Environment and Time as Constraints on the Biogeographical Distribution of Gibbons

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Highlights:

- Time budgets limit the biogeographic distribution of gibbons and siamang
- Time that has to be allocated to resting seems to be the main constraint
- The limits on group size are not so restrictive as to make monogamy obligatory

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2 # Environment and Time as Constraints on the Biogeographical

3 ## Distribution of Gibbons

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5 by

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Abstract

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28   We develop a time budget model for the hylobatid family with the aim of assessing

29   the extent to which their contemporary and historical biogeographic distributions

30   might be explained by ecological constraints. The model uses local climate to predict

31   time budgets, and from this the limiting size of social group that animals could

32   manage at a given location. The model predicts maximum group sizes that vary

33   between 3-15 within the taxon's current distribution, indicating that the combination

34   of their dietary and locomotor styles with the kinds of habitats they inhabit radically

35   constrain group size. Beyond the edges of their current distribution, sustainable group

36   size rapidly tends to zero, although if they had been able to bypass some of these

37   areas, they would have found very suitable habitats in southern India and beyond the

38   Wallace Line. While travel time would be a major constraint on group size at larger

39   group sizes, as it is in great apes, the main factor limiting the gibbon's current

40   distribution is the time they need to spend resting that is imposed on them by the

41   environment. The model also indicates that gibbons would not now be able to survive

42   in regions of central and southeastern China where they are known to have occurred

43   within historical times, perhaps because historical climate change following the Little

44   Ice Age of the C18th made these regions uninhabitable for them. Finally, our results

45   indicate that gibbons have the ecological capacity to live in larger groups than they

46   do, making it unlikely that their adoption of monogamy reflects purely ecological

47   constraints.

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50   *Key Words*: climate, biogeographic distribution, group size, foraging ecology

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54   *Highlights:*
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56   • Time budgets limit the biogeographic distribution of gibbons and siamang
57   • Time that has to be allocated to resting seems to be the main constraint
58   • The limits on group size are not so restrictive as to make monogamy
59   obligatory
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63   **Introduction**

64   Inability to satisfy nutritional and other demands within a defined time period

65   (usually a 24-hr physiological cycle: Peters 1983) limits both where a species can live

66   and, in social species, how big their groups can be (Dunbar, Korstjens & Lehmann

67   2009). In this respect, time is a central problem for all animals, especially for those

68   that, like primates, are not active 24 hours a day. This insight has led to the

69   development of a series of time budget models designed to understand the role of time

70   in limiting the biogeographical distributions of individual taxa (Dunbar et al. 2009).

71   These models are premised on the fact that behaviour (as the outcome of core

72   physiological processes) is the interface between the environment and the animals'

73   ability to survive (with the latter indexed by the size of group they can maintain as an

74   ecological entity) (see also Marshall, Carter, Rowcliffe & Cowlishaw 2013). There is

75   little that most animals can do to alter their physiology in response to changes in

76   climate or vegetation, but they can, and do, adjust their behaviour and group sizes in

77   response to changing environmental conditions.

78   For species like anthropoid primates that (with obvious the exception *Aotus*)

79   are strictly diurnal due to poor night vision, the time available for satisfying their

80   nutritional requirements is further reduced to the ~12 hours of tropical daylight. In

81   addition, climate and the species' dietary adaptations may force animals to rest

82   ('enforced rest' *sensu* Korstjens et al. 2010), thereby further reducing the length of

83   their active day. The climatic component of this relationship reflects the fact that

84   almost all tropical animals are obliged to cease being active and seek shelter during

85   the hottest part of the day when ambient temperatures in open sunlight exceed their

86   thermoneutral zone (Mount 1979; Dàvid-Barrett & Dunbar 2016); the dietary aspect is

87   mainly a problem for folivores for whom gut fermentation of foliage is incompatible

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88   with any form of activity because the microbial activity on which it depends is

89   extremely sensitive to the heat generated by physical activity (van Soest 1982).

90   Intensely social species, like most anthropoid primates, face an additional

91   problem because the functionality of their relationships, as well as the cohesion of

92   their social groups, depends directly on the time they invest in grooming with each

93   other (Lehmann, Korstjens & Dunbar 2007a; Pollard & Blumstein 2008; Dunbar &

94   Shultz 2010; Dunbar & Lehmann 2013; Sutcliffe, Dunbar, Binder & Arrow 2012;

95   Dunbar 2018a,b). For both humans and nonhuman primates, an individual's

96   willingness to give coalitionary aid to another depends directly on the amount of time

97   they spend engaged in affiliative interaction (Seyfarth & Cheney 1984; Dunbar 1980,

98   2018a,b; Burton-Chellew & Dunbar 2015). At some point, animals will inevitably run

99   out of time, and this will ultimately limit where they can live. Ecologists tend to see

100   this as a constraint imposed by nutrient availability, but the real constraint is time.

101   In effect, animals face a three-way optimization problem in which they have to

102   offset the costs of living in social groups (i.e. time investment in social grooming) and

103   the acquisition of nutrients (specified by the ecological determinants of time required

104   for feeding and travel, including the fixed costs of fuelling fertility and lactation)

105   against the benefits of group size as their principal form of predator deterrence.

106   Predation risk will depend on whether the animal is terrestrial or arboreal and on the

107   hunting styles of the local predators (the 'landscape of fear') (Shultz et al. 2004;

108   Shultz & Finlayson 2010; Tolon et al. 2009; Laundré et al. 2014; Coleman & Hill

109   2014; Riginos 2015; Gallagher, Creel, Wilson & Cooke 2017), but within these

110   constraints the prey species can adjust group size to offset local predation risk.

111   Focusing on time budgets is not an alternative to the more conventional socio-

112   ecological approach that typically seeks to identify correlations between, on the one

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113   hand, group size, population density or behaviour and, on the other hand, climate or

114   forage quality/distribution (Wrangham, Gittleman & Chapman 1993; Chapman &

115   Chapman 2000; Snaith & Chapman 2007; Clutton-Brock & Janson 2012), or

116   apraoches that explore the efficiency of foraging (e.g. Brockleman et al. 2014).

117   Rather, a focus on time budgets provides a way of looking at the mechanisms that

118   underpin the correlations between environmental variables and population level

119   outcomes (group size and dispersion). Time budgets thus provide insights into the

120   mechanisms involved in these processes, and, in doing so, emphasize a component of

121   that mechanism (namely, time) that is invariably overlooked in most studies.

122   In addition, a time budget approach allows us to incorporate, directly or

123   indirectly, a range of other factors and causal relationships that bear on animals'

124   ability to survive in particular locations. This is important because biological

125   phenomena are naturally systems-based, with most causal relationships being

126   multivariate and subject to the influence of feedback loops. Failure to incorporate

127   these more complex relationships may result in the over- or underestimation of the

128   importance of particular causal relationships. Time budget models allow us to

129   integrate a range of variables and relationships into a single, coherent model.

130   The formal structure of time budget models is that of a causal chain. The

131   climate at a given location directly or (via its effect on vegetation quality) indirectly

132   determines the three core elements of the time budget (feeding, moving and minimum

133   or 'enforced' rest time). These in turn determine the limiting size of group that

134   animals can maintain at that location because this depends on how much

135   'uncommitted' time the animals then have available and the parametric relationship

136   between time invested in relationships and the resulting quality of those relationships

137   (as reflected in group coherence). The maximum possible group size predicted by the

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138 time budget ultimately determines whether or not the species can live at that location

139 (Dunbar et al. 2009). It is important to appreciate that these models are location-

140 specific: they make explicit predictions about the behaviour of the animals at a given

141 location, subject to the particular climatic and vegetational conditions pertaining at

142 that location. They are also necessarily genus-specific because they reflect the

143 particular dietary, physiological and body size adaptations that characterize a genus.

144 In this respect, a genus is an ecological species.

145   Climate variables affect both energy demand (energy animals need to maintain

146 thermoregulation: Mount 1979) and the rate of nutrient intake through the effect

147 climate has on the quality and digestibility of forage (and hence the amount of forage

148 that has to be eaten to extract the required amount of nutrients: van Soest 1982). They

149 also influence both travel time (through their effect on foraging patch size and inter-

150 patch distances: Janson & van Schaik 1988; Chapman & Chapman 2000) and rest

151 time (when animals are forced to seek shelter because ambient temperatures exceed

152 their thermoneutral zone: Mount 1979; Dàvid-Barrett & Dunbar 2016). In addition,

153 the feedback loops that are invariably an important component of biological systems

154 can arise when group size, in particular, influences travel time (because larger groups

155 have to travel further if feeding patches are depleted easily – typically, less of a

156 problem for folivores), and this in turn requires more time to be devoted to foraging to

157 replace the energy consumption of the additional travel.

158   Time budget models consist of a set of simultaneous equations (one for each

159 of the core time budget variables) and thus have the form of a linear programming (or

160 linear optimization) model (Dunbar 2002). The intersection of these equations defines

161 the limit on group size that a species can maintain at a given location. If any of these

162 relationships involve nonlinear components, solving the simultaneous equation set to

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163 find the maximum group size usually requires numerical rather than analytical

164 methods (Dunbar 1992a; Dunbar et al. 2009). However, if all relationships are linear,

165 analytical methods based on substitution can be used. The set of climatic parameters

166 (the primary drivers, with each as a separate X, Z,…. axis) and group size (the final

167 output, or Y-axis, variable) define a multidimensional state space. The isosurface

168 defined by points where maximum group size drops below the minimum required for

169 demographic viability and/or predator defence defines the taxon's biogeographic

170 distribution. So long as the predicted maximum group size lies above these minima,

171 the species can survive at those locations, with the magnitude of the difference

172 between the maximum and these minima providing a measure of how much

173 ecological stress the animals are under.

174   These mechanism-based models are at least as successful as conventional

175 climate (or niche) envelope models at predicting the geographical distributions of the

176 taxa concerned (Willems & Hill 2009; Korstjens et al. 2010). In addition, they provide

177 insights into which aspects of the animals' ecology are responsible for limiting their

178 distribution (Dunbar et al. 2009). This is because time budget models allow us (a) to

179 determine how much ecological and demographic stress a particular population is

180 under (Dunbar 1992a; Lehmann et al. 2007a) and (b) to specify why the species is

181 unable to occupy particular habitats in ways that directly reflect their physiological

182 adaptations (Dunbar et al. 2009). Neither of these is possible with conventional

183 climate envelope models, which are essentially simple correlational relationships. In

184 addition, because time budget models are driven directly by climate, the models can

185 be used to explore the consequences of past and future climate change for a taxon's

186 biogeographic distribution in a more nuanced way (Lehmann et al. 2010; Kortsjens,

187 Lehmann & Dunbar 2010; Bettridge & Dunbar 2012).

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188   Time budget models have been published for a number of terrestrial (gelada:

189   Dunbar 1992a; baboons: Dunbar 1992b; Bettridge, Lehmann & Dunbar 2010; African

190   great apes: Lehmann, Korstjens & Dunbar 2007b, 2008a) and arboreal (spider

191   monkeys: Korstjens, Verhoeckx & Dunbar 2006; colobins: Korstjens & Dunbar 2007;

192   vervets: Willems & Hill 2009; forest guenons: Korstjens, Lehmann & Dunbar 2018;

193   orangutans: Carne, Semple & Lehmann 2012) primates, as well as one ungulate (feral

194   goats: Dunbar & Shi 2013).

195   Although models have been developed for all three great apes (Lehmann et al.

196   2007b, 2008a,b; Carne et al. 2012), the small apes (the gibbon family, Hylobatidae)

197   have yet to be considered. The small apes offer a particularly interesting challenge for

198   several reasons. First, for an ape, they occupy an unusually exclusive arboreal

199   terminal branch niche otherwise occupied only by some of the smaller monkeys.

200   Second, they typically live in very small groups (monogamous pairs plus dependent

201   offspring) and this raises questions as to why they have opted for this form of social

202   system. One classic explanation for monogamy in mammals is that females are forced

203   by their ecology to forage on their own in territories that are too large for a male to be

204   able to successfully defend more than one female (Wittenberg & Tilson 1980; van

205   Schaik & Dunbar 1990; Komers & Brotherton 1997; Lukas & Clutton-Brock 2013);

206   an alternative suggestion is that male parental support allows females to reduce their

207   energy demand, especially in strongly seasonal habitats or where locomotor costs

208   prohibit fat storage (Dunbar 1995a,b; Heldstab, van Schaik & Isler 2017). Third, they

209   have a relatively limited distribution in southeast Asia, and this limited range begs

210   explanation.

211   Historically, hylobatids probably evolved as a distinct family in southwest

212   China in the late Miocene, and gradually extended their range down through the

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213   Malay peninsula into the islands of the Sunda shelf as opportunities provided by

214   changes in climate and sea level allowed (Jablonski & Chaplin 2009). Turvey, Crees

215   & Di Fonzo (2015) found that, as late as the 1800s, gibbons were still being recorded

216   in historical documents as widely distributed throughout southern and central China as

217   far north as latitude ~35°N (Shanxi, Shaanxi and Shangdong provinces). What

218   explains their disappearance from all but the southwest corner of China (Yunnan and

219   Hainan) remains to be determined, though anthropogenic factors have been suspected

220   (Fan 2017).

