

1 Machine learning pipeline extracts biologically significant data automatically from avian monitoring  
2 videos

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8

## 9 **Abstract**

10 Measuring parental care behaviour in the wild is central to the study of animal life-history trade-offs,  
11 but is often labour and time-intensive. More efficient machine learning-based video processing tools  
12 have recently emerged that allow parental nest visit rates to be measured using video cameras and  
13 computer processing. Here, we used open-source software to detect movement events from videos  
14 taken at the nest box of a wild passerine bird population. We show that visit numbers from our  
15 automatic data collection pipeline strongly correlate with human observations and predicts an  
16 increase in brood fitness. Using a machine learning-assisted annotation approach on a subset of 18  
17 videos, we show that the accuracy largely increased and cut annotation time by an average of 5.5x  
18 compared to that of a cohort of undergraduate students. Since our automatic pipeline collected  
19 biological-meaningful data that would have taken approximately 800 days by human observers, we  
20 encourage more researchers to apply existing open-source tools to assist data collection in animal  
21 behaviour studies.

22 **Keywords:** Parental Care, Computer Vision, Deep Meerkat, House Sparrow, Machine Learning,

23

24

## 25 **Introduction**

26 Parental care behaviour is a life-history trait that is commonly studied in a wide range of animals (Royle  
27 et al., 2012). Parental care is defined as any behaviour that increases the fitness of offspring (Clutton-  
28 Brock, 1991; Royle et al., 2012), but often decreases the survival probability of parents (Trivers, 1972)  
29 presenting as a life history trade-off (Stearns, 1992). While there are many forms of parental care (*e.g.*,  
30 nest building, predator defence, incubation, and feeding; see Royle et al., 2012), parental care in the  
31 form of feeding of the young is traditionally considered as very costly for parents (Owens and Bennett,  
32 1994; Winkler and Wilkinson, 1988). As such, a large body of literature focuses on how nest visit  
33 frequency with dependent young is associated with aspects of an animal's life history, with a focus on  
34 birds, where 90% of species engage in parental care (Cockburn, 2006). For example, work on life  
35 history trade-off in terms of parent and offspring fitness (Schroeder et al., 2013), parent coordination  
36 (see Ihle et al., 2019; Wojczulanis-Jakubas et al., 2018), parent-offspring conflict (Estramil et al., 2013),  
37 or ageing (Wilcoxon et al., 2010) all used the frequency of parental visits to nests (or provisioning) as  
38 a proxy of parental investment.

39

40 However, measuring parental visitation rates in the wild is labour intensive and time-consuming. Data  
41 is traditionally collected by direct observations (*e.g.* Dunn and Cockburn, 1996), which can be invasive  
42 by disturbing animals in the vicinity of their nest (Rose, 2009). Less invasive methods include video  
43 recording (*e.g.*, García-Navas and Sanz, 2010; Nakagawa et al., 2007), radio tracking (see Mitchell et  
44 al., 2012), and the use of radio-frequency tags and antennas at the nest (RFID; Mariette et al., 2011;  
45 Ringsby et al., 2009). While automatic methods allow visitation rates to be quantified over long  
46 periods of time, the technology is prone to error (up to 20%; see Mariette et al., 2011). On the other  
47 hand, video analysis is more flexible and allows for other behaviours to be quantified, like nest defence,  
48 copulations or feeding load (Lendvai et al., 2015). Maybe most importantly, using video cameras also  
49 allows for visual confirmation of individual identity if colour rings are used. However, manual

50 annotation of video data is time-consuming and error-prone (Tuytens et al., 2014), calling for more  
51 effective ways to extract data from videos (Conway et al., 2021; Weinstein, 2018a).

52

53 Recent advances in deep learning (Borowiec et al., 2021) and computer vision (Weinstein, 2018a)  
54 allow quick and reliable information to be extracted from field data (Valletta et al., 2017). For example,  
55 machine learning methods have been successfully applied to solve problems with species  
56 identification (see Wäldchen and Mäder, 2018), bird song complexity measurement (see Pearse et al.,  
57 2018; Priyadarshani et al., 2018), social behaviour measurement (see Robie et al., 2017), and individual  
58 identification (Bogucki et al., 2019; Ferreira et al., 2020; Körschens et al., 2018; Schofield et al., 2019).  
59 Since computing resources are much cheaper than human-labour, such approaches have immense  
60 potential in reducing the financial and time costs of data collection, evidenced by a recent increase in  
61 popularity for ecological applications (Borowiec et al., 2021; Tuia et al., 2022).

