

This is an Open Access document downloaded from ORCA, Cardiff University's institutional repository: <https://orca.cardiff.ac.uk/id/eprint/137380/>

This is the author's version of a work that was submitted to / accepted for publication.

Citation for final published version:

Shewring, Mike, Jenks, Paddy, Cross, Anthony V., Vaughan, Ian P. and Thomas, Robert J. 2020. Testing for effects of tail mounted radio tags and environmental variables on European Nightjar *Caprimulgus europaeus* nest survival. *Bird Study* 67 (4) , pp. 429-439. 10.1080/00063657.2021.1927979

Publishers page: <https://doi.org/10.1080/00063657.2021.1927979>

Please note:

Changes made as a result of publishing processes such as copy-editing, formatting and page numbers may not be reflected in this version. For the definitive version of this publication, please refer to the published source. You are advised to consult the publisher's version if you wish to cite this paper.

This version is being made available in accordance with publisher policies. See <http://orca.cf.ac.uk/policies.html> for usage policies. Copyright and moral rights for publications made available in ORCA are retained by the copyright holders.



Testing for effects of tail mounted radio tags and environmental variables on European Nightjar (*Caprimulgus europaeus*) nest survival

Mike Shewring^{a*}, Paddy Jenks^b, Anthony V. Cross^c, Ian P. Vaughan^a and Robert J. Thomas^{a,d}

^a School of Biosciences, Cardiff University, The Sir Martin Evans Building, Museum Avenue, Cardiff, CF10 3AX

^b Aderyn Ecology Ltd, Brynelwyn, Hebron, Whitland, Carmarthenshire, SA34 0JS

^c Samaria, Nantmel, Llandrindod Wells, Powys, LD1 6EN

^d Eco-explore Community Interest Company www.eco-explore.co.uk

* Correspondence author: Mike.shewring@gmail.com

Keywords: Radio tag, Nightjar, Nest survival

Short Title: Effects of radio tags on Nightjar nest survival

19 **Summary**

20 Capsule – Monitoring of European Nightjar *Caprimulgus europaeus* nest sites over
21 multiple years (2013-2019) produced no evidence of a negative effect of tail-mounted
22 radio tag deployment on nest success.

23 Aims – To test whether nest success of European Nightjar was affected by radio tag
24 deployment.

25 Methods – The breeding parameters of European Nightjar were monitored at the
26 Brechfa West Wind Farm, Carmarthenshire, Wales, from 2013 to 2019. A total of 85
27 nests were located through a combination of capture and radio tracking of breeding
28 individuals, and direct observation combined with focused searching. All located nests
29 were subsequently monitored thorough a combination of visual checks and trail camera
30 deployment until their natural conclusion.

31 Results – No evidence was identified to support a negative effect of tail mounted radio
32 tag deployment on the nest success of European Nightjar. However, nesting success (1
33 or more chicks fledged) was positively associated with mean temperature during the
34 nesting period, although the strength of this effect varied through time.

35 Conclusion – The use of tail mounted radio tags on European Nightjar has no negative
36 effect on nest survival.

Introduction

The marking and tagging of birds are widespread and important methods that have informed studies of many aspects of animal ecology, including migration, foraging behaviour and physiological ecology (Bodey et al. 2017). The techniques used for such marking are continuously evolving, and have been used in some form for many decades. The extra mass that these devices impose, the tag configuration and attachment method used has, however, been a cause for concern, especially for relatively heavy devices such as radio tags, GPS devices and geolocators (e.g. Bowlin et al. 2014). The deployment of such devices has been shown in some cases to reduce survival, inhibit parental care (Bodey et al. 2017), induce potentially costly behavioural modifications (Vandenabeele et al. 2014), or reduce the probability of nesting (Barron et al. 2010). Several mechanisms for such effects have been identified including; increased energetic costs of flight through drag (Bowlin et al. 2010), reduced foraging success (Wanless et al. 1988), impacts on young through reduced provisioning (Robert et al. 2006) and increased thermoregulatory costs due to feather loss and skin damage (Hines and Zwickle 1985). Although it is likely that such affects are in many cases species specific with other studies identifying few, if any effects (e.g. Bell et al. 2017, Brlik et al. 2020).

In an attempt to overcome such device effects, the research community has adopted rules of thumb for the design of tagging studies, such as the ‘5% rule’. This dictates a maximum tag mass limit of 5% of a bird’s body mass (Brander & Cochran 1969). The figure of 5% has been considered too high by some authors or for some taxa; for example Kenward (2001) suggested a limit of no more than 3%, supported by studies of nest abandonment in albatross and petrel species (Phillips et al. 2003, Casper 2009).

In recent years, further research has shown a simple percentage mass rule of thumb is

likely to be over-simplified. For example, various studies have shown that factors such as device-induced drag (Vandenabeele et al. 2013), tag shape and attachment location (Kay et al. 2019) are also critical considerations. These considerations, coupled with the apparently species-specific nature of tag effects, highlight the importance of testing for tagging impacts on individual species.

European Nightjar *Caprimulgus europaeus* (henceforth “nightjar”) breeding in Welsh upland coniferous forest, are difficult to monitor using conventional survey techniques due to their crepuscular nature, cryptic camouflage, and low density population (Cross et al. 2005, Gilbert et al. 1998). Therefore, a combination of radio tracking and observational nest finding methods have generally been utilised together for such studies at upland sites in Wales.

Radio transmitters and GPS devices suitable for deployment on nightjar have been available for some time, and have been widely used in breeding studies, most often as tail mounted devices (e.g. Alexander et al. 1990, Cross et al. 2005, Evens et al. 2018). Despite their widespread use in studies of breeding nightjars (e.g. Sharps et al. 2015, Evens et al. 2017) there is to our knowledge no published study of the effects of such tag deployment on breeding success. It is, however, critical that such effects should be investigated so that risks can be evaluated and minimised (Wilson et al. 2006, Casas et al. 2015).

An additional challenge in evaluating tag effects is to distinguish them from environmental impacts on survival or breeding success due to factors such as habitat quality or weather. Previous studies on nest survival in nightjars have identified probable effects of weather on nest survival (English et al. 2018) and similar effects are widely documented from studies in other species (e.g. Miller et al. 2017, Martin et al.

2017). As such, it is critical in studies of tag effects to account for such variables to accurately gauge any evidence of effects. In the present study, we therefore considered tag effects together with a set of environmental variables that we hypothesised may influence nightjar breeding success.

The present study directly compares observed nesting success of tagged birds and untagged birds, in order to investigate the potential effects of tag deployment and environmental variables on nesting nightjars. These data have been collected as part of on-going ecological impact monitoring requirements associated with the Brechfa West Wind Farm development. The data set includes nest record data from the study site during the pre-development, construction and operational phases of the wind farm.

Methods

Study Species

Nightjars are ground nesting birds that typically lay two eggs (occasionally one egg) and usually produce two broods per breeding season (Holyoak et al. 2001). The nightjar is usually thought of as a heathland species, but in Wales they mainly breed in clear-fell forestry (i.e. recently felled forestry, before substantial re-planting / re-growth), check coupes (i.e. stands of stunted tree growth) and recently restocked conifer plantations (Conway et al. 2007). Male nightjars establish breeding territories within the study area in May; females arrive in mid-May and subsequently become paired with established territorial males.

Nightjars are of conservation concern due to historic population declines and range contraction (Balmer et al. 2013, Hagemeijer & Blair 1997). The nightjar is an Annex 1 species in the EU (Council Directive 2009/147/EC), has Amber status in the UK (Bird

of Conservation Concern; Eaton et al. 2015) and is listed under Section 7 of the Environment (Wales) Act 2016. The nightjar population in Wales has been increasing since at least 1981 (Morris et al. 1994), possibly due to increased habitat availability following the maturation and felling of plantations that were planted in the 1950's.

Study Site

This study utilises nest data from Brechfa Forest (South Wales, UK – Latitude 51.967432, Longitude -4.1964175), a commercial plantation forestry managed by Natural Resources Wales on behalf of Welsh Government. The forest is dominated by dense Sitka Spruce *Picea stitchensis* forest blocks (coupes), interspersed with recently felled areas around wind turbines, and with semi-natural woodland along watercourses.

Topography and forest age at this site has enabled observational nest finding to be relatively successful during recent commercial ecological monitoring work.

Nest data collection

The inclusion of nightjar in species protection legislation ensures that nightjar nest locations are protected from damage/ destruction under the Wildlife and Countryside Act (1981). Suitably licensed and experienced individuals undertook all tagging and nest monitoring visits completed in this study.

Territory identification

Active territories were located by systematic searches in areas of suitable habitat, and were confirmed by observation of pairs or of displaying males, which produce a distinctive “churring” call (Ferguson-Lees et al. 2011).

Observational nest location

Nest searching commenced annually in late May, and continued until August. Active territories were systematically watched on multiple occasions by multiple observers at dusk, and visual cues were used to guide follow up nest searches (Langston et al. 2009). Subsequent nest searches consisted of detailed visual inspection in areas of observed nightjar activity during dusk watches, with searchers aiming to pass within 3-4 metres of any point within the search area.

