1 A GENERALISED RANDOM ENCOUNTER MODEL FOR ESTIMATING 2 ANIMAL DENSITY WITH REMOTE SENSOR DATA

- 3 Running title: A generalised random encounter model for animals.
- 4 Word count:
- 5 Authors:
- 6 Tim C.D. Lucas^{1,2,3}, Elizabeth A. Moorcroft^{1,4,5}, Robin Freeman⁵, Marcus J. Rowcliffe⁵,
- 7 Kate E. Jones^{2,5}
- 8 Addresses:
- 9 1 CoMPLEX, University College London, Physics Building, Gower Street, Lon-
- 10 don, WC1E 6BT, UK
- 11 2 Centre for Biodiversity and Environment Research, Department of Genetics,
- 12 Evolution and Environment, University College London, Gower Street, London,
- 13 WC1E 6BT, UK
- 3 Department of Statistical Science, University College London, Gower Street,
- 15 London, WC1E 6BT, UK
- ¹⁶ 4 Department of Computer Science, University College London, Gower Street,
- 17 London, WC1E 6BT, UK
- 5 Institute of Zoology, Zoological Society of London, Regents Park, London, NW1
- 19 4RY, UK
- 20 Corresponding authors:
- 21 Kate E. Jones,
- 22 Centre for Biodiversity and Environment Research,
- 23 Department of Genetics, Evolution and Environment,
- 24 University College London,
- 25 Gower Street,
- 26 London,
- 27 WC1E 6BT,
- 28 UK

- 29 kate.e.jones@ucl.ac.uk
- 30
- Marcus J. Rowcliffe,
- Institute of Zoology,
- 33 Zoological Society of London,
- 34 Regents Park,
- 35 London,
- 36 NW1 4RY,
- 37 UK
- marcus.rowcliffe@ioz.ac.uk

1. Abstract

39

1: Wildlife monitoring technology has advanced rapidly and the use of remote sensors such as camera traps, and acoustic detectors is becoming common in both the terrestrial and marine environments. Current capture-recapture or distance methods to estimate abundance or density require individual recognition of animals or knowing the distance of the animal from the sensor, which is often difficult. A method without these requirements, the random encounter model (REM), has been successfully applied to estimate animal densities from count data generated from camera traps. However, count data from acoustic detectors do not fit the assumptions of the REM due to the directionality of animal signals.

2: We developed a generalised REM (gREM), to estimate absolute animal density
from count data from both camera traps and acoustic detectors. We derived the
gREM for different combinations of sensor detection widths and animal signal
widths (a measure of directionality). We tested the accuracy and precision of this
model using simulations of different combinations of sensor detection widths and
animal signal widths, number of captures, and models of animal movement.

3: We find that the gREM produces accurate estimates of absolute animal density
for all combinations of sensor detection widths and animal signal widths. However, larger sensor detection and animal signal widths were found to be more precise. While the model is accurate for all capture efforts tested, the precision of the
estimate increases with the number of captures. We found no effect of different
animal movement models tested on the accuracy and precision of the gREM.

4: We conclude that the gREM provides an effective method to estimate absolute animal densities from remote sensor count data over a range of sensor and animal signal widths. The gREM is applicable for use for count data obtained in both marine and terrestrial environments, visually or acoustically (e.g., big cats, sharks, birds, bats and cetaceans). As sensors such as camera traps and acoustic detectors become more ubiquitous, the gREM will be increasingly useful for monitoring animal populations across broad spatial, temporal and taxonomic scales.

1.1. **Keywords.** Acoustic detection, Camera traps, Marine, Population monitoring, Simulations, Terrestrial

2. Introduction

70

Animal population density is one of the fundamental measures needed in ecol-71 ogy and conservation. The density of a population has important implications for 72 a range of issues such as sensitivity to stochastic fluctuations (Richter-Dyn & Goel, 73 1972; Wright & Hubbell, 1983) and risk of extinction (Purvis et al., 2000). Monitor-74 ing animal population changes in response to anthropogenic pressure is becoming 75 increasingly important as humans modify habitats and change climates as never before (Everatt et al., 2014). Sensor technology, such as camera traps (Rowcliffe & 77 Carbone, 2008; Karanth, 1995) and acoustic detectors (O'Farrell & Gannon, 1999; 78 Clark, 1995; Acevedo & Villanueva-Rivera, 2006) are becoming increasingly used to monitor changes in animal populations (Rowcliffe & Carbone, 2008; Kessel et al., 2014), as they are efficient, relativity cheap and non-invasive (Cutler & Swann, 1999), allowing for surveys over large areas and long periods. However, the problem of converting sampled count data to estimates of density remains as efforts 83 must be made to account for detectability of the animals (Anderson, 2001). 84

Methods do already exist for estimating animal density if the distance between 85 the animal and the sensor can be estimated (e.g., capture-mark recapture meth-86 ods (Karanth, 1995) and distance sampling (Harris et al., 2013)). However, these 87 methods often require additional information that may not be available. For example, capture-mark-recapture methods (Karanth, 1995; Trolle & Kéry, 2003; Soisalo & Cavalcanti, 2006; Trolle et al., 2007) require recognition of individuals; distance methods require a distance estimation of how far away individuals are from the sensor barlow2005estimates, marques2011estimating. The development of the ran-92 dom encounter model (REM) (a modification of a gas model) enabled animal den-93 sities to be estimated from unmarked individuals of a known speed, and sensor detection parameters (Rowcliffe et al., 2008). The REM method has been success-95 fully applied to estimate animal densities from camera trap surveys (Manzo et al., 96 2012; Zero et al., 2013). However, extending the REM method to other types of sensors (for example acoustic detectors) is more problematic, because the original

derivation assumes a relatively narrow sensor width (up to $\pi/2$ radians) and that the animal is equally detectable irrespective of its heading (Rowcliffe *et al.*, 2008).