62

63 Simultaneously, open-source tools were developed to aid data collection from video data. Examples  
64 include software on tracking animals and behaviours in captive settings (Harmer and Thomas, 2019;  
65 Pennington et al., 2019; Sridhar et al., 2019; Walter and Couzin, 2021), pose estimation (Graving et al.,  
66 2019; Lauer et al., 2021; Pereira et al., 2019), or species identification (Falzon et al., 2020; Van Horn  
67 et al., 2015). Particularly, Weinstein (2018b) developed an open-source tool named Deep Meerkat  
68 which uses convolutional neural networks (CNNs) to capture movement events from wildlife  
69 monitoring videos. Despite the name, the software was initially designed for use with a hummingbird  
70 population (Marcot et al., 2019; Weinstein, 2018b), but the software has been adapted for use in  
71 marine (Sheehan et al., 2020) and insect (Mertens et al., 2021; Pegoraro et al., 2020) systems. To the  
72 best of our knowledge, no literature exists that documents the use of the software in avian systems  
73 other than the original hummingbird population.

74

75 Here, we applied Deep Meerkat (Weinstein, 2018b) to automatically process and extract visitation  
76 rates from parental provisioning videos collected in the house sparrow (*Passer domesticus*) population  
77 on Lundy Island, UK since 2004 (see Nakagawa et al., 2007). We first compared the results to manually-  
78 collected data by trained individuals. Next, we compared the outputs from the automatic processing  
79 with alternative methods of annotation, including a machine learning-assisted approach and  
80 crowdsourced annotation by a cohort of undergraduate students. Lastly, we determined the biological  
81 relevance of the automatic data by testing the hypothesis that increased rates of feeding would lead  
82 to increased annual reproductive fitness of the parents – a classic life-history theory hypothesis from  
83 behavioural ecology (Trivers, 1972).

84

85

86 **Materials and Methods**

87 **Study System**

88 Data were collected from a population of house sparrows (*Passer domesticus*) on Lundy Island (51° 10'  
89 N, 4°40' W) located on the Bristol Channel, UK. The population is part of a long-term study and is  
90 monitored systematically since 2000, with >99% of individuals marked with a unique combination of  
91 coloured metal rings by the British Trust for Ornithology (Cleasby et al., 2011), and unique passive-  
92 integrated transponders (Schroeder et al., 2011). Since house sparrows rarely fly over large bodies of  
93 water (Magnussen and Jensen, 2017) very little immigration or emigration has taken place in the  
94 population (Schroeder et al., 2015). As a result, the population has high recapture rates with no  
95 trapping bias (Simons et al. 2015) and reliable life history data for every individual (Schroeder et al.,  
96 2015).

97

98 **Parental provisioning videos**

99 The Lundy sparrow population is situated within a 0.2km<sup>2</sup> area around a small village, since that is the  
100 only viable habitat on the island (Schroeder et al., 2011). Nest boxes were put up for the sparrows,  
101 and every box is checked systematically to detect all breeding attempts throughout the summer  
102 breeding season (Cleasby et al., 2011). After eggs were found and the identity of the parents  
103 confirmed by their colour ring combination, 90-minute videos were recorded on the 7<sup>th</sup> and 11<sup>th</sup> day  
104 after egg hatching, with a video camera 2-5m away from the nest box and a field of view of 30cm  
105 radius (see Nakagawa et al., 2007a for detailed procedure) to measure parental visitations.

106

107 When annotating the videos, birds were considered as feeding the young when entering and exiting  
108 the nest box, and when feeding behaviour could be seen through the nest box hole without entering  
109 the nest box. We recorded when birds are perching outside the nest box but as a separate behaviour  
110 such that it did not count towards feeding rate estimates. Feeding rates were then calculated from  
111 the first visit of either parent until the end of the video, or until 90 minutes had elapsed from the first

112 visit, whichever came first. We started counting from the first visit, and not the beginning of the video,  
113 to allow time for the birds to adjust to the disturbance possibly caused by the camera setup. The  
114 resulting time during which visits were scored was termed the effective observational time (Nakagawa  
115 et al., 2007a). The total number of visits by both parents was divided by the effective observational  
116 time to obtain the manual feed rate (feeds per hour) as a measure of parental provisioning.