Radio tracking nest location

Where observation of active nightjar territories yielded little information, or nest searches were unsuccessful, or where pairs were considered likely to attempt a 2nd brood, then these territories/ pairs were targeted for radio tagging effort. Mist nets were set up in the vicinity of identified territories, and male nightjars were then tape lured into the mist nets by playing the species' typical territorial song (Squire and Alexander 1981). Tape luring proved less effective at attracting incubating females. Females were caught by mist-netting at favoured feeding sites, or by trapping at known 1st brood nests (found by field observation) to allow radio tracking to 2nd brood nests.

Captured birds were fitted (under licence) with PIP-3 radio-transmitters (from Lotek Ltd – as per Alexander and Cresswell (1990)), attached to the base of one of the central tail feathers. Attaching the radio-transmitters in this way ensures that they are shed during post-breeding moult in the wintering grounds, and thus does not affect the birds during their spring migration. The tags used in this study each weighed 1.2g, male nightjars weighed between 60.2–87.0 g (n=34), and females weighed between 69.0–100.8 (n=23) - so tags weighed 1.34–1.99% of male body weight, and 1.19–1.72% of female body weight.

Tags were deployed across the breeding season, with tagging dates ranging between the 3rd of June and 24th of July. The median tagging date was the 25th of June; the mean tagging date was the 25th of June for females and 27th of June for males. Tags were deployed both prior to and after nests were located; 19 of the 39 tagged females were tagged after their nest was located, as were 11 of the 25 tagged males.

Following the identification of active nests through either observation or radio tracking, all nests were monitored to their natural completion (fledging or nest failure) by an experienced nightjar fieldworker, using regular (~weekly) nest site visits. Nests were classified as either successful or failed, based on a combination of the timing of nest visit records and available evidence at the nest site and within the territory (i.e. flying young present).

Weather data

In order to account for the influence of weather on nesting success, data from the closest available weather station (Pembrey; 51.7144117°N, -4.366197°E, approximately 30km south of the study site) was obtained using the GSODR package (Sparks, Hengl, and Nelson 2017) using R software version 3.6.1 (R Core Team 2019), implemented via R Studio (RStudio team 2018). The GSODR package provides automated downloading, parsing and cleaning of Global Surface Summary of the Day (GSOD) (United States National Oceanic and Atmospheric Administration National Climatic Data Center) weather data. This provided daily rainfall (mm) and mean temperature (T_m, °C). Data manipulation and visualisation was undertaken using the R libraries tidyverse (Wickham et al. 2019), lubridate (Groelmund & Wickham 2011) and ggplot2 (Wickham, 2016). Mean temperature and mean precipitation were calculated for the

active period of each nest (laying date to last known presence) and utilised in subsequent analysis.

Statistical analysis

We performed all statistical analyses in R 3.6.0 (R Core Team 2019). In order to account for the inherent bias in nest studies arising from the lower detection probability of failed nests (due to their shorter time available for potential observation), we estimated daily nest survival rates - DSR (Mayfield 1975, Dinsmore et al. 2002) using RMark version 2.2.7 and MARK (Laake 2013, White and Burnham 1999).

Daily nest survival rates were estimated and modelled with selected covariates using the R package RMark version 2.2.7 (Laake 2013). We undertook model selection of nest survival models using an information theoretical approach based on the second-order Akaike information criterion for small sample sizes (AICc; Burnham and Anderson 2002).

A set of 193 biologically plausible models was derived, including additive effects of Julian day, nest age (as estimated based on hatch date, if available, or if not then using estimates based on egg floatation (Westerskov 1950) or observational information), brood (1st, 2nd, 3rd), year, mean rainfall within the relevant active nest period, mean temperature (Tm) within the relevant active nest period, the presence of windfarm construction activity (binary yes/no – nest active in year of construction activity), adult male tag status (tag status of the male associated with nest - binary yes/ no), adult female tag (tag status of the female associated with nest - binary yes/ no) and combined adult tag status (tag status of both adults associated with nest - binary yes/ no - i.e. both birds tagged or not). The candidate models also included the interaction between mean temperature and date, to help distinguish the effect of temperature from seasonality. The

combined adult tag status variable was included to account for potential synergistic effects of tagging both parents. All covariates were scaled prior to analysis, to have a mean of zero and a standard deviation of one. The set of candidate models also included a global model (containing all candidate independent variables) and a null model (containing no independent variables). Co-linearity between variables was determined using Pearson's correlation coefficient, and this identified low levels of correlation between candidate model variables. No candidate model variables exceeded the threshold correlation of 0.7 (Dorman et al. 2013) and all candidate variables were thus included in the analysis.

Models were ranked using AICc, and the Δ AICc values and Akaike weights (w_i) were used to infer support for each of the candidate models (Appendix A). In our model selection analysis, no single model was clearly better than all others, and to account for model selection uncertainty, models within two AICc units of the top model, were selected for model averaging, as this can provide a robust means of obtaining parameter estimates in such scenarios (Burnham & Anderson 2002, Grueber et al. 2011, Harrison et al. 2018). A weighted average of the parameter estimates (and 95% confidence limits) was calculated for all of the variables contained in the top models, using the package MuMIn (Grueber et al. 2011, Barton 2018, Mwangi et al. 2018) (Table 2). Parameters were considered statistically significant where their model-averaged 95% confidence limits did not span zero.

Overall nest survival was calculated from predictions daily of nest survival rate (DSR) made by the final, averaged model. These were converted to the overall nest success by assuming a 36 day standard nesting period (DSR^{36}) from the median nest initiation date. Variance in the nest survival estimates were obtained using the delta method (Powell 2007).

The same suite of models was also re-run using a subset of the data representing the egg stage and chick stage respectively. Whilst this reduced the sample size for these models, it was considered to potentially provide greater insights into potential tag effects during the two different breeding stages, given the likely different energetic demands and behaviours associated with each stage. Due to convergence problems, because of small sample sizes, the chick stage models were run without the year parameter.

Results

Nest finding and monitoring

Eighty five nightjar nests were located over the course of the study (2013-2019); sixty-one of these were located through direct observation of adult behaviour, and twenty four were located using radio tracking. Median nest initiation date was 16th June (range = 27th May – 27th July). In total, 59 nests were confirmed first brood nests and 13 confirmed second brood nests. Two nest attempts were also recorded as ‘third brood’ nests, although these were a result of early failure of previous nesting attempts (1st or 2nd brood) and thus are replacement clutches; they have nevertheless been referred to as third brood nests for the ease of reference. Brood number could not be confirmed at 11 of the located nests.

We found nests at different stages of development: 52 (61.1%) during incubation and 33 (38.8%) were found during the nestling period. From all of the nests, 52 fledged at least one chick, whilst the remainder (33) failed, with 15 at the egg stage and 18 failing at the chick stage. A summary of nest success and the number of nests with attending tagged adults is provided in Table 1, whilst Table 2 details the breakdown of nests attended by tagged adults, by adult sex, and brood number.

Nest survival

In our model selection analysis, there were three models within 2 AIC units and they contained the following variables – nest age, female tag status, adult tag status, temperature, precipitation and Julian day (Table 1). In order to account for model selection uncertainty, a conditional weighted average (averaged over only the models containing those parameters) and a full weighted average (all models using zero value for parameters not present) of the parameter estimates and 95% confidence limits was calculated for all of the variables contained in the top three models (conditional weighted averages in Table 4, and full weighted averages in Table 5). Full weighted model average parameter estimates are reported below, along with the standard error (SE).

Estimated average daily nest survival (\pm SE), across all years and tag treatments, was 0.986 (\pm 0.008). This extrapolates over the 36-d nesting cycle to an average annual nest success rate of 0.63 (\pm 0.18).

The same suite of models run on subsets of the full data set for the egg stage of the nesting cycle failed to identify any parameters as having an important effect on DSR and identified no detectable difference between DSR for tagged nests vs. untagged nests at either stage. Top selected models and model averaged coefficients for the identified top models are presented in supplementary materials Appendix B – Table B1 to Table B3. The same suite of models for the chick stage of the nesting cycle failed to converge due to low sample sizes.

Radio tag effects

There was no evidence for tags reducing nesting success. Although two of the three top models of daily nest survival rate included either female tag status or adult tag status variables, these all indicated a positive relationship that was not significant: a result confirmed by the averaged model ($\beta_{fm_tag} = +0.158 \pm 0.429$; $\beta_{f_tag} = +0.445 \pm 0.499$).

Overall DSR rates for untagged female attended nests and tagged female attended nests were $0.984 (\pm 0.010 [SE])$ and $0.990 (\pm 0.006 [SE])$ respectively (Figure 1). Estimated DSR for untagged and tagged adult attended nests (male or female) were very similar, at $0.986 (\pm 0.010)$ and $0.991 (\pm 0.006)$ respectively (Figure 2).

Nest age and Julian day

The top model of daily survival rate included significant effects of Julian day and nest age (initiation date; Table 3). Nest survival rate of nightjar decreased as the season progressed (model-averaged parameter \pm SE; $\beta_{\text{Julian day}} = -0.07 \pm 0.023$) but increased with the age of the nest ($\beta_{\text{nest initiation date}} = +0.072 \pm 0.028$). Over the nesting season, model averaged DSR ranged from $0.988 (\pm 0.012)$ on day 1 of the nesting season (28th May), to $0.986 (\pm 0.013)$ on day 81 (17th August).