Whilst these restrictions are not problematic for most camera trap makes (e.g. Reconyx, Cuddeback), the REM could not be used to estimate densities from cam-era traps with a wider sensor width (e.g. canopy monitoring with fish eye lens (Brusa & Bunker, 2014)). Additionally, the REM method would not be useful in estimating densities from acoustic survey data as the acoustic detector angles are often wider than $\pi/2$ radians. Acoustic detectors are designed for a range of di-verse tasks and environments (Kessel et al., 2014), which will naturally lead to a wide range of sensor detection widths and detection distances. In addition to this, calls emitted by many animals are directional (breaking the assumption of the REM method).

There has been a sharp rise in interest around passive acoustic detectors in recent years, with a 10 fold increase in publications in the decade between 2000 and 2010 (Kessel *et al.*, 2014). Acoustic monitoring is being developed to study many aspects of ecology, including the interactions of animals and their environments (Blumstein *et al.*, 2011; Rogers *et al.*, 2013), the presence and relative abundances of species (Marcoux *et al.*, 2011), and biodiversity of an area (Depraetere *et al.*, 2012).

Acoustic data suffers from many of the problems associated with data from camera trap surveys in that individuals are often unmarked so capture-make-recapture methods cannot be used to estimate densities. In some cases the distance between the animal and the sensor is known, for example when an array of sensors and the position of the animal is estimated by triangulation (Lewis *et al.*, 2007). In these situations distance-sampling methods can be applied, a method typically used for marine mammals (Rogers *et al.*, 2013). However, in many cases distance estimation is not possible, for example when single sensors are deployed, a situation typical in the majority of terrestrial acoustic surveys (Elphick, 2008; Buckland *et al.*, 2008). In these cases, only relative measures of local abundance can be calculated, and not absolute densities. This means that comparison of populations between species and sites is problematic without assuming equal detectability (Schmidt, 2003). Equality detectability is unlikely because of differences in environmental conditions, sensor type, habitats, species biology.

In this study we create a generalised REM (gREM), as an extension to the cam-131 era trap model of (Rowcliffe et al., 2008), to estimate absolute density from count 132 data from acoustic detectors, or camera traps, where the sensor width can vary 133 from 0 to 2π radians, and the signal given off from the animal can be directional. 134 We assessed the accuracy and precision of the gREM within a simulated environ-135 ment, by varying the sensor detection widths, animal signal widths, number of 136 captures and models of animal movement. We use the simulation results to rec-137 ommend best survey practice for estimating animal densities from remote sensors. 138

3. Methods

139

158

159

160

3.1. Analytical Model. The REM presented by (Rowcliffe et al., 2008) adapts the 140 gas model to model count data from camera trap surveys. The REM is derived as-141 suming a stationary sensor with a detection width less than $\pi/2$ radians. However, 142 in order to apply this approach more generally, and in particular to acoustic de-143 tectors, we need both to relax the constraint on sensor detection width, and allow 144 for animals with directional signals. Consequently, we derive the gREM for any 145 detection width, θ , between 0 and 2π with a detection distance r giving a circular sector within which animals can be captured (the detection zone)(Figure 1). Additionally, we model the animal as having an associated signal width α between 148 0 and 2π (Figure 1, see Appendix S1 for a list of symbols). We start deriving the 149 gREM with the simplest situation, the gas model where $\theta = 2\pi$ and $\alpha = 2\pi$. 150

3.1.1. *Gas Model.* Following Yapp (1956), we derive the gas model where sensors can capture animals in any direction and animal's signal is detectable from any direction($\theta = 2\pi$ and $\alpha = 2\pi$). We assume that animals are in a homogeneous environment, and move in straight lines of random direction with velocity v. We allow that our stationary sensor can capture animals at a detection distance r and that if an animal moves within this detection zone they are captured with a probability of one, while animals outside the zone are never captured.

In order to derive animal density, we need to consider relative velocity from the reference frame of the animals. Conceptually, this requires us to imagine that all animals are stationary and randomly distributed in space, while the sensor moves with velocity v. If we calculate the area covered by the sensor during the

survey period we can estimate the number of animals the sensor should capture.

As a circle moving across a plane, the area covered by the sensor per unit time is 2rv. The number of expected captures, z, for a survey period of t, with an animal

density of D is z = 2rvtD. To estimate the density, we rearrange to get D = z/2rvt.

3.1.2. gREM derivations for different detection and signal widths. Different combina-166 tions of θ and α would be expected to occur (e.g., sensors have different detection 167 widths and animals have different signal widths). For different combinations θ 168 and α , the area covered per unit time is no longer given by 2rv. Instead of the size 169 of the sensor detection zone having a diameter of 2r, the size changes with the 170 approach angle between the sensor and the animal. For any given signal width 171 and detector width and depending on the angle that the animal approaches the 172 sensor, the width of the area within which an animal can be detected is called the 173 profile, p. The size of the profile (averaged across all approach angles) is defined 174 as the average profile \bar{p} . However, different combinations of θ and α need different 175 equations to calculate \bar{p} . 176

We have identified the parameter space for the combinations of θ and α for which the derivation of the equations are the same (defined as sub-models in the gREM) (Figure 2). For example, the gas model becomes the simplest gREM sub-model (upper right in (Figure 2) and the REM from (Rowcliffe *et al.*, 2008) is another gREM sub-model where $\theta < \pi/2$ and $\alpha = 2\pi$. We derive one gREM sub-model SE2 as an example below (where $4\pi - 2\alpha < \theta < 2\pi$, $0 < \alpha < \pi$) (see Appendix S2 for other gREM sub-models).