117

118 Between 2004 and 2015, videos (N=2112) were manually annotated by postgraduate students and  
119 researchers, which contributed to multiple publications (see Ihle et al., 2019; Nakagawa et al., 2007;  
120 Schroeder et al., 2019, 2016, 2013, 2012). We used these data as a baseline – the ‘manual feed rate’  
121 – to be compared against the automatically-collected data (‘automatic feed rate’) outlined below.

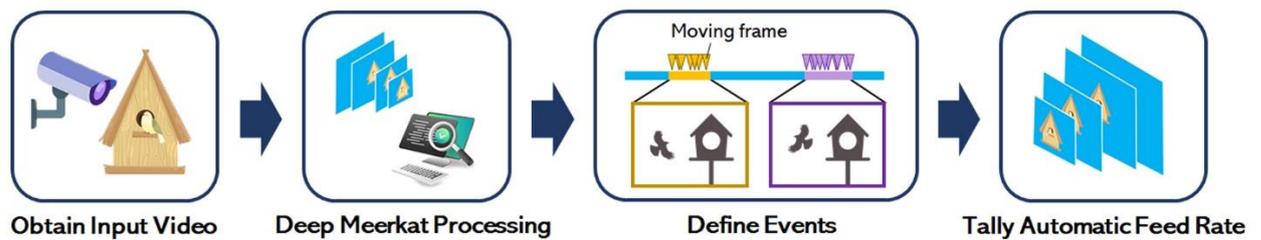
122

### 123 **Automatic Video Processing Pipeline**

124 We processed videos collected between 2011 and 2019 (N=2629) through the open-source  
125 programme Deep Meerkat, which uses convolution neural networks (CNNs) to classify moving frames  
126 and identify moving objects in wildlife monitoring videos (Weinstein, 2018b). Since the performance  
127 of the default model was satisfactory from initial tests, we did not further fine-tune the model using  
128 video frames from our study. We processed the output from Deep Meerkat to merge movement  
129 events detected fewer than 40 frames apart and more than 2 frames in length into the same event.  
130 We then further grouped the events into 7-second video clips, to allow effective post-processing  
131 annotation and matching with manual data. We then tallied the number of events for each video, then  
132 divided the tally by the effective observational time (see above) to obtain the automatic feed rate (in  
133 events per hour; Figure 1). Since most of the feeding events on Lundy sparrows were done by parents  
134 entering the nest box, we assume each feeding event broadly corresponds to 2 detected movement  
135 events by the pipeline (entering and exiting the nest box), hence the automatic feed rate was further  
136 divided by 2. Finally, since certain videos produced massively over-inflated feed rate measures due to  
137 filming environment (*e.g.* camera shaking, continuous background movement), we removed any

138 videos that has an automatic feed rate of 72.7 events/hour or more (1.31% videos removed), since  
139 that is the max feed rate we have ever recorded from previous manual annotation, assuming any rate  
140 more extreme will be an anomaly.

141



142

143 **Figure 1: Automatic data collection pipeline for processing parental provisioning videos of house**

144 **sparrows on Lundy Island, UK.** Videos were processed using open source software Deep Meerkat

145 (Weinstein, 2018b), outputting frames with movement detected. Events were then defined by

146 grouping movement frames that were less than 40 frames apart and at least 2 frames long. Lastly,

147 events for each provisioning video were tallied to obtain an automatic feed rate

148

### 149 **Validation**

150 After processing all videos using the automatic pipeline, we first tested the association between the

151 automatic and manual feed rates using Pearson's correlation test. We then chose 18 random videos

152 to be used as a validation dataset, to compare the performance of three different methods of data

153 collection. The first was the automatic feed rate as described above, where videos were processed

154 through the pipeline (Figure 1) without human intervention. The second was a machine learning (ML)-

155 assisted approach, where video outputs of events from the pipeline were further annotated by hand

156 to filter out non-feeding visits. The third was a crowd-sourcing approach, where a cohort of 36

157 undergraduate students were given full-length videos to annotate from scratch. To compare the

158 annotation time between the latter two methods, we also measured the annotation time per video

159 for the ML-assisted and crowd-sourcing approaches.