Weather effects

Initial data exploration of weather data identified a weak positive correlation between relative humidity (surrogate for cloud cover) and minimum temperature ($\tau = 0.177$), with a similar positive correlation noted between relative humidity (surrogate for cloud cover) and minimum temperature ($\tau = 0.219$). As such, weather effects should be interpreted in this context.

The top models together provide good evidence that temperature has an important effect on nest success, as temperature was consistently selected in top models. Alternative models without this variable did not receive strong statistical support and were at least 2.7 AICc units from the top model.

Average temperatures during active nest periods over the study years ranged from 12.8 to 19.5 °C, and model predictions showed a positive relationship with temperature ($\beta_{m_temp} = +2.501 \pm 1.083$; Table 5). As confidence intervals did not include zero, this is considered a statistically significant effect. The top models also consistently incorporated an interactive effect between temperature and Julian day on DSR, and this interaction term appeared in all top models.

Model estimates show a negative parameter for the temperature x Julian day interaction term ($\beta_{m_temp: Time} = -0.035 \pm 0.019$; Table 5). As confidence intervals include zero this is however not considered to be a statistically significant interaction. Despite this, the important effects of temperature on DSR must be viewed in the context of its relationship with time, as its inclusion in top models suggests that the magnitude of the positive effect temperature is potentially conditioned on Julian day. This interaction term describes how the effect of temperature varies through time, and indicates that the positive effect of temperature on DSR depends on the Julian day and decreases through the breeding season. This may be due to threshold effects of temperature, as temperature exhibits a non-linear relationship with time through the season, or could be due to further interactions with the stage of nest development – i.e. nests are more likely to have chicks later in the season.

Predicted DSR increased from 0.36 (95% CI 0.03 to 0.920) to 0.999 (95% CI 0.994 to 0.999) over the recorded temperature range (12.8 to 19.5 °C), for a nest initiated on the

316 16th June (median date of nest initiation) assuming average values for the other
317 covariates (Figure 3).

318 Mean daily rainfall during the active nest periods ranged from 0 mm to 10.55 mm, with
319 a mean of 2.10 mm. No significant effect of precipitation on DSR was detected (β
320 m_prcp 0.509 ± 0.382 , Table 5); confidence intervals for this estimate spanned zero,
321 suggesting a lack of any statistically significant effect.

322

Discussion

Mean temperature and nest age were identified as important factors associated with annual reproductive success of nightjars at the study site (see Table 2 and Table 3). No evidence for a negative effect of tagging was identified by the models of nest survival, and this is consistent with the raw data, where mean nest success across the seven years of the study was 61% for nests attended by one or two tagged parents, and 62% for nests attended by untagged parents. This provides good evidence that the continued use of radio tagging to facilitate nest finding is unlikely to impact nest survival.

Models identified no evidence that any of the other candidate variables affected nesting success, with no statistically significant effect noted for Julian day, precipitation, brood or year of construction. Previous studies of nightjar nest success have focused on the effects of recreational disturbance (e.g. Langston et al. 2007; Lowe et al. 2014) and in general have identified a negative effect of such disturbance, but have not investigated relationships with tagging, time or weather. Langston et al. (2007) estimated overall nesting success to be 39% in the Dorset heathlands, whereas Lowe et al. (2014) estimated success at 53% in Nottinghamshire plantation forestry sites. Overall nest success estimates of 61-62% from the upland forestry habitats of the Brechfa Forest study site thus compare favourably with reported nest success rates from other studies.

A significant effect of nest age on daily survival rate was identified, with DSR increasing with nest age within individual breeding attempts. Similar variation in chick/nest survival with age has been observed in other species (e.g. Grant et al. 2005, McDonald et al. 2016, English et al. 2018, Maziarz et al. 2019, Zhao et al. 2020). The positive pattern noted here could be due to older chicks having greater resilience to poor weather and being more able to overcome the nutritional and thermoregulatory burden

of poor weather, as has been suggested for Northern Bobwhite chicks (*Colinus virginianus* – Terhune et al. 2019).

The identified positive association between temperature and nest survival is unsurprising, as during periods of low temperature nests can fail due to chick starvation (pers, obs) and similar positive effects of temperature have been made in North American nightjar species - whip-poor-will (*Antrostomus vociferous* - English et al. 2018). In general, young, downy chicks are likely to be less able to thermally regulate effectively (Du Rant et al. 2001, Newberry et al. 2018), and thus may be particularly vulnerable to adverse weather and predation. Young chicks will repeatedly call when chilled; this advertisement is likely to increase predation risk as has been observed in other bird species (e.g. Deardon 1999, Briskie et al. 1999, Ibanez-Alamo et al. 2012, Husby 2019, Gonchorova et al. 2019), and may form part of the mechanism by which low temperature leads to nest failure. In addition, moth activity is generally positively correlated with temperature (Holyoak et al. 1997), so a direct negative effect of cold weather on nest success through reduced food availability, would be expected though direct impacts on provisioning at the chick stage, or indirectly through reduced incubation intensity at the egg stage. Similar effects of temperature on chick survival have also been noted in a North American nightjar species (the Whip-Poor-Will - *Antrostomus vociferus*, English et al. 2018) with higher chick survival recorded on warmer nights.

It is surprising, however, that rainfall did not show a negative effect on nest survival, as nest failure due to hypothermia/starvation has previously been recorded following protracted heavy rain (pers. obs), and moth activity is generally negatively correlated with rainfall (Holyoak et al. 1997). One explanation may be the presence of a positive correlation between the minimum daily temperature (likely at night) and rainfall ($\tau =$

0.177), as during cloudy conditions night-time temperatures are usually higher than under clear skies. This may be particularly relevant for the dawn foraging period for nightjars, when at 300m elevation (as at the study site), the temperature is often below 10°C following a night of clear skies during the main breeding season (See Appendix C – Figure C1 and C2). Hence it may be that extreme rainfall events have a negative effect by causing direct chick mortality, as has been shown in White Stork (*Ciconia ciconia* - Tobolka et al. 2015) and Northern Wheatear (*Oenanthe oenanthe* - Oberg et al. 2015), but food availability is perhaps increased both when evenings are warm following sunny weather, and during cloudy, drizzly conditions, when both dusk and dawn foraging periods are relatively mild. This increase in food availability may lead to improved nest survival, as has been noted in other species (White Ibis *Eudocimus albus* - Herring et al. 2011, and Eurasian reed warbler *Acrocephalus scirpaceus* - Vafidis et al. 2016). However, more work is needed in this area, including collecting insect abundance data, to try to unpick the relationships between weather, insect abundance and nest survival (Shewring et al. in prep.).

Wind farm construction had no observable effect on the daily nest survival rate, and the year of construction variable was not selected in any of the top models. It is, however, worth noting that any effects of construction disturbance are likely to be influenced by the proximity of individual nests to construction activity. Such detailed data were not available to inform the current study, but would certainly be recommended in future studies focused on the effects of construction disturbance. In addition, there were deliberate attempts to limit construction effects on nightjar at the Brechfa windfarm (e.g. by using disturbance exclusion buffers around located nests) and as such, this conclusion is only relevant to construction where such mitigation procedures are implemented. In light of this, we would advise that this aspect of the analysis be treated

with the appropriate caution when interpreting the sensitivity of nightjar to construction disturbance.

It should however be noted that nest survival is a single metric for impact identification of tagging, and other effects of tag deployment on nightjar cannot be discounted based on the current study. It is certainly possible that tagging has affected foraging success and ranging behaviour, as has been noted in other species (e.g. Taylor et al 2001, Phillips et al. 2003), but any such effects have not fed through to detectable effects on nest survival. As such, we would recommend further study of tag effects in nightjar, especially where tagging is proposed for longer durations or where heavier tags are proposed.

In conclusion, the current study confirms the importance of weather effects on nightjar nest survival, particularly the positive effect of temperature. It also confirms the lack of observable tagging effects on nest survival when using tail mounted radio tags, and indicates that their continued use in nest finding studies is unlikely to have a negative impact on nest survival. Integrating these two conclusions leads us to recommend that future tagging studies adequately consider potentially confounding weather effects.

414 **Acknowledgements**

415 The authors would like to thank RWE renewables UK Ltd for the help and support
416 during fieldwork at Brechfa West Wind Farm and two anonymous reviewers for their
417 insightful and helpful comments. Mike Shewring would also like to thank the
418 Knowledge Economy Skills Scholarship (KESS2) and Wildlife Trust of South and West
419 Wales for their financial support.