3.1.3. Example derivation of SE2. In order to calculate \bar{p} , we have to integrate over the focal angle, x_1 (Figure 3a). This is the angle taken from the centre line of the sensor. Other focal angles are possible (x_2, x_3, x_4) and are used in other gREM sub-models (see Appendix S2). As the size of the profile depends on the approach angle, we present the derivation across all approach angles. When the sensor is directly approaching the animal $x_1 = \pi/2$.

Starting from $x_1 = \pi/2$ until $\theta/2 + \pi/2 - \alpha/2$, the size of the profile is $2r \sin \alpha/2$ (Figure 3b). During this first interval, the size of α limits the width of the profile.

When the animal reaches $x_1 = \theta/2 + \pi/2 - \alpha/2$ (Figure 3c), the size of the profile is

 $r \sin(\alpha/2) + r \cos(x_1 - \theta/2)$ and the size of $\theta/$ and α both limit the width of the profile (Figure 3c). Finally, at $x_1 = 5\pi/2 - \theta/2 - \alpha/2$ until $x_1 = 3\pi/2$, the width of the profile is again $2r \sin \alpha/2$ (Figure 3d) and the size of α again limits the width of the profile. The profile width p for π radians of rotation (from directly towards the sensor to directly behind the sensor) is completely characterised by the three intervals (Figure 3b–d). Average profile width \bar{p} is calculated by integrating these profiles over their appropriate intervals of x_1 and dividing by π which gives

$$\bar{p} = \frac{1}{\pi} \left(\int_{\frac{\pi}{2}}^{\frac{\alpha}{2} + \frac{\theta}{2} - \frac{\alpha}{2}} 2r \sin\frac{\alpha}{2} dx_1 + \int_{\frac{\pi}{2} + \frac{\theta}{2} - \frac{\alpha}{2}}^{\frac{5\pi}{2} - \frac{\theta}{2} - \frac{\alpha}{2}} r \sin\frac{\alpha}{2} + r \cos\left(x_1 - \frac{\theta}{2}\right) dx_1 + \int_{\frac{5\pi}{2} - \frac{\theta}{2} - \frac{\alpha}{2}}^{\frac{3\pi}{2}} 2r \sin\frac{\alpha}{2} dx_1 \right)$$

$$= \frac{r}{\pi} \left(\theta \sin\frac{\alpha}{2} - \cos\frac{\alpha}{2} + \cos\left(\frac{\alpha}{2} + \theta\right) \right)$$

$$= qn \ 1$$

$$= qn \ 2$$

We then, as with the gas model, use this expression to calculate density

200

$$D = z/vt\bar{p}.$$
 eqn 3

Rather than having one equation that describes \bar{p} globally, the gREM must be 202 split into submodels due to discontiunous changes in p as α and β change. These 203 discontinuities can occur for a number of reasons such as a profile switching be-204 tween being limited by α and θ , the difference between very small profiles and 205 profiles of size zero and the fact that the width of a sector stops increasing once 206 the central angle reaches π radians (i.e., a semi circle is just as wide as a full circle.) 207 As a visual example, if α is small, there is an interval between Fig. 3c and 3d 208 where the 'blind spot' would prevent animals being detected at all giving p = 0. 209 This would require an extra integral in our equation as simply putting our small value of α into eqn 1 would not give us this integral of p = 0. gREM submodel specifications were done by hand, and the integration was 212 done using SymPy (SymPy Development Team, 2014) in Python (Appendix S3). 213 The gREM submodels were checked by confirming that: 1) submodels adjacent 214 in parameter space were equal at the boundary between them; 2) submodels that 215 border $\alpha = 0$ had p = 0 when $\alpha = 0$; 3) average profile widths \bar{p} were between 0 and 216

2r and; 4) each integral, divided by the range of angles that it was integrated over, was between 0 and 2r. The scripts for these tests are included in Appendix S3 and 218 the R (R Development Core Team, 2010) implementation of the gREM is given in 219 Appendix S4. 220