221   In this paper, we have five main aims. The first is to develop a model of

222   hylobatid time budgets in terms of climatic variables. Second, we test the model's

223   validity by using it to predict the taxon's presence and absence in various locations

224   within continental Asia and the islands of the Sunda Shelf and New Guinea. Third, we

225   use the model to determine which time budget variables are most responsible for

226   limiting the taxon's biogeographic distribution. Fourth, we use the model to determine

227   the possible causes of the historical extinction of gibbon populations in China.

228   Finally, we use the results to assess how plausible it is that female spacing best

229   explains hylobatid monogamy.

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231   **Methods**

232   *Gibbon data*

233   A database was compiled from the gibbon literature to provide quantitative

234   data on key demographic, environmental and ecological variables. We identified 77

235   studies that provide data on at least some of these (see online *Dataset S1*). Of these,

236   59 provide data on mean group size, 41 provide data on group density or biomass, and

237   29 provide data on activity budgets and other behavioural or demographic variables.

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238   In respect of activity budgets, we identify five mutually exclusive states (feeding,

239   moving or travel, resting, social interaction – mainly social grooming – and singing)

240   which, in this sample, between them account for ~100% of time.

241   Inevitably, a number of issues arise when comparing data from different

242   studies. One is that studies vary in the definitions they use for individual behaviour

243   categories; a second is that methods of collecting time budget data vary, both in

244   intensity (e.g. some researchers use group scans, others focal individual sampling) and

245   in coverage (the hours of daylight may not be sampled evenly). While these are

246   certainly issues, past experience developing time budget models suggests that the

247   impact of these methodological issues is at best modest (see Dunbar & Shi 2013).

248   More importantly, the main consequence of variations in definition and procedure is

249   that they increase the error variance in parameter estimates; increased error variance

250   will simply make it harder to obtain significant results, and will therefore bias the

251   statistical analysis in favour of the null hypothesis (i.e. no relationship). Ultimately, of

252   course, the real test is whether our models predict what we see on the ground *despite*

253   these methodological flaws. We should be less concerned with standardisation of

254   definitions or methods (though these should always be encouraged) than with whether

255   models based on them correctly predict what we know to be the case.

256   Where these were given by the original field sources, we also extracted data

257   on a number of ecological and behavioural variables: the percentage of leaf in the diet

258   (%Leaf), percentage of fruit in the diet (%Fruit), the density of gibbon groups

259   (groups/ha), gibbon population biomass (kg/km^2), mean day journey length (km),

260   mean territory size (ha), density of fig trees (Figdens, indexed as trees/ha) and number

261   of sympatric primate species. Mean day journey length and time spent moving

262   (transformed into hours spent moving per day) were used to calculate travel speed

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263 (m/hr). Together, these comprise the set of ecological and covariate variables we will

264 use in model-building.

265   Data for all sites are provided in online *Dataset S1*.

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267 *Climatic and geophysical data*

268   Rainfall, altitude and latitude for each study site are those provided by the

269 cited publications, where these are given. Since latitudinal effects should be

270 symmetrical about the equator, we transform all latitudes into absolute latitude. Since

271 individual studies often do not provide all the indices we need, we have sourced all

272 temperature variables (mean annual temperature, TEMP; mean minimum temperature,

273 $TEMP_{min}$; mean maximum temperature, $TEMP_{max}$; mean daily temperature variation,

274 $TEMP_{var}$ [difference between mean minimum and mean maximum temperature]; and

275 the standard deviation of mean monthly temperature, T_{moSD} [an index of seasonality])

276 from either http://www.globalspecies.org/weather_stations/ (which provides climate

277 data for individual wildlife reserves) or http://en.climate-data.org (which provides

278 climate data for civic weather stations). Wherever possible, we gave preference to the

279 first. These climatic variables and the two geographical variables (i.e. latitude and

280 altitude, both of which are determinants of climate) constitute the set of climatic

281 variables that we will use as the main independent variables in model-building.

282   Previous analyses of weather station data for sub-Saharan Africa have

283 demonstrated that only three climatic variables are needed to predict evapo-

284 transpiration (the principal predictor of tropical primary productivity: Rosenzweig

285 1968; Le Houérou & Hoste 1977; Lo Seen Chong, Mougin & Gastellu-Etchegorry

286 1993). These are mean ambient temperature, total annual rainfall and an index of

287 seasonality (Williamson & Dunbar 1998; see also Hill 1999). As in our previous

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models for African primates, the standard deviation of mean monthly temperature (TmoSD) is our main index of seasonality. This also allows us to include any effects due to the fact that, in some contexts, certain times of the year when resource availability is poor or climate especially challenging may impose limits on what animals can do (see, for example, Dunbar & Shi 2013).

Collinearity between the climate variables in the field site dataset is, in general, low and within conventionally acceptable levels, the only exception being that between rainfall and temperature (mainly due to the effect of the cold, dry habitats in southwest China disproportionately influencing what is otherwise a much less clearcut relationship). At best, this relationship explains only 20% of the variance, and as little as 10% if the Chinese habitats are excluded.

Model-building procedure

For the model-building analysis, we excluded time budget data for one study on a heavily logged habitat (Johns 1986) because it had an implausibly low estimate for time spent feeding (8%, >3 standard deviations below the mean for gibbons, and almost a full SD below the next lowest value) and an implausibly high value for time spent resting (62%, 2.5 SDs above the overall mean, and a full SD above the next highest value), two studies (West Garo Hills, NE India: Alfred & Sati 1986, 1990; Ujung Kulon, Indonesia: P. Kappeler 1984a,b) that had very high values for time spent feeding (>60%, 3.1 and 3.9 SDs above the mean for gibbons, and >1 SD above the next nearest value) and unusually low values for resting time, and one (Tanjong Triang: Ellefson 1974) that had an unusually high value for moving (47%) and low value for rest (7%). Such high values for feeding and moving and low values for resting are indicative of gibbons that were not fully habituated.

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In developing any model, we face a choice between detail and generality. It is always possible to build a model that takes every conceivable environmental variable into account, and hence is 100% accurate in its fit to the data. But in doing so, we inevitably lose generality: in order to make predictions about where the taxon can live, we will need to know much more about the particular environmental parameters of an individual location (e.g. soil type and acidity, inclination and tree composition, as well as all the larger scale variables like rainfall and temperature). Using general climate as the basis for a model allows us, at the expense of some loss of accuracy, to be more general, since relevant climate values can be mapped geographically on a continental scale from climate models, as well as being projected backwards and forwards in time.

For the purposes of building a time budget model, we need to determine taxon-specific equations for just two key variables (feeding and moving time). The other two main components (enforced resting time and social time) use general primate equations given, respectively, by Korstjens et al. (2010) and Lehmann et al. (2007a). For the purposes of the present model, we shall take time devoted to singing as a constant (at the mean observed value of 4.96%).

Enforced resting time, as defined by Korstjens et al. (2010), specifies the minimum amount of time that has to be spent resting as a consequence of high ambient temperatures (to avoid thermal overload or excessive heat loss) and for the digestion of leaves (as a function of climatic conditions and the species' dietary physiology). Enforced resting time differs from observed resting time (time spent inactive, as recorded by field observers) in that observed rest time consists of enforced resting time plus uncommitted time ("free rest") (Korstjens et al. 2010).

The time during the day that animals have left over after making their habitat-

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338   specific allocations to feeding, moving, enforced rest and singing (subject to any

339   feedback loops in the equation set) gives us the amount of time in the day that could

340   be devoted to social interaction (grooming). Because grooming time correlates with

341   group size (Dunbar 1992; Lehman et al. 2007; Dunbar & Lehmann 2014), we can use

342   this relationship to specify the maximum size of group that the animals could maintain

343   as a coherent social entity at that location. This does not mean the animals *have* to

344   devote all this time to social interaction: it simply sets the upper limit, and hence the

345   upper limit on social group size.

346   The social time equation reflects the fact that social grooming is the principal

347   bonding mechanism for primate social groups and increases linearly with group size

348   across primates as a whole (Dunbar 1991; Lehmann et al. 2007; Dunbar & Lehmann

349   2014). As such, it represents the time investment in social interaction that is necessary

350   to maintain a group's cohesion through time so as to prevent it disintegrating (Dunbar

351   1992a, 2012). It is worth noting that this increase in time devoted to grooming with

352   group size does not mean that animals in large groups groom with more individuals.

353   Typically they do not. Rather, as group size increases, animals invest increasingly in

354   the handful of core grooming partners that act as their primary coalition partners

355   within the group because these act as their primary defence against the stresses of

356   living in a group (Kudo & Dunbar 2001; Dunbar 2012, 2018a).

357   As in previous models, we first examined all relevant pairwise plots visually to

358   determine whether any factors correlate nonlinearly with the main time budget

359   variables. There were no nonlinear relationships. We then ran backward stepwise

360   linear regression models with the full set of climatic and ecological variables as

361   predictors of each of the two dependent variables (feeding and moving time). The

362   criterion used in selecting the final equation in each case was based on maximising

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363   the proportion of variance explained, subject to the requirement that the overall

364   equation and all its individual predictor variables were significant.

365   Because siamang are more than twice as large as other gibbons, siamang

366   feeding time was first corrected to a gibbon-equivalent (i.e. what it would need to do

367   if it was a conventional gibbon) by rescaling it by the ratio of the metabolic body

368   weights of the two taxa (i.e. $5.5^{0.75}/10.75^{0.75}=0.605$). This allows us to use all the data

369   without having to include an additional factor for taxon or body mass. None of the

370   other gibbon species differed significantly in body weight (see *Dataset S1*).

371   In extracting these equations from the data, we have ignored the possible

372   influence of phylogeny. As in previous analyses (see Dunbar et al. 2009), we do so

373   principally because almost all of the variables we are concerned with are behavioural

374   rather than anatomical, and are likely to be influenced more heavily by local

375   environmental conditions than by biological inheritance, and thus typically have

376   phylogenetic signals that are close to zero (see Kamilar & Cooper 2013). In effect, we

377   treat all the hylobatids as belonging to a single ecological species. Inter-population

378   analyses of just this kind have previously been successfully used to explore aspects of

379   the behaviour of gibbons (song function: Cowlishaw 1992, 1996). It is perhaps

380   important to note that we are not seeking to describe the "typical" behaviour of

381   individual gibbon species: rather, our concern is to understand the overall pattern of

382   behavioural flexibility and the effects of climate on biogeography across the whole

383   taxon. For convenience, we adopt the convention of referring to the entire group as

384   hylobatids, and differentiate between siamang and all other gibbon species only on the

385   basis of body size.

386   We then use this set of equations to determine maximum group size for any

387   given location. Since all the equations in the present case are linear, we use analytical

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388   methods and solve directly by substitution:

389         $N_{Max} = (100 - (Feed + Move + Rest_{Enf})) / (grooming\ equation)$

390   This value is the limiting group size (the maximum ecologically tolerable group size,

391   N_{max}).

392   In order to exhibit the biogeographical implications of the model in graphable

393   form, we present the main results as 3-dimensional surface plots of maximum group

394   size plotted against (a) absolute latitude and altitude and (b) mean annual rainfall and

395   mean annual temperature. Because a relatively large number of environmental and

396   behavioural variables are involved in the basic model, we need to convert all other

397   variables into functions of just the two variables used for each graph. We do this by

398   using backwards stepwise regression modelling to derive equations for these variables

399   from the data given in online *Dataset S1*. The resulting equations are given in Table 2.

400   Of these, latitude, altitude and temperature are well known to be systematically

401   related.

402   All statistical analyses and graphs were executed in SPSS v.23.

403

404   *Testing the model*

405   We test the model by evaluating its ability to predict the presence versus

406   absence of gibbons at a range of locations across the Indian subcontinent, the Indo-

407   China peninsula, Indonesia and the Malay Archipelago, New Guinea and mainland

408   China. For these purposes, we selected a number of locations of known altitude and

409   latitude, which had climatic data available in www.en.climatedata.org. The locations

410   and their climatic and geographical data are given in online *Dataset S2*. Since these

411   analyses predict where hylobatids can and cannot survive, they also allow us to make

412   inferences about the biogeographic distribution of the taxon. In most cases, we

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413　selected major cities for these purposes, since our question is whether or not gibbons

414　could live in the general area, not whether they live at a particular location. These data

415　also allow us to assess whether gibbons could now live in those provinces of China

416　where they were recorded historically by Turvey et al. (2015), and hence determine

417　whether anthropogenic factors or climate change might have been responsible for

418　their extinction. For these purposes we selected one site at random roughly in the

419　center of each of the provinces of modern China.