References

- Alexander, I. and Cresswell, B. 1990. Foraging by Nightjars *Caprimulgus europaeus* away from their nesting areas. *Ibis*, 132: 568-574. doi:10.1111/j.1474-919X.1990.tb00280.x
- Balmer, D.E., Gillings, S., Caffrey, B.J., Swann, R.L., Downie, I.S. & Fuller, R.J. (eds). 2013. Bird Atlas 2007–11: the breeding and wintering birds of Britain and Ireland. BTO Books, Thetford. Banks, A.N., Coombes, R.H. & Crick, H.Q.P.
- Barron, D.G., Brawn, J.D. and Weatherhead, P.J. 2010. Meta-analysis of transmitter effects on avian behaviour and ecology. *Methods in Ecology and Evolution*, 1: 180-187. doi:10.1111/j.2041-210X.2010.00013.x
- Bodey, T.W., Cleasby, I.R., Bell, F., Parr, N., Schultz, A., Votier, S.C. and Bearhop, S.. 2018. A phylogenetically controlled meta-analysis of biologging device effects on birds: Deleterious effects and a call for more standardized reporting of study data. *Methods in Ecology and Evolution*, 9: 946– 955. <https://doi.org/10.1111/2041-210X.12934>
- Bowlin, M.S., Henningsson, P., Muijres, F.T., Vleugels, R.H.E., Liechti, F. and Hedenström, A. 2010. The effects of geolocator drag and weight on the flight ranges of small migrants. *Methods in Ecology and Evolution*, 1: 398-402. doi:10.1111/j.2041-210X.2010.00043.x
- Brander, R.B. & Cochran, W.W. 1969. Radio location telemetry. *Wildlife Management Techniques* (ed. R.H. Giles Jr), pp. 95–103. The Wildlife Society, Washington, DC.
- Brlík, V, Koleček, J, Burgess, M, et al. Weak effects of geolocators on small birds: A meta-analysis controlled for phylogeny and publication bias. *J Anim Ecol.* 2020; 89: 207– 220. <https://doi.org/10.1111/1365-2656.12962>

443 Briskie, J. V., Martin, P. R., & Martin, T. E. (1999). Nest predation and the evolution of
 444 nestling begging calls. *Proceedings of the Royal Society of London. Series B:*
 445 *Biological Sciences*, 266(1434), 2153-2159.

446 Burnham, K.P. & Anderson, D.R. 2002. *Model Selection and Multimodel Inference: A*
 447 *Practical Information-Theoretic Approach*, 2nd edition. Springer, Berlin.

448 Casas, F., Benítez-López, A., García, J.T., Martín, C. A., Viñuela, J., & Mougeot, F.
 449 2015. Assessing the short-term effects of capture, handling and tagging of sandgrouse.
 450 *Ibis*, 157(1): 115-124.

451 Casper, R. 2009. Guidelines for the instrumentation of wild birds and mammals. *Animal*
 452 *Behaviour*, 78(6): 1477-1483.

453 Conway, G., Wotton, S., Henderson, I., Langston, R., Drewitt, A. and Curriem, F. 2007.
 454 Status and distribution of European Nightjars *Caprimulgus europaeus* in the UK in
 455 2004, *Bird Study*, 54:1, 98-111. DOI: 10.1080/00063650709461461

456 Dinsmore, S.J., White, G.C. and Knopf, F.L. 2002. Advanced techniques for modeling
 457 avian nest survival. *Ecology*, 83: 3476-3488. doi:10.1890/0012-
 458 9658(2002)083[3476:ATFMAN]2.0.CO;2

459 Dearborn, D. C. (1999). Brown-headed cowbird nestling vocalizations and risk of nest
 460 predation. *The Auk*, 116(2), 448-457.

461 Dormann, C.F., Elith, J., Bacher, S., Buchmann, C., Carl, G., Carré, G., Marquéz,
 462 J.R.G., Gruber, B., Lafourcade, B., Leitão, P.J., Münkemüller, T., McClean, C.,
 463 Osborne, P.E., Reineking, B., Schröder, B., Skidmore, A.K., Zurell, D. and Lautenbach,
 464 S. (2013), *Collinearity: a review of methods to deal with it and a simulation study*

465 evaluating their performance. *Ecography*, 36: 27-46. doi:10.1111/j.1600-
466 0587.2012.07348.x

467 DuRant, S. E., Hopkins, W. A., Wilson, A. F., & Hepp, G. R. (2012). Incubation
468 temperature affects the metabolic cost of thermoregulation in a young precocial bird.
469 *Functional Ecology*, 26(2), 416-422.

470 Eaton, M., Aebischer, N., Brown, A., Hearn, R., Lock, L., Musgrove, A. & Gregory, R.
471 2015. Birds of Conservation Concern 4: the population status of birds in the UK,
472 Channel Islands and Isle of Man. *British Birds*, 108(12), 708-746.

473 English, P. A., Nocera, J. J., & Green, D. J. (2018). Nightjars may adjust breeding
474 phenology to compensate for mismatches between moths and moonlight. *Ecology and*
475 *evolution*, 8(11), 5515-5529.

476 Evens, R., Beenaerts, N., Witters, N. & Artois, T. (2017). Study on the foraging
477 behaviour of the European nightjar *Caprimulgus europaeus* reveals the need for a
478 change in conservation strategy in Belgium. *Journal of Avian Biology*, 48: 1238-1245.
479 doi:10.1111/jav.00996

480 Evens, R., Beenaerts, N., Ulenaers, E., Witters, N., & Artois, T. (2018). An effective,
481 low-tech drop-off solution to facilitate the retrieval of data loggers in animal-tracking
482 studies. *Ringling & Migration*, 33(1), 10-18.

483 Gilbert G, Gibbons D. W. and Evans J. 1998. Bird Monitoring Methods. RSPB
484 Sandy.

485 Goncharova, M. V., & Klenova, A. V. (2019). Siberian crane chick calls reflect their
486 thermal state. *Bioacoustics*, 28(2), 115-128.

487 Grant, T. A., Shaffer, T. L., Madden, E. M., & Pietz, P. J. (2005). Time-specific
 488 variation in passerine nest survival: new insights into old questions. *The Auk*, 122(2),
 489 661-672.

490 Grolemond, G., and Wickham, H. 2011. Dates and Times Made Easy with lubridate.
 491 *Journal of Statistical Software*, 40(3): 1-25. URL <http://www.jstatsoft.org/v40/i03/>.

492 Grueber, C.E., Nakagawa, S., Laws, R.J. and Jamieson, I.G. (2011). Multimodel
 493 inference in ecology and evolution: challenges and solutions. *Journal of Evolutionary*
 494 *Biology*, 24: 699-711. doi:10.1111/j.1420-9101.2010.02210.x

495 Hagemeyer, W. J., & Blair, M. J. (1997). *The EBCC atlas of European breeding birds*.
 496 Poyser, London, 479.

497 Harrison X.A., Donaldson L., Correa-Cano M.E., Evans J., Fisher D.N., Goodwin
 498 C.E.D., Robinson B.S., Hodgson D.J. & Inger R. (2018). A brief introduction to mixed
 499 effects modelling and multi-model inference in ecology. *PeerJ* 6:e4794
 500 <https://doi.org/10.7717/peerj.4794>

501 Hines, J. E., & Zwickel, F. C. (1985). Influence of
 502 radio packages on young blue grouse. *The Journal of wildlife Management*, Vol. 49,
 503 No. 4, 1050-1054.

504 Hensler, G. L., & Nichols, J. D. (1981). The Mayfield method of estimating nesting
 505 success: a model, estimators and simulation results. *The Wilson Bulletin*, 42-53.

506 Herring, G., Cook, M. I., Gawlik, D. E., & Call, E. M. (2011). Food availability is
 507 expressed through physiological stress indicators in nestling white ibis: a food
 supplementation experiment. *Functional Ecology*, 25(3), 682-690.

508 Holyoak, D. T. (2001). Nightjars and their allies: the Caprimulgiformes (Vol. 7).
509 Oxford University Press.

510 Holyoak, M., Jarosik, V., & Novak, I. (1997). Weather-induced changes in moth
511 activity bias measurement of long-term population dynamics from light trap samples.
512 *Entomologia Experimentalis et Applicata*, 83(3), 329-335.

513 Husby, M. (2019). Nestling begging calls increase predation risk by corvids. *Animal*
514 *Biology*, 69(2), 137-155.

515 Ibáñez-Álamo, J. D., Arco, L., & Soler, M. (2012). Experimental evidence for a
516 predation cost of begging using active nests and real chicks. *Journal of ornithology*,
517 153(3), 801-807.

518 Kay, W.P., Naumann, D.S., Bowen, H.J., Withers, S., Evans, B., Wilson, R., Stringell,
519 T., Bull, J., Hopkins, P., & Börger, L. (2019). Minimizing the impact of biologging
520 devices: Using computational fluid dynamics for optimizing tag design and positioning.
521 *Methods in Ecology and Evolution*, 10: 1222– 1233. [https://doi.org/10.1111/2041-](https://doi.org/10.1111/2041-210X.13216)
522 [210X.13216](https://doi.org/10.1111/2041-210X.13216)

523 Kenward, R.E. (2001). A manual for wildlife radio tagging. London, UK: Academic
524 Press.

525 Langston, R. H. W.; Liley, D.; Murison, G., Woodfield, E. & Clarke, R.T. (2007). What
526 effects do walkers and dogs have on the distribution and productivity of breeding
527 European Nightjar *Caprimulgus europaeus*? *IBIS*. Volume: 149, Supplement: 1, Pages:
528 27-36.

529 Lowe, A., A. C. Rogers, and K. L. Durrant. (2014). Effect of human disturbance on
530 long-term habitat use and breeding success of the European Nightjar, *Caprimulgus*
531 *europaeus*. *Avian Conservation and Ecology* 9(2): 6.

