

PHILOSOPHICAL TRANSACTIONS OF THE ROYAL SOCIETY B

BIOLOGICAL SCIENCES

Complementarity, completeness and quality of long-term faunal archives in an Asian biodiversity hotspot

Journal:	<i>Philosophical Transactions B</i>
Manuscript ID	RSTB-2019-0217.R1
Article Type:	Research
Date Submitted by the Author:	n/a
Complete List of Authors:	Turvey, Samuel; Zoological Society of London, Institute of Zoology Walsh, Connor; Imperial College London Hansford, James; Zoological Society of London, Institute of Zoology; Northern Illinois University Crees, Jennifer; Natural History Museum, Department of Earth Sciences Bielby, Jon; Liverpool John Moores University Faculty of Science, School of Natural Sciences and Psychology Duncan, Clare; Zoological Society of London, Institute of Zoology Hu, Kaijin; Sun Yat-Sen University, School of Life Sciences Hudson, Mike; Zoological Society of London, Institute of Zoology; Durrell Wildlife Conservation Trust
Issue Code (this should have already been entered but please contact the Editorial Office if it is not present):	FOSSIL
Subject:	Ecology < BIOLOGY, Palaeontology < BIOLOGY, Environmental Science < BIOLOGY
Keywords:	extinction, China, Hainan, historical baseline, Holocene, zooarchaeology

SCHOLARONE™
Manuscripts

Author-supplied statements

Relevant information will appear here if provided.

Ethics

Does your article include research that required ethical approval or permits?:

Yes

Statement (if applicable):

Interviews were only conducted following informed consent of all respondents. Project design was approved by the Zoological Society of London's Ethics Committee, and complied with protocols approved by the Hainan Provincial Forestry Department, the Hainan Provincial Government, and the management office of each protected area where fieldwork was conducted.

Data

It is a condition of publication that data, code and materials supporting your paper are made publicly available. Does your paper present new data?:

Yes

Statement (if applicable):

The datasets supporting this paper are available in the supplementary materials.

Conflict of interest

I/We declare we have no competing interests

Statement (if applicable):

CUST_STATE_CONFLICT :No data available.

Authors' contributions

This paper has multiple authors and our individual contributions were as below

Statement (if applicable):

S.T.T. designed research; C.W., J.J.C., K.H. and S.T.T. coordinated data collection; C.W., J.H., J.B., C.D., M.A.H., K.H. and S.T.T. interpreted and analysed data; and S.T.T. wrote the paper with support from other authors.

1
2
3
4 1 **Complementarity, completeness and quality of long-term faunal**
5
6 2 **archives in an Asian biodiversity hotspot**
7
8
9 3

10
11 4 Samuel T. Turvey^{1,*}, Connor Walsh², James P. Hansford^{1,3}, Jennifer J. Crees⁴, Jon
12
13 5 Bielby⁵, Clare Duncan¹, Kaijin Hu⁶, Michael A. Hudson^{1,7}
14
15 6

16
17
18 7 *¹Institute of Zoology, Zoological Society of London, Regent's Park, London NW1*
19
20 8 *4RY, UK*

21
22
23 9 *²Imperial College London, Silwood Park Campus, Ascot, Berkshire SL5 7PY, UK*
24

25 10 *³Department of Biological Sciences, Northern Illinois University, DeKalb, Illinois*
26
27 11 *60115, USA*

28
29
30 12 *⁴Department of Earth Sciences, Natural History Museum, London SW7 5BD, UK*
31

32 13 *⁵School of Natural Sciences and Psychology, Liverpool John Moores University,*
33
34 14 *Liverpool L3 8PY, UK*

35
36
37 15 *⁶School of Life Sciences, Sun Yat-sen University, Guangzhou, Guangdong 510275,*
38
39 16 *China*

40
41
42 17 *⁷Durrell Wildlife Conservation Trust, Les Augrès Manor, Trinity, Jersey JE3 5BP,*
43
44 18 *Channel Islands*

45
46 19
47
48 20 *Corresponding author: samuel.turvey@ioz.ac.uk
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 21 **Abstract.** Long-term baselines on biodiversity change through time are crucial to
4
5 22 inform conservation decision-making in biodiversity hotspots, but
6
7 23 environmental archives remain unavailable for many regions. Extensive
8
9 24 palaeontological, zooarchaeological and historical records and indigenous
10
11 25 knowledge about past environmental conditions exist for China, a megadiverse
12
13 26 country experiencing large-scale biodiversity loss, but their potential to
14
15 27 understand past human-caused faunal turnover is not fully assessed. We
16
17 28 investigate a series of complementary environmental archives to evaluate the
18
19 29 quality of the Holocene-historical faunal record of Hainan Island, China's
20
21 30 southernmost province, for establishing new baselines on postglacial
22
23 31 mammalian diversity and extinction dynamics. Synthesis of multiple archives
24
25 32 provides an integrated model of long-term biodiversity change, revealing that
26
27 33 Hainan has experienced protracted and ongoing human-caused depletion of its
28
29 34 mammal fauna from prehistory to the present, and that past baselines can inform
30
31 35 practical conservation management. However, China's Holocene-historical
32
33 36 archives exhibit substantial incompleteness and bias at regional and country-
34
35 37 wide scales, with limited taxonomic representation especially for small-bodied
36
37 38 species, and poor sampling of high-elevation landscapes facing current-day
38
39 39 climate change risks. Establishing a clearer understanding of the quality of
40
41 40 environmental archives in threatened ecoregions, and their ability to provide a
42
43 41 meaningful understanding of the past, is needed to identify future conservation-
44
45 42 relevant historical research priorities.
46
47 43
48
49 44 **Key words:** extinction, China, Hainan, historical baseline, Holocene,
50
51 45 zooarchaeology
52
53
54
55
56
57
58
59
60

1. INTRODUCTION

Effective conservation of threatened global biodiversity hotspots urgently requires scientific evidence to inform and guide management [1]. However, whereas biodiversity richness is greatest in the tropics, biodiversity data richness is skewed towards the poles, especially for long-term datasets needed to understand population dynamics, responses to potential threats, and biodiversity change through time [2-3]. Gaps in conservation-relevant data availability are of particular concern in decision-making and prioritisation for eastern and southeast Asian terrestrial ecosystems, which are experiencing extreme anthropogenic pressure and contain the world's highest numbers of threatened vertebrates and plants [4,5]. Identifying the different types of environmental data that exist for these biodiversity hotspots, and determining their information-content and conservation usefulness, is therefore a vital conservation research priority [6].

Conservation planning typically uses modern-day ecological data, with very limited use of longer-term records [6,7]. However, there is increasing recognition that long-term environmental archives, including fossil, zooarchaeological and historical records, can contribute to conservation research, policy and practice by providing unique insights about diversity and composition of past ecosystems, biotic responses to environmental change, species and ecosystem vulnerability to past stressors, and extinction rates and dynamics [8,9]. Many ecosystems, particularly those with long histories of human presence, are likely to have experienced an "extinction filter" whereby biodiversity that was vulnerable to past human pressures has already been lost, making assumptions about ecology, biogeography and extinction risk based only on modern-day data

1
2
3 71 potentially incomplete or misleading [10]. Approaches for integrating past and
4
5 72 present environmental data are now explored by the emerging disciplines of
6
7 73 conservation palaeobiology and historical ecology, which aim to model future
8
9 74 biodiversity scenarios and identify management tools and restoration targets
10
11 75 based on historical baselines [6-9].
12
13
14

15 76 Unfortunately, assessing the information-content of environmental archives
16
17 77 and incorporating historical baselines into conservation planning remains
18
19 78 challenging for biodiversity hotspots due to limited availability of relevant
20
21 79 archives for many regions, especially in the tropics [6,11]. China represents an
22
23 80 important exception. This huge (9.6 million km²), 'megadiverse' country contains
24
25 81 >10% of global mammal species and covers a diverse range of habitats [12], but
26
27 82 has experienced human overpopulation, resource overexploitation and habitat
28
29 83 modification throughout the climatically stable postglacial Holocene Epoch and
30
31 84 historical period [13,14], leading to catastrophic ongoing biodiversity loss
32
33 85 including ecosystem functional and compositional collapse, population
34
35 86 extirpations and species extinctions [15-17]. Multiple environmental archives,
36
37 87 spanning different temporal depths and spatio-temporal resolutions across the
38
39 88 Holocene-historical period, are available to investigate postglacial human-
40
41 89 ecosystem interactions and impacts in China, including palaeontological and
42
43 90 zooarchaeological records [11,17,18], a written record going back over two
44
45 91 millennia with abundant information on past environmental conditions [16], and
46
47 92 a rich body of indigenous knowledge about past and present biodiversity held by
48
49 93 China's large rural population [19].
50
51
52
53
54
55

56
57 94 China's long-term archives have been used to reconstruct regional ecological
58
59 95 histories and investigate historical and prehistoric human-environmental
60

1
2
3 96 interactions [13]. They also have the potential to provide important insights into
4
5 97 the changing status of China's biodiversity and make predictive hypotheses to
6
7
8 98 guide conservation management. However, although 253 mainland Chinese
9
10 99 Holocene archaeological and palaeontological sites contain identified wild
11
12 100 mammal species (figure 1a) [17], the ability of these archives to define past
13
14
15 101 biodiversity baselines and faunal responses to human activities has not been
16
17 102 fully assessed. Previous Chinese Holocene-historical faunal studies have focused
18
19 103 on using specific archives to identify extinct species [20] and reconstruct the past
20
21 104 ecology and distribution of threatened species [16,21]. However, important
22
23
24 105 wider questions remain unexplored about the quality of China's environmental
25
26 106 archives, the relative contribution of different historical baselines for
27
28 107 understanding patterns and processes of biodiversity change, and the ability of
29
30 108 long-term datasets to provide a meaningful understanding of the past of use for
31
32
33 109 conservation.