160

161 To compare the methods quantitatively, we calculated three return rates corresponding to the three  
162 methods - the automatic pipeline, machine learning-assisted, and undergraduate return rates. We  
163 took the manual feed rate as a baseline value, then calculated return rates by dividing the number of  
164 events detected using each method over the manual feed rate for each video (*e.g.* for a given video,  
165 if manual feed rate has 10 feeding events and undergraduates found 7, the undergraduate return rate  
166 is 0.7). Since a return rate of 1 shows that a method detected the exact number of events as the  
167 manual feed rate baseline, we did one sampled t-tests to see whether each return rate was  
168 significantly different from this baseline (by setting the theoretical mean ( $\mu$ ) to 1). Lastly, we also  
169 compared all methods using pairwise t-tests between the return rates for each annotation method.

170

#### 171 **Case study**

172 Finally, we used the automatic feed rate in a case study to test for the first part of the life-history  
173 trade-off predicted by Trivers (1972). We tested whether broods whose parents visited the nest more  
174 often had a higher fledgling and recruitment success, using the automatic feed rate. Since each brood  
175 was measured on the 7<sup>th</sup> and 11<sup>th</sup> day after hatching, we averaged both values to obtain an average  
176 feed rate for each brood. However, since feeding rates increases as a brood age (see Schroeder et al.,  
177 2019), we also ran the models separately using Day 7 and 11 rates to ensure results are consistent.

178

179 To test the prediction that increased automatic feed rate was associated with higher chick survival in  
180 a brood, we fitted generalised linear mixed models with the number of fledglings and the number of  
181 recruits for each brood as respective response variables, against automatic feed rates (events per hour)  
182 as explanatory variables, using a Poisson link function. A fledgling was defined as a sparrow chick that  
183 successfully fledged; recruits were defined as fledglings that produced at least one genetic offspring  
184 in their lifetime. To allow effect sizes to be compared, we z-transformed the feed rates. To control for  
185 other effects that might affect the fitness metrics, we added the age of the mother and father (see

186 Wiebe, 2018), and hatch date (Days after April 1<sup>st</sup>) as fixed effects. Since breeding success usually  
187 corresponds to peak food abundance (see Cresswell and McCleery, 2003; Lack, 1968), we also added  
188 a quadratic fixed-effect term for hatch date. Next, the population has undergone routine cross-  
189 fostering, which is associated with increased survival (see Winney et al., 2015), hence we added a fixed  
190 factor for fostered status (yes/no) in all models, however, note that the feed rate we used were always  
191 from the parents that did the actual feeding, which are not always the genetic parents of the young.  
192 We added the social parent IDs and the year as random effects to control for environmental effects  
193 (see Rose et al., 1998) and repeatable feeding rates by individual parents (Nakagawa et al., 2007). The  
194 location of the nest box was added as a random effect to control for known environmental effects  
195 (Schroeder et al., 2012).

196

197 We ran all models using the R packages MCMCglmm (Hadfield, 2010) on R version 3.6.1 (R Core Team,  
198 2013). The posterior distributions and autocorrelations were checked following Hadfield (2014) to  
199 ensure all fixed and random effects converged without violating any model assumptions. We defined  
200 a parameter estimate as statistically significant if the 95% credible interval did not overlap with zero.

201

## 202 **Results**

### 203 **Automatic Video Processing Pipeline**

204 Using the automatic pipeline, we processed a total of 2629 videos and so approximately 3900 hours  
205 (160 days) of recordings. We found a significant positive correlation between automatic and manual  
206 feed rates ( $r=0.62$ , 95%CI: 0.58-0.66,  $p < 0.001$ ; Figure 2A).