532 Laake, J.L. (2013). RMark: An R Interface for Analysis of Capture-Recapture Data with
533 MARK. AFSC Processed Rep 2013-01, 25p. Alaska Fish. Sci. Cent., NOAA, Natl. Mar.
534 Fish. Serv., 7600. Sand Point Way NE, Seattle WA 98115.

535 MacDonald, E. C., Camfield, A. F., Martin, M., Wilson, S., & Martin, K. (2016). Nest-
536 site selection and consequences for nest survival among three sympatric songbirds in an
537 alpine environment. *Journal of Ornithology*, 157(2), 393-405.

538 Martin, K., Wilson, S., MacDonald, E. C., Camfield, A. F., Martin, M., & Trefry, S. A.
539 (2017). Effects of severe weather on reproduction for sympatric songbirds in an alpine
540 environment: Interactions of climate extremes influence nesting success. *The Auk:*
541 *Ornithological Advances*, 134(3), 696-709.

542 Mayfield, H. (1975). Suggestions for Calculating Nest Success. *The Wilson Bulletin*,
543 87(4): 456-466.

544 Maziarz, M., Grendelmeier, A., Wesołowski, T., Arlettaz, R., Broughton, R. K., &
545 Pasinelli, G. (2019). Patterns of predator behaviour and Wood Warbler *Phylloscopus*
546 *sibilatrix* nest survival in a primaeval forest. *Ibis*, 161(4), 854-866.

547 Miller, M. W., Leech, D. I., Pearce-Higgins, J. W., & Robinson, R. A. (2017). Multi-
548 state, multi-stage modeling of nest-success suggests interaction between weather and
549 land-use. *Ecology*, 98(1), 175-186.

550 Morris, A., Burges, D., Evans A., Smith, K., & Fuller, R. (1994). The status and
551 distribution of nightjars *Caprimulgus europaeus* in Britain in 1992. A report to the
552 British Trust for Ornithology. *Bird Study*, 41(3): 181-191.

553 Mwangi, J., Ndithia, H.K., Kentie, R., Muchai, M. & Tieleman, B.I. (2018). Nest
554 survival in year-round breeding tropical red-capped larks *Calandrella cinerea* increases
555 with higher nest abundance but decreases with higher invertebrate availability and
556 rainfall. *Journal of Avian Biology*, 49: e01645. doi:10.1111/jav.01645

557 Newberry, G. N., & Swanson, D. L. (2018). Elevated temperatures are associated with
558 stress in rooftop-nesting Common Nighthawk (*Chordeiles minor*) chicks. *Conservation*
559 *physiology*, 6(1), coy010.

560 Öberg, M., Arlt, D., Pärt, T., Laugen, A. T., Eggers, S., & Low, M. (2015). Rainfall
561 during parental care reduces reproductive and survival components of fitness in a
562 passerine bird. *Ecology and Evolution*, 5(2), 345-356.

563 Powell, L.A. (2007). Approximating Variance of Demographic Parameters Using the
564 Delta Method: A Reference for Avian Biologists, *The Condor*, 109(4): 949–954.
565 <https://doi.org/10.1093/condor/109.4.949>

566 Phillips, R. A., Xavier, J. C., & Croxall, J. P. (2003). Effects of satellite transmitters on
567 albatrosses and petrels. *The Auk*, 120(4), 1082-1090.

568 RStudio Team. (2018). RStudio: Integrated Development for R. RStudio, Inc., Boston,
569 MA, USA. <http://www.rstudio.com/>.

570 Robert, M., Drolet, B., & Savard, J.P.L. (2006). Effects of backpack radio-transmitters
571 on female Barrow's Goldeneyes. *Waterbirds*, 29(1): 115-120.

572 Sparks, A, Hengl, T & Nelson, A. (2017). GSODR: Global Summary Daily Weather
573 Data in R. *Journal of Open Source Software*, 2(10), 177, doi:10.21105/joss.0017

574 Squire, T. & Alexander, I. 1981. Capture techniques for full grown Nightjars. *Ringers’*
575 *Bulletin*, 5: 132.

576 Taylor, S. S., Leonard, M. L., Boness, D. J., & Majluf, P. (2001). Foraging trip duration
577 increases for Humboldt penguins tagged with recording devices. *Journal of Avian*
578 *Biology*, 32(4), 369-372.

579 Taylor, G. & Thomas, A. (2002). Animal flight dynamics II. Longitudinal stability in
580 flapping flight. *Journal of Theoretical Biology*, 214: 351–370.

581 Terhune, T. M., Palmer, W. E., & Wellendorf, S. D. (2019). Northern bobwhite chick
582 survival and effects of weather. *The Journal of Wildlife Management*, 83(4), 963-974.

583 Tobolka, M., Zolnierowicz, K. M., & Reeve, N. F. (2015). The effect of extreme
584 weather events on breeding parameters of the White Stork *Ciconia ciconia*. *Bird Study*,
585 62(3), 377-385.

586 Vafidis, J. O., Vaughan, I. P., Jones, T. H., Facey, R. J., Parry, R., & Thomas, R. J.
587 (2016). The effects of supplementary food on the breeding performance of Eurasian
588 reed warblers *Acrocephalus scirpaceus*; Implications for climate change impacts. *PLoS*
589 *One*, 11(7), e0159933.

590 Vandenabeele, S.P., Shepard, E.L.C., Grogan, A. & Wilson, R.P. (2012). When three
591 per cent may not be three per cent; device-equipped seabirds experience variable flight
592 constraints. *Marine Biology*, 159: 1–14.

593 Vandenabeele, S.P., Grundy, E., Friswell, M.I., Grogan, A., Votier, S.C. & Wilson, R.P.
594 (2014). Excess baggage for birds: inappropriate placement of tags on gannets changes
595 flight patterns. PLoS One, 9(3): e92657. doi:10.1371/journal.pone.0092657

596 Weidinger, K. (2008). Nest Monitoring Does Not Increase Nest Predation in Open-
597 Nesting Songbirds: Inference from Continuous Nest-Survival Data, The Auk, 125(4):
598 859–868. <https://doi.org/10.1525/auk.2008.07016>

599 Westerskov, K. (1950). Methods for determining the age of game bird eggs. Journal of
600 Wildlife Management 14 : 56 – 67 .

601 White, G.C. & Burnham, K.P. (1999). Program MARK: survival estimation from
602 populations of marked animals, Bird Study, 46:sup1, S120-S139, DOI:
603 10.1080/00063659909477239

604 Wilson, R. P., & McMahon, C. R. (2006). Measuring devices on wild animals: What
605 constitutes acceptable practice? Frontiers in Ecology and the Environment, 4(3): 147–
606 154. [https://doi.org/10.1890/1540-9295\(2006\)004\[0147:MDOWAW\]2.0.CO;2](https://doi.org/10.1890/1540-9295(2006)004[0147:MDOWAW]2.0.CO;2)

607 Zhao, J. M., Yang, C., Lou, Y. Q., Shi, M., Fang, Y., & Sun, Y. H. (2020). Nesting
608 season, nest age, and disturbance, but not habitat characteristics, affect nest survival of
609 Chinese grouse. Current Zoology, 66(1), 29-37.

610

Appendix A

Table A1. All candidate models of nightjar daily nest survival rates, for a set of independent variables comprising: total rainfall (s_prcp), average temperature (m_temp), nest age (NestAge), time, construction year (ycons), adult female tag status (f_tag), adult male tag status (m_tag), both adult tag status (fm_tag), adult male or female tag status (f_m_tag) and year (2013 to 2019).

Model	nPar	AICc	DeltaAICc	weight	Deviance
S(~NestAge + f_tag + m_temp * Time + m_prcp2)	7	170.11	0.00	0.12	156.02
S(~NestAge + fm_tag + m_temp * Time + m_prcp2)	7	171.27	1.15	0.07	157.17
S(~NestAge + m_temp * Time + m_prcp2)	6	171.86	1.75	0.05	159.79

S(~NestAge + f_tag + m_temp * Time)	6	172.46	2.35	0.04	160.39
S(~NestAge + f_tag + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	10	172.96	2.84	0.03	152.77
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time + m_prcp2)	14	173.35	3.23	0.02	144.99
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 +	14	173.35	3.23	0.02	144.99

y19 + m_prcp2 + Time * m_temp)					
S(~NestAge + fm_tag + m_temp * Time)	6	173.40	3.29	0.02	161.33
S(~NestAge + f_tag + brood1 + brood2 + brood3 + Time * m_temp)	9	173.53	3.42	0.02	155.38
S(~NestAge + f_tag + m_temp + m_prcp2 + Time)	6	173.58	3.47	0.02	161.51
S(~NestAge + f_tag + Time)	4	173.60	3.49	0.02	165.57
S(~NestAge + m_temp * Time)	5	173.66	3.55	0.02	163.61

S(~f_tag + Time)	3	173.80	3.69	0.02	167.78
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time + m_prdp2)	14	173.80	3.69	0.02	145.45
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_prdp2 + Time * m_temp)	14	173.80	3.69	0.02	145.45
S(~NestAge + m_tag + m_temp * Time + m_prdp2)	7	173.88	3.76	0.02	159.78