34
35
36 110 In this study, we assess the extent to which long-term faunal archives can
37
38 111 contribute unique conservation-relevant information on Chinese biodiversity in
39
40 112 two ways. We use a series of environmental archives available for a regional
41
42 113 Chinese study system to define successive Holocene faunal baselines, identify
43
44 114 differences in species composition between past and present, reconstruct the
45
46 115 timing and drivers of past biodiversity loss, and determine the extent that
47
48 116 different archives can complement each other to reconstruct faunal dynamics
49
50 117 through time. We also identify and quantify patterns of incompleteness and bias
51
52 118 in Chinese faunal archives at both regional and country-wide scales. These
53
54
55 119 analyses establish a new framework for assessing the unique opportunities and
56
57
58
59
60

1
2
3 120 inherent limitations in using environmental archives to inform conservation
4
5 121 planning.

6
7
8 122

9
10 123 **2. MATERIAL AND METHODS**

11
12 124

13
14
15 125 **(a) Regional study system**

16
17 126 Hainan Island, China's southernmost province, is a 33,920km² subtropical-
18
19 127 tropical continental-shelf island in the South China Sea (figure 1b). Hainan
20
21 128 probably became isolated from mainland China through marine transgression
22
23 129 during the early Holocene between 7,100-10,500 yr BP [22]. Its current-day land
24
25 130 mammal fauna contains 83 recorded species and is relatively depauperate,
26
27 131 lacking numerous species known from mainland China and southeast Asia [12]
28
29 132 (electronic supplementary material, table S1). This fauna includes the Hainan
30
31 133 gibbon (*Nomascus hainanus*), one of the world's rarest mammals, with a global
32
33 134 population of only 27 surviving individuals [23]. Ancient DNA analysis of
34
35 135 historical museum collections has recently shown that the last population of Père
36
37 136 David's deer or milu (*Elaphurus davidianus*), which has been extinct in the wild
38
39 137 for over a century, occurred on Hainan during the mid-1800s [24]. The current
40
41 138 depauperate state of Hainan's mammal fauna may therefore represent depletion
42
43 139 following additional past extinctions. Because faunal turnover on an island is not
44
45 140 influenced by population migration, Hainan constitutes a "closed system" for
46
47 141 investigating extinction dynamics. However, the magnitude, timing, and drivers
48
49 142 of any such events have not been investigated.

50
51
52 143 In addition to occasional historical accounts by visiting naturalists [e.g. 25],
53
54
55 144 three main temporally non-overlapping Holocene-historical faunal archives are

1
2
3 145 available for Hainan: **(1)** A rich fossil deposit from Luobidong Cave, dated to
4
5 146 10,642±207 yr BP [26], containing abundant mammal material [27] and
6
7
8 147 therefore providing a faunal baseline approximately at the point when Hainan
9
10 148 became an island (figure 1b); **(2)** Hainan's gazetteer record, covering the late
11
12 149 Ming Dynasty, Qing Dynasty and Republican Period, and containing considerable
13
14 150 local environmental data including animal records [28]; **(3)** Indigenous
15
16 151 knowledge about past and present biodiversity, possessed by Li and Miao ethnic
17
18 152 communities in Hainan's forested interior [19].
19
20
21
22 153

23 24 154 **(b) Hainan fossil data**

25
26 155 The Luobidong cave fauna contains 38 identifiable mammal species (corrected to
27
28 156 taxonomy in [12]), including 12 unknown on Hainan today in Proboscidea,
29
30 157 Perissodactyla, Artiodactyla, Carnivora, Rodentia and Chiroptera: Asian elephant
31
32 158 (*Elephas maximus*), tapir (*Tapirus* sp.), buffalo (*Bubalus* sp.), serow (*Capricornis*
33
34 159 sp.), tufted deer (*Elaphodus cephalophus*), Reeves' muntjac (*Muntiacus reevesi*),
35
36 160 tiger (*Panthera tigris*), dhole (*Cuon alpinus*), hog badger (*Arctonyx collaris*),
37
38 161 greater bandicoot rat (*Bandicota indica*), vole (*Microtus* sp.), hairy-winged bat
39
40 162 (*Harpiocephalus* sp.). Regional mammal extinctions are unlikely to be associated
41
42 163 with Holocene climate change, which was very limited compared to Late
43
44 164 Pleistocene change [29,30]. We identify two competing hypotheses: (1) human-
45
46 165 caused extinction; (2) stochastic extinction on a closed island system [31,32]. We
47
48 166 tested between these hypotheses by assessing likelihood of stochastic extinction
49
50 167 across the Holocene for regionally extirpated megafaunal mammals. We
51
52 168 conducted population viability analysis (PVA) for the three largest regionally
53
54 169 extinct herbivores (Asian elephant, buffalo, tapir) and largest regionally extinct
55
56
57
58
59
60

1
2
3 170 carnivore (tiger) in the Luobidong fauna, as these species are most vulnerable to
4
5 171 stochastic extinction due to low population densities and large spatial
6
7
8 172 requirements [33]. We ran PVA base models in Vortex v.10 [34] over an 8,000-
9
10 173 year period, in 50 evenly-spaced survival-level increments between 5% and 95%
11
12 174 inclusive, including catastrophes as a stochastic extinction driver and with Latin
13
14 175 hypercube sensitivity analysis of different variables (electronic supplementary
15
16 176 material, text S1, table S2).
17
18
19
20
21

22 178 **(c) Hainan historical data**

23
24 179 We surveyed 44 gazetteer volumes dating between 1521-1935 AD from Hainan
25
26 180 [28]. We recorded animals listed in the 'beasts' (*shou*) sections and/or that
27
28 181 accompanied separate reports of gibbons, excluding obviously mythical or
29
30 182 fantastical reports (electronic supplementary material, table S3). Hainan was
31
32 183 periodically administered with nearby mainland provinces, so we confirmed that
33
34 184 records referred to Hainan from supporting information. For selected animal
35
36 185 records (see Results), we conducted optimal linear estimation (OLE), a
37
38 186 probabilistic approach that uses temporal distributions of independent sighting
39
40 187 events to estimate an extinction date [35], implemented using the "sExtinct"
41
42 188 package [36] in R [37].
43
44
45
46
47
48
49

50 190 **(d) Hainan local ecological knowledge**

51
52 191 We conducted interviews in January-April 2015 in villages close to seven
53
54 192 Hainanese protected areas (Bawangling, Diaoluoshan, Jianfengling, Wuzhishan
55
56 193 and Yinggeling National Nature Reserves; Jiayi and Limushan Provincial Nature
57
58 194 Reserves; figure 1b). Local people use animal and plant resources collected from
59
60

1
2
3 195 inside these protected areas [23]. We randomly selected 10 villages around each
4
5 196 reserve and aimed to conduct 10 interviews per village. We used a standard
6
7
8 197 anonymous questionnaire for all interviews, which took up to 1 hour to
9
10 198 complete, with interviews mainly conducted in Mandarin or Hainanese and
11
12 199 recorded in Chinese (electronic supplementary material, text S2). Respondent
13
14 200 selection criteria/methods and interview protocols are given in ref. 19. Project
15
16 201 design was approved by the Zoological Society of London's Ethics Committee.