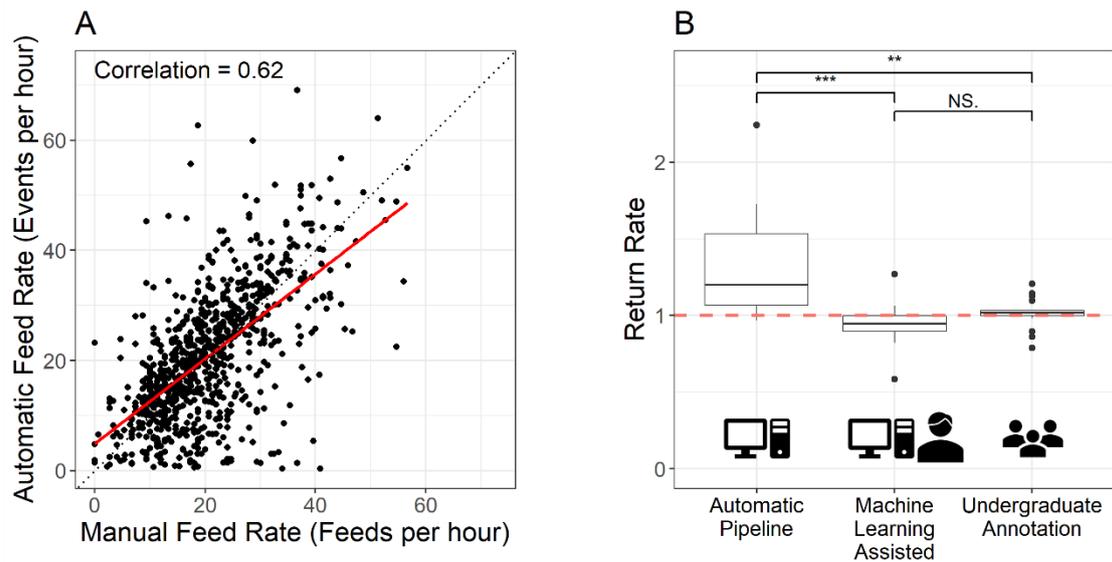
207

### 208 **Validation**

209 From the 18 validation videos, the automatic pipeline return rate obtained without human  
210 intervention was significantly larger than 1 (Table 1), showing that the measure was inflated compared  
211 to the baseline set by manual feed rates (Figure 2B). However, the ML-assisted return rate showed  
212 that once the pipeline outputs were manually annotated to eliminate non-feeding events, the rate  
213 dropped and became not significantly different from 1 and comparable to the baseline (Table 1). The  
214 undergraduates performed similarly well, with return rates not significantly different from 1 (Table 1)  
215 and not significantly different from the ML-assisted return rate (Figure 2B).

216

217 On average, undergraduate students took 65.4 minutes (min 25 minutes, max 100 minutes) to  
218 annotate each video, whereas the ML-assisted approach took an average of 12.0 minutes per video  
219 (min 4.6 minutes, max 31.0 minutes), equating to an average of 53.4 minutes saved per video by first  
220 processing the videos using the pipeline.



221

222 **Figure 2: Validation results of automatic video processing pipeline and alternative annotation**  
 223 **methods for data collection on house sparrow (*Passer domesticus*) parental provisioning videos on**  
 224 **Lundy Island UK. A) Positive correlation between the automatic feed rate collected using the pipeline**  
 225 **and manual feed rate collected manually. The red line shows a significant positive correlation, and the**  
 226 **dotted line shows a 1:1 line if both rates were equal. B) Comparison of return rates for three separate**  
 227 **data collection methods using return rate measures of 18 validation videos. 1) Automatic pipeline:**  
 228 **derived from automatic feed rate 2) Machine learning-assisted: further manual annotation of output**  
 229 **clips from the pipeline 3) Undergraduate annotation: annotation by a cohort of undergraduate**  
 230 **students. The red dotted line represents the reference level of 1, brackets and labels represent the**  
 231 **significance of t-test between all groups (N.S: not significant, \*:  $p < 0.05$ , \*\*:  $p < 0.01$ , \*\*\*:  $p < 0.001$ ).**

232

233 **Table 1: Comparison of three annotation methods using a validation dataset of 18 provisioning**  
 234 **videos of house sparrows (*Passer domesticus*) on Lundy Island, UK. The return rates were calculated**  
 235 **by dividing the number of detected events using each method by the manual feed rate. Test statistics**  
 236 **were obtained from a one-sampled t-test, with the theoretical mean ( $\mu$ ) set to 1.**