221

3.2. **Simulation Model.** We tested the accuracy and precision of the gREM by developing a spatially explicit simulation of the interaction of sensors and animals 222 using different combinations of sensor detection widths, animal signal widths, 223 number of captures, and models of animal movement. 100 simulations were run 224 where each consisted of a 7.5 km by 7.5 km square (with periodic boundaries). A 225 stationary sensor of radius r was set up in the exact centre of each simulation, cov-226 ering 7 sensor detection widths θ between 0 and 2π (2/9 π , 4/9 π , 6/9 π , 8/9 π , 10/9 π , 227 $14/9\pi$, 2π). Each simulation was populated with a density of 70 animals km⁻², cal-228 culated from the equation in Damuth (1981) as the expected density of mammals of weighing 1 g. This density therefore represents the highest likely density of in-230 divudals, given that the smallest mammal is around 2 g Jones et al. (2009). A total 231 of 3937 individuals per simulation were created which were placed randomly at 232 the start of the simulation. Individuals were assigned 11 signal detection widths 233 α between 0 and π (1/11 π , 2/11 π , 3/11 π , 4/11 π , 5/11 π , 6/11 π , 7/11 π , 8/11 π , 9/11 π , 234 $10/11\pi, \pi$). 235 Each simulation lasted for N steps (14400) of duration T (15 minutes) giving a 236 total duration of 150 days. The individuals moved within each step with a distance 237 d, with an average speed, v. d, was sampled from a normal distribution with 238 mean distance, $\mu_d = vT$, and standard deviation $\sigma_d = vT/10$. An average speed, 239 $v = 40 \,\mathrm{km} \,\mathrm{days}^{-1}$, was chosen as this represents the largest day range of terrestrial 240 animals (Carbone et al., 2005), and represents the upper limit of realistic speeds. 241 At the end step, individuals were allowed to either remain stationary for a time 242 step (with a given probability, S), change direction (with a maximum angle, A) 243 between 0 and π . This resulted in 7 different movement models where: (1) simple 244 movement, where S and A = 0; (2)stop-start movement, where (i) S = 0.25, A = 0, 245 (ii) S = 0.5, A = 0, (iii) S = 0.75, A = 0; (3) random walk movement, where (i) S = 0.5246

, $A = \pi/3$, (ii) S = 0, $A = 2\pi/3$, iii) S = 0, $A = \pi$. Individuals were counted as they moved in and out of the detection zone of the sensor per simulation.

We calculated the estimated animal density from the gREM by summing the number of captures per simulation and inputting these values into the correct gREM submodel. gREM accuracy was determined by comparing the density in the simulation with the estimated density. High accuracy is indicated by the mean difference between the estimated and actual values not being significantly different from zero (Wilcoxon signed-rank test). gREM precision was determined by the standard deviation of estimated densities. We used this method to compare the accuracy and precision of all the gREM submodels. As these submodels are derived for different combinations of α and θ , the accuracy and precision of the submodels was used to determine the impact of different values of α and θ .

The influence of the number of captures and animal movement models on accuracy and precision was investigated using 4 different gREM submodels representative of the range α and θ values (submodels NW1, SW1, NE1, and SE3, Figure 2). Using these four submodels, we calculated how long the simulation needed to run to generate a range of different capture numbers (from 10 to 100 captures in 10 unit intervals), and estimated animal density. These estimated densities were compared to the real density to assess the impact on the accuracy and precision on the gREM of different simulation lengths. We also used these four submodels to compare the accuracy and precision of a simple movement model, to stop-start movement models and random walk movement models. The gREM assumes that individuals move continuously with straight-line movement (simple movement model) and we therefore assessed the impact of breaking the gREM assumptions.

271 4. RESULTS

4.1. **Analytical model.** The equation for \bar{p} has been newly derived for each submodel in the gREM, except for the gas model and REM which have been calculated previously. However, many models, although derived separately, have the same expression for \bar{p} . Figure 4 shows the expression for \bar{p} in each case. The general equation for density, using the correct expression for \bar{p} is then substituted into eqn 3. Although more thorough checks are performed in Appendix S3, it can be seen that all adjacent expressions in Figure 4 are equal when expressions for the boundaries between them are substituted in.

280 4.2. Simulation model.

4.2.1. gREM submodels. All gREM submodels showed a high accuracy, i.e., the 281 mean difference between the estimated and actual values was not significantly 282 different from zero across all models, corrected for multiple tests (all gREM sub 283 models Wilcoxon signed-rank test, p >0.002)(Figure 5). However, the precision of 284 the submodels do vary, where the gas model is the most precise and the SW7 sub 285 model the least precise, having the smallest and the largest interquartile range, re-286 spectively (Figure 5). The standard deviation of the error between the estimated 287 and true densities is strongly related to both the sensor and signal widths (Fig-288 ure 6), such that larger widths have lower standard deviations (greater precision). 289 However, even smaller sensor and signal widths have a relativity high level of 290 precision. 291

4.2.2. *Number of captures*. Within the four gREM submodels tested (NW1, SW1, SE3, NE1), the accuracy was not affected by the number of captures, where the mean difference between the estimated and actual values was not significantly different from zero across all capture rates, corrected for multiple tests (all gREM sub models Wilcoxon signed-rank test, p > 0.008)(Figure 7). However, the precision was dependent on the number of captures across all four of the gREM submodels, where precision increases as number of captures increases(Figure 7). For all gREM submodels, the the coefficient of variation falls to 10% at 100 captures.

4.2.3. Movement models. Within the four gREM submodels tested (NW1, SW1, SE3, NE1), neither the accuracy or precision was affected by the amount of time spent stationary. The mean difference between the estimated and actual values was not significantly different from zero for each category of stationary time (0, 0.25, 0.5 and 0.75), corrected for multiple tests (all gREM sub models Wilcoxon signed-rank test, p >0.12)(Figure 8a). Altering the maximum change in direction in each step (0, pi/3, 2pi/3, and pi) did not affect the accuracy or precision of the four gREM

submodels tested (all gREM sub models Wilcoxon signed-rank test, p >0.05)(Figure 8b). 308

5. Discussion

309

310

311

312

315

316

321

We have developed the gREM such that it can be used to estimate density from acoustic sensors and camera traps. This has entailed a generalisation of the gas model and the REM in (Rowcliffe et al., 2008) to be applicable to any combination of sensor width and signal directionality. We have used simulations to show, as a proof of principle, that these models are accurate and precise. The precision of the gREM was found to be dependent on the width of the sensor and the call, and the number of captures.