19 202 In addition to other data presented in ref. 19,38,39, we collected data on
20
21 203 respondent awareness and experience of nine mammal species: wild pig (*Sus*
22
23 204 *scrofa*), rhesus macaque (*Macaca mulatta*), Hainan gibbon, clouded leopard
24
25 205 (*Neofelis nebulosa*), Asian black bear (*Ursus thibetanus*), Chinese pangolin (*Manis*
26
27 206 *pentadactyla*), binturong (*Arctictis binturong*), sambar deer (*Rusa unicolor*), giant
28
29 207 anteater (*Myrmecophaga tridactyla*). Most of these species are known or
30
31 208 suspected to occur in Hainan [12,40]; giant anteaters are native to the Neotropics
32
33 209 and were a negative control to check response accuracy. We showed colour
34
35 210 photographs of these mammals (sourced from www.arkive.org and the
36
37 211 Zoological Society of London), shown in the same order given above in all
38
39 212 interviews, and asked respondents to name species and provide further
40
41 213 ecological/morphological details to confirm recognition. If respondents did not
42
43 214 recognize photographs, we used standard Chinese names to prompt recall. We
44
45 215 asked if respondents had encountered animals in the photographs (including
46
47 216 sightings, hearing gibbon song, and diagnostic footprints/sign), and if so where
48
49 217 and how recently. We also asked if respondents knew about any animals that had
50
51 218 existed in the past but no longer occurred locally, and whether they knew any
52
53 219 old stories that described animals that had only existed in the past.
54
55
56
57
58
59
60

1
2
3 220 Encounter records were converted to direct calendar years for analysis,
4
5 221 following ref. 39; encounter data reported below represent converted data.
6
7 222 Differences in species last-encounter histories for 1990-2015 were analysed
8
9 223 using generalised linear models (GLM) in R. Frequency of last-encounter dates
10
11 224 per species per year was expressed as a proportion of total number of
12
13 225 observations for each species encounter-history dataset, and regressed on year
14
15 226 (predictor) [41]. We used a binomial error structure unless data showed
16
17 227 overdispersion, when a quasibinomial error structure was used. Last-encounter
18
19 228 history trajectories between species over time were considered significantly
20
21 229 different if confidence intervals of regression slopes did not overlap; 83%
22
23 230 confidence intervals were used for comparison because these give an
24
25 231 approximate $\alpha=0.05$ test, whereas comparisons using 95% confidence intervals
26
27 232 are too conservative [42]. Lower encounter-history slopes indicate fewer
28
29 233 encounters have occurred close to the present. The oldest 5% of records for each
30
31 234 species all date from before 1990, so there was no need to further exclude these
32
33 235 data from analysis to reduce the effect of long encounter data “tails” (which
34
35 236 produce flatter overall encounter-history slopes that are harder to differentiate
36
37 237 statistically) [41].
38
39
40
41
42
43
44
45
46

47 239 **(e) Bias in China’s Holocene record**

48
49 240 We investigated whether representation of past mammalian diversity in China’s
50
51 241 Holocene faunal record is biased by exploring whether biological/ecological
52
53 242 traits other than abundance can predict the number of Holocene site records for
54
55 243 mainland Chinese species. We considered body mass and geographic range,
56
57 244 which are both predictors of fossil species occurrence in other systems [43,44].
58
59
60

1
2
3 245 We used phylogenetic comparative methods to account for biases associated
4
5 246 with shared evolutionary history, and ran all models using the `ppls` function in
6
7 247 the R package “`caper`” [45], using the dated mammal supertree of ref. 46 and with
8
9 248 taxonomy standardized between datasets (electronic supplementary material,
10
11 249 text S1). We first investigated the relationship between body mass and site
12
13 250 records for 493 species, using log-transformed body mass estimates (electronic
14
15 251 supplementary material, text S1, table S4). This dataset contained a high
16
17 252 proportion of species with 0 site records ($n=377$); because zero-inflated datasets
18
19 253 can create problems for quantifying relationships between variables, we
20
21 254 conducted bootstrapping to understand how removal of different proportions of
22
23 255 species with 0 site records affected parameter estimates and robustness of
24
25 256 model inferences (electronic supplementary material, text S1). We then
26
27 257 investigated the predictive power of both body mass and geographic range. Many
28
29 258 Chinese mammal ranges have decreased over the Holocene, making modern-day
30
31 259 distributions inappropriate proxies for past distributions [16,17]. Standardised
32
33 260 Holocene range estimates are available for 34 species [17], so we analysed this
34
35 261 reduced species subset using both body mass and Holocene range as predictors
36
37 262 of site records (electronic supplementary material, table S4).

38
39 263 We also investigated whether China’s Holocene faunal record is spatially
40
41 264 biased and representative of past ecological diversity, using two approaches. We
42
43 265 used nearest-neighbour analysis in ArcGIS Pro v.2.3.0 [47] to test whether
44
45 266 Holocene sites are spatially clustered, by measuring mean nearest-neighbour
46
47 267 distance between sites and comparing this with expected mean nearest-
48
49 268 neighbour distances for a point set with a random distribution. We also carried
50
51 269 out chi-squared tests in R on number of sites present in each mainland Chinese
52
53
54
55
56
57
58
59
60

1
2
3 270 ecoregion as defined in the Terrestrial Ecosystems of the World dataset [48], to
4
5 271 test whether spatial distribution of sites shows biogeographic bias. We
6
7
8 272 calculated expected values manually by multiplying mean site density across
9
10 273 China by total area of each ecoregion, excluding all ecoregions under 15,000km²
11
12 274 (size of smallest ecoregion containing at least one site) to reduce the number of
13
14
15 275 low expected counts.
16
17 276

19 277 **3. RESULTS**

24 279 **(a) Hainan's long-term archives**

26 280 PVA base models for elephant, buffalo, tapir and tiger populations on Hainan
27
28 281 quickly grew to their carrying capacities and remained stable with no incidences
29
30
31 282 of extinction in the absence of catastrophes. Modelled stochastic catastrophes
32
33 283 had to be severe to drive populations to extinction (*Elephas maximus*: all extinct
34
35 284 at $\leq 82\%$ survival, all survive at $\geq 88\%$ survival; *Bubalus* sp.: all extinct at $\leq 44\%$
36
37 285 survival, all survive at $\geq 64\%$ survival; *Tapirus* sp.: all extinct at $\leq 64\%$ survival, all
38
39 286 survive at $\geq 84\%$ survival; *Panthera tigris*: all extinct at $\leq 31\%$ survival, all survive
40
41 287 at $\geq 58\%$ survival).

45 288 Hainan's gazetteer record contains 104 land mammal "types" (excluding
46
47 289 bats, which are usually classified separately as "flying creatures" or "insects" and
48
49 290 were not catalogued here). Of these, 84 do not obviously correspond with
50
51 291 domestic taxa (electronic supplementary material, table S3). Interpretation and
52
53 292 identification of records, although often aided by accompanying brief
54
55 293 descriptions, is inevitably subjective (e.g. "cat" may refer to domestic or wild
56
57 294 taxa). We are able to identify 15 recognisable species of Artiodactyla, Carnivora,

1
2
3 295 Erinaceomorpha, Lagomorpha, Pholidota and Primates: wild pig, sambar, red
4
5 296 muntjac (*Muntiacus muntjak*), Eld's deer (*Rucervus eldii*), Asian black bear,
6
7
8 297 clouded leopard, leopard cat (*Prionailurus bengalensis*), wolf (*Canis lupus*), dhole,
9
10 298 yellow-throated marten (*Martes flavigula*), Hainan gymnure (*Neohylomys*
11
12 299 *hainanensis*), Hainan hare (*Lepus hainanus*), Chinese pangolin, rhesus macaque,
13
14 300 Hainan gibbon. Some species are referenced with multiple historical names (to a
15
16 301 maximum of five for black bear). Other types refer to wider species groups (e.g.
17
18 302 "porcupines", "squirrels") or cannot be identified beyond a broad taxonomic
19
20 303 category (e.g. 16 small carnivore types cannot be identified beyond Viverridae,
21
22 304 Herpestidae or Mustelidae). Records of "wild cattle" may refer to gaur (*Bos*
23
24 305 *gaurus*), which are not otherwise recorded from Hainan, but also possibly to
25
26 306 other wild/domestic ungulates. Deer referred to as "milu" are reported until
27
28 307 1917, but these cannot definitely be identified as Père David's deer because
29
30 308 reported deer nomenclature is confused; 14 deer types are recorded, some with
31
32 309 similar names (e.g. "mi" [elk]), and including other species otherwise unknown
33
34 310 from Hainan (e.g. water deer, musk deer) that probably represent
35
36 311 misidentifications. Occasional mentions of other regionally extinct or otherwise
37
38 312 unknown species are clearly allegorical or poetic (e.g. 1908: "The wind brings
39
40 313 the stink of a crouching tiger"), so are ignored here.