420

421　**Results**

422　*The model*

423　　　　Table 1 lists the best fit equations for feeding and moving that were generated

424　by the data, as well as the generic primate equations for enforced resting time and

425　social time. The causal relationships involved are summarized in Fig. 1. For reference,

426　alternative "next best" significant equations for feeding and moving time selected by

427　the backwards stepwise regression are given in the *Online Supplementary Material*

428　(Table S1). Both feeding and moving time increase as habitats become climatically

429　more stressful (low rainfall, low temperature, high altitude, high latitude). Although

430　group size has no effect on feeding time, it has a weak positive effect on moving time

431　even despite the extremely limited variation in hylobatid group size (Fig. 2). Since

432　moving time is the main limiting factor for great apes, and severely limits their

433　biogeographic distribution and group size (Lehmann et al. 2007b, 2008a,b), we ran a

434　separate regression model with just latitude and group size as predictors (Eqn. 2a). It

435　is of significance that, despite an order of magnitude difference in the range of group

436　sizes, the slope coefficient for group size in gibbons is very similar to that for the

437　chimpanzee time budget model (3.08 vs 2.59, respectively).

17

438   Across the sample of study populations, percentage of fruit in the diet is

439 determined mainly by ambient temperature (Table 2). In contrast, percentage of leaf

440 in diet increases with the level of environmental stress as reflected in rainfall, altitude

441 and temperature variance (Table 2). Although day journey length was not predicted by

442 any of the climatic variables or by fig tree density, territory size is predicted with a

443 very high r^2 by a complex of climatic variables (Table 2), suggesting that territory

444 sizes get larger as environmental stress increases.

445   The next step is to use the time budget equations to predict maximum

446 ecologically possible (i.e. limiting) group size, *Nmax*, for the sampled populations. To

447 do this, we interpolate the time available for social interaction (i.e that remaining after

448 removing the time predicted for feeding, moving and enforced rest at the site, plus the

449 constant for singing) into Eqn. (5). For this analysis, siamang predicted feeding time

450 is rescaled back to siamang body mass by reversing the transformation used to convert

451 their feeding time to gibbon-equivalents.

452   Fig. 3(a) plots maximum tolerable group size, N_{max}, against observed mean

453 population group size for all the gibbon and siamang populations in our sample. The

454 horizontal line demarcates a group size of 2.96 individuals, representing the minimum

455 group size for demographic viability (i.e. demographic stability defined by a

456 population growth rate of r=0: this requires two surviving offspring over an average

457 gibbon 17-year reproductive lifespan, and would equate to a group with two adults

458 plus 4/17 = 0.12 births per year, hence an average of ~0.96 dependent offspring aged

459 <8 years at any given time, taking 8 to be the age at dispersal and thus the range of

460 individuals who would be observed in a group). (This value assumes minimal

461 mortality in immature gibbons, which seems broadly to be the case; assuming 50%

462 mortality up to puberty, a figure typical of terrestrial baboons, would only increase

18

463   minimum viable group size to 2 adults + 1.5*0.96 living offspring = 3.44.) The

464   dashed diagonal line marks the line of equivalence (N_{max} is equal to observed mean

465   population size).

466   Three points may be noted. First, time budget models specify that group sizes

467   should not exceed the predicted maximum size (hence all data points in Fig. 3a should

468   lie above the dashed line). In the present case, 80.5% of the 41 populations have a

469   predicted maximum group size that is larger than their observed mean group size.

470   This is broadly encouraging as far as model fit is concerned, but suggests that

471   something else is needed to account for the exceptions. Second, no population has

472   N_{max}>15 individuals. This suggests that hylobatids are under rather greater ecological

473   constraint than many other Old and New World monkeys (and African great apes),

474   where limiting group sizes are typically >20. Third, notwithstanding this, it is clear

475   that many (but not all) populations could, on ecological grounds, live in much larger

476   groups than they actually do. Such groups would inevitably be multi-female.

477   One likely reason why some populations might have maximum predicted

478   group sizes below their observed group sizes is the length of the active day. Gibbons

479   are well known to retire early, often several hours before dusk (Raemaekers 1979;

480   Chivers 1984; Palombit 1997; Fei, Zhang, Yuan, Zhang & Fan 2017), a behaviour

481   that is quite unusual for most primates. As a result, the sampled activity budgets are

482   based on an active day that averages 9.3±0.89 hrs (range 8.1-10.6, N=9 sites), despite

483   the fact that daylength in tropical habitats is ~12 hrs. Progessively extending the

484   length of the active day in units of 5% suggests that an active day that is 15% (84

485   min) longer than the observed mean would be enough to lift all but the three lowest

486   siamang populations above the demographic viability threshold (Fig. 3b). (This would

487   increase maximum N_{max} for the other populations only to ~20.) Unfortunately, data on

19

the length of the active day are not available for any of the populations with N_{max}<2.96. However, an extra 84 min would only increase the mean value of day length to 10.7 hrs, virtually identical to the longest observed day length in our dataset (10.6 hrs in a siamang population: *Dataset S1*).

We ran a sensitivity analysis of the model using the alternative equations for feeding and moving time from Table S1. To do this, we altered one of the equations at a time in the original model from Table 1, keeping all the other equations as in the original model. We also ran a model combining the alternative moving time equation with the two most extreme alternative feeding time equations. Predicted maximum group size for the individual populations in the study site database determined using these alternative equations are highly correlated with those determined using the original model (Table S2; Fig. S1). The alternative moving time equation increases the largest maximum group size from ~15 to ~25, but does not substantively change any of the results. This suggests that the Table 1 model is quite robust. Similarly, decreasing the slope coefficient for group size in the moving time equation to 2.59 (the value for the chimpanzee model) increases the largest N_{max} to ~18, while increasing it to 5.0 decreases N_{max} to ~9, but does not alter the broad pattern or the fact that most mean population group sizes are comfortably below N_{max}.

To provide some indication as to how environmental parameters influence hylobatid biogeography, Fig. 4 plots predicted maximum group size as a function of (a) latitude and altitude and (b) rainfall and temperature. To produce these graphs, we have used the equations given in Table 2 to reduce all the climatic variables in the model to the two indices of interest in each case. This can be expected to increase error variance, with a consequential tendency for reversion to the mean, but the results provide us with an indication of how maximum group size is likely to vary across

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513   habitats. The results suggest that hylobatids do best (i.e. are able to sustain larger

514   groups) in cooler, high altitude, high rainfall habitats at low latitudes (i.e. near the

515   equator). When rainfall is less than ~1500mm per annum, or at latitudes above ~20°

516   (in effect, outside the Tropics), hylobatids are unable to maintain minimally viable

517   groups (N≈3 individuals) unless they are living at altitudes above ~1000m (i.e. under

518   cooler conditions) (see also Turvey et al. 2015). This last prediction is confirmed by

519   the fact that the northern Chinese populations are at significantly higher altitudes than

520   all the other hylobatid populations (mean altitudes: 1750.0±783.8SD m vs

521   401.2±409.5SD m asl, N=14 and 50 respectively; $F_{1,62}$=76.2, p<0.0001).

522

Testing the model

524   To explore the model's ability to predict hylobatid biogeography, we used a

525   sample of locations within the current hylobatid biogeographic range in southeast

526   Asia and a sample of locations on the Indian subcontinent, mainland China and the

527   islands of the Malay archipelago outside the current hylobatid range. In addition, we

528   have included one representative site in each of the Chinese provinces where gibbons

529   were recorded as occurring prior to 1800 AD (see Turvey et al. 2015). Note that, for

530   the latter cases, the climate data are current, not historical, values. In each case, we

531   predict maximum group size using current local climate

532   Fig. 5 plots the mean and range of maximum group size predicted by the

533   model for each of these sites grouped by geographical location. Predicted maximum

534   group sizes average 6.6±4.0SD for 18 sites within the current hylobatid range. In

535   contrast, sites in northeast India and Bangladesh to the west of the Brahmaputra

536   (which forms the northwest boundary of the current gibbon range) average 0.3±0.5SD

537   (N=7 sites), as is the case for locations further west in northwest India (mean

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538   1.6±2.2SD, N=7 sites) (see also Fig. S2). This suggests that, even in the absence of

539   the physical barrier of the Brahmaputra river system, the taxon's ability to expand its

540   range westwards has been limited by substantial tracts of land it would have been

541   unable to cross. Had gibbons been able to bypass these two barriers, they would have

542   found very congenial habitats in the southern parts of the subcontinent (e.g. the Nilgiri

543   Hills and adjacent ranges in the Deccan, and in Sri Lanka), where they would have

544   been able to support groups as large as 8.7±5.1SD (N=7 sites). The island habitats

545   across the Wallace Line (specifically, the Celebes and Papua New Guinea) provide

546   equally gibbon-friendly habitats (mean group size 10.3±1.1SD, N=2 sites), but it

547   seems that, as with most other primates (Brandon-Jones 1998), the gibbons were

548   never able to traverse this major sea barrier. Increasing the length of the available day

549   by 15% (as in Fig. 3b) does not change the overall pattern, or make regions such as

550   northern India or central China any more habitable for gibbons.

551   More puzzling, however, is the fact that central and east Java (including

552   nearby islands like Bali) provide very suitable habitats (mean N_{max} 10.4±1.2SD, N=7

553   sites) yet have no gibbon populations. Climatically at least, these habitats seem to be

554   at least as suitable as those in west Java where gibbons do occur (mean N_{max}

555   10.2±1.9SD, N=7 sites). In contrast, gibbons would now find it impossible to survive

556   in either the central and southeastern provinces of China where they were recorded as

557   living prior to 1800 AD (mean 0.0±0.0SD, N=14 sites) or, perhaps less surprisingly,

558   the provinces of northern China where they did not occur historically (mean

559   1.3±3.6SD, N=7). This contrasts with locations in southwest China (Yunnan province

560   and Hainan Island) that currently do support gibbon populations, for which the model

561   predicts viable group sizes (mean N_{max} = 4.3±1.7SD, N=3 sites).

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563   *What limits hylobatid distribution?*

564   An important feature of time budget models is that they allow us to determine

565   which aspect of the animals' biology is the principal constraint on their ability to

566   occupy habitats, and hence what actually limits their biogeographic distribution. In

567   turn, this tells us something about the taxon's risk of extinction under climate change.

568   To explore this, we plotted the predicted time required for feeding, moving and

569   enforced rest for the *Dataset S2* locations (Fig. 6). Neither feeding nor moving differ

570   consistently between regions where hylobatids are present vs not present, suggesting

571   that neither of these is a major constraint (except, in the limit, the effect of group size

572   on moving time at much larger group sizes); in contrast, resting is significantly higher

573   in regions where they are absent and seems to account for the fact that they do not

574   have sufficient time for social interaction to maintain groups of any significant size.

575   It thus seems that the main problem lies in the extent to which climatic

576   variables oblige them to rest. This is confirmed by an analysis of observed versus

577   predicted time budget allocations in the sample study sites. Fig. 7a plots the ratio of

578   observed feeding and resting time divided by the values predicted by the model

579   equations in Table 1 for those populations in the field study dataset (*Dataset S1*) for

580   whom predicted maximum ecologically tolerable group size is either less than or

581   greater than 5 (the upper limit for observed mean population group size). For these

582   purposes, predicted rest is enforced rest time, while observed rest is total rest time.

583   Feeding time is close to that predicted in both cases, as should be the case. In contrast,

584   observed rest time is significantly less than the minimum predicted ('enforced' rest) in

585   populations where *Nmax* is less than 5, whereas populations where *Nmax* exceeds 5

586   typically have surplus rest time capacity. The strength of this effect is clear from Fig.

587   7b which plots the difference between observed and predicted rest against the

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588 difference between observed group size versus predicted *Nmax* for the populations.

589 The correlation is highly significant (r=-0.847, p<0.0001).

590

591 **Discussion**

592 The time budget model for hylobatids that we develop here is broadly similar

593 in form to the models developed for 10 other primate genera. It works well for

594 contemporary populations, predicting presence in most cases where gibbons occur and

595 absence where they do not (Fig. 5). In other cases, it is clear that physical barriers

596 (e.g. habitat conditions in northeast India or the Wallace Line to the southeast) have

597 prevented gibbons expanding into habitats where they could do well. It is important to

598 remember that time budget models predict the largest groups that a taxon can maintain

599 at a given site (in effect, the carrying capacity), not necessarily the actual size of

600 group. They simply set the upper limit defined by local environmental conditions.

601 Since, for primates in general, fertility is almost always adversely affected by

602 increasing group size (van Schaik 1982; Dunbar 2018a; Dunbar, MacCarron & Shultz

603 2018a), animals will generally try to minimize group size in any given location to the

604 extent that this is compatible with the constraints imposed by the local predation risk

605 (Dunbar et al. 2009; Dunbar et al. 2018a).