S(~NestAge + fm_tag + Time)	4	173.94	3.82	0.02	165.90
S(~fm_tag + Time)	3	173.96	3.85	0.02	167.94
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time)	13	174.19	4.08	0.02	147.88
S(~NestAge + fm_tag + m_temp + m_prcp2 + Time)	6	174.20	4.09	0.02	162.13
S(~NestAge + m_temp + m_prcp2 + Time)	5	174.34	4.23	0.01	164.29

S(~f_tag + m_temp * Time)	5	174.41	4.29	0.01	164.35
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time)	13	174.49	4.38	0.01	148.18
S(~NestAge + fm_tag + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	10	174.50	4.38	0.01	154.31
S(~NestAge + fm_tag + brood1 + brood2 + brood3 + Time * m_temp)	9	174.82	4.70	0.01	156.66
S(~fm_tag)	2	175.00	4.89	0.01	170.99

S(~f_tag + NestAge * m_temp * Time)	9	175.01	4.90	0.01	156.86
S(~fm_tag + m_temp * Time)	5	175.09	4.97	0.01	165.04
S(~NestAge + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	9	175.29	5.17	0.01	157.13
S(~f_tag + m_prcp2 + m_temp * Time)	6	175.37	5.26	0.01	163.30
S(~NestAge * m_temp * Time)	8	175.45	5.34	0.01	159.33
S(~f_tag)	2	175.48	5.36	0.01	171.47
S(~NestAge * Time + m_temp + m_prcp2)	6	175.50	5.39	0.01	163.43

S(~NestAge + brood1 + brood2 + brood3 + Time * m_temp)	8	175.55	5.43	0.01	159.42
S(~NestAge + m_tag + m_temp * Time)	6	175.68	5.57	0.01	163.61
S(~NestAge + Time)	3	175.70	5.58	0.01	169.68
S(~fm_tag + NestAge * m_temp * Time)	9	175.72	5.61	0.01	157.57
S(~NestAge * m_temp * Time + fm_tag)	9	175.72	5.61	0.01	157.57
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp +	13	175.87	5.75	0.01	149.55

m_prcp2 + Time)					
S(~Time)	2	175.89	5.77	0.01	171.88
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time + m_prcp2)	13	175.92	5.80	0.01	149.61
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_prcp2 + Time * m_temp)	13	175.92	5.80	0.01	149.61
S(~m_temp * Time)	4	176.09	5.98	0.01	168.06
S(~fm_tag + m_prcp2 + m_temp * Time)	6	176.15	6.04	0.01	164.08

S(~NestAge + fm_tag)	3	176.32	6.21	0.01	170.30
S(~NestAge + m_tag + m_temp + m_prcp2 + Time)	6	176.36	6.25	0.01	164.29
S(~NestAge * Time + m_temp + brood1 + brood2 + brood3)	8	176.48	6.37	0.01	160.36
S(~f_tag + m_temp + m_prcp2 + Time)	5	176.54	6.42	0.00	166.49
S(~NestAge + f_tag + m_temp + m_prcp2 + Time + brood1 +	9	176.55	6.44	0.00	158.40

brood2 + brood3)					
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time)	12	176.62	6.50	0.00	152.35
S(~1)	1	176.65	6.54	0.00	174.65
S(~fm_tag + m_temp + m_prcp2 + Time)	5	176.91	6.80	0.00	166.86
S(~NestAge + f_tag)	3	176.91	6.80	0.00	170.89
S(~NestAge + f_tag + Time + brood1 + brood2 + brood3)	7	177.04	6.93	0.00	162.95

S(~NestAge * Time)	4	177.12	7.01	0.00	169.09
S(~NestAge + fm_tag + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	9	177.14	7.03	0.00	158.99
S(~NestAge + m_tag + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	10	177.25	7.14	0.00	157.07
S(~m_prcp2 + m_temp * Time)	5	177.31	7.19	0.00	167.25
S(~NestAge + m_temp + m_prcp2 + Time + brood1 +	8	177.37	7.25	0.00	161.24

brood2 + brood3)					
S(~f_tag + m_temp * Time + m_prcp2 + ycons)	7	177.38	7.26	0.00	163.28
S(~f_tag + m_prcp2 + ycons + Time * m_temp)	7	177.38	7.26	0.00	163.28
S(~NestAge + m_tag + brood1 + brood2 + brood3 + Time * m_temp)	9	177.39	7.27	0.00	159.24
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp +	12	177.39	7.28	0.00	153.13

m_prcp2 + Time)					
S(~m_tag + NestAge * m_temp * Time)	9	177.48	7.36	0.00	159.33
S(~NestAge * m_temp * Time + m_tag)	9	177.48	7.36	0.00	159.33
S(~ycons + NestAge * m_temp * Time)	9	177.48	7.37	0.00	159.33
S(~NestAge + fm_tag + Time + brood1 + brood2 + brood3)	7	177.53	7.42	0.00	163.44
S(~fm_tag + m_temp * Time + m_prcp2 + ycons)	7	177.62	7.50	0.00	163.52

S(~fm_tag + m_prcp2 + ycons + Time * m_temp)	7	177.62	7.50	0.00	163.52
S(~ycons + Time)	3	177.66	7.54	0.00	171.64
S(~NestAge + m_tag + Time)	4	177.67	7.56	0.00	169.64
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time + m_prcp2)	14	177.74	7.62	0.00	149.38
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 +	14	177.74	7.62	0.00	149.38

y19 + m_prcp2 + Time * m_temp)					
S(~m_tag + Time)	3	177.88	7.77	0.00	171.86
S(~f_tag + brood1 + brood2 + brood3 + Time * m_temp)	8	177.95	7.84	0.00	161.83
S(~NestAge)	2	177.97	7.85	0.00	173.96
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	17	178.09	7.97	0.00	143.56
S(~ycons + m_temp * Time)	5	178.10	7.98	0.00	168.05

S(~m_tag + m_temp * Time)	5	178.11	7.99	0.00	168.06
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp * Time)	13	178.24	8.12	0.00	151.93
S(~f_tag + m_temp + m_prcp2 + ycons + Time)	6	178.37	8.25	0.00	166.29
S(~f_tag + Time + brood1 + brood2 + brood3)	6	178.44	8.32	0.00	166.36
S(~fm_tag + m_temp + m_prcp2)	4	178.53	8.42	0.00	170.50

S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	17	178.55	8.43	0.00	144.02
S(~fm_tag + Time + brood1 + brood2 + brood3)	6	178.58	8.46	0.00	166.51
S(~m_tag)	2	178.66	8.54	0.00	174.65
S(~ycons)	2	178.66	8.54	0.00	174.65
S(~fm_tag + m_temp + m_prcp2 + ycons + Time)	6	178.91	8.79	0.00	166.83

S(~f_tag + m_temp + m_prcp2)	4	178.95	8.84	0.00	170.92
S(~m_temp * Time + m_prcp2 + ycons)	6	179.10	8.98	0.00	167.02
S(~m_temp + m_prcp2 + ycons + Time * m_temp)	6	179.10	8.98	0.00	167.02
S(~NestAge + m_tag + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	9	179.31	9.19	0.00	161.15
S(~m_tag + m_prcp2 + m_temp * Time)	6	179.32	9.21	0.00	167.25

S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	16	179.35	9.24	0.00	146.89
S(~NestAge * Time + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19)	12	179.37	9.25	0.00	155.10
S(~f_tag + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	9	179.51	9.39	0.00	161.36
S(~NestAge + f_tag + y13 + y14 + y15 + y16	16	179.75	9.64	0.00	147.29

+ y17 + y18 + y19 + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)					
S(~+m_temp + m_prcp2 + ycons + Time)	5	179.77	9.65	0.00	169.72
S(~m_tag + m_temp + m_prcp2 + Time)	5	179.77	9.66	0.00	169.72
S(~f_tag + brood1 + brood2 + brood3)	5	179.81	9.70	0.00	169.76
S(~NestAge + Time + brood1 + brood2 + brood3)	6	179.81	9.70	0.00	167.74

S(~NestAge + m_tag)	3	179.98	9.86	0.00	173.96
S(~NestAge + fm_tag + m_temp + m_prdp2)	5	180.08	9.96	0.00	170.03
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time)	11	180.09	9.98	0.00	157.87
S(~m_temp + m_prdp2)	3	180.10	9.99	0.00	174.08
S(~fm_tag + Time * m_temp + m_prdp2 + brood1 + brood2 + brood3)	9	180.47	10.35	0.00	162.31
S(~fm_tag + m_temp +	5	180.55	10.44	0.00	170.50

m_prcp2 + ycons)					
S(~NestAge + fm_tag + brood1 + brood2 + brood3)	6	180.58	10.46	0.00	168.50
S(~NestAge + f_tag + m_temp + m_prcp2)	5	180.58	10.47	0.00	170.53
S(~f_tag + m_temp + m_prcp2 + ycons)	5	180.58	10.47	0.00	170.53
S(~brood1 + brood2 + brood3 + m_temp * Time)	7	180.84	10.73	0.00	166.75
S(~Time * m_temp +	7	180.84	10.73	0.00	166.75

brood1 + brood2 + brood3)					
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3 + Time * m_temp)	15	180.86	10.74	0.00	150.45
S(~f_tag + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	8	181.01	10.90	0.00	164.89
S(~m_tag + m_temp * Time + m_prcp2 + ycons)	7	181.12	11.00	0.00	167.02
S(~m_tag + m_prcp2 + ycons	7	181.12	11.00	0.00	167.02