41
42
43 314 Two well-described species from Hainan's gazetteer record (wolf, 7 records,
44
45 315 1618-1931; dhole, 15 records, 1521-1935) are not present today on Hainan,
46
47 316 although they still occur in nearby mainland China and southeast Asia [12,40].
48
49 317 We conducted OLE on dated records for each species, giving estimated extinction
50
51 318 dates of 1941 for wolf (95% CI=1931-2079), and 1942 for dhole (95% CI=1935-
52
53 319 1993).
54
55
56
57
58
59
60

1
2
3 320 We interviewed 709 respondents in villages across Hainan (mean age=50.1,
4
5 321 range=20-94, male:female=83:17%), who reported past encounter data for
6
7 322 seven of our eight target Chinese mammals. We excluded reports that were
8
9 323 obviously not of wild animals (e.g. “on television”, “in a market”), and data from
10
11 324 two respondents who claimed to have seen giant anteaters. Six respondents
12
13 325 reported possible old sightings (20-60 years ago) of binturong, a species not
14
15 326 confirmed from Hainan [40], but only provided basic descriptions and did not
16
17 327 differentiate it from other regionally occurring civets, so we do not consider
18
19 328 these uncertain reports further. Our interview dataset shows substantial
20
21 329 between-species variation in numbers of respondents reporting encounters and
22
23 330 last-encounter dates, interpreted as reflecting variation in species’ regional
24
25 331 abundance and recent survivorship (table 1). Pig and macaque have the highest
26
27 332 encounter-history slopes, followed by gibbon, bear, clouded leopard, sambar and
28
29 333 pangolin; pig, macaque and gibbon all have significantly higher encounter-
30
31 334 history slopes compared to bear, clouded leopard, sambar and pangolin (figure 2,
32
33 335 table 1). Only 20 respondents named specific animals they thought had existed in
34
35 336 the past but were now locally extinct; these included pangolin (n=6), bear (n=3),
36
37 337 parrot (n=3), snake/python (n=3), turtle (n=3), gibbon (n=2), tiger (n=1), wild
38
39 338 pig (n=1), and muntjac (n=1).

339

340 **(b) Quality of China’s Holocene record**

341 In full analysis of 493 species, body mass was a significant predictor of site
342 records for Chinese mammals (est=1.829, S.E.=0.488, t-value=3.746, $p<0.001$;
343 $df=369$, $R^2=0.03$, $\lambda=0.596$), with larger-bodied species present in more
344 sites (figure 3). Bootstrapping treatments yielded a positive significant

1
2
3 345 relationship between log-transformed body mass and site number, showing that
4
5 346 our results are robust to the proportion of zeroes present in the dataset
6
7
8 347 (electronic supplementary material, text S1, figure S1). In reduced analysis of 34
9
10 348 species, neither body mass nor geographic range were significant predictors of
11
12 349 site number (*body mass*: est=-0.0007, S.E.=0.011, t-value=-0.071, p=0.944;
13
14 350 *geographic range*: est=-0.000002, S.E.<0.001, t-value=-0.627, p=0.535; df=31,
15
16
17 351 $R^2 < 0.001$, lambda=0.099), probably representing a Type II error associated with
18
19 352 small sample size.

20
21
22 353 Holocene sites are significantly more clumped than expected under a
23
24 354 random distribution (z-score=-14.61, p<0.0001), with mean expected and
25
26 355 observed nearest-neighbour distances showing a ratio of 0.52
27
28
29 356 (50.65km:97.38km). Site density differs significantly across mainland Chinese
30
31 357 ecoregions (chi-sq=494.35, df=47, p<0.0001; electronic supplementary material,
32
33 358 figure S2, table S5). The three ecoregions with the greatest number of sites
34
35
36 359 compared with expected values are the Yellow River Plain mixed forest (O=69,
37
38 360 E=11.77), Dabashan evergreen forest (O=22, E=4.56) and Yangtze Plain
39
40
41 361 evergreen forest (O=25, E=11.87), and the three ecoregions with the lowest
42
43 362 number of sites compared with expected values are the Taklimakan desert (O=2,
44
45 363 E=20.16), Central Tibetan Plateau alpine steppe (O=2, E=17.04) and Alashan
46
47 364 Plateau semi-desert (O=2, E=12.39).

48
49
50 365

51 52 366 **4. DISCUSSION**

53
54 367 Our investigation of multiple Chinese long-term environmental archives,
55
56 368 spanning different temporal resolutions and spatial scales, provides a new
57
58
59 369 assessment of the conservation-relevant information-content of different
60

1
2
3 370 historical datasets that can potentially inform evidence-based management in a
4
5 371 biodiversity hotspot. Through the use of diverse analyses, we highlight the types
6
7
8 372 of novel insights provided by long-term faunal records on historical
9
10 373 environmental baselines and patterns and dynamics of biodiversity change, but
11
12 374 also key issues regarding quality and completeness of faunal records, and the
13
14
15 375 extent to which data incompleteness and bias might limit integration of
16
17 376 environmental archives into conservation decision-making in global-priority
18
19
20 377 regions.

21
22 378 Our analyses of Holocene-historical faunal records from Hainan demonstrate
23
24 379 how regional archives can contribute unique new perspectives that improve our
25
26 380 understanding of biogeography, ecosystem composition and extinction
27
28
29 381 vulnerability, with direct relevance for conservation research and practical
30
31 382 management. Firstly, the long-term archives available for Hainan provide a new
32
33 383 baseline on the island's past postglacial species diversity that is unattainable
34
35
36 384 from modern-day data. The Holocene fossil and historical records, and additional
37
38 385 insights from ancient DNA analysis of museum archives [24], together reveal that
39
40 386 the current depauperate state of Hainan's mammal fauna is a historically recent
41
42
43 387 rather than a long-term "natural" ecological condition, with 14 species in six
44
45 388 orders (17% of Hainan's present-day mammalian species richness) recorded in
46
47 389 postglacial faunal archives but unknown from Hainan today. It is possible that
48
49
50 390 some bats and rodents recorded at Luobidong might be undetected rather than
51
52 391 extinct today, as Hainan's small mammal fauna remains relatively understudied
53
54
55 392 [49]. However, Holocene-historical archives also demonstrate that Hainan
56
57 393 formerly contained a typical southeast Asian large mammal fauna comprising a
58
59
60

1
2
3 394 diverse assemblage of megaherbivores, megacarnivores, mesoherbivores and
4
5 395 mesocarnivores, which have now largely disappeared from the island.
6
7

8 396 Hainan's long-term archives also provide a baseline for reconstructing
9
10 397 relative extinction timings for different components of the island's large mammal
11
12 398 fauna over time. The largest-bodied herbivores and carnivores present at
13
14 399 Luobidong (elephants, tapirs, buffalos, tigers) had already disappeared by the
15
16 400 time historical accounts of Hainan's biodiversity and natural resources were first
17
18 401 recorded. Although the exact timing of extinctions remains unclear, these species
19
20 402 are not referenced in Hainan's gazetteer archive dating from the 1500s onwards,
21
22 403 and Hainan was described as "without horses or tigers" as early as 80 CE [13].
23
24 404 Smaller-bodied mesoherbivores and mesocarnivores survived until much more
25
26 405 recently. Père David's deer persisted on Hainan until at least the nineteenth
27
28 406 century [24] and possibly until 1917 based on gazetteer records; and OLE
29
30 407 analysis of gazetteer records for wolf and dhole together with indigenous
31
32 408 knowledge from local respondents demonstrates these species persisted into the
33
34 409 mid-twentieth century but apparently disappeared before living memory.
35
36 410 Comparison of different probabilistic methods for inferring extinction using
37
38 411 sighting records suggests OLE is more robust than other approaches, and
39
40 412 generally gives accurate predictions when applied to >5 records [50], although
41
42 413 use of >10 records is recommended by ref. 51, a condition fulfilled by dhole but
43
44 414 not wolf. We also note that estimated extinction dates for wolf and dhole
45
46 415 represent last-occurrence dates only, because OLE relies on the implicit
47
48 416 assumption that recording effort never falls to zero [35], but Hainan's gazetteer
49
50 417 record stops at the end of the Nationalist Era.
51
52
53
54
55
56
57
58
59
60

1
2
3 418 These archives also demonstrate that the decline of Hainan's mammal fauna
4
5 419 is ongoing. Our large-scale dataset of species sightings over recent decades
6
7
8 420 shows that bears, clouded leopards, sambar and pangolins all have lower
9
10 421 encounter-history slopes in comparison to encounter data for the Hainan gibbon,
11
12 422 one of the world's rarest mammals, with fewer sightings of these species close to
13
14 423 the present suggestive of steeply declining populations. Although this pattern
15
16 424 might be explained partly by differing species detectabilities, with gibbons
17
18 425 potentially easier to detect due to their diurnal activity and singing behaviour,
19
20 426 we consider it sadly likely that remnant populations of some or all of these other
21
22 427 species are now on the verge of extinction if not already gone, especially because
23
24 428 there are currently no species-specific conservation programmes to help
25
26 429 safeguard any mammal species on Hainan other than Hainan gibbon or Eld's
27
28 430 deer [23,52].
29
30
31
32