5.1. Analytical model. The gREM was derived for different combinations of α 317 and θ resulting in 25 different submodels, the expression for \bar{p} are equal for many of these submodels resulting in 8 different equations including the previously derived gas model and REM. These submodels were tested for consistency with adja-320 cent expressions being equal at their boundaries. These new submodels will allow researchers to evaluate the absolute density of animals that have previously been 322 difficult to study with noninvasive methods such as remove sensors. The gREM 323 allows the data from acoustic detectors to be used where an animal has a direc-324 tional calls, this could be used for a range of animals including bats, songbirds, 325 Cetaceans and forest primates. 326

There are a number of positive extensions to the gREM which could be devel-327 oped in the future. The original gas model was formulated for the case where both 328 subjects, either animal and detector, or animal and animal, are moving (Hutchinson & Waser, 2007). Indeed any of the models with animals that are equally de-330 tectable in all directions ($\alpha = 2\pi$) can be trivially expanded for moving by substi-331 tuting the sum of the average animal velocity and the sensor velocity for v as used 332 here. However, when the animal has a directional call, the extension becomes less 333 simple. The approach would be to calculate again the mean profile width. How-334 ever, for each angle of approach, one would have to average the profile width for 335 an animal facing in any direction (i.e. not necessarily moving towards the sen-336 sor) weighted by the relative velocity of that direction. There are a number of

situations where a moving detector and animal could occur and as such may be 338 advantage to have a method of estimating densities from the data collected, e.g. an 339 acoustic detector based off a boat when studying Cetacea or sea birds (Yack et al., 340 2013). Another interesting, and so far unstudied problem, is edge effects caused 341 by trigger delays (the delay between sensing an animal and attempting to record 342 the encounter) and time expansion acoustic detectors which repeatedly turn on 343 an off during sampling. Both of these have potential biases as animals can move 344 through the detection zone without being detected. The models herein are formu-345 lated assuming constant surveillance and so the error quickly becomes negligible. 346 For example, if it takes longer for the recording device to be switched on than the 347 length of some animal calls there could be a systematic underestimation of density.

5.2. Accuracy and Precision. We tested each of the gREM submodels for accuracy and precision through a simulation. All the submodels produced estimated den-350 sities that were not significantly different from the true density of the simulation. 351 Therefore based on these simulations we believe that the gREM has the poten-352 tial to produce accurate estimates for many different species, using either camera 353 traps or acoustic detectors. However the precision of the gREM differed between 354 submodels. For example, when the sensor and signal width were smaller then the 355 precision of the model was reduced, so when choosing a sensor for use in a gREM 356 study the detection width should be maximised, and if the study species has a 357 narrow signal directionality other aspects of the study protocol should be used to 358 compensate.

The precision of the gREM is greatly affected by the number of captures that are 360 collected, the coefficient of variation falls dramatically between 10 and 60 captures 361 and then after this continues to slowly reduce. At 100 captures the submodels 362 reach 10% coefficient of variation, and therefore we believe at this point the models 363 are precise. The length of surveys in the field will need to be adjusted so that 364 enough data is collected to reach this level of precision, populations of fast moving 365 animals or populations with large densities will require less survey effort than 366 those with slow moving or low densities. 367

The gREM was both accurate and precise for all the movement models we 368 tested against, stop-start movement and correlated random walks. However these 369 movement models are still simple representations of true animal movement which 370 often consist of multiple be dependent on multiple factors such as behavioural 371 state and and existence of home ranges (?). The accuracy of the gREM may be 372 affected by the interaction between the movement model and the size of the detec-373 tion radius. In figure 8b we studied a relatively long step length compared to the 374 size of the detection radius, and therefore the chance of catching the same animal 375 multiple times within a short space of time was reduced. However if the ratio of 376 step length to detection radius was smaller then this may decrease the precision of 377 the model, however this should not decrease its accuracy. 378

Although we have used simulations to validate the gREM submodels, much 379 more robust testing is needed. Although difficult, proper field test validation 380 would be required before the models could be fully trusted. The REM (Rowcliffe et al., 2008) has already been field tested, and both Rowcliffe et al. (2008) and Zero 382 et al. (2013) both found that the REM was an effective manner of estimating animal 383 densities (Rowcliffe et al., 2008; Zero et al., 2013). In some taxa gold standard meth-384 ods of estimating animal density exist, such as capture mark recapture. Where 385 these gold standard exist, and have been proved to work, a simultaneous gREM 386 study could be completed to test the accuracy under field conditions. An eas-387 ier way to continue to evaluate the models is to run more extensive simulations 388 which break the assumptions of the analytical models. The main element that 389 cannot be analytically treated is the complex movement of real animals. There-390 fore testing these methods against true animal traces, or more complex movement models would be required. 392

381

391

393

394

395

396

397

398

Within the simulation we have assumed an equal density across the entire world, however in a field environment the situation would be much more complex, with additional variation coming from local changes in density between camera sites. In the simulation we ran the speed of the animal as 40 km days⁻¹, the largest day range of terrestrial animals (Carbone et al., 2005), other speed values should not alter the accuracy or the precision of the gREM. We also assume perfect knowledge of the average speed of an animal and size of the detection zone, and instant

triggering of the camera. All of which may lead to possible bias or a decrease in precision.