237

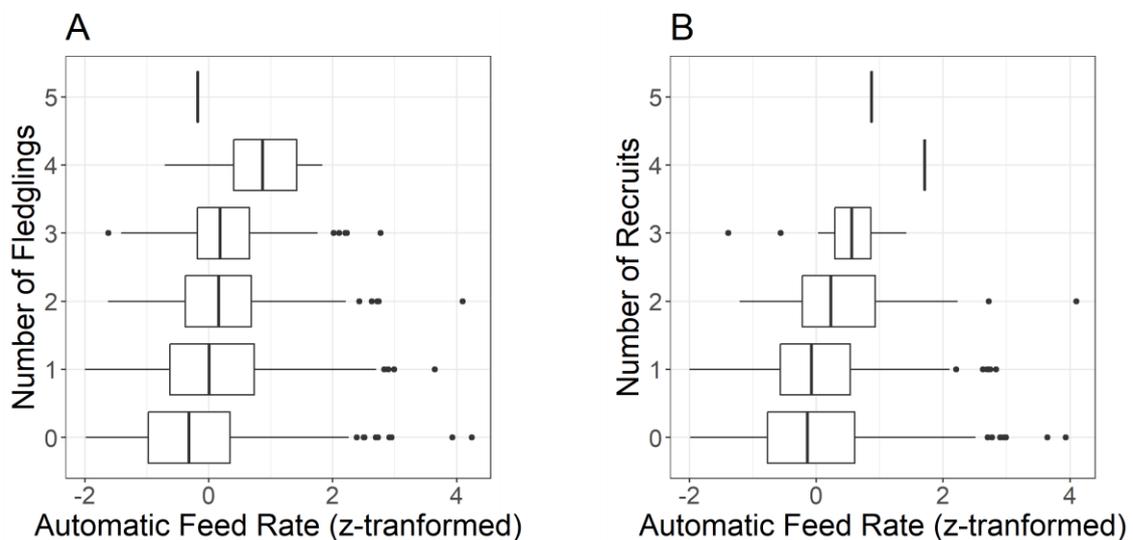
Metrics	Mean	95% Confidence Interval	t	p-value
A) Automatic pipeline return rate	1.39	1.17-1.63	3.68	0.002
B) Machine learning-assisted return rate	0.94	0.88 – 1.01	-1.78	0.10
C) Undergraduate return rate	1.01	0.97 – 1.06	1.13	0.52

238

239 **Case study**

240 The data that we used contained 658 unique broods, of 2116 individual chicks, of which 728 (34%)  
 241 fledged and 328 (16%) were recruited. Automatic feed rates significantly predict an increase in the  
 242 number of fledgelings and recruits for all models fitted (Figure 4; Supporting Information 1). We found  
 243 a noticeable change in brood survival over the breeding season, with a significant negative quadratic  
 244 effect for hatch date, showing that fitness is the highest in mid-breeding season (Supporting Table S1).

245



246

247 **Figure 4: Increase in the number of (A) fledglings and (B) recruits as average automatic feed rates**  
 248 **increase in house sparrows (*Passer domesticus*) on Lundy Island, UK** Automatic feed rates were  
 249 obtained from the pipeline without human intervention. Feed rates were further z\_transformed to  
 250 allow effect sizes to be comparable, with 0 representing the mean and each unit representing one  
 251 standard deviation away from the mean.

252 **Discussion**

253 Using an automatic data collection pipeline based on Deep Meerkat (Weinstein, 2018b), we extracted  
254 visitation rates from parental provisioning videos of house sparrows and found that automatic feed  
255 rates positively correlated with manual annotation, and can reproduce biological results, equivalent  
256 to ~800 hours (~100 8-hour workdays) of human labour work. Even though the computational time  
257 for Deep Meerkat is 1:1 (1 hour video takes ~1 hour to process; see Marcot et al., 2019), computing  
258 time is much cheaper than human labour time, especially when techniques like parallel computing  
259 were used to further speed up processing. As such, we processed a huge backlog of unprocessed  
260 videos from the Lundy sparrow system that would have been infeasible without the use of machine  
261 learning methods.