33 431 Further analysis of baseline data from long-term archives also identifies the
34
35 432 likely driver of extinctions in Hainan's postglacial mammal fauna. Our PVA
36
37 433 results show that, unlike some other Late Quaternary island systems with
38
39 434 isolated large-bodied mammal populations [32,53], Hainan is large enough to
40
41 435 support long-term viable populations of megaherbivores and megacarnivores
42
43 436 that were present when the island became isolated in the early Holocene, with
44
45 437 natural catastrophes required to cause stochastic extinction in model
46
47 438 simulations considered too severe to be ecologically plausible (i.e. requiring
48
49 439 destruction of $\geq 12\%$ of the island's carrying capacity). We can therefore exclude
50
51 440 the hypothesis of stochastic extinction, and identify human activity as the only
52
53 441 plausible driver of Holocene mammalian losses on Hainan. Prehistoric human-
54
55 442 environment interactions and demographic changes on Hainan are poorly
56
57
58
59
60

1
2
3 443 understood, making it difficult to determine specific activities that caused past
4
5 444 extinctions. Hainan's ecosystems were being heavily exploited for natural
6
7
8 445 resources by the 18th century for trade with mainland China [13], but the
9
10 446 island's megafauna was already extinct by this point. Neolithic cultures have
11
12 447 been present on Hainan since at least 6,000 yr BP [54], and aboriginal peoples
13
14
15 448 are known to have transformed Hainan's environment to some degree through
16
17 449 hunting and agricultural conversion during recent millennia, but prehistoric
18
19 450 human populations have generally been assumed to be too small to cause much
20
21 451 environmental impact [13]. However, evidence for heavy metal pollution from
22
23 452 around 4,000 yr BP, associated with appearance of abundant archaeological sites
24
25 453 on Hainan, indicates intensification of regional human activities (e.g. agricultural
26
27 454 development, deforestation, metal utilization) that could have been associated
28
29 455 with mammal extinctions [55].
30
31
32

33 456 Investigation of multiple complementary faunal archives, stretching from the
34
35 457 living memory of local inhabitants back to the early Holocene, thus enables
36
37 458 development of an integrated model of long-term mammalian biodiversity
38
39 459 change for Hainan (figure 4). This overview of faunal dynamics is only possible
40
41 460 through synthesis of different archives, and makes it possible to answer key
42
43 461 questions that cannot be addressed using modern-day data: was Hainan's fauna
44
45 462 different in the past, and what happened to this fauna between past and present?
46
47 463 Long-term archives reveal that rather than having been a naturally depauperate
48
49 464 system or having lost biodiversity in a punctuated ancient or recent event,
50
51 465 Hainan has experienced protracted and ongoing human-caused depletion of its
52
53 466 mammal fauna from prehistory to the present, with its largest-bodied species
54
55 467 lost first and followed by progressive loss of smaller-bodied species. This pattern
56
57
58
59
60

1
2
3 468 is similar to the staggered extinction dynamics seen in several continental
4
5 469 mammal faunas across the Holocene-historical period [56,57]. These findings
6
7
8 470 reveal that Hainan is now experiencing “empty forest syndrome” [58], and with
9
10 471 the Hainan gibbon “merely” the latest of Hainan’s mammals to be sliding towards
11
12 472 extinction. They also raise key questions for future investigation: why has
13
14 473 Hainan’s mammal fauna been so vulnerable to extinction, given that its forest
15
16 474 cover remained fairly extensive until the twentieth century [13,59]; how did
17
18 475 gibbons manage to survive; and what does this long-term perspective suggest
19
20
21 476 about the future of Hainan’s biodiversity?
22
23

24 477 This new baseline on past diversity and faunal turnover provides a practical
25
26 478 framework for conservation managers to understand the extent of human-
27
28 479 caused biodiversity loss on Hainan, and emphasizes the urgent need for active
29
30 480 regional conservation programmes for many more species. Our new model of
31
32 481 Hainan’s long-term extinction dynamics can be compared and contrasted with
33
34 482 data for other Asian regions, for example islands that have experienced either
35
36 483 survival or extinction of species formerly present on Hainan (e.g. tigers, clouded
37
38 484 leopards), to identify intrinsic or extrinsic correlates of species vulnerability and
39
40
41 485 resilience and make predictive hypotheses to inform conservation planning
42
43 486 [31,60,61]. Integrated faunal archives can also inform direct conservation
44
45 487 management, for example to set new restoration or rewilding targets (e.g.
46
47 488 reintroduction of extirpated species, such as Père David’s deer to Hainan’s
48
49 489 wetlands; management of disrupted forest regeneration processes requiring
50
51 490 mammalian dispersers), or to forecast potential faunal responses to future
52
53 491 environmental change scenarios and develop appropriate mitigation strategies
54
55 492 against ongoing biodiversity loss.
56
57
58
59
60

1
2
3 493 However, despite the invaluable new insights about the status of regional
4
5 494 Chinese biodiversity provided by these long-term perspectives, China's
6
7
8 495 environmental archives exhibit extensive problems with incompleteness,
9
10 496 representativeness and bias in the information they contain about past faunal
11
12 497 baselines, cautioning against their use at face value. Each archive we investigated
13
14 498 for Hainan contains only a small percentage of the mammal species occurring on
15
16 499 the island today (fossil record=31%, gazetteer record=16%), and only one
17
18 500 regionally extinct species (dhole) is definitely included in more than one archive
19
20 501 (electronic supplementary material, table S1), suggesting that many more
21
22 502 species, potentially including numerous regionally extinct species of unknown
23
24 503 identity, remain undocumented. The usefulness of the gazetteer record and
25
26 504 potentially also the indigenous knowledge record are limited further due to
27
28 505 problems with accurate species identification by untrained observers. We also
29
30 506 demonstrate that, in contrast to some other social-ecological systems [19,62],
31
32 507 Hainan's indigenous knowledge record is an extremely poor source of
33
34 508 information on past extinctions, with almost no local awareness of formerly
35
36 509 occurring species, and most responses likely "cued" from previous interview
37
38 510 questions about named animals (pangolin, bear, gibbon, wild pig).

39
40
41 511 Our analyses of mainland China's faunal record demonstrate similar
42
43 512 incompleteness and bias, with only 22% of extant Chinese mammal species
44
45 513 represented in Holocene sites, and a strong effect of body mass on likelihood of
46
47 514 species representation. Similar biases are observed in many Quaternary and
48
49 515 older faunal assemblages, and likely reflect multiple burial and post-burial
50
51 516 processes including preferential past human hunting of larger-bodied animals,
52
53 517 greater survival of robust skeletal elements, biased excavation procedures,
54
55
56
57
58
59
60

1
2
3 518 and/or limited osteological information for species identification of many small-
4
5 519 bodied mammals [44,63], with few studies available on Chinese Quaternary small
6
7
8 520 mammal assemblages [64]. Although it is not possible to quantify how these
9
10 521 different processes have biased data for our regional study system, it is therefore
11
12 522 likely that Hainan's large-bodied Holocene mammal fauna is better understood
13
14
15 523 than its small-bodied fauna, with potential faunal turnover in Chinese small
16
17 524 mammal assemblages more challenging to identify. Distribution of Holocene
18
19 525 sites across China is also spatially uneven, and with very different representation
20
21 526 of different ecoregions. Far more excavations have been conducted in regions
22
23 527 with higher historical human populations (e.g. Yellow River and Yangtze plains)
24
25 528 [18], constituting an important target for archaeologists but not representative
26
27 529 of past human-environmental interactions and impacts across China as a whole,
28
29 530 and providing very different power to understand past environments and
30
31 531 biodiversity change in different landscapes, notably high-elevation Asian
32
33 532 ecosystems facing increased climate change risks today.

34
35
36
37
38 533 Our analyses of the quality of China's Holocene faunal record provide a new
39
40 534 baseline for assessing the insights that historical data can provide for
41
42 535 conservation, as well as the challenges that necessitate caution and care in
43
44 536 interpreting these data, which prevent long-term archives from ultimately being
45
46 537 able to answer many questions of importance to conservation biologists.
47
48 538 Establishing a clearer understanding of patterns of incompleteness and bias in
49
50 539 the faunal record can help identify future research priorities, including increased
51
52 540 sampling in understudied ecoregions, or extrapolations to estimate numbers of
53
54 541 regionally extinct species that remain unidentified [65]. Unfortunately, other
55
56 542 biodiversity hotspots do not have the range and resolution of long-term archives
57
58
59
60

1
2
3 543 available for China, so any region-specific conservation insights from the past
4
5 544 will be even more limited for these systems, especially when coupled with
6
7
8 545 problems of specimen preservation exacerbated in tropical environments such
9
10 546 as thermal degradation of ancient biomolecules [24]. Long-term environmental
11
12 547 records provide windows into the past that are essential for understanding
13
14 548 environmental baselines and biodiversity change, and enable development of
15
16 549 more inclusive decision-making frameworks, but incorporating these records
17
18 550 into conservation planning requires careful and nuanced interpretation.
19
20
21

22 551

23
24 552 **Data accessibility.** The datasets supporting this paper are available in the
25
26 553 supplementary materials.

27
28 554 **Authors' contributions.** S.T.T. designed research; C.W., J.J.C., K.H. and S.T.T. coordinated
29
30 555 data collection; C.W., J.H., J.B., C.D., M.A.H., K.H. and S.T.T. interpreted and analysed data;
31
32 556 and S.T.T. wrote the paper with support from other authors.