606 The fact that hylobatids commonly live in groups that are smaller than those

607 allowed by the local ecology has two important implications. First, it implies that,

608 while the model certainly tells us that gibbons are under greater ecological constraint

609 than most Old World monkeys and African great apes, the level of ecological stress is

610 not sufficiently high to force them to live in groups as small as those they actually live

611 in. While it is true that some populations have time budgets with little or no spare

612 capacity, many do not (Fig. 7). This conclusion is supported by the fact that, unlike

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613 most other monkeys and apes, hylobatids commonly go to their night rest mid-

614 afternoon (Raemaekers 1979; Srikosamatara 1984; Palombit 1999; Fei et al. 2017),

615 implying that they are not under significant time pressure. Second, it suggests that

616 predation risk must be low for hylobatids; group size evidently does not provide

617 gibbons with a significant anti-predator advantage, so they can afford to minimize

618 group size in order to maximize fertility (see Dunbar et al. 2009; Bettridge, Lehmann

619 & Dunbar 2010; Dunbar, MacCarron & Robertson 2018b). Indeed, in comparison to

620 almost all other primates, cases of predation are conspicuous by their absence from

621 the gibbon literature (Reichard 2003).

622   This calls into question the claim that gibbons are monogamous because

623 females are forced by their ecology to forage on their own rather than in groups (van

624 Schaik & van Hooff 1983; Rutberg 1985; Komers & Brothertpn 1997; Brotherton &

625 Komers 2003; Lukas & Clutton-Brock 2013). Contrary to this claim, it seems that

626 most populations could in fact maintain larger groups (range 5-12 animals), especially

627 in the more southerly parts of the Malay archipelago. Groups of this size could easily

628 include 2-4 reproductive females (across all primates, reproductive females form a

629 very consistent 30-35% of the group: Dunbar et al. 2018a,b). Even if females lived

630 alone, it would be possible to have more of them within a gibbon group's current

631 territory, and males would have access to more than one reproductive female if they

632 pursued a roving male strategy (much as orang utan males do) (see also van Schaik &

633 Dunbar 1990; Bartlett 2009). This implies, as suggested by van Schaik & Dunbar

634 (1990) and Opie, Atkinson, Dunbar & Shultz (2014), that monogamy must have

635 evolved in response to factors other than ecological constraints.

636   One of the main benefits of the time budget model approach is that it provides

637 insights into the behavioural and physiological constraints that limit a taxon's ability

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638   to successfully occupy different kinds of habitats. As with previous primate time

639   budget models, feeding time is strongly influenced by predictors of habitat quality: in

640   the hylobatid case, the main determinants of feeding time (rainfall, altitude and

641   latitude, with the latter two both being core predictors of temperature) are all variables

642   that influence both tree species composition and the nutritional quality of vegetation

643   (see also Marshall & Leighton 2006). As food quality falls, animals will need to spend

644   longer feeding to meet their nutritional requirements.

645   In contrast, time spent moving remains largely independent of the climatic

646   variables, being affected only by absolute latitude and group size. Models for African

647   genera have also noted that moving time tends to be independent of environmental

648   variables and is often close to being a constant (Dunbar et al. 2009), mainly because,

649   in order to save time for other more pressing activities, animals respond by increasing

650   travel speed rather than increasing travel time when they need to travel further

651   (baboons: Dunbar 1992a). There is some suggestion that gibbons also do this: travel

652   speed increases as climatic conditions deteriorate (i.e. when temperatures are high and

653   latitude is low, although the effect is weak and not significant (Fig. S3). However,

654   since there is inevitably a limit on the speed at which animals can travel, at some point

655   animals simply have to devote more time to moving. The substantive issue for day

656   journey length (and hence moving time) is, at least for non-folivores (Snaith &

657   Chapman 2007), group size: the area that has to be covered to allow every animal in

658   the group to meet its nutritional requirements is inevitably a linear function of the

659   number of animals in the group, forcing the group to travel further each day (Dunbar

660   et al. 2009). The great apes are especially susceptible to the effects of group size on

661   moving time, and it is this that is mainly responsible for chimpanzees' fission-fusion

662   form of sociality (Lehmann et al. 2007b, 2008a,b). It is significant that, despite the

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663   very limited variance in the size of their groups, the group size slope parameter for the

664   gibbons is similar to that for chimpanzees. For groups as small as those found in

665   gibbons, the impact of this effect will, of course, be modest; but it does mean that

666   when groups are larger than ~10 the impact will rapidly becomes prohibitive (as is the

667   case for African great apes: Lehmann et al. 2007b [Fig. 7]).

668   However, it seems to be enforced resting time that is the main constraint for

669   hylobatids (Figs. 6 and 7). Since this is mainly affected by ambient temperature, it

670   might explain why social groups get larger in northern populations irrespective of

671   species (Fig. 8). Ultimately, however, the northern extension of hylobatids is

672   constrained by the fact that feeding and moving time are also positive functions of

673   latitude (and hence, in effect, declining temperatures). A comparison of the slope

674   parameters for latitude in Tables 1 and 2 indicates that the additive effects of the

675   latitude coefficients for feeding and moving time increase at a combined rate that is

676   ~4 times the savings in terms of the slope for enforced rest ($0.363 + 0.462 = 0.825$ vs

677    $1.33*0.158 = 0.210$), so that time budgets become progressively squeezed as animals

678   occupy increasingly high latitude habitats. At the latitude of Mt Wuliang in Yunnan

679   Province, latitude has added a net 15 percentage points to the time budget.

680   There is a widespread perception that *Nomascus* populations live in larger

681   groups than all other hylobatids, hence the fact that they have polygamous groups. Up

682   to a point, this is true; however, this is not true for all *Nomascus* populations (Fig. 8).

683   Moreover, *Hoolock* populations also have large groups at high latitudes without these

684   being polygamous. Mean population group size is in fact a cubic function of absolute

685   latitude (Fig. 8). Group size does increase steadily up to around 20°N (roughly the

686   latitude of Chaing Mai in northern Thailand), but then it levels off only to drop

687   precipitately after latitude 25°N (roughly the latitude of Mt Wuliang in southwestern

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China). This decline at very high latitudes is suggestive of populations living at the limits of their range, and indeed mean group size at these highest latitude populations is only just above the minimum for demographic viability.

It is evident from Fig. 3(b) that siamang incur an additional cost due to the fact that their larger body size imposes a higher feeding time demand. A convincing explanation for their larger body size remains elusive, especially given the fact that their range overlaps that of other gibbons and the two taxa are often sympatric. As a result, however, they are often under significant time budget pressure. There is some evidence to suggest that they make time budgeting adjustments that might be sufficient to reduce this pressure: for the limited sample available, it seems that siamang devote less time each day to both singing and social interaction than gibbons do, and they have a longer active day (Table 3). For this limited sample, the net gain for siamang is equivalent to increasing the length of the active day by ~17% (the combined effect of a longer active day adjusted for less time spent socialising and singing) while only having to increase the actual length of the active day by ~10%. That is sufficient to just lift the lower siamang populations in Fig. 3(b) into the minimum viability zone. Nonetheless, siamang populations are clearly under more ecological pressure than is typical of the other gibbons (Fig. 3).

Gibbons share with the great apes a dietary physiology specialized for frugivory; indeed, their feeding time equation is very similar to that of the gorilla, though their size precludes their being able to survive on low quality folivorous fallback foods in the way gorillas can. Although the larger-bodied siamang exhibit some capacity in this direction (Raemaekers 1979; Palombit 1995), their ability to resort to a heavily folivorous diet is likely to be considerably less than the gorilla's simply because of their smaller body size. Mast years aside, fruits invariably have a

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713   more patchy distribution than leaf, and the travel demands imposed by this may be

714   part of the problem that affects gibbon time budgets, and hence limits group sizes.

715   Gibbon ecology appears to be quite tightly defined: their geographical range is

716   surprisingly limited. They are currently confined to the Indochina peninsula (bridging

717   out into southwest China) and the associated islands of the Malay archipelago

718   (Sumatra, Java and Borneo). They appear to have been prevented from extending

719   southeastwards by the Wallace Line (despite the fact that the islands beyond would be

720   perfectly habitable for them: Fig. 5), while their capacity to encroach into the Indian

721   subcontinent has been limited by the climate west of the Brahmaputra (Fig. 6).

722   Nonetheless, within their core distribution they seem to be under much less ecological

723   constraint than we might have anticipated given the female-dispersion explanation for

724   their adoption of a monogamous mating/social system.

725   Historically, gibbons are known to have occurred at much higher latitudes in

726   China than they do now. The model suggests that current climate in these areas makes

727   these habitats completely unsuitable for gibbons. Although the historical extinction of

728   these northern populations has been attributed to anthropogenic factors (Fan 2017), in

729   fact these habitats would not support gibbons now even if there were no humans

730   living there. There are only three plausible explanations for the disappearance of these

731   populations. One is that the populations in question had evolved novel adaptations to

732   these environments, such that the slope parameters on their feeding and/or moving

733   equations were radically different from those for gibbons from further south, but that

734   humans caused their extermination nonetheless. However, to suggest that these

735   northern gibbons were doing something completely different to all other New and Old

736   World anthropoid primates as well as other gibbons implies that they could not have

737   been conventional primates, and that should be, at best, an explanation of last resort.

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738 A second possibility is that humans had released animals in these areas in order to

739 have them as exotics (something that humans seem to be especially prone to do), even

740 though the habitats were not really suitable for them; as a result, the populations went

741 extinct once they were no longer being replenished (a situation not too dissimilar to

742 that of *Macaca sylvanus* on Gibraltar). This is a possibility, but the fact that gibbons

743 seem to be unable to survive there at all would imply the need for constant

744 replenishment, and this seems unlikely over such a wide area and such a long time

745 period. The third option is that the climate has changed in the past 300 years in ways

746 that would have led to the inevitable extinction of these populations (with or without

747 the assistance of humans).

748 A potential culprit in the latter respect is the Little Ice Age of 1645-1715,

749 which resulted in average world temperatures falling by ~1°C. In China, this resulted

750 in a significant increase in climatic seasonality. The cultivation of Mediterranean-type

751 citrus crops was abandoned in Jiangxi Province (one of the provinces that historically

752 harboured gibbons: Turvey et al. 2015) after the 1750s, despite their having been

753 cultivated there for many centuries (Reiter 2000). Guangdong (another of the

754 provinces where gibbons have been documented) experienced a particularly cold, dry

755 spell, accompanied by an unusually high frequency of typhoon strikes, after the Little

756 Ice Age (Liu, Shen & Louie 2001). Ameca y Juárez, Mace, Cowlishaw, Cornforth &

757 Pettorelli (2013) have shown, for mammals as a whole and primates in particular, that

758 high frequencies of cyclones and droughts correlate with elevated extinction risk

759 (indexed by the number of taxa classified as "threatened"), with the southeast corner

760 of China being especially prone to this effect. Turvey et al. (2015) noted that there

761 was increasing fragmentation of these populations from 1700 onwards (with a marked

762 upturn in fragmentation from around 1900), with a 50-100 year lag to last reported

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763   occurrence. This is indicative of populations struggling to survive.

764   A comparison of the distribution of climatic variables in southwest China

765   (where gibbons live now) and central and southeast China (where they lived

766   historically) suggests that the only climate variables in which these two regions differ

767   significantly are $TEMP_{moSD}$ (standard deviation of mean monthly temperature across

768   the year, an index of seasonality) and annual rainfall (Table 4). (We do not consider

769   latitude or altitude since these cannot have changed historically.) This may well

770   reflect shifting patterns in the latitudinal distribution of the monsoon circulation,

771   allowing an extension of a more seasonal monsoon climate further into mainland

772   China after the Little Ice Age. A 65% increase in rainfall and a dramatic reduction in

773   seasonality would be required to allow gibbons to survive in the southeastern

774   provinces as well as they currently do in the southwest.

775   The bottom line seems to be that even if anthropogenic factors have been

776   important in the final demise of these populations (as both Turvey et al. 2015 and Fan

777   2017 have suggested), these populations' sensitivity to anthropogenic factors is likely

778   to have been exacerbated by environmental effects on the animals' increasing inability

779   to cope as climate changed. The results in Fig. 4, for example, imply that, historically,

780   declining populations at high latitudes are likely to have become locked into

781   mountain-top retreats as climate deteriorated, leading to small, isolated pockets that

782   are inevitably more vulnerable to extinction in the face of environmental shocks

783   (Dunbar 1998; Cowlishaw & Dunbar 2000).

784   One final puzzle is the fact that gibbons are not found east of the Dieng

785   Mountains in central Java. Kappeler (1984a,b) attributed this to the kinds of forests

786   found in the eastern half of the island, which tend to be more deciduous. Fossil

787   gibbons have been recorded at Trinil, central Java (in the Lower/Middle Pleistocene

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788 deposits that produced the hominins: Ingicco, Vos & Hoffman 2014) and at the

789 Ngandong and Gunung Dawung hominin sites in east Java (Storm & Vos 2006) (see

790 also Jablonski & Chapin 2009), so this area clearly did once support gibbon

791 populations. The fact that west Java was one of the refugia for gibbons during the dry

792 phases of the late Pleistocene Ice Ages (Brandon-Jones 1998) makes this all the more

793 puzzling. It may be that anthropogenic factors resulted in their demise, although these

794 are unlikely to have kicked in until historical times.

