hedgerow characteristics and b) Hedgerow management analyses at
625 the egg and chick stage separately. Terms with the variable name in bold were included
626 in interactions to examine the potential for interactions between nest visibility and
627 predator metrics. Terms where the statistics are highlighted in bold are those that were
628 highlighted as potentially important at $p < 0.1$ by univariate testing and were thus
629 included in subsequent multivariate analyses.

630

631 Table 3. Results from a conditional logistic regression determining which features
632 differed between nest sites and random sites within the same hedgerow. η^2 values are
633 from comparison of the final model with and without the term.

634

635 Table 4. Factors affecting the probability of nest predation at the egg and chick stages. a)
636 lists variables retained in multivariate GLMMs (at $P < 0.01$) while b) lists variables that
637 were tested but failed to achieve this level of statistical significance. See section 2.5 for
638 further details of model selection.

639

640 Table 5. Predicted daily nest survival rates (DSR) and whole nest survival rates at the
641 egg and chick stages separately, and combined, assuming 13-day incubation and chick-
642 rearing periods. Predictions are for combinations of high and low vegetation access
643 (Small ball accessibility: Y = high access; N = low access) and high and low corvid
644 exposure (corvid distance and distance to vantage point: near = high corvid; far = low
645 corvid), predicted from the final models (Table 4). Continuous variables are predicted at
646 levels of the 10th and 90th percentiles from the raw data. To test the sensitivity of our
647 nest survival predictions to the potentially counterintuitive effect of vantage distance,
648 we re-ran our predictions from the models excluding this variable (figures in brackets).

649

650 Table 6. Hedgerow management terms affecting the probability of nest predation at the
651 egg and chick stages. a) lists variables retained in multivariate GLMMs (at $P < 0.01$) while
652 b) lists variables that were tested but failed to achieve this level of statistical
653 significance. See section 2.5 for further details of model selection.

654

655 Table 7. Summary statistics (Mean \pm SE) and GLMMs comparing vegetation structure
656 between a) nest sites and b) random locations within the same hedgerow in high (Leggy,
657 recently laid and remnant) and low (mechanically cut, stock-proof and trimmed; see
658 Figure 2) predation risk hedgerow managements. For a) GLMMs contain hedgerow
659 within farm as random terms, and for b) GLMMs contain an additional nested random
660 term of nest ID (to control for multiple random points per stretch of hedgerow). Small
661 ball accessibility was not measured at random locations.

662

663

664 Table 1

665

666 a)

Variable	Description	Median (Range) or levels (for factors)
Nest height	Height of the rim of the nest cup above ground level (m)	1.350 (0.300 – 2.300)
Horizontal depth	Shortest horizontal distance of the edge of the nest cup to the nearest hedgerow edge (m). A hedgerow edge is defined as the beginning of dense thick twigs, as determined by using the weight of a cricket ball to move any light vegetation out of the way and determine where the hedgerow ends. Thus, widely spaced, thin twigs do not constitute the edge, whereas dense thick twigs do.	0.550 (0.050 – 1.800)
Vertical depth	Shortest vertical distance of the nest rim to the top of the hedgerow (m)	1.194 (0.200 – 3.650)
Nest volume	Maximum vertical x horizontal dimensions (cm ³)	1040.0 (117.8 – 5542.0)
Primary species ^a	Primary supporting woody plant species	Blackthorn, Bramble, Hawthorn, Rose and Other
Surrounding species ^a	Surrounding vegetation not supporting the nest but offering protection	Bramble, Ivy, Rose, None and Other
Cross-sectional area	Hedgerow height at the nest x hedgerow width at the nest (each \pm 5 cm; area in m ²). Height and width were measured so as to include woody hedgerow vegetation and recent growth, but exclude trees	5.57 (1.23 – 22.5)
Light penetration	Assessed using two light meters (Wavetek Metreman LM631; range 0.01 – 20,000	0.011 (0.001 – 0.226)