5.3. **Implications for conservation.** The gREM is therefore available for the esti-402 mation of density of a number of taxa of importance to conservation, zoonotic dis-403 eases and ecosystem services. The models provided are suitable for certain groups 404 for which there are currently no, or few, effective methods for density estimation. 405 Any species that would be consistently recorded at least once when within range 406 of a detector would be a suitable subject for the gREM, such as bats (Kunz et al., 407 2009), songbirds (Buckland & Handel, 2006), Cetaceans (Marques et al., 2009) or forest primates (Hassel-Finnegan et al., 2008). Within increasing technological ca-409 pabilities, this list of species is likely to increase dramatically. 410 Importantly the methods are noninvasive and do not require human marking or 411 naturally identifying marks (as required for mark-recapture models). This makes 412 them suitable for large, continuous monitoring projects with limited human re-413 sources. It also makes them suitable for species that are under pressure, species 414 that cannot naturally be individually recognised or species that are difficult or 415 dangerous to catch.

6. ACKNOWLEDGMENTS

418 REFERENCES

417

Acevedo, M.A. & Villanueva-Rivera, L.J. (2006) Using automated digital recording systems as effective tools for the monitoring of birds and amphibians. *Wildlife Society Bulletin*, **34**, 211–214.

Anderson, D.R. (2001) The need to get the basics right in wildlife field studies.

Wildlife Society Bulletin, pp. 1294–1297.

Blumstein, D.T., Mennill, D.J., Clemins, P., Girod, L., Yao, K., Patricelli, G., Deppe, J.L., Krakauer, A.H., Clark, C., Cortopassi, K.A. *et al.* (2011) Acoustic monitoring in terrestrial environments using microphone arrays: applications, technological considerations and prospectus. *Journal of Applied Ecology*, **48**, 758–767.

- Brusa, A. & Bunker, D.E. (2014) Increasing the precision of canopy closure es-
- timates from hemispherical photography: Blue channel analysis and under-
- exposure. Agricultural and Forest Meteorology, **195**, 102–107.
- Buckland, S.T. & Handel, C. (2006) Point-transect surveys for songbirds: robust
- methodologies. *The Auk*, **123**, 345–357.
- Buckland, S.T., Marsden, S.J. & Green, R.E. (2008) Estimating bird abundance:
- making methods work. *Bird Conservation International*, **18**, S91–S108.
- Carbone, C., Cowlishaw, G., Isaac, N.J. & Rowcliffe, J.M. (2005) How far do ani-
- mals go? Determinants of day range in mammals. The American Naturalist, 165,
- 437 290–297.
- ⁴³⁸ Clark, C.W. (1995) Application of US Navy underwater hydrophone arrays for
- scientific research on whales. Reports of the International Whaling Commission, 45,
- 440 210–212.
- Cutler, T.L. & Swann, D.E. (1999) Using remote photography in wildlife ecology:
- a review. Wildlife Society Bulletin, pp. 571–581.
- Damuth, J. (1981) Population density and body size in mammals. Nature, 290,
- 444 699–700.
- Depraetere, M., Pavoine, S., Jiguet, F., Gasc, A., Duvail, S. & Sueur, J. (2012) Mon-
- itoring animal diversity using acoustic indices: implementation in a temperate
- woodland. Ecological Indicators, 13, 46–54.
- Elphick, C.S. (2008) How you count counts: the importance of methods research
- in applied ecology. *Journal of Applied Ecology*, **45**, 1313–1320.
- Everatt, K.T., Andresen, L. & Somers, M.J. (2014) Trophic scaling and occupancy
- analysis reveals a lion population limited by top-down anthropogenic pressure
- in the limpopo national park, mozambique. *PloS one*, **9**, e99389.
- 453 Harris, D., Matias, L., Thomas, L., Harwood, J. & Geissler, W.H. (2013) Applying
- distance sampling to fin whale calls recorded by single seismic instruments in
- the northeast atlantic. The Journal of the Acoustical Society of America, 134, 3522–
- 456 3535.
- 457 Hassel-Finnegan, H.M., Borries, C., Larney, E., Umponjan, M. & Koenig, A. (2008)
- 458 How reliable are density estimates for diurnal primates? International Journal of
- 459 *Primatology*, **29**, 1175–1187.