33
34 557 **Competing interests.** We have no competing interests.

35
36 558 **Funding.** Funding was provided by a Royal Society University Research Fellowship to
37
38 559 S.T.T. (UF130573) and the Arcus Foundation.

39
40 560 **Acknowledgements.** We thank the indigenous communities of Hainan for sharing their
41
42 561 indigenous knowledge. Data collection was supported by Michelle Wong, Hongyu Fu,
43
44 562 Qiaomei Li, Yuegui Lin, Cong Zhou, Hao Zhang, Jinghua Zhang, Helen Nash, Peng Zhang,
45
46 563 the management offices and reserve staff of Hainan's protected areas, and the Hainan
47
48 564 Provincial Forestry Department. We thank Hannah O'Regan and Claire Risley for advice
49
50 565 on analysis.
51

52
53 566

54
55 567 **REFERENCES**
56
57
58
59
60

- 1
2
3 568 1. Segan DB, Bottrill MC, Baxter PWJ, Possingham HP. 2010 Using conservation
4
5 569 evidence to guide management. *Cons. Biol.* **25**, 200-202.
- 6
7
8 570 2. Collen B, Ram M, Zamin T, McRae L. 2008 The tropical biodiversity data gap:
9
10 571 addressing disparity in global monitoring. *Trop. Conserv. Sci.* **1**, 75-88.
- 11
12 572 3. Boakes EH, McGowan PJK, Fuller RA, Ding C, Clark NE, O'Connor K, Mace GM.
13
14 573 2010 Distorted views of biodiversity: spatial and temporal bias in species
15
16 574 occurrence data. *PLoS Biol.* **8**, e1000385.
- 17
18 575 4. Schipper J, *et al.* 2008 The status of the world's land and marine mammals:
19
20 576 diversity, threat, and knowledge. *Science* **322**, 225-230.
- 21
22 577 5. Sodhi NS, Posa MRC, Lee TM, Bickford D, Koh LP, Brook BW. 2010 The state
23
24 578 and conservation of southeast Asian biodiversity. *Biodiv. Cons.* **19**, 317-328.
- 25
26 579 6. Willis KJ, Araújo MB, Bennett KD, Figueroa-Rangel B, Froyd CA, Myers N.
27
28 580 2007 How can a knowledge of the past help to conserve the future?
29
30 581 Biodiversity conservation and the relevance of long-term ecological studies.
31
32 582 *Phil. Trans. Roy. Soc. B* **362**, 175-186.
- 33
34 583 7. Davies AL, Colombo S, Hanley N. 2014 Improving the application of long-
35
36 584 term ecology in conservation and land management. *J. Appl. Ecol.* **51**, 63-70.
- 37
38 585 8. Dietl GP, Kidwell SM, Brenner M, Burney DA, Flessa KW, Jackson ST, Kock PL.
39
40 586 2015 Conservation paleobiology: leveraging knowledge of the past to inform
41
42 587 conservation and restoration. *Annu. Rev. Earth Planet Sci.* **43**, 79-103.
- 43
44 588 9. Barnosky AD, *et al.* 2017 Merging paleobiology with conservation biology to
45
46 589 guide the future of terrestrial ecosystems. *Science* **355**, eaah4787.
- 47
48 590 10. Balmford A. 1996 Extinction filters and current resilience: the significance
49
50 591 for past selection pressures for conservation biology. *Trends Ecol. Evol.* **11**,
51
52 592 193-196.

- 1
2
3 593 11. Louys J, Curnoe D, Tong H. 2007 Characteristics of Pleistocene megafauna
4
5 594 extinctions in southeast Asia. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* **243**,
6
7 595 152-173.
8
9
10 596 12. Smith AT, Xie Y (Eds.). 2013 *Mammals of China*. Princeton and Oxford:
11
12 597 Princeton University Press.
13
14 598 13. Marks RB. 2017 *China: an environmental history*. Second edition. Lanham,
15
16 599 MD: Rowman & Littlefield.
17
18
19 600 14. Shapiro J. 2001 *Mao's war against nature: politics and the environment in*
20
21 601 *revolutionary China*. Cambridge, UK: Cambridge University Press.
22
23
24 602 15. Dudgeon D. 2010 Requiem for a river: extinctions, climate change and the
25
26 603 last of the Yangtze. *Aquatic Conserv. Mar. Freshw. Ecosyst.* **20**, 127-131.
27
28
29 604 16. Turvey ST, Crees JJ, Di Fonzo MMI. 2015 Historical data as a baseline for
30
31 605 conservation: reconstructing long-term faunal extinction dynamics in Late
32
33 606 Imperial-modern China. *Proc. R. Soc. B* **282**, 20151299.
34
35
36 607 17. Turvey ST, Crees JJ, Li Z, Bielby J, Yuan J. 2017 Long-term archives reveal
37
38 608 shifting extinction selectivity in China's postglacial mammal fauna. *Proc. R.*
39
40 609 *Soc. B* **284**, 20171979.
41
42
43 610 18. Liu L, Chen X. 2012 *The archaeology of China: from the Late Paleolithic to the*
44
45 611 *early Bronze Age*. Cambridge, UK: Cambridge University Press.
46
47
48 612 19. Turvey ST, Bryant JV, McClune KA. 2018 Differential loss of components of
49
50 613 traditional ecological knowledge following a primate extinction event. *R. Soc.*
51
52 614 *Open Sci.* **5**, 172352.
53
54
55 615 20. Turvey ST, Bruun K, Ortiz A, Hansford J, Hu S, Ding Y, Zhang T, Chatterjee HJ.
56
57 616 2018 New genus of extinct Holocene gibbon associated with humans in
58
59 617 Imperial China. *Science* **360**, 1346-1349.
60

- 1
2
3 618 21. Han H, *et al.* 2019. Diet evolution and habitat contraction of giant pandas via
4
5 619 stable isotope analysis. *Curr. Biol.* **29**, 664-669.
6
7
8 620 22. Zhao H, Wang L, Yuan J. 2007 Origin and time of Qiongzhou Strait. *Mar. Geol.*
9
10 621 *Quat. Geol.* **27**, 33-40.
11
12 622 23. Turvey ST, Traylor-Holzer K, Wong MHG, Bryant JV, Zeng X, Hong X, Long Y
13
14 623 (Eds). 2015 *International conservation planning workshop for the Hainan*
15
16 624 *gibbon: final report*. London: Zoological Society of London / Apple Valley,
17
18 625 MN: IUCN SSC Conservation Breeding Specialist Group.
19
20 626 24. Turvey ST, Barnes I, Marr M, Brace S. 2017 Imperial trophy or island relict? A
21
22 627 new extinction paradigm for Père David's deer: a Chinese conservation icon.
23
24 628 *R. Soc. Open Sci.* **4**, 171096.
25
26 629 25. Swinhoe R. 1870 On the mammals of Hainan. *Proc. Zool. Soc. Lond.* **1870**,
27
28 630 224-239.
29
30 631 26. Yan JA. 2006 Paleontology and ecologic environmental evolution of the
31
32 632 Quaternary in Hainan Island. *J. Palaeogeogr.* **8**, 103-115.
33
34 633 27. Hao S, Huang W. 1998 *Sanya Luobidong cave site*. Haikou: Nanfang
35
36 634 Publishing House.
37
38 635 28. Hong S. 2003 *Hainan difangzhi congkan*. Haikou: Hainan Publishing House.
39
40 636 29. Zhang Z, Liu R. 1991 The Holocene along the coast of Hainan Island, China.
41
42 637 *Chinese Geogr. Sci.* **1**, 188-196.
43
44 638 30. Roberts N. 1998 *The Holocene: an environmental history*. Oxford: Blackwell.
45
46 639 31. Wilkinson DM, O'Regan HJ. 2003 Modelling differential extinctions to
47
48 640 understand big cat distribution on Indonesian islands. *Global Ecol. Biogeogr.*
49
50 641 **12**, 519-524.
51
52
53
54
55
56
57
58
59
60