Lux) to assess the amount of light penetration above the nest relative to light levels outside the hedge. Only nests for which simultaneous measures from inside and outside the hedgerow (egg n=158; chick n=147) were included in analyses as the variable used was the ratio of light level above the nest to light level outside the hedgerow

Horizontal visibility ^{ab}	Maximum value of a measure from each side of the hedgerow. Assessed using a 14cm x 14cm black card containing a 5 x 5 regular grid of white circles (diameter 12 mm). The card was placed adjacent to each side of the nest (parallel to the hedgerow side) and viewed from three different angles (-45°, 90° and +45° to the nest). The number of circles that were at least 75% visible at each angle was summed for each side of the hedgerow, to give a measure (range 1-75) of horizontal visibility on each side of the hedgerow.	4.11 (0-23)
Vegetation cover ^a	% vegetation cover above the nest assessed using a digital photograph taken by placing a Casio EX-Z3 digital camera (set at the widest field of view) on the base of the nest facing vertically upwards. Photoshop software (v 7.0.1) was used to determine the proportion of pixels of sky in the image and thus the % vegetation cover above the nest	96.23 (70.40-99.99)
Small ball accessibility	Accessibility assessed by attempting to manoeuvre a baseball (circumference 30.5 cm) from the edge of the hedgerow to the nest by any route above or level with the nest without breaking any woody vegetation. We assessed whether or not the ball could reach the nest	Yes or No

Large ball accessibility	Accessibility assessed by attempting to manoeuvre a football (circumference 69 cm) from the edge of the hedgerow to the nest by any route above or level with the nest without breaking any woody vegetation. We assessed whether or not the ball could reach the nest	Yes or No
Corvid distance	Distance to nearest corvid nest (m)	270 (40 – 1350)
Wood distance	Distance to nearest wood, copse, spinney or woodland strip (at least 5m wide)	170 (5 – 645)
Vantage distance	Distance to nearest tree, pylon, telegraph pole or any other vantage point >5m in height (m)	40 (3-300)
Corvid abundance	Mean abundance of corvids per km, per site per year.	7.167 (0 – 28.830)

667

668 ^a variable included in nest site selection analysis

669 ^b visibility on each side of hedgerow included in nest site selection analysis as two separate variables but combined (as maximum visibility from
670 either side of the hedgerow) for nest failure analyses

671

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673

674

b)

Variable	Description	Median (Range)
Aspect	Aspect of hedgerow	E-W, N-S, NE-SW and SE-NW
Hedgerow management	Category: remnant (REMN), recently laid (RECE), mechanically cut (MECH), trimmed but dense (TRIM), overgrown and leggy (lacking branches and foliage in the bottom 1 m of the hedgerow; LEGG) or uncut and stock-proof (STOC) (from Bickmore, 2002; see Table 7 for further details of structure)	Leggy, Mechanically trimmed Remnant, Recently laid, Trimmed but dense, Stock-proof
Cutting style	Hedgerow shape. Categorised as: 'A' shaped (at least 2 cuts), Chamfered (at least 4 cuts), box (at least 3 cuts), free growth on top (only two sides cut), or free growth all round (uncut)	'A' shaped, Chamfered, Box, Free growth on top, or Free growth all round
% gaps	The % gaps ($\pm 5\%$) within the 30m hedgerow section	0 (0 – 30)
Number of trees	Number of trees >10m in height within 30m hedgerow section	0 (0 – 6)
Recent trim	Whether or not a hedgerow had been trimmed in the last 5 years	Trimmed or not
Last cut	Years since last cut	1 (1 – 4)
Margin width	Average width of herbaceous vegetation on both sides of the hedgerow (± 1 m)	2.0 (1.0 – 3.5)