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1064   **Legends to Figures**

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1066   Fig. 1. Flow chart for the gibbon time budget model, summarising the causal

1067   relationships between climatic and behavioural variables based on the

1068   equations given in Table 1. Solid lines: positive effects; dashed lines: negative

1069   effects.

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1071   Fig. 2. Time spent moving plotted against mean population group size. Dashed line is

1072   least squares regression line. Filled symbols: gibbons; unfilled symbols:

1073   siamang. Source: online *Dataset S1*.

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1075   Fig. 3. (a) Maximum ecologically tolerable group size predicted by the time budget

1076   model, plotted against observed mean group size for different gibbon (filled

1077   symbols) and siamang (unfilled symbols) populations. (b) Predicted maximum

1078   group size allowing for a 15% increase in the length of the active day. The solid

1079   line demarcates the minimum group size for demographic viability (2 adults plus

1080   0.96 immatures: see text for details); dashed line in (a) is the line of equilibrium

1081   (*Nmax* = observed mean).

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1083   Fig. 4. Maximum group size predicted by the time budget model for different

1084   combinations of (a) latitude and altitude and (b) annual rainfall and mean

1085   temperature.

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1087   Fig. 5. Mean (±95% CI) maximum group size (*Nmax*) predicted by the time budget

1088   model for habitats within different geographic regions. Filled circles: regions

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1089   where gibbons currently live; unfilled circles: regions where gibbons do not

1090   currently live; grey circle: gibbons historically present before 1800 AD. Dashed

1091   line: minimum group size for demographic viability (2.96 individuals: see text).

1092   Present: 18 sites within the current biogeographic distribution of gibbons within

1093   Indo-China, Sumatra and Borneo. W Java: 3 sites in west Java where gibbons

1094   occur; E Java: 8 sites in Java east of the Dieng Mountains where gibbons do not

1095   currently occur; E Indonesia: 2 sites on Celebes and western Papua New Guinea,

1096   east of the Wallace Line where gibbons do not occur; NW and NE India: 8 and 7

1097   sites, respectively, in northwest and northeast India (including Bangladesh) where

1098   gibbons do not occur; S India: 7 sites in southern India plus Sri Lanka; SW

1099   China: 3 sites within current gibbon range in Yunnan and Hainan. SE China: one

1100   site from each province in southern China where gibbons were recorded as being

1101   present in historical documents before 1800 AD (N=14 site). N China: one site

1102   from each province in northern China (including Tibet) where gibbons were not

1103   recorded historically (N=7 sites). Locations and climate data from online *Dataset*

1104   *S2*.

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1106   Fig. 6. Mean (±95% CI) (a) feeding time, (b) moving time and (c) enforced rest time

1107   predicted by the time budget model for the regional sites shown in Fig. 5. Moving

1108   time does not include an adjustment for group size. Regions and symbols as for

1109   Fig. 5.

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1111   Fig. 7. (a) Mean (±95% CI) ratio of observed to predicted time spent feeding and

1112   resting for populations for which predicted *Nmax* <5 (unfilled symbols) or *Nmax*

1113   >5 (filled symbols). Predicted rest time is enforced rest time. (b) Difference in

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1114   rest time (observed minus predicted by model) for individual hylobatid

1115   populations plotted against difference in group size (observed minus *Nmax*).

1116   Source: Dataset S1.

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1118   Fig. 8. Mean observed group size for individual populations plotted against absolute

1119   latitude for the population. The best fit equation (dashed line) is Group size =

1120 $3.71 - 0.014*Lat + 0.012*Lat^2 - 0.0004*Lat^3$ (r^2=0.159, $F_{3,50}$=3.15, p=0.033).

1121   Source: Dataset S1.

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Table 1. Regression equations for the hylobatid model.

Variable	Equation	r^2	F	df	p
(1) Feed time (%)	$41.634 - 0.003*RAIN - 0.009*ALT + 0.363* LAT_{abs}$ $^{\Omega}$	0.501	5.03	3,15	0.013
(2) Moving time (%)	$17.316 + 0.462*LAT_{abs}$	0.166	3.59	1,18	0.074
(2a) Moving time (%)	$5.297 + 0.290*LAT_{abs} + 3.080*N$	0.220	1.98	2,14	0.175
(3) $Rest_{ENF}$ time (%)¶	$-23.24 + 1.33*TEMP + 0.259*LEAF + 6.12*T_{moSD}$	from Korstjens et al. (2010)			
(4) Social(%)†	$2.968 + 0.109N$	0.440	29.90	1,38	<0.0001
(5) Time budget	$100 = FEED + MOVE + REST_{ENF} + SING + SOCIAL$				
(5a)	$100 = FEED + (5.297 + 0.290*LATabs + 3.080*N) + (REST_{ENF} + REST_{FREE}) + SING + (2.968 + 0.109*N)$				
(5b) Nmax§	$= ((100 - FEED - (5.297 + 0.290*LATabs) - REST - 4.96‡) + 2.968)/(0.109 + 3.080)$ by substitution				

Ω Absolute latitude

¶ Rest time consists of two components: enforced rest ($REST_{ENF}$, imposed on the animal by climatic conditions and its dietary strategy) and free

  rest ($REST_{FREE}$) that represents uncommitted time that can be allocated to any other activity when required (see Korstjens et al. 2010).

† Recalculated from Lehmann et al (2007a)

§ Maximum ecologically tolerable group size (the maximum group size that will allow the time budget to be balanced)

‡ 4.96 = average percentage of day devoted to singing by gibbons

Table 2. Supplementary equations

Variable	Equation	r^2	F	df	p
TmoSD **	$10.440 + 0.093*LAT_{Abs} - 0.001*ALT - 0.821*TEMP$				
	$+ 0.588*TEMP_{min}$	0.597	17.75	4,48	<0.0001
Mean temperature (°C)	$27.651 - 0.003*ALT - 0.158*LAT_{Abs}$	0.672	60.33	2,59	0.0001
Altitude (m)	$4145.354 - 147.236*TEMP + 0.023*RAIN$	0.575	37.14	2,55	<0.0001
$Temp_{var}$ †	10.196				
Absolute Latitude	$48.732 - 0.004*RAIN - 1.142*TEMP$	0.580	35.23	2,51	<0.0001
Rain	$3279.523 - 53.609*LAT_{Abs} + 0.025*ALT$	0.286	10.60	2,52	<0.0001
$Temp_{min}$ (°C)	$-5.776 + 1.033*TEMP$	0.922	623.63	1,53	<0.0001
Territory size (ha)	$2498.88 + 0.077*ALT - 477.830*TEMP_{min} + 401.718*TEMP_{max}$				
	$- 477.03*TEMP_{var} - 32.260*_{TEMPmoSD} - 0.032*RAIN$	0.894	19.60	6,14	<0.0001
Leaf in diet (%)	$99.430 - 1.000*LAT_{abs} + 0.009*ALT + 7.012* TEMP_{var} +$				
	$7.879* TEMP_{min} - 9.779* TEMP_{max}$	0.611	5.04	5,16	0.006
Fruit in diet (%)	$-51.27 + 10.65*TEMP - 7.71*TEMP_{min}$	0.382	6.19	2,20	0.008
Active day (hr) $^{\Omega}$	$11.273 - 0.001*RAIN$	0.905	19.15	1,2	0.048

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** Standard deviation of monthly mean temperatures across the year

† Difference between mean monthly maximum and mean monthly minimum temperature

Ω Gibbons only

3

Table 3. Time budget differences between siamang and gibbons

Variable	Siamang		Gibbons*	
	Mean±SD	N§	Mean±SD	N§
% fruit in diet	40.7±16.1	6	57.9±18.7	22
% leaf in diet	34.6±11.6	4	33.7±30.4	22
Feed (%)	49.6±6.8	5	34.4±8.5	19
Move (%)	18.0±5.8	5	21.4±7.9	16
Rest (%)	34.4±10.1	5	34.7±9.8	18
Sing (%)	2.0	1	4.96±3.1	10
Social time (%)	3.0	1	5.87±3.5	12
Active day (hr)	10.1±0.3	2	9.06±0.9	7
Net difference in available time†	16.8%			

* sites used in time budget analyses only

§ Number of studies

† Equivalent change in gross time budget achieved by the siamang compared to gibbons
 (~3% by reducing singing, ~3% by reducing social time and ~11% by increasing length
 of active day)

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Table 4. Comparison of climatic variables between southwest China (where gibbons currently exist) and southeast China (where they were present historically, but no longer are).

Provinces of China	Gibbons present	TEMP	$TEMP_{min}$	$TEMP_{max}$	$TEMP_{var}$	$TEMP_{moSD}$	RAIN
Southwest	current	20.6	16.5	24.6	8.1	3.8	1973
Southeast	historical	16.6	12.5	20.9	9.7	8.1	1163
	$F_{1,15}$	2.50	1.88	2.61	0.21	19.97	9.22
	p	0.135	0.191	0.127	0.657	0.0005	0.008

5

Environment and Time as Constraints on the Biogeographical Distribution of Gibbons

R.I.M. Dunbar, Susan M. Cheyne, Daoying Lan, Amanda Korstjens, Julia Lehmann & G. Cowlishaw

Online Supplementary Material

Alternative model equations

Table S1 provides alternative multivariate equations for feeding and moving time generated by the backwards regression model. Most of the slope coefficients are similar across the alternative equations. However, note that the magnitude of the effect of group size on moving time for Equation (2c) is considerably less. Nonetheless, the values for both Equations (2a) and (2c) straddle the observed value for the chimpanzee model.

Table S2 gives the correlation values for maximum ecologically tolerable group size predicted by the original model based on Table 1 in the main text and the various alternative feeding and moving time equations given in Table S1. All alternative statistical models yield predicted maximum group sizes that correlate significantly with those predicted by the original model based on the equations of Table 1.

Fig. S1 plots predicted maximum group size for individual study sites in the main dataset (online *Dataset S1*) against the equivalent value predicted by substituting the various alternative feeding and moving time equations. Alternative feeding time equations do not have a significant effect on predicted maximum group size. Note that the alternative moving time equation predicts slightly larger group sizes than those predicted by the original model of Table 1 because the effect of group size on moving time in Equation 2c is lower (1.72 vs 3.08), yielding slightly larger group sizes. Nonetheless, the difference in the size of the largest groups predicted is modest (25 vs 15), the outcomes do not differ significantly (Table S2). Hence, using these alternative equations does not change any of the main conclusions.

Table S1. Alternative regression equations for feeding and moving time for the hylobatid model.

Variable	Equation	r^2	F	df	p
(1) Feed time (%)	$41.634 + 0.363* LAT_{abs} - 0.009*ALT - 0.003*RAIN$	0.501	5.03	3,15	0.013
(1a)	$53.955 + 0.243*LAT - 0.010*ALT - 0.472*TEMP + 0.243*TEMPVAR - 0.004*RAIN$				
		0.340	2.86	5,13	0.059
(1b)	$55.879 + 0.279*LATtabs - 0.010*ALT -0.485*TEMP -.004*RAIN$				
		0.520	3.80	4,14	0.027
(1c)	$33.230 + 0.469*LATabs - 0.009*ALT$	0.461	6.85	2,16	0.007
(2) Moving time (%)	$17.316 + 0.462*LAT_{abs}$	0.273	5.25	1,14	0.038
(2a)	$5.297 + 0.290*LAT_{abs} + 3.080*N$	0.220	1.98	2,14	0.175
(2b)	$29.196 + 0.306*LAT - 0.365*TEMPmax$	0.294	2.71	2,13	0.104
(2c)	$19.409 + 0.303*LATabs - 0.275*TEMPmax + 1.718*N$	0.312	1.81	3,12	0.199

Table S2. Correlations between maximum group size predicted by the original model of Table 1 and a model using the different equations for feeding and moving time given in Table S1, for the actual hylobatid study sites (as listed in online *Dataset S1*). In each case, one equation is substituted, with all the remaining equations as in the original model of Table 1.