- 1
2
3 642 32. Leonard SA, Risley CL, Turvey ST. 2013 Could brown bears (*Ursus arctos*)
4
5 643 have survived in Ireland during the Last Glacial Maximum? *Biol. Lett.* **9**,
6
7 644 20130281.
8
9
10 645 33. Cardillo M, Mace GM, Jones KE, Bielby J, Bininda-Emonds ORP, Sechrest W,
11
12 646 Orme CDL, Purvis A. 2005 Multiple causes of high extinction risk in large
13
14 647 mammal species. *Science* **309**, 1239-1241.
15
16
17 648 34. Lacy RC, Pollak JP. 2014 *Vortex: A Stochastic Simulation of the Extinction*
18
19 649 *Process. Version 10.0*. Brookfield, IL: Chicago Zoological Society.
20
21
22 650 35. Solow AR. 2005 Inferring extinction from a sighting record. *Math. Biosci.*
23
24 651 **195**, 47-55.
25
26
27 652 36. Clements C. 2013 *sExtinct*. R package version 1.0.1.
28
29 653 37. R Development Core Team. 2015 *R: a language and environment for*
30
31 654 *statistical computing*. Vienna: R Foundation for Statistical Computing.
32
33
34 655 38. Nash HC, Wong MHG, Turvey ST. 2016 Determining status and threats of the
35
36 656 Critically Endangered Chinese pangolin (*Manis pentadactyla*) in Hainan,
37
38 657 China, using local ecological knowledge. *Biol. Cons.* **196**, 189-195.
39
40
41 658 39. Turvey ST, *et al.* 2017 How many remnant gibbon populations are left on
42
43 659 Hainan? Testing the use of local ecological knowledge to detect cryptic
44
45 660 threatened primates. *Am. J. Primatol.* **79**, e22593.
46
47
48 661 40. Lau MW, Fellowes JR, Chan BPL. 2010 Carnivores (Mammalia: Carnivora) in
49
50 662 South China: a status review with notes on the commercial trade. *Mamm.*
51
52 663 *Rev.* **40**, 247-292.
53
54
55 664 41. Turvey ST, *et al.* 2015 Interview-based sighting histories can inform regional
56
57 665 conservation prioritization for highly threatened cryptic species. *J. Appl. Ecol.*
58
59 666 **52**, 422-433.
60

- 1
2
3 667 42. Payton ME, Greenstone MH, Schrenker N. 2003 Overlapping confidence
4
5 668 intervals or standard error intervals: what do they mean in terms of
6
7
8 669 statistical significance? *J. Insect Sci.* **3**, 34.
9
10 670 43. Valentine JW, Jablonski D, Kidwell S, Roy K. 2006 Assessing the fidelity of the
11
12 671 fossil record by using marine bivalves. *Proc. Natl Acad. Sci. USA* **103**, 6599-
13
14 672 6604.
15
16
17 673 44. Turvey ST, Blackburn TM. 2011 Determinants of species abundance in the
18
19 674 Quaternary vertebrate fossil record. *Paleobiol.* **37**, 537-546.
20
21
22 675 45. Orme D. 2013 *Caper: comparative analyses of phylogenetics and evolution in*
23
24 676 *R*. R package version 0.5.2.
25
26
27 677 46. Bininda-Emonds ORP, Cardillo M, Jones KE, MacPhee RDE, Beck RMD,
28
29 678 Grenyer R, Price SA, Vos RA, Gittleman JL, Purvis A. 2007 The delayed rise of
30
31 679 present-day mammals. *Nature* **446**, 507-512.
32
33
34 680 47. ESRI. 2018 *ArcGIS Pro: Version 2.3.0*. Redlands, CA: Environmental Systems
35
36 681 Research Institute.
37
38
39 682 48. Olson DM, *et al.* 2001 Terrestrial ecoregions of the world: a new map of life
40
41 683 on Earth. *BioScience* **51**, 933-938.
42
43
44 684 49. Lu L, *et al.* 2012 Small mammal investigation in spotted fever focus with
45
46 685 DNA-barcoding and taxonomic implications on rodents species from Hainan
47
48 686 of China. *PLoS ONE* **7**, e43479.
49
50
51 687 50. Boakes EH, Rout TM, Collen B. 2015 Inferring species extinctions: the use of
52
53 688 sighting records. *Methods Ecol. Evol.* **6**, 678-687.
54
55
56 689 51. Clements CF, Collen B, Blackburn TM, Petchey OL. 2014 Recent
57
58 690 environmental change may affect accurate inference of extinction. *Conserv.*
59
60 691 *Biol.* **28**, 971-981.

- 1
2
3 692 52. Zeng Z, Song Y, Li J, Teng L, Zhang Q, Guo F. 2005 Distribution, status and
4
5 693 conservation of Hainan Eld's deer (*Cervus eldi hainanus*) in China. *Folia Zool.*
6
7 694 **54**, 249-257.
- 8
9
10 695 53. Guthrie RD. 2004 Radiocarbon evidence of mid-Holocene mammoths
11
12 696 stranded on an Alaskan Bering Sea island. *Nature* **429**, 746-749.
- 13
14
15 697 54. Hao S, Wang D. 2003 Retrospection and prospection of archaeology in
16
17 698 Hainan. *Archaeology (China)* **4**, 291-299.
- 18
19
20 699 55. Xu F, Hu B, Dou Y, Song Z, Liu X, Yuan S, Sun Z, Li A, Yin X. 2018 Prehistoric
21
22 700 heavy metal pollution on the continental shelf off Hainan Island, South China
23
24 701 Sea: from natural to anthropogenic impacts around 4.0 kyr BP. *Holocene* **28**,
25
26 702 455-463.
- 27
28
29 703 56. Woinarski JCZ, Burbidge AA, Harrison PL. 2015 Ongoing unraveling of a
30
31 704 continental fauna: decline and extinction of Australian mammals since
32
33 705 European settlement. *Proc. Natl Acad. Sci. USA* **112**, 4531-4540.
- 34
35
36 706 57. Crees JJ, Carbone C, Sommer RS, Benecke N, Turvey ST. 2016 Millennial-scale
37
38 707 faunal record reveals differential resilience of European large mammals to
39
40 708 human impacts across the Holocene. *Proc. R. Soc. B* **283**, 20152152.
- 41
42
43 709 58. Corlett RT. 2007 The impact of hunting on the mammalian fauna of tropical
44
45 710 Asian forests. *Biotropica* **39**, 292-303.
- 46
47
48 711 59. Zhang Y, Uusivuori J, Kuuluvainen J. 2000 Econometric analysis of the causes
49
50 712 of forest land use changes in Hainan, China. *Can. J. For. Res.* **30**, 1913-1921.
- 51
52
53 713 60. Diamond JM. 1989 Quaternary megafaunal extinctions: variations on a
54
55 714 Theme by Paganini. *J. Archaeol. Sci.* **16**, 167-175.
- 56
57
58
59
60

- 1
2
3 715 61. Chiang P, Pei KJ, Vaughan MR, Li C, Chen M, Liu J, Lin C, Lin L, Lai Y. 2015 Is
4
5 716 the clouded leopard *Neofelis nebulosa* extinct in Taiwan, and could it be
6
7 717 reintroduced? An assessment of prey and habitat. *Oryx* **49**, 261-269.
8
9
10 718 62. Wehi PM, Cox MP, Roa T, Whaanga H. 2018 Human perceptions of
11
12 719 megafaunal extinction events revealed by linguistic analysis of indigenous
13
14 720 oral traditions. *Hum. Ecol.* **46**, 461-470.
15
16
17 721 63. Behrensmeyer AK, Fürsich FT, Gastaldo RA, Kidwell SM, Kosnik MA,
18
19 722 Kowalewski M, Plotnick RE, Rogers RR, Alroy J. 2005 Are the most durable
20
21 723 shelly taxa also the most common in the marine fossil record? *Paleobiol.* **31**,
22
23 724 607-623.
24
25
26 725 64. Jin JJH, Jablonski NG, Flynn LJ, Chaplin G, Ji X, Li Z, Shi X, Li G. 2012
27
28 726 Micromammals from an early Holocene archaeological site in southwest
29
30 727 China: paleoenvironmental and taphonomic perspectives. *Quat. Int.* **281**, 58-
31
32 728 65.
33
34
35 729 65. Curnutt J, Pimm S. 2001 How many bird species in Hawai'i and the central
36
37 730 Pacific before first contact? *Stud. Avian Biol.* **22**, 15-30.
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 731 **FIGURE LEGENDS**
4

5 732
6

7
8 733 **Figure 1.** (a) Distribution of 253 Holocene zooarchaeological and
9
10 734 palaeontological sites across mainland China with wild mammal records. (b) Map
11
12 735 of Hainan, showing location of Luobidong Cave (star) and villages where
13
14 736 interviews were conducted (circles). B, Bawangling; D, Diaoluoshan; JN,
15
16 737 Jianfengling; JX, Jiaxi; L, Limushan; W, Wuzhishan; Y, Yinggeling.
17
18
19

20 738

21
22 739 **Figure 2.** Slopes and 83% CIs of local respondent encounter-history data for
23
24 740 seven Hainanese mammal species. Left to right: wild pig, rhesus macaque,
25
26 741 Hainan gibbon, Asian black bear, clouded leopard, sambar deer, Chinese
27
28 742 pangolin.
29

30 743
31

32
33 744 **Figure 3.** Box plot of body masses for mainland Chinese mammal species that
34
35 745 are present or absent in the Holocene zooarchaeological and palaeontological
36
37 746 record.
38