675

676 Table 2.

a <i>Base model</i>	Egg stage					Chick stage				
	Slope	SE	df	χ^2	p	Slope	SE	df	χ^2	p
First egg date/Hatch date	-0.014	0.007	1	4.529	0.038	-0.013	0.006	1	5.395	0.020
First egg date ² /Hatch date ²	-0.001	0.001	1	0.509	0.476	-0.001	0.001	1	1.753	0.186
Clutch size/Brood size	-0.892	0.168	1	28.617	<0.001	0.261	0.172	1	2.459	0.117
Species			5	6.356	0.273			4	1.039	0.904
Year	0.066	0.351	1	0.036	0.850	-0.078	0.337	1	0.053	0.818
<i>Univariate tests</i>										
Nest height	0.034	0.555	1	0.004	0.950	-0.620	0.561	1	1.226	0.268
Horizontal depth	-0.228	0.465	1	0.239	0.625	0.678	0.793	1	0.713	0.399
Vertical depth	-0.209	0.251	1	0.716	0.398	0.046	0.277	1	0.027	0.870
Nest volume	0.092	0.217	1	0.177	0.674	0.063	0.254	1	0.063	0.802
Primary species			4	8.461	0.076			4	2.887	0.577
Surrounding species			4	0.657	0.957			4	1.618	0.806
Cross sectional area	-0.012	0.043	1	0.074	0.786	-0.062	0.274	1	0.051	0.821
Light penetration	4.552	4.318	1	1.024	0.312	-3.760	5.898	1	0.443	0.506
Horizontal visibility	0.162	0.170	1	0.883	0.347	-0.041	0.187	1	0.049	0.825
Vegetation cover	-0.886	1.318	1	0.445	0.505	0.987	1.375	1	0.533	0.466
Small ball accessibility	0.095	0.413	1	0.054	0.817	0.942	0.541	1	3.627	0.057
Large ball accessibility	-0.062	0.327	1	0.036	0.850	-0.225	0.365	1	0.392	0.531

Corvid distance	-0.462	0.265	1	2.965	0.085	0.077	0.231	1	0.111	0.739
Vantage distance	-0.002	0.003	1	0.691	0.406	0.290	0.132	1	4.680	0.031
Corvid abundance	-0.014	0.029	1	0.248	0.619	-0.010	0.268	1	0.002	0.969

677

<i>Base model</i>	Egg stage					Chick stage				
	Slope	SE	df	χ^2	p	Slope	SE	df	χ^2	p
First egg date/Hatch date	-0.013	0.006	1	0.860	0.354	-0.014	0.006	1	6.395	0.011
First egg date ² /Hatch date ²	-0.001	0.001	1	1.093	0.296	-0.001	0.001	1	0.366	0.545
Clutch size/Brood size	-0.877	0.169	1	25.176	<0.001	0.334	0.174	1	3.977	0.046
Year	0.411	0.369	1	1.666	0.197	-0.492	0.319	1	2.330	0.127
Species			5	4.469	0.484			5	3.411	0.637
<i>Univariate tests</i>										
Aspect			3	4.095	0.251			3	0.120	0.989
Hedgerow management			5	3.481	0.626			5	19.132	0.002
Cutting style			4	3.324	0.505			4	7.587	0.108
% gaps	-0.031	0.035	1	0.887	0.346	0.021	0.026	1	0.654	0.419
Number of trees	0.116	0.320	1	0.131	0.718	-0.643	0.363	1	3.444	0.063
Recent trim	0.881	0.656	1	2.108	0.147	-1.047	0.441	1	4.990	0.025
Last cut	-0.400	0.228	1	3.265	0.071	0.259	0.149	1	2.772	0.096
Margin width	0.081	0.250	1	0.105	0.746	-0.059	0.262	1	0.051	0.821

678

679 Table 3.

680

Variable	df	χ^2	p
Vegetation cover	1	31.189	<0.001
Horizontal visibility (non-nest side)	1	18.568	<0.001
Horizontal visibility (nest side)	1	17.751	<0.001
Surrounding species	5	20.773	<0.001
Primary species	4	9.845	0.043