Pearson Correlations

		Feed Eqn 1a	Feed Eqn 1b	Feed Eqn 1c	Move Eqn 2c	Feed Eqn 1a with Move Eqn 2c	Feed Eqn 1c with Move Eqn 2c
Original model	Correlation	0.994	0.995	0.988	0.996	0.987	0.986
	p (2-tailed)	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001
	N	51	51	51	51	51	51
Feed Eqn 1a	Correlation		0.999	0.974	0.996	0.997	0.978
	p (2-tailed)		<0.0001	<0.0001	<0.0001	<0.0001	<0.0001
	N		51	51	51	51	51
Feed Eqn 1b	Correlation			0.975	0.997	0.997	0.980
	p (2-tailed)			<0.0001	<0.0001	<0.0001	<0.0001
	N			51	51	51	51
Feed Eqn 1c	Correlation				0.984	0.966	0.997
	p (2-tailed)				<0.0001	<0.0001	<0.0001
	N				51	51	57

Move Eqn 2c	Correlation					0.995	0.989
	p (2-tailed)					<0.0001	<0.0001
	N					51	51
Feed Eqn 1a with	Correlation						0.977
Move Eqn 2c	p (2-tailed)						<0.0001
	N						51

Figure S1

Maximum group size predicted by the original model using the Table 1 equations plotted against maximum group size for all populations in the sample of studies (*Dataset S1*), using the alternative Feed and Move equations given in Table S1. Except for the last pair of graphs, each alternative equation is substituted individually, and all other equations are held constant as in the original model of Table 1. Filled symbols: gibbon populations; unfilled symbols: siamang populations. Dashed line: line of equivalence (the predicted maximum group size is the same for both models).
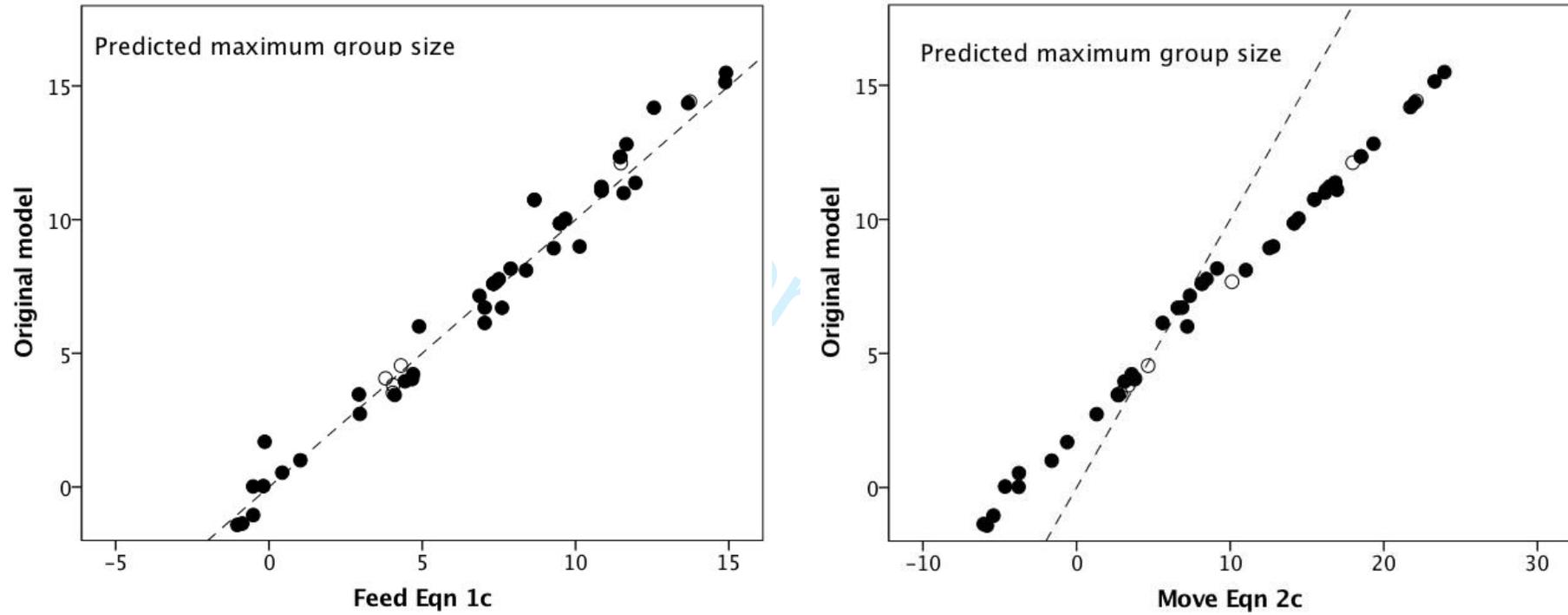
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Fig. S1 (contd)

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Fig. S1 (contd)
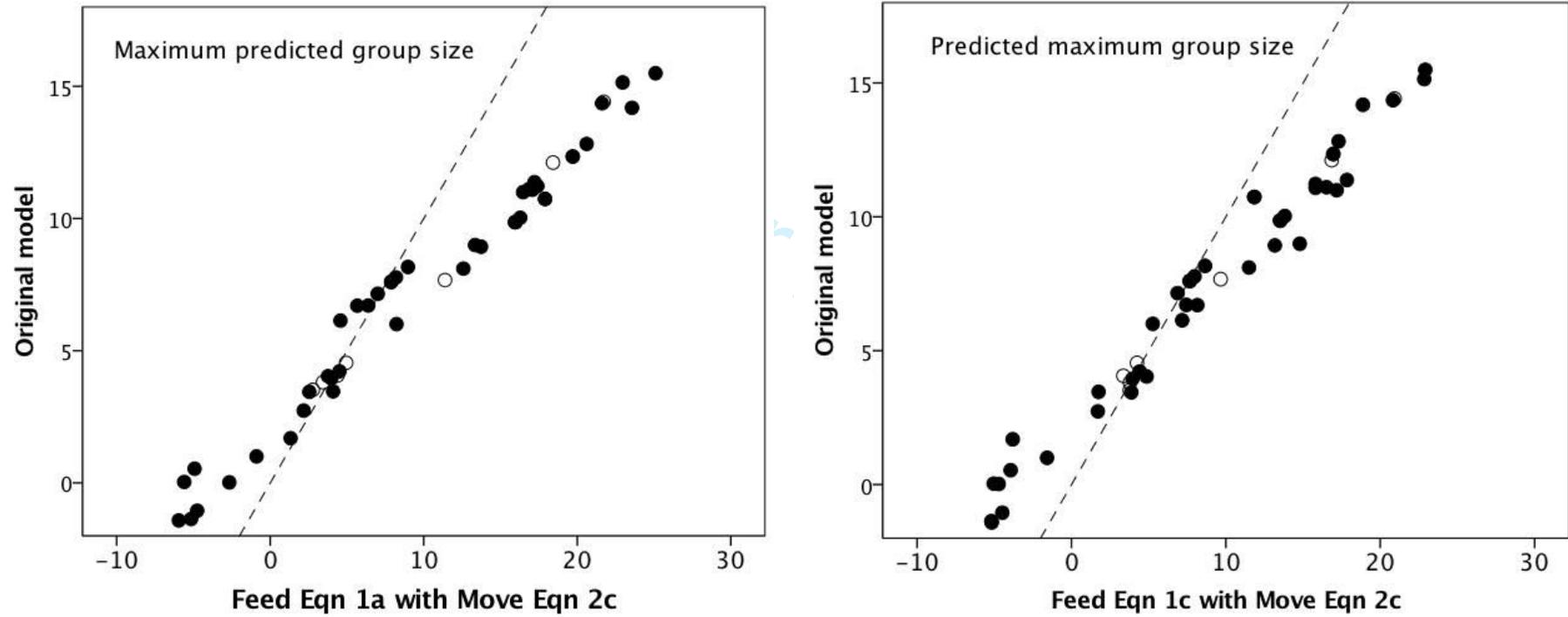
Model predictions

Fig. S2 plots the mean (±95% CI) predicted maximum group size (*Nmax*) for sites in countries where hylobatids are known to live and those in adjacet geographical regions (which should be accessible to them) where they do not. These values are calculated from the climate data in *Dataset S2* for those regions listed as within their current geographic range, plus SE China (essentially the provinces where gibbons occurred prior to 1800), NE India (the area north and west of the Brahmaputra in Bangladesh and northeastern India), and central and eastern Java. Indonesia east of the Wallace Line is excluded.

The difference is highly significant (means 2.92 vs 6.8, N=31 and 23, respectively; $F_{1,52}$=10.71, p=0.002). Note that, in areas where gibbons do not live, half the *Nmax* values are below the group size required for demographic viability (2.96 individuals), and most are below the observed mean size of 4.01 individuals. In contrast, all values for locations where gibbons live are above the observed mean group size.

Figure S2

Mean (±95 CI) predicted maximum group size (*Nmax*) for sites in countries where hylobatids are currently found, compared to that for geographically adjacent areas where they do not now occur (see main text for details). The solid line is the calculated minimum group size for demographic viability (2.96 individuals of all ages: see main text); the dashed line is the observed mean group size across all gibbon populations. Source: *Dataset S2*
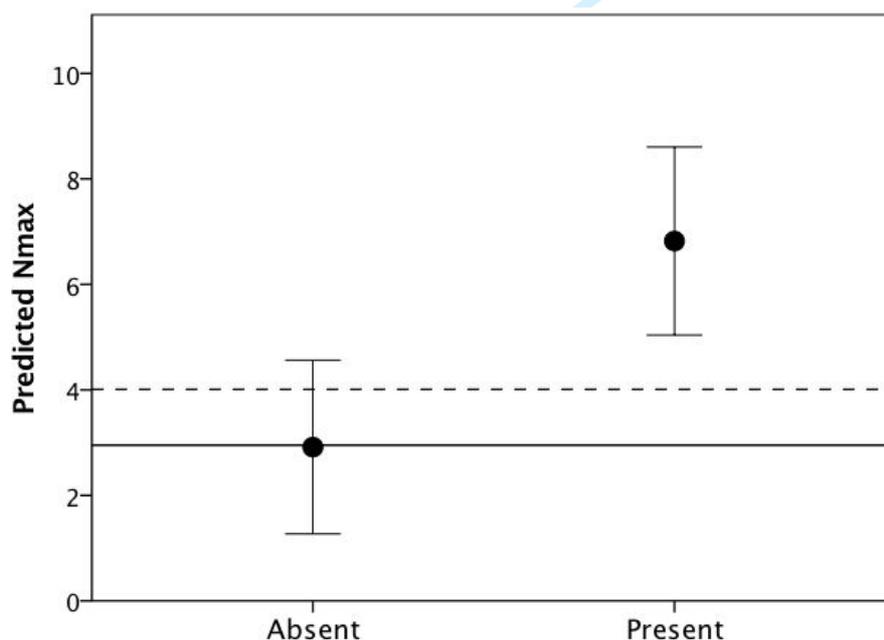
Speed of travel

We calculate average speed of travel across the day as day journey length (in metres) divided by the number of hours in the day (12) multiplied by the proportion of the day devoted to moving. Speed of travel is weakly predicted by latitude, altitude and temperature:

$$SPEED \ (m/hr) = -621.2 - 25.7* \ LAT_{Abs} + 0.4*ALT + 46.5*TEMP$$

(r^2=0.674, $F_{3,7}$=1.94, p=0.211).

Fig. S3(a) plots speed against absolute latitude and mean temperature. There is a very slight suggestion that the relationship might be quadratic (U-shaped) in temperature, with speed increasing when ambient temperatures are both low and high, with a minimum at temperatures around 20°C. Alternatively, the relationship may be better described by a power curve in which speed increases dramatically above ~22°C at latitudes <5° either side of the equator.

Figure S3(a)

Mean speed for individual gibbon (filled symbols) and siamang (unfilled symbols) plotted against mean temperature and absolute latitude of the study site, assuming a 12-hour tropical day.

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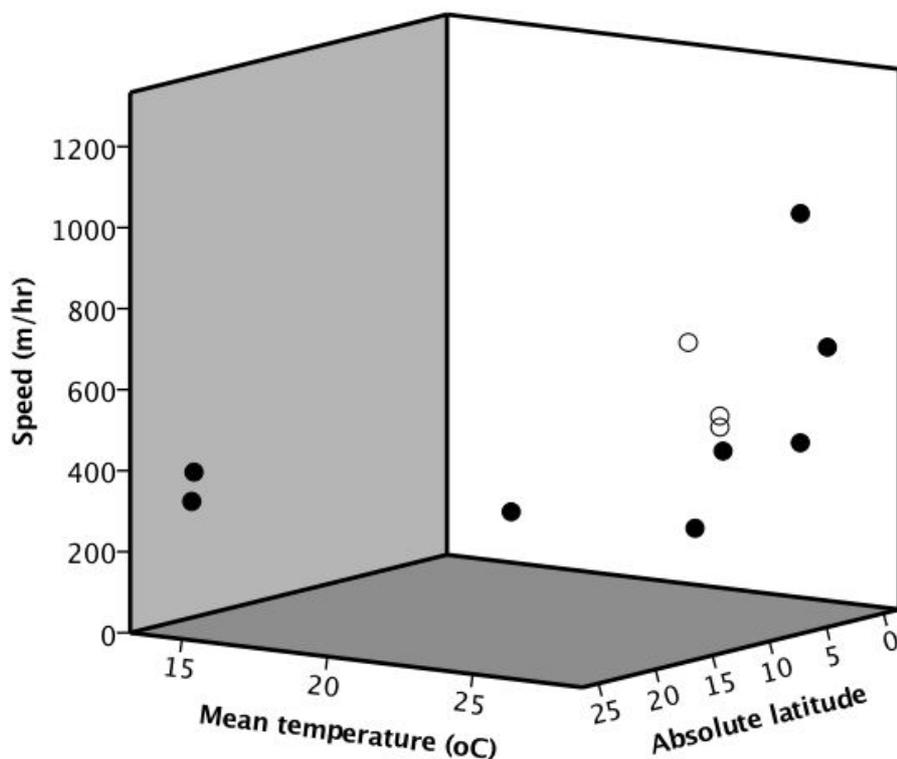
Many gibbon populations retire to nest several hours before sunset, with length of active daylength best predicted by mean rainfall (Table 2). To check whether a shorter active day makes any difference to these results, we recalculated speed as a function of actual length of active day using the equation given in Table 2 to predict length of active day from rainfall for each site. Aside from a slight uniform increase in speed, the transformation has no effect (Fig. S3b).