39 747
40

41
42
43 748 **Figure 4.** Integrated model of long-term mammalian biodiversity change on
44
45 749 Hainan based on synthesis of multiple environmental archives, showing
46
47 750 progressive depletion of regional mammal fauna across the Holocene to the
48
49 751 present as evidenced by different species-specific data sources on temporal
50
51 752 patterns of population persistence.
52
53
54
55
56
57
58
59
60

1
2
3 **753 Table 1.** Summary of respondent encounter history data for seven Hainanese
4
5 **754** mammals, and species last-encounter history regression slopes with 83%
6
7 **755** confidence interval upper and lower bounds (df for all regressions=24).
8
9
10 **756**

species	no. of encounter records	mean last-encounter date	% encounters in past 10 yrs (2006-2015)	slope	SD	lower bound (8.5%)	upper bound (91.5%)
wild pig	549	2012	59.6	0.158	0.035	0.113	0.209
rhesus macaque	432	2010	54.9	0.125	0.030	0.086	0.168
Hainan gibbon	187	1983	21.9	0.065	0.027	0.028	0.103
Asian black bear	193	1987	13.0	-0.015	0.026	-0.051	0.020
clouded leopard	125	1980	6.4	-0.023	0.027	-0.061	0.014
sambar deer	359	1993	15.3	-0.025	0.017	-0.049	-0.001
Chinese pangolin	495	1993	11.9	-0.031	0.021	-0.061	-0.002

23 **757**

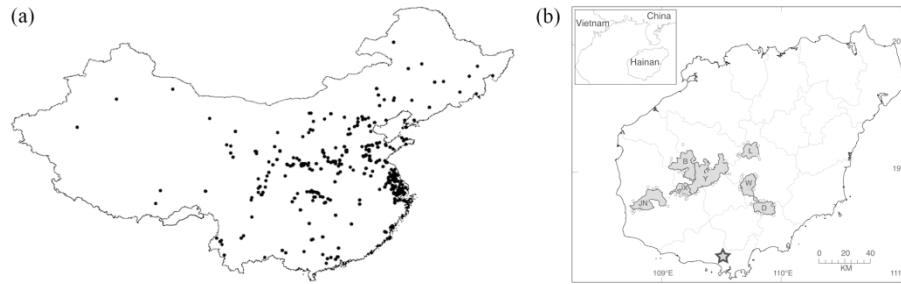


Figure 1. (a) Distribution of 253 Holocene zooarchaeological and palaeontological sites across mainland China with wild mammal records. (b) Map of Hainan, showing location of Luobidong Cave (star) and villages where interviews were conducted (circles). B, Bawangling; D, Diaoluoshan; JN, Jianfengling; JX, Jiayi; L, Limushan; W, Wuzhishan; Y, Yinggeling.

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

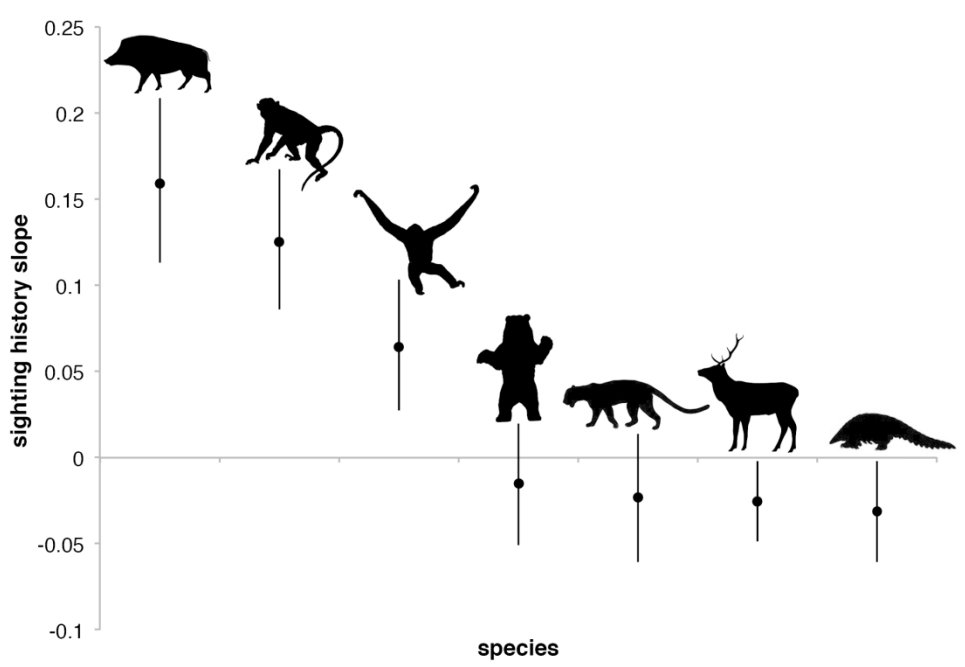


Figure 2. Slopes and 83% CIs of local respondent encounter-history data for seven Hainanese mammal species. Left to right: wild pig, rhesus macaque, Hainan gibbon, Asian black bear, clouded leopard, sambar deer, Chinese pangolin.

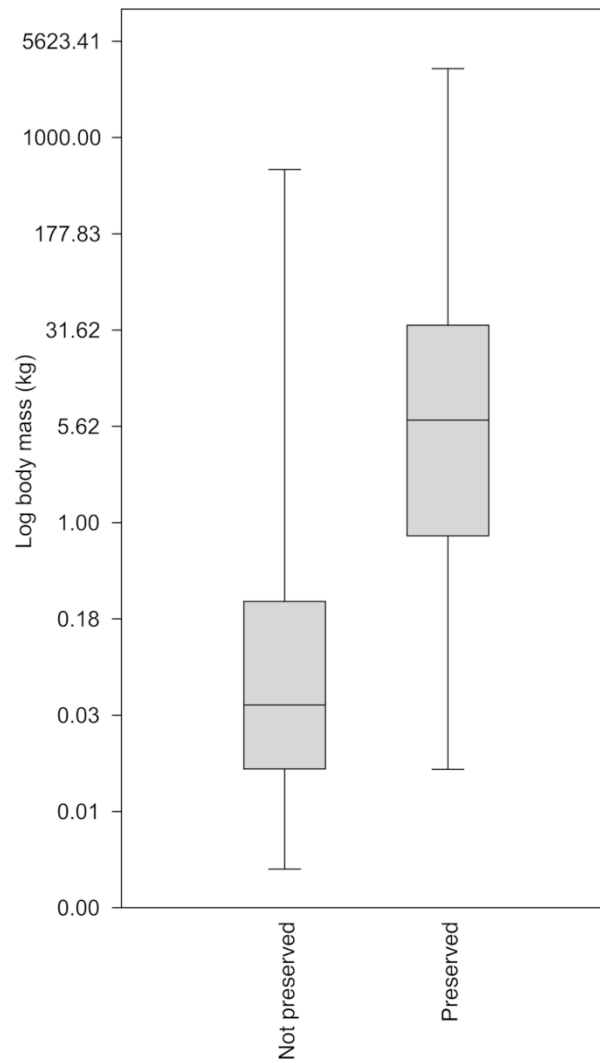


Figure 3. Box plot of body masses for mainland Chinese mammal species that are present or absent in the Holocene zooarchaeological and palaeontological record.

100x161mm (300 x 300 DPI)

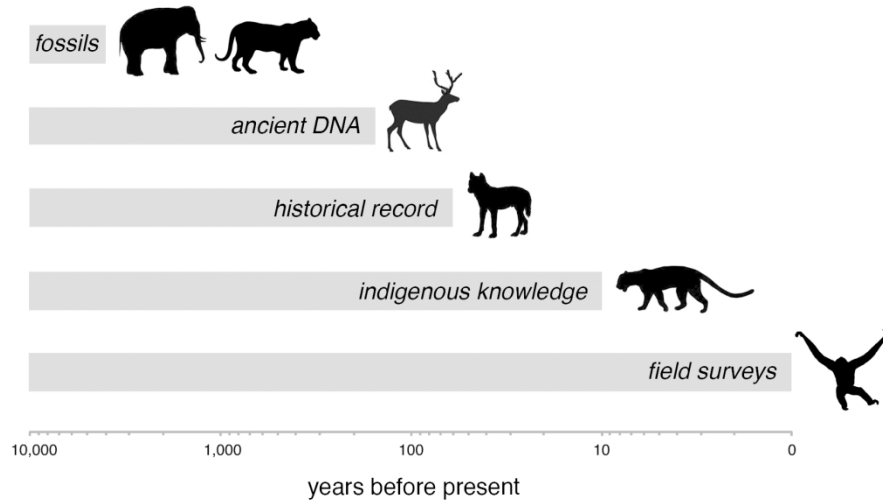


Figure 4. Integrated model of long-term mammalian biodiversity change on Hainan based on synthesis of multiple environmental archives, showing progressive depletion of regional mammal fauna across the Holocene to the present as evidenced by different species-specific data sources on temporal patterns of population persistence.

167x102mm (300 x 300 DPI)