- 460 Hutchinson, J.M.C. & Waser, P.M. (2007) Use, misuse and extensions of "ideal gas"
- models of animal encounter. Biological Reviews of the Cambridge Philosophical So-
- 462 *ciety*, **82**, 335–359.
- Jones, K.E., Bielby, J., Cardillo, M., Fritz, S.A., O'Dell, J., Orme, C.D.L., Safi, K.,
- 464 Sechrest, W., Boakes, E.H., Carbone, C. et al. (2009) PanTHERIA: a species-level
- database of life history, ecology, and geography of extant and recently extinct
- mammals: Ecological archives e090-184. *Ecology*, **90**, 2648–2648.
- 467 Karanth, K. (1995) Estimating tiger (Panthera tigris) populations from camera-trap
- data using capture–recapture models. *Biological Conservation*, **71**, 333–338.
- 469 Kessel, S., Cooke, S., Heupel, M., Hussey, N., Simpfendorfer, C., Vagle, S. & Fisk, A.
- 470 (2014) A review of detection range testing in aquatic passive acoustic telemetry
- studies. *Reviews in Fish Biology and Fisheries*, **24**, 199–218.
- Kunz, T.H., Betke, M., Hristov, N.I. & Vonhof, M. (2009) Methods for assessing
- colony size, population size, and relative abundance of bats. Ecological and be-
- 474 havioral methods for the study of bats (TH Kunz and S Parsons, eds) 2nd ed Johns
- 475 Hopkins University Press, Baltimore, Maryland, pp. 133–157.
- 476 Lewis, T., Gillespie, D., Lacey, C., Matthews, J., Danbolt, M., Leaper, R.,
- 477 McLanaghan, R. & Moscrop, A. (2007) Sperm whale abundance estimates from
- acoustic surveys of the ionian sea and straits of sicily in 2003. Journal of the Ma-
- rine Biological Association of the United Kingdom, **87**, 353–357.
- 480 Manzo, E., Bartolommei, P., Rowcliffe, J.M. & Cozzolino, R. (2012) Estimation of
- population density of european pine marten in central italy using camera trap-
- 482 ping. *Acta Theriologica*, **57**, 165–172.
- 483 Marcoux, M., Auger-Méthé, M., Chmelnitsky, E.G., Ferguson, S.H. & Humphries,
- 484 M.M. (2011) Local passive acoustic monitoring of narwhal presence in the cana-
- dian arctic: a pilot project. Arctic, pp. 307–316.
- 486 Marques, T.A., Thomas, L., Ward, J., DiMarzio, N. & Tyack, P.L. (2009) Estimating
- cetacean population density using fixed passive acoustic sensors: An example
- with Blainville's beaked whales. The Journal of the Acoustical Society of America,
- **125**, 1982–1994.
- 490 O'Farrell, M.J. & Gannon, W.L. (1999) A comparison of acoustic versus capture
- techniques for the inventory of bats. *Journal of Mammalogy*, pp. 24–30.

- Purvis, A., Gittleman, J.L., Cowlishaw, G. & Mace, G.M. (2000) Predicting extinc-
- tion risk in declining species. Proceedings of the Royal Society of London Series B:
- 494 Biological Sciences, **267**, 1947–1952.
- 495 R Development Core Team (2010) R: A Language And Environment For Statistical
- 496 Computing. R Foundation For Statistical Computing, Vienna, Austria. ISBN 3-
- 497 900051-07-0.
- ⁴⁹⁸ Richter-Dyn, N. & Goel, N.S. (1972) On the extinction of a colonizing species. *The*-
- oretical Population Biology, **3**, 406–433.
- Rogers, T.L., Ciaglia, M.B., Klinck, H. & Southwell, C. (2013) Density can be mis-
- leading for low-density species: benefits of passive acoustic monitoring. *Public*
- Library of Science One, 8, e52542.
- Rowcliffe, J.M. & Carbone, C. (2008) Surveys using camera traps: are we looking
- to a brighter future? *Animal Conservation*, **11**, 185–186.
- Rowcliffe, J., Field, J., Turvey, S. & Carbone, C. (2008) Estimating animal density
- using camera traps without the need for individual recognition. Journal of Ap-
- plied Ecology, **45**, 1228–1236.
- 508 Schmidt, B.R. (2003) Count data, detection probabilities, and the demography, dy-
- namics, distribution, and decline of amphibians. Comptes Rendus Biologies, 326,
- 510 119–124.
- 511 Soisalo, M.K. & Cavalcanti, S. (2006) Estimating the density of a jaguar population
- in the Brazilian Pantanal using camera-traps and capture-recapture sampling in
- combination with GPS radio-telemetry. *Biological Conservation*, **129**, 487–496.
- 514 SymPy Development Team (2014) SymPy: Python library for symbolic mathematics.
- Trolle, M. & Kéry, M. (2003) Estimation of ocelot density in the Pantanal using
- capture-recapture analysis of camera-trapping data. Journal of mammalogy, 84,
- 517 607-614.
- Trolle, M., Noss, A.J., Lima, E.D.S. & Dalponte, J.C. (2007) Camera-trap studies of
- maned wolf density in the Cerrado and the Pantanal of Brazil. Biodiversity and
- 520 *Conservation*, **16**, 1197–1204.
- Wright, S.J. & Hubbell, S.P. (1983) Stochastic extinction and reserve size: a focal
- species approach. Oikos, pp. 466–476.

Lucas et al.: A generalised random encounter model for animals

- Yack, T.M., Barlow, J., Calambokidis, J., Southall, B. & Coates, S. (2013) Passive
- acoustic monitoring using a towed hydrophone array results in identification of
- a previously unknown beaked whale habitat. The Journal of the Acoustical Society
- of America, **134**, 2589–2595.
- Yapp, W. (1956) The theory of line transects. *Bird study*, **3**, 93–104.
- ⁵²⁸ Zero, V.H., Sundaresan, S.R., O'Brien, T.G. & Kinnaird, M.F. (2013) Monitoring
- an endangered savannah ungulate, Grevy's zebra (Equus grevyi): choosing a
- method for estimating population densities. *Oryx*, **47**, 410–419.

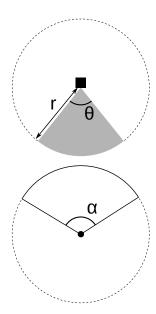


FIGURE 1. Representation of sensor detection width and animal signal width. The filled square and circle represent a sensor and an animal, respectively; θ , sensor detection width (radians); r, sensor detection distance; dark grey shaded area, sensor detection zone; α , animal signal width (radians). Dashed lines around the filled square and circle represents the maximum extent of θ and α , respectively.