Figure S3(b)

Mean speed for individual gibbon (filled symbols) and siamang (unfilled symbols) plotted against mean temperature and absolute latitude of the study site, with speed calculated for active day length (estimated using the equation in rainfall from Table 2).
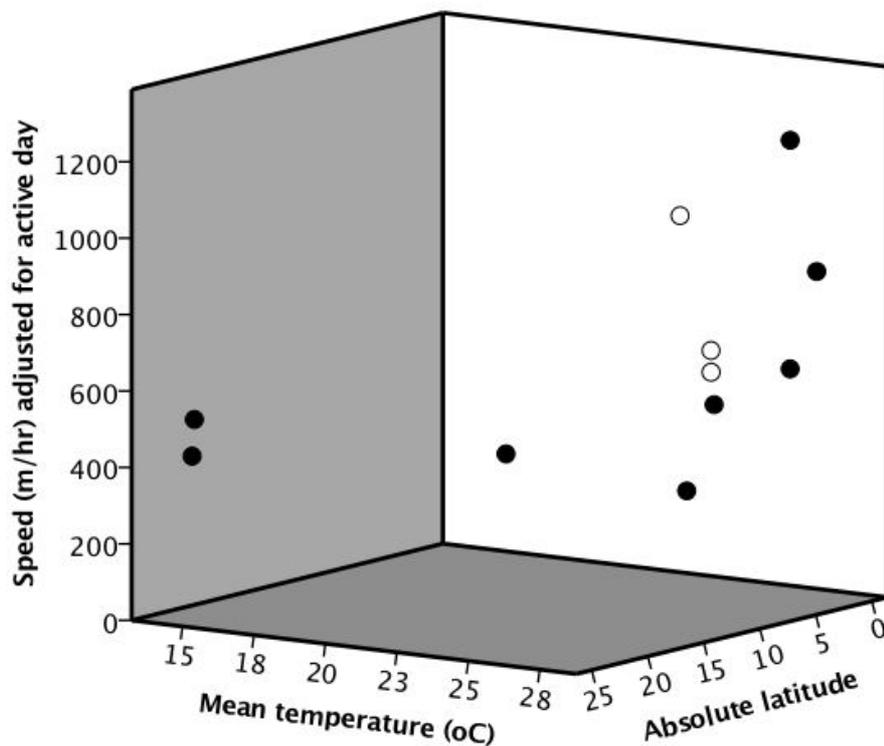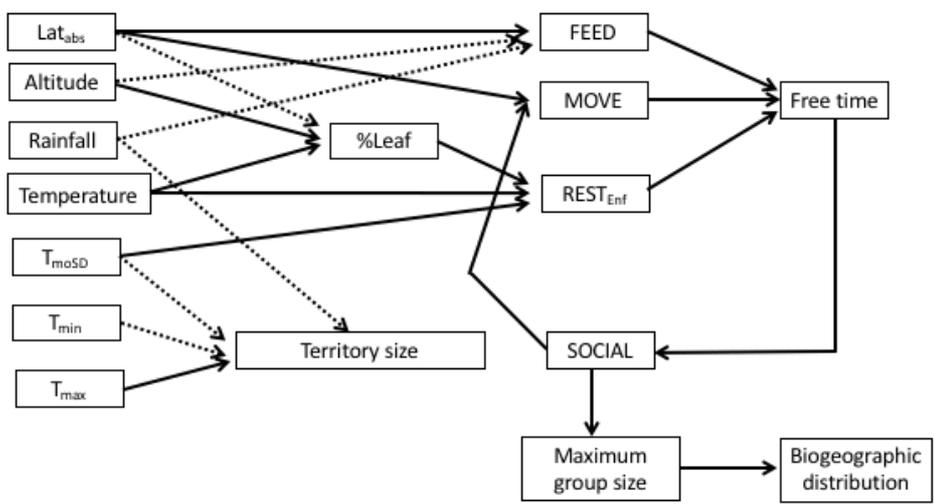
Fig. 2

165x131mm (300 x 300 DPI)

FIG. 3A
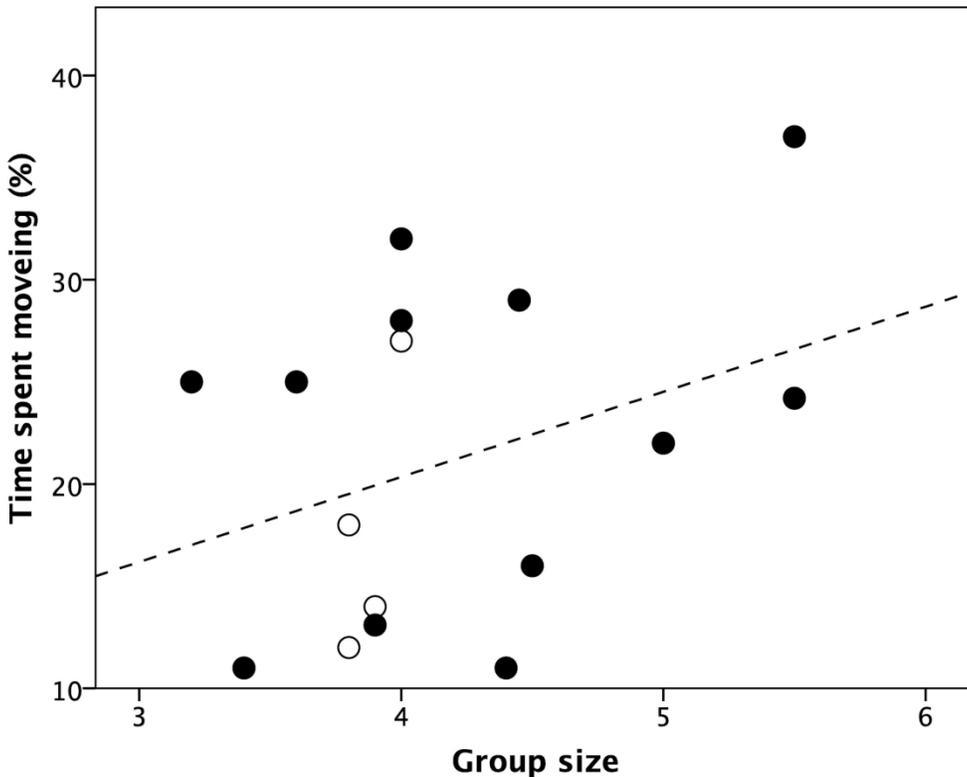
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FIG. 3B

165x131mm (300 x 300 DPI)

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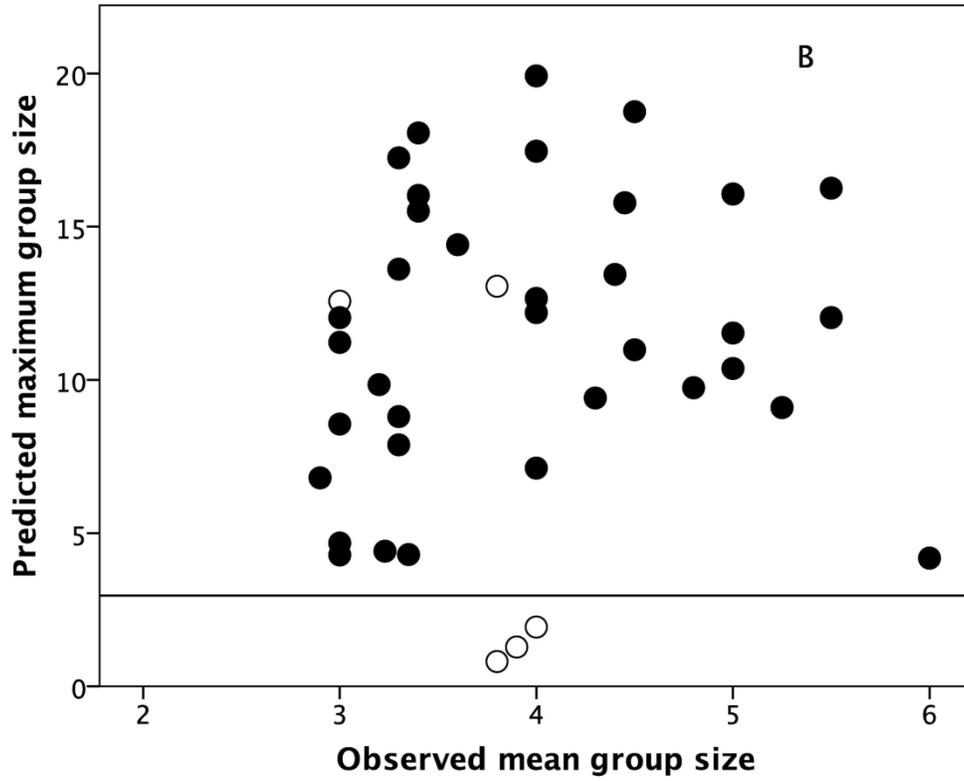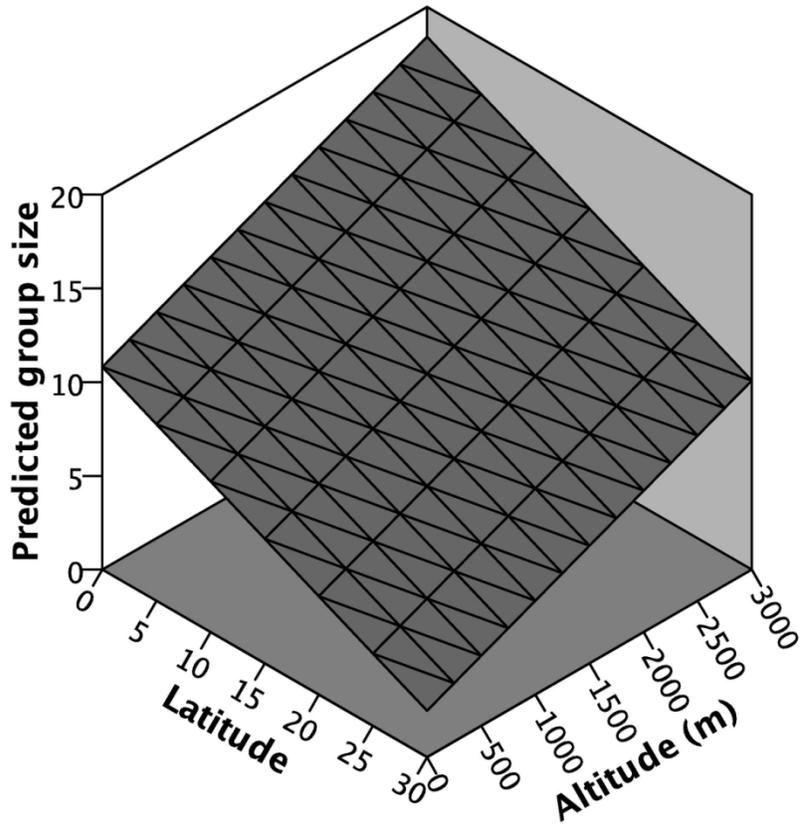
Fig. 4A