681

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685

686 Table 4.

687

a Variable	df	Egg stage		Direction of effect	df	Chick stage		Direction of effect
		χ^2	p			χ^2	p	
First egg date/Hatch date	1	0.534	0.465	-ve	1	7.737	0.005	-ve
Clutch size/Brood size	1	23.462	<0.001	-ve	-	-	-	N/A
Corvid distance	1	2.965	0.085	-ve	-	-	-	N/A
Vantage distance	-	-	-	N/A	1	4.52	0.034	+ve
Small ball accessibility	-	-	-	N/A	1	3.656	0.056	See Fig 2

688

b Variable	df	Egg stage		Chick stage		
		χ^2	p	df	χ^2	p
Primary species	4	6.094	0.192	-	-	-
Vegetation cover x Corvid abundance	1	0.461	0.497	1	0.515	0.473
Vegetation cover x Corvid distance	1	0.001	0.972	1	0.334	0.563

689

690

691 Table 5.

	High access, high corvid	Low access, high corvid	High access, low corvid	Low access, low corvid
Predicted egg stage DSR	0.953 (0.953)	0.953 (0.953)	0.981 (0.981)	0.981 (0.981)
Predicted chick stage DSR	0.977 (0.963)	0.991 (0.985)	0.942 (0.963)	0.976 (0.985)
Predicted egg-stage survival	0.533 (0.533)	0.533 (0.533)	0.781 (0.781)	0.781 (0.781)
Predicted chick stage survival	0.743 (0.609)	0.886 (0.824)	0.461 (0.609)	0.728 (0.824)
Predicted overall nest survival	0.396 (0.325)	0.472 (0.439)	0.360 (0.476)	0.569 (0.644)

692

693

694 Table 6.

a

Variable	Egg stage			Direction of effect	Chick stage			Direction of effect
	df	χ^2	p		df	χ^2	p	
First egg date/Hatch date	-	-	-	N/A	1	10.155	0.001	-ve
Clutch size/Brood size	1	37.636	<0.001	-ve	1	1.388	0.239	+ve
Hedgerow management	-	-	-	N/A	5	12.200	0.032	See Fig 3
Last cut	1	3.265	0.071	-ve	-	-	-	N/A

695

b

Variable	Egg stage			Chick stage		
	df	χ^2	p	df	χ^2	p
Recent trim	-	-	-	1	0.744	0.389
Last cut	-	-	-	1	0.148	0.700
Number of trees	-	-	-	1	0.868	0.352

696

697

698

699 Table 7.

700

701 a)

Variable	High predation risk	Low predation risk	χ^2	p
Vegetation cover	93.77 ± 0.62	93.85 ± 0.41	0.15	0.70
Horizontal visibility (nest side)	3.97 ± 0.53	3.72 ± 0.28	1.38	0.24
Horizontal visibility (non nest side)	1.10 ± 0.22	0.68 ± 0.10	0.01	0.95
Horizontal visibility	2.54 ± 0.32	2.20 ± 0.16	0.53	0.47
Small ball accessibility (%)	91.3 ± 0.03	81.5 ± 0.03	3.03	0.08

702

703 b)

Variable	High predation risk	Low predation risk	χ^2	p
Vegetation cover	90.72 ± 0.42	91.79 ± 0.23	0.01	0.96
Horizontal visibility (nest side)	4.15 ± 0.21	3.08 ± 0.10	3.13	0.08
Horizontal visibility (non nest side)	2.02 ± 0.15	0.98 ± 0.06	18.67	<0.01
Horizontal visibility	3.08 ± 0.16	2.03 ± 0.07	6.63	0.01

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706 Appendix 1. Correlation matrix for all continuous variables highlighted as potentially important by univariate analysis. Figures represent the
707 correlation co-efficient from a Pearson's product moment test. Correlations significant at $p < 0.05$ are highlighted in bold.