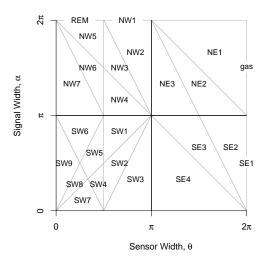


FIGURE 2. Locations where derivation of the average profile \bar{p} is the same for different combinations of sensor detection width and animal signal width. Symbols within each polygon refer to each gREM submodel named after their compass point, except for Gas and REM which highlight the position of these previously derived models within the gREM. Symbols on the edge of the plot are for submodels with $\alpha, \theta = 2\pi$

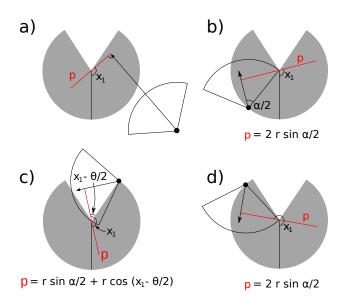


FIGURE 3. An overview of the derivation of SE2. The filled circles represent animals, with the animal signal shown as a unfilled sector and the direction of movement shown as an arrow. The detection zone of the sensors are shown as filled grey sectors with a detection distance of r. The SYMBOL shows the direction the sensor is facing; θ , sensor detection width; α , animal signal width. The profile p (the line an animal must pass through in order to be captured) is shown in red and x_1 is the focal angle, where (a) shows the location of x_1 . The derivation of p changes as the animal approaches the sensor from different directions where (b) is the derivation of p when p is in the interval $\left[\frac{\pi}{2}, \frac{\pi}{2} + \frac{\theta}{2} - \frac{\alpha}{2}\right]$, (c) p when p is in the interval $\left[\frac{\pi}{2}, \frac{\pi}{2} - \frac{\theta}{2} - \frac{\alpha}{2}\right]$ and (d) p when p is in the interval $\left[\frac{\pi}{2}, \frac{\pi}{2} - \frac{\theta}{2} - \frac{\alpha}{2}\right]$. The resultant equation for p is shown beneath each figure.

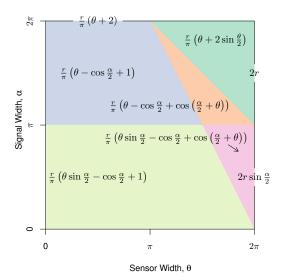


FIGURE 4. Expressions for the average profile width, \bar{p} , given sensor and signal widths. Despite independent derivation within each block, many models result in the same expression. These are collected together and presented as one block of colour. Expressions on the edge of the plot are for submodels with α , $\theta = 2\pi$.

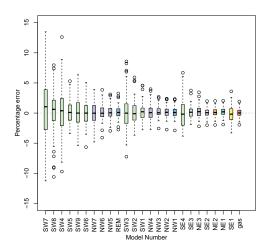


FIGURE 5. Simulation model results of the accuracy and precision for gREM submodels. The precentage error between estimated and true density for each gREM sub model is shown within each box plot, where the black line represents the median percentage error across all simulations, boxes represent the the middle 50% of the data. Box colours correspond to the expressions for average profile width \bar{p} given in f:equalModelResults.

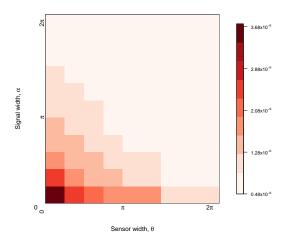


FIGURE 6. Simulation model results of the gREM precision given a range of sensor and signal widths, shown by the standard deviation of the error between the estimated and true densities. Standard deviations are shown from deep red to pink, representing high to low values between 0.483×10^-6 to 3.74×10^-6 .

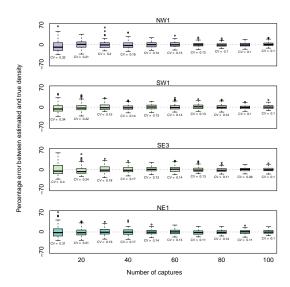


FIGURE 7. Simulation model results of the accuracy and precision of four gREM submodels (NW1, SW1, SE3 and NE1) given different numbers of captures. The percentage error between estimated and true density within each gREM sub model for capture rate is shown within each box plot. Sensor and signal widths vary between submodels. The colour of each box plot corresponds to the expressions for average profile width \bar{p} given in Figure 4

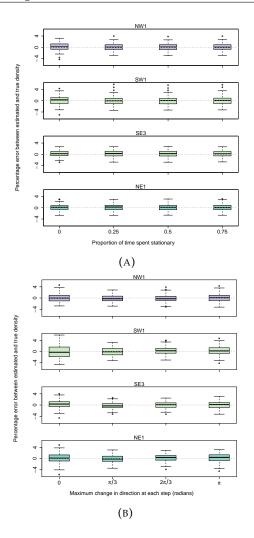


FIGURE 8. Simulation model results of the accuracy and precision of four gREM submodels (NW1, SW1, SE3 and NE1) given different movement models where (A) amount of time spent stationary (stop-start movement) and (B) maximum change in direction at each step (correlated random walk model). The percentage error between estimated and true density within each gREM sub model for the different movement models is shown within each box plot. The simple model is represented where time and maximum change in direction equals 0. The colour of each box plot corresponds to the expressions for average profile width \bar{p} given in 4.