+ Time * m_temp)					
S(~Time + brood1 + brood2 + brood3)	5	181.15	11.03	0.00	171.09
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	16	181.17	11.06	0.00	148.71
S(~fm_tag + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	8	181.27	11.15	0.00	165.14
S(~NestAge + f_tag + brood1 +	6	181.41	11.30	0.00	169.34

brood2 + brood3)					
S(~brood1 + brood2 + brood3)	4	181.42	11.30	0.00	173.38
S(~f_tag + Time * m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	10	181.50	11.38	0.00	161.31
S(~NestAge + m_temp + m_prcp2)	4	181.55	11.43	0.00	173.52
S(~NestAge + m_tag + Time + brood1 + brood2 + brood3)	7	181.61	11.50	0.00	167.52
S(~m_tag + m_temp +	6	181.78	11.66	0.00	169.71

m_prcp2 + ycons + Time)					
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3 + NestAge * Time * m_temp)	18	181.81	11.69	0.00	145.22
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time)	11	181.92	11.81	0.00	159.70
S(~NestAge * Time + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19)	12	181.99	11.87	0.00	157.72

S(~m_temp + m_prcp2 + ycons)	4	182.07	11.95	0.00	174.03
S(~m_tag + m_temp + m_prcp2)	4	182.11	12.00	0.00	174.08
S(~fm_tag + Time * m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	10	182.26	12.14	0.00	162.07
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	15	182.32	12.21	0.00	151.91

S(~Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	8	182.35	12.23	0.00	166.22
S(~f_tag + m_temp + m_prcp2 + ycons + Time + brood1 + brood2 + brood3)	9	182.37	12.26	0.00	164.22
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3 + Time * m_temp)	16	182.57	12.45	0.00	150.10
S(~m_tag + brood1 + brood2	8	182.79	12.68	0.00	166.67

+ brood3 + Time * m_temp)					
S(~NestAge + brood1 + brood2 + brood3)	5	182.82	12.70	0.00	172.76
S(~fm_tag + m_temp + m_prcp2 + brood1 + brood2 + brood3)	7	182.82	12.70	0.00	168.72
S(~m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	7	182.82	12.71	0.00	168.73
S(~ycons + brood1 + brood2 + brood3 + Time * m_temp)	8	182.87	12.75	0.00	166.75

S(~ycons + Time + brood1 + brood2 + brood3)	6	182.99	12.88	0.00	170.92
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	17	183.01	12.90	0.00	148.49
S(~m_tag + Time + brood1 + brood2 + brood3)	6	183.03	12.91	0.00	170.96
S(~NestAge + fm_tag + y13 + y14 + y15 + y16	10	183.18	13.06	0.00	162.99

+ y17 + y18 + y19)					
S(~fm_tag + m_temp + m_prcp2 + ycons + Time + brood1 + brood2 + brood3)	9	183.28	13.16	0.00	165.12
S(~m_tag + brood1 + brood2 + brood3)	5	183.35	13.24	0.00	173.30
S(~f_tag + m_temp + m_prcp2 + brood1 + brood2 + brood3)	7	183.43	13.31	0.00	169.33
S(~ycons + brood1 + brood2 + brood3)	5	183.43	13.32	0.00	173.38

S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time)	10	183.48	13.37	0.00	163.29
S(~NestAge + m_tag + m_temp + m_prcp2)	5	183.57	13.45	0.00	173.52
S(~NestAge * Time + y13 + y14 + y15 + y16 + y17 + y18 + y19)	11	183.65	13.53	0.00	161.42
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 + Time + brood1 +	16	183.72	13.61	0.00	151.26

brood2 + brood3)					
S(~m_tag + m_temp + m_prcp2 + ycons)	5	184.08	13.97	0.00	174.03
S(~Time * m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	9	184.29	14.17	0.00	166.13
S(~m_tag + Time * m_temp + m_prcp2 + brood1 + brood2 + brood3)	9	184.32	14.21	0.00	166.17
S(~NestAge + fm_tag + m_temp + m_prcp2 +	8	184.37	14.26	0.00	168.25

brood1 + brood2 + brood3)					
S(~f_tag + m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	8	184.70	14.59	0.00	168.58
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time + brood1 + brood2 + brood3)	14	184.73	14.61	0.00	156.37
S(~NestAge + m_tag + brood1 + brood2 + brood3)	6	184.73	14.62	0.00	172.66
S(~m_temp + m_prcp2 + ycons	8	184.77	14.65	0.00	168.64

+ Time + brood1 + brood2 + brood3)					
S(~m_tag + m_temp + m_prcp2 + Time + brood1 + brood2 + brood3)	8	184.80	14.69	0.00	168.68
S(~fm_tag + m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	8	184.81	14.70	0.00	168.69
S(~m_temp + m_prcp2 + brood1 + brood2 + brood3)	6	184.96	14.85	0.00	172.89
S(~NestAge + f_tag + m_temp	8	185.13	15.02	0.00	169.01

+ m_prcp2 + brood1 + brood2 + brood3)					
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time)	11	185.16	15.04	0.00	162.93
S(~NestAge * Time + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19)	12	185.19	15.07	0.00	160.92
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time + brood1 + brood2 + brood3)	14	185.57	15.45	0.00	157.21

S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2)	12	185.99	15.88	0.00	161.72
S(~m_tag + Time * m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	10	186.24	16.13	0.00	166.06
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19)	10	186.26	16.15	0.00	166.07
S(~NestAge + m_temp + m_prcp2 +	7	186.37	16.26	0.00	172.28

brood1 + brood2 + brood3)					
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3)	13	186.37	16.26	0.00	160.06
S(~m_tag + m_temp + m_prcp2 + ycons + Time + brood1 + brood2 + brood3)	9	186.77	16.65	0.00	168.61
S(~m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	7	186.86	16.75	0.00	172.77

S(~m_tag + m_temp + m_prcp2 + brood1 + brood2 + brood3)	7	186.93	16.82	0.00	172.84
S(~NestAge + fm_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 + brood1 + brood2 + brood3)	15	187.39	17.28	0.00	156.98
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3 + NestAge * Time)	14	187.95	17.83	0.00	159.59
S(~NestAge + y13 + y14 + y15	9	187.99	17.88	0.00	169.84

+ y16 + y17 + y18 + y19)					
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time + brood1 + brood2 + brood3)	13	188.26	18.14	0.00	161.95
S(~NestAge + m_tag + m_temp + m_prcp2 + brood1 + brood2 + brood3)	8	188.33	18.22	0.00	172.21
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2)	12	188.61	18.50	0.00	164.35

S(~m_tag + m_temp + m_prcp2 + ycons + brood1 + brood2 + brood3)	8	188.86	18.75	0.00	172.74
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19)	10	189.57	19.46	0.00	169.38
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + Time + brood1 + brood2 + brood3)	14	190.06	19.95	0.00	161.70
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 +	11	190.60	20.49	0.00	168.38

m_temp + m_prcp2)					
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3)	13	190.71	20.59	0.00	164.40
S(~NestAge + f_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 + brood1 + brood2 + brood3)	15	191.95	21.84	0.00	161.54
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 +	12	192.13	22.01	0.00	167.86

y19 + m_temp + m_prcp2)					
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3)	12	192.78	22.66	0.00	168.51
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + brood1 + brood2 + brood3)	13	194.09	23.98	0.00	167.78
S(~NestAge + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 +	14	194.49	24.37	0.00	166.13

brood1 + brood2 + brood3)					
S(~NestAge + m_tag + y13 + y14 + y15 + y16 + y17 + y18 + y19 + m_temp + m_prcp2 + brood1 + brood2 + brood3)	15	195.38	25.26	0.00	164.96

618

619

Appendix B

Table B1. Top models (i.e. models within 2 AICc units of the top model) of nightjar daily nest survival rates during the egg stage, for a set of models including mean rainfall (m_prcp2), average temperature (m_temp), time (Julian day) and adult female tag status (f_tag).

Model	nPar	AICc	DeltaAICc	Weight	Deviance
S(~f_tag + m_prcp2 + m_temp * Time)	6	66.22	0	0.07	53.99
S(~f_tag + Time)	3	66.75	0.53	0.06	60.689
S(~f_tag)	2	66.94	0.72	0.05	62.909
S(~f_tag + m_temp * Time)	5	67.18	0.95	0.04	57.01
S(~f_tag + m_temp + m_prcp2 + Time)	5	67.54	1.32	0.04	57.37

Table B2. Full model averaged estimates (\pm SE) of the effects of mean rainfall, mean temperature, Julian day (Time) and adult female tag status, on daily nest survival rates (DSR) of egg stage nightjar nests at Brechfa Forest. Model averaged parameter estimates were derived by weighted averaging across all models within 2 AICc units of the top model (Table B1).