708

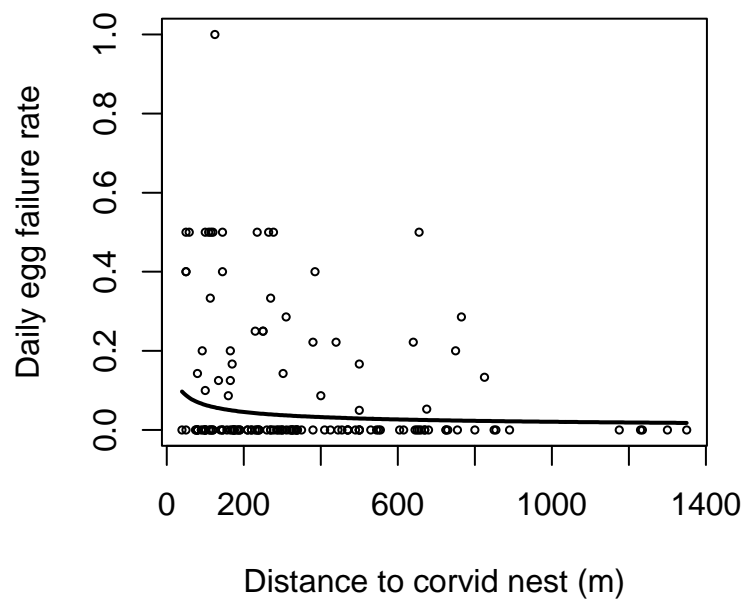
709 1)

	Horizontal depth	Vegetation cover	Horizontal visibility	Corvid abundance	Corvid distance
Vegetation cover	-0.04	-	-	-	-
Horizontal visibility	0.05	-0.35	-	-	-
Corvid abundance	-0.02	0.01	-0.05	-	-
Corvid distance	-0.07	-0.02	-0.06	-0.18	-
Vantage distance	-0.03	-0.02	-0.02	-0.10	0.04

710

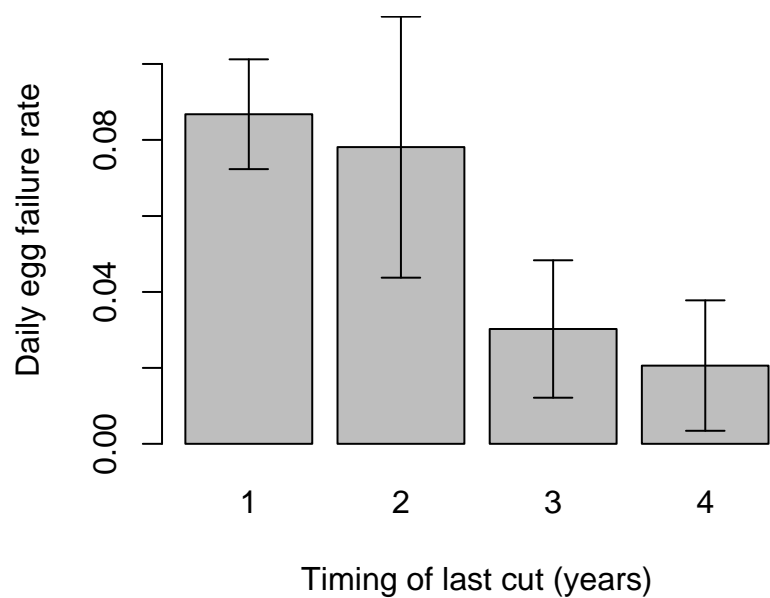
711

712 Appendix 2. Effect of distance from the nearest corvid nest on egg stage failure rate. Points show raw data; line is predicted from the final model
713 (Table 4) assuming mean hatch date and brood size.
714



715
716

717 Appendix 3. Effect of timing of last cut on egg stage nest failure rate. Bars show raw means \pm 1 SE; last cut is analysed as a continuous variable but
718 displayed categorically for clarity
719



720
721