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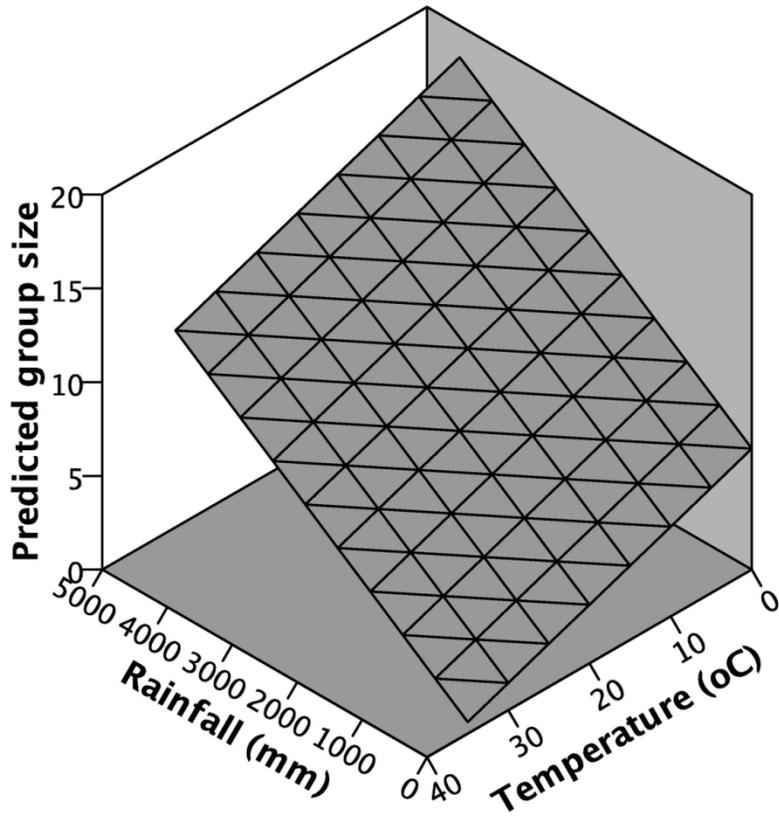
Fig. 4B

123x123mm (300 x 300 DPI)
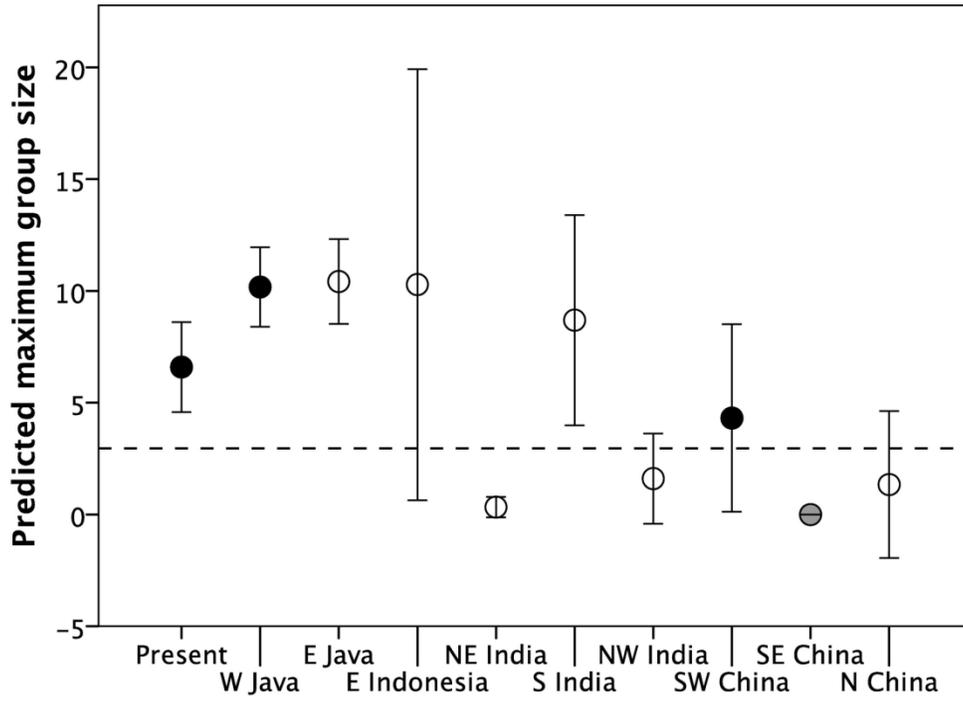
FIG. 5

165x131mm (300 x 300 DPI)
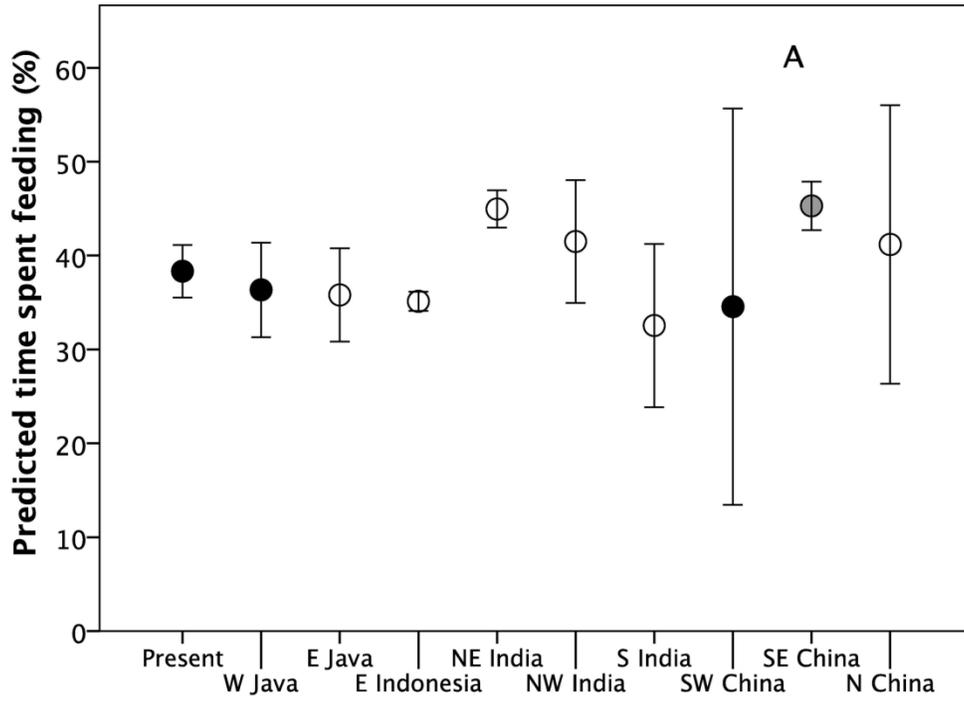
Fig. 6A

165x131mm (300 x 300 DPI)
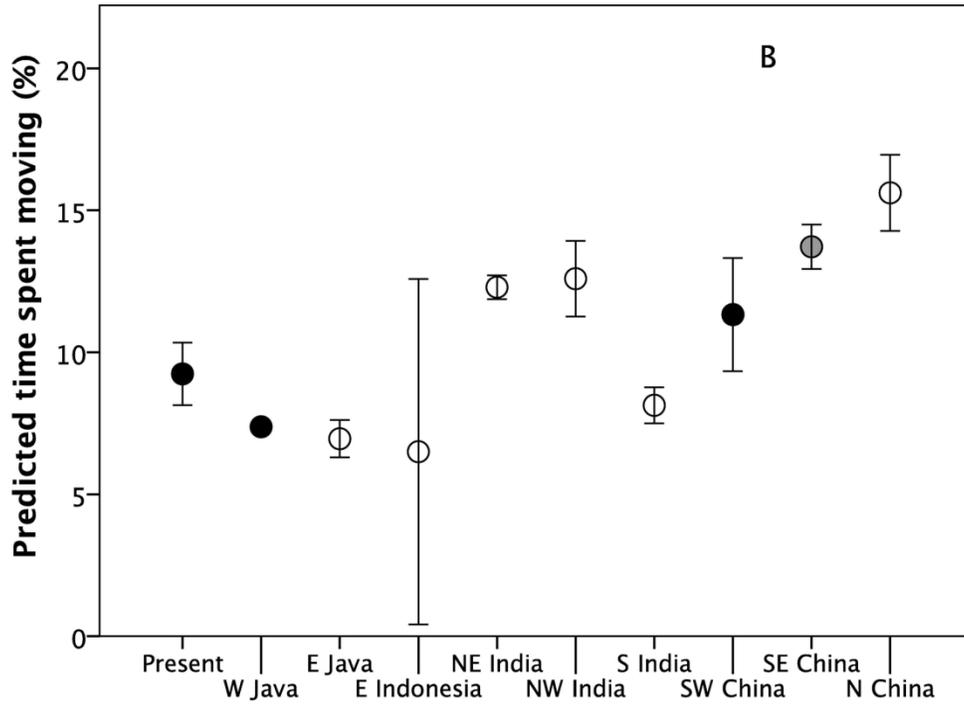
Fig. 6B

165x131mm (300 x 300 DPI)
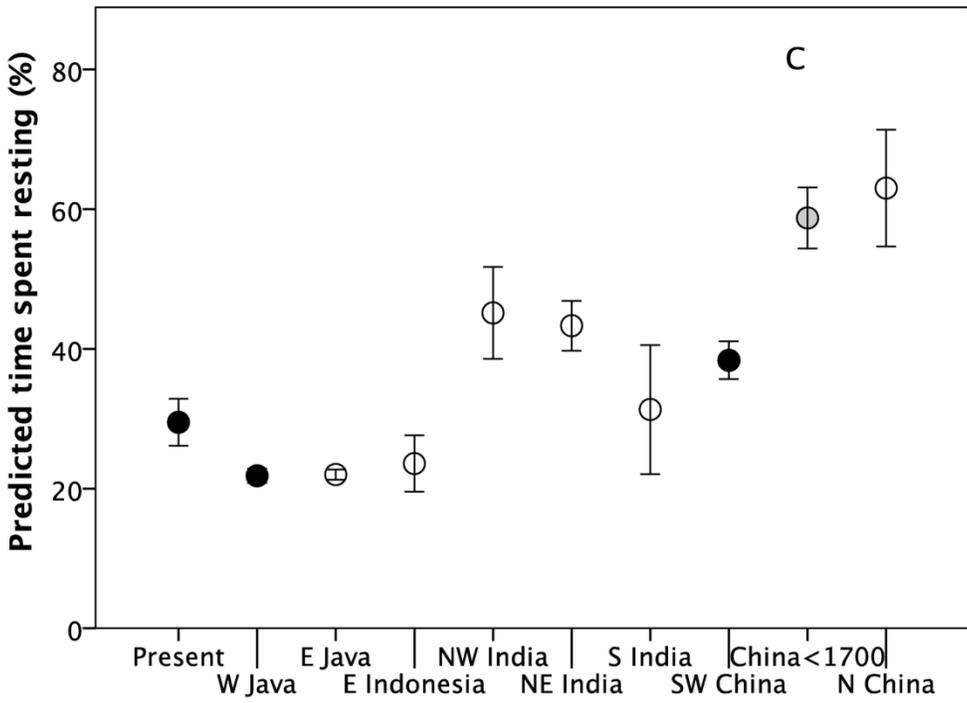
Fig. 6C

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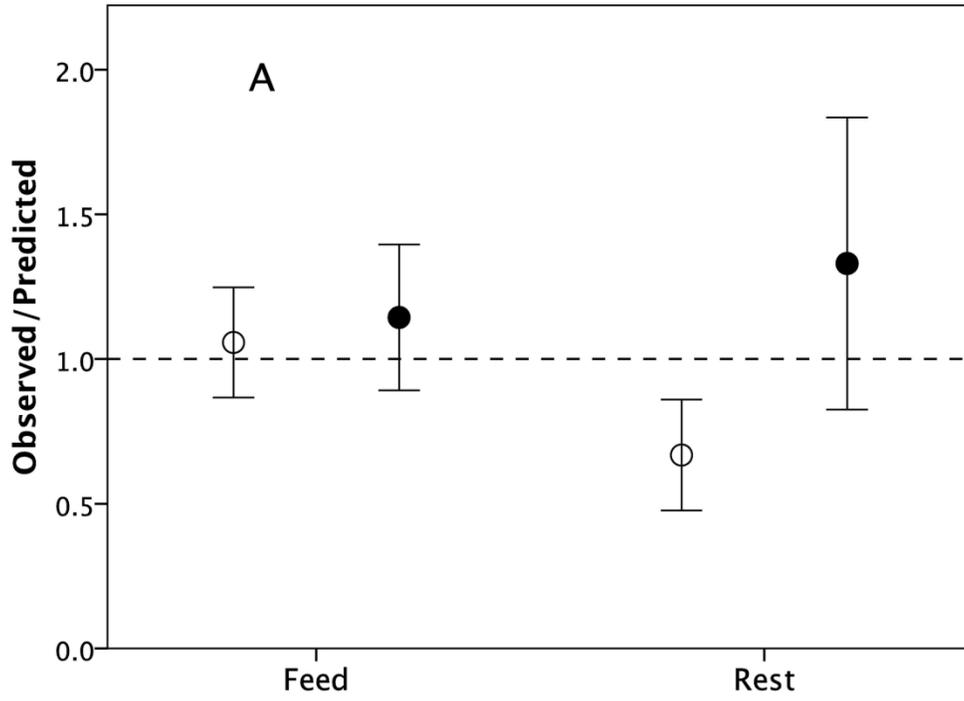
Fig. 7A

165x131mm (300 x 300 DPI)

Fig. 7B

165x131mm (300 x 300 DPI)

FIG. 8

165x131mm (300 x 300 DPI)