262

263 When applying machine learning methods, there is often a trade-off between the variance and bias of  
264 a trained model (Geman et al., 1992; but see Neal, 2019). Here, the 18 test videos showed that the  
265 pipeline produced biased results, since the obtained automatic feed rate produced overinflated  
266 estimates compared to manual feed rates, likely due to over-detection of non-feeding events when  
267 birds are still present within the video frame. However, the bias was complemented with low variance  
268 (or high consistency), with automatic feed rate correlating well with manual feed rates and predicting  
269 an increase in fledgelings and recruits within broods in our case study. Although not comparable with  
270 manual feed rates, the automatic feed rate can hence be considered as another proxy of parental  
271 investment (see Schroeder et al., 2013; Trivers, 1972) allowing more biological hypotheses to be  
272 tested with increased sample size and statistical power. Moreover, the motion detection algorithm of  
273 Deep Meerkat is not species-specific, allowing the same method to be applied to other study systems  
274 or species, opening a new avenue of research for newly established systems or large video datasets.

275

276 In this study, we also tested other methods for data collection, including crowdsourcing  
277 undergraduate students and adopting a ML-assisted approach. Manual annotation by undergraduate

278 students was accurate, with mean event return rates slightly higher than the manual feed rates,  
279 possibly due to the multiple observer effect (*e.g.* Guay et al., 2013), but can also be due to inflation in  
280 detected events due to misidentifications. We also showed that there were no significant differences  
281 between the undergraduate and ML-assisted return rates, highlighting the value of using an ML-  
282 assisted approach, which was 5.5x faster. Given sex-specific hypotheses on parental investment are  
283 often tested in the population (*e.g.* Schroeder et al., 2016), adopting a ML-assisted approach can be  
284 useful for collecting additional sex or behavioural data within the feeding clips. Further development  
285 using deep learning and computer vision techniques can also classify the sex (see Ferreira et al., 2020)  
286 and behaviour (see Conway et al., 2021; Ditria et al., 2021) automatically, which has potential to fully  
287 replace human annotation in the future.

288

289 Machine learning approaches are becoming widely used in ecology in recent years (Borowiec et al.,  
290 2021). Here, we presented an excellent case study of machine learning assisted data collection that  
291 can obtain biologically meaningful results in existing datasets without further manual intervention.  
292 With the increase in open-source tools that are being developed in the field to reduce manual  
293 annotation efforts (*e.g.* Van Horn et al., 2015; Walter and Couzin, 2021; Weinstein, 2018b), and the  
294 increase in computing literacy of ecology graduates (Farrell and Carey, 2018), we encourage  
295 researchers to make use of such tools to be adapted to their own existing datasets. This would not  
296 only unlock the bottleneck of unanalysed data that would otherwise go to waste, but also allow more  
297 interesting hypotheses to be tested.

298

## 299 **Conclusion**

300 In this study, we used a data collection pipeline based on Deep Meerkat to automatically process  
301 parental provisioning videos on Lundy sparrows. Automatic feed rates collected from the machine  
302 learning pipeline correlated well with manual annotation, as well as predicting recruitment and  
303 fledging success in broods. The pipeline also acts as an effective pre-processing step to allow

304 annotation time to be largely reduced, as we encourage researchers to adopt similar methods in their  
305 study systems. Machine learning techniques and open-source tools are becoming widespread in  
306 ecology, we provide here a successful case study of the use of such tools to eliminate the bottleneck  
307 of laborious manual data collection.

308

#### 309 **Significance Statement**

310 Collecting data on avian parental provisioning rates are often time and labour intensive. Recent  
311 advancements in open-source machine learning software allow such data to be effectively collected  
312 without human annotation. This paper provides a case-study of the validity and potential of such  
313 software to easily collect parental provisioning data in the wild.

314

#### 315 **Conflict of Interest Statement**

316 None

317

#### 318 **Ethics Statement**

319 The study complied with all relevant legislation and guidelines, and had the appropriate ethical  
320 approval.

321

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327

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331

332 **Author's Contributions**

333 Conception: TB, JS, AC, Funding: TB, JS, Analysis: AC, WP, JS, Writing: AC, TB, JS, WP

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