Parameter	Estimate	SE	95% Confidence limits
S((Intercept))	4.48	2.08	0.41 to 8.55
S(f_tag1)	1.56	0.84	-0.08 to 3.19
S(m_prcp2)	0.24	0.46	-0.66 to 1.14
S(m_temp)	0.78	2.37	-3.87 to 5.43
S(Time)	-0.04	0.05	-0.13 to 0.06
S(m_temp:Time)	-0.01	0.05	-0.1 to 0.09
S(NestAge)	0.07	0.22	-0.36 to 0.51
S(m_temp:NestAge)	0.09	0.27	-0.45 to 0.62
S(NestAge:Time)	0	0	-0.01 to 0.01

S(m_temp:NestAge:Time)	0	0.01	-0.01 to 0.01
------------------------	---	------	---------------

632

633

Table B3. Nest survival rate (DSR¹⁸) estimates for **egg stage** nests at Brechfa Forest using predicted DSR from model averaged top models for nests initiated on day 20 (16th June – median nest initiation date).

Tag status	Sample Size	NSR Estimate	95% Confidence limits
Female tagged	27	0.82	0.56 to 1.00
Female untagged	23	0.55	0.37 to 0.72
Adult tagged	33	0.91	0.82 to 0.99
Adult untagged	17	0.87	0.77 to 0.98

Appendix C

Figure C1: Boxplot of minimum temperature vs. relative humidity at Brechfa Forest, Carmarthenshire, Wales, 2013–2019 for data split into low temperature ($<10^{\circ}\text{C}$) and high temperature groups ($>10^{\circ}\text{C}$).

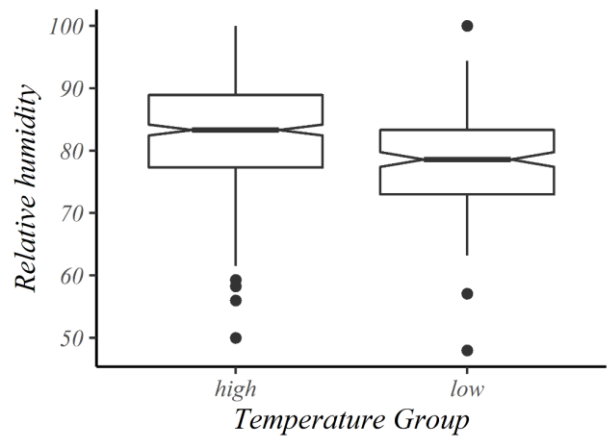
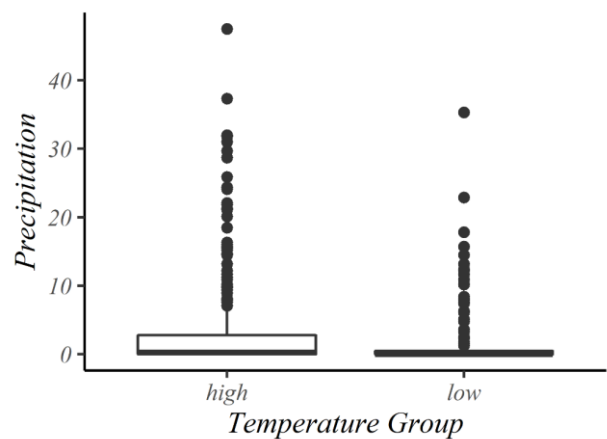


Figure C2: Boxplot of minimum temperature vs. precipitation (mm) at Brechfa Forest, Carmarthenshire, Wales, 2013–2019 for data split into low temperature ($<10^{\circ}\text{C}$) and high temperature groups ($>10^{\circ}\text{C}$).



648 **Tables**

649 **Table 1.** Summary of nest monitoring results (total no. of nests fledging one chick or
650 more, and percentage success rates) with a breakdown by tag status of the attending
651 adults and brood number.

	Total No. nests	No. Succes sful	Overall 1 % succes s	% succes s 1 st Brood	% succes s 2 nd Brood	% succes s 3 rd Brood	% succes s unkno wn
All nests	85	52	61.2	69.5	46.2	50	36.4
Untagged nests	34	21	61.8	70.4	33.3	100.0	0
Nests attended by at least 1 tagged adult	51	31	60.8	68.8	50.0	0.0	50.0
Nests attended by tagged adult male	25	16	64.0	66.7	60.0	NA	60.0
Nests attended by tagged adult female	39	26	66.7	75.0	55.6	0.0	60.0

Nests attended by tagged adult male and female	13	11	84.6	85.7	75.0	NA	100.0
--	----	----	------	------	------	----	-------

652

653 **Table 2.** Summary of number of nests attended tagged parents, broken down by brood
654 status.

	Total	1st brood	2nd brood	3rd brood	unknown
No. of nests	85	59	13	2	11
No. attended by tagged adult	51	32	10	1	8
No. attended by tagged adult male	25	15	5	0	5
No. attended by tagged adult female	39	24	9	1	5
No. attended by 2 tagged adults	13	7	4	0	2
% attended by tagged adult	60.0	54.2	76.9	50.0	72.7
% attended by tagged adult male	29.4	25.4	38.5	0.0	45.5

% attended by tagged adult female	45.9	40.7	69.2	50.0	45.5
% attended by 2 tagged adults	15.3	11.9	30.8	0.0	18.2

655

656 **Table 3.** Top models (i.e. models within 2 AICc units of the top model) of nightjar daily
657 nest survival rates, for a set of models including mean rainfall (m_prdp2), average
658 temperature (m_temp), nest age (NestAge), time (Julian day), adult female tag status
659 (f_tag), year (2013 to 2019) and adult tag status (fm_tag).

Model	nPar	AICc	DeltaAICc	Weight	Deviance
S(~NestAge + f_tag + m_temp * Time + m_prdp2)	7	170.12	0	0.1227	156.02
S(~NestAge + fm_tag + m_temp * Time + m_prdp2)	7	171.27	1.15	0.072	157.17
S(~NestAge + m_temp * Time + m_prdp2)	6	171.86	1.75	0.05	159.79

660

661 **Table 4.** Conditional model averaged estimates (\pm SE) of the effects of mean rainfall,
662 mean temperature, nest age, time (days from 28th of May), construction year, adult
663 female tag status and adult male or female tag status, on daily nest survival rates (DSR)
664 of nightjars at Brechfa Forest. Model averaged parameter estimates were derived by

665 weighted averaging across all models within 2 AICc units of the top model (Table 1).
 666 Parameters in bold are considered to have an important effect based on 95% CL.

	Estimate	SE	95% Confidence limits
Intercept	5.7157	0.99284	3.770 to 7.662
Nest age	0.07146	0.02756	0.017 to 0.125
Female adult tag status (tagged)	0.76275	0.43062	-0.081 to 1.607
Mean temperature	2.50182	1.08336	0.378 to 4.625
Mean precipitation	0.61952	0.33058	-0.028 to 1.268
Time	-0.07332	0.0233	-0.119 to -0.028
Adult tag status (tagged)	0.80302	0.6496	-0.470 to 2.077
Mean Temperature: Time	-0.03543	0.01924	-0.073 to 0.002

667

668

Table 5. Full model averaged estimates (\pm SE) of the effects of total rainfall, mean temperature, nest age, time (days from 28th of May), construction year, adult female tag status and adult male or female tag status, on daily nest survival rates (DSR) of nightjars at Brechfa Forest. Model averaged parameter estimates were derived by weighted averaging across all models within 2 AICc units of the top model (Table 1). Parameters in bold are considered to have an important effect based on 95% CL.

	Estimate	SE	95% Confidence limits
Intercept	5.7157	0.99284	3.770 to 7.661
Nest age	0.07146	0.02756	0.017 to 0.126
Female adult tag status (tagged)	0.44476	0.49955	-0.534 to 1.424
Mean temperature	2.50182	1.08336	0.378 to 4.625
Mean precipitation	0.50877	0.38222	-0.240 to 1.258
Time	-0.07332	0.0233	-0.119 to -0.028
Adult tag status (tagged)	0.15784	0.42985	-0.685 to 1.000

Mean Temperature: Time	-0.03543	0.01924	-0.073 to 0.002
---------------------------	----------	---------	-----------------

Legends to figures

Figure 1: Relationship between daily survival rate (DSR) and radio tag deployment status of parental adult nightjar at Brechfa Forest, Carmarthenshire, Wales, 2013–2019. Daily survival results are based on 85 nests pooled across 2013–2019. The points represent the estimated mean DSR values, and the bars represent the 95% confidence intervals.

Figure 2: Relationship between daily survival rate (DSR) and radio tag deployment status of parental female adult nightjar at Brechfa Forest, Carmarthenshire, Wales, 2013–2019. Daily survival results are based on 85 nests pooled across 2013–2019. The points represent the estimated mean DSR values, and the bars represent the 95% confidence intervals.

Figure 3: Model averaged predicted daily nest survival rate in relation to mean temperature during the nightjar nesting period in Brechfa Forest, Carmarthenshire, Wales, 2013–2019. Estimates (lines) and 95% confidence bands (shaded) are shown for day 1 of the season (28th May), day 20 (16th June – median nest initiation date), and day 46 (12th July – median hatch date), with other covariates fixed at mean values.

Figures

