

1 **Drivers of taxonomic bias in ecology and evolution: insights from ethologists and behavioural**
2 **ecologists**

3 Pietro Pollo*, Michael M. Kasumovic

4 Evolution & Ecology Research Centre, School of Biological, Earth & Environmental Sciences,
5 University of New South Wales, Kensington, NSW, 2052, Australia

6 * Corresponding author: pietro_pollo@hotmail.com

7

8 **ORCID**

9 Pietro Pollo: <https://orcid.org/0000-0001-6555-5400>

10 Michael M. Kasumovic: <https://orcid.org/0000-0003-0158-5517>

11

12 **Abstract**

13 Taxonomic bias (i.e. the disproportionate attention given to some taxa relative to their diversity)
14 remains a major barrier to achieving generality in ecology and evolution, yet its underlying causes
15 are poorly understood. We propose a framework explaining taxonomic bias along three major axes,
16 supported by evidence from a survey of 868 researchers' experiences. First, rational
17 considerations, such as logistical ease and societal relevance, were associated with the choice of
18 research organisms within major animal groups but rarely across them. Second, emotional factors,
19 including taxonomic affinities, closely mirrored taxonomic patterns in the literature. Third,
20 contextual factors, like the prominence of certain organisms within peer networks or early-career
21 exposure to specific taxa, were also associated with which taxa are chosen as study systems. Based
22 on these findings, we suggest actions to mitigate taxonomic bias, including promoting (i) outreach
23 initiatives featuring neglected taxa, (ii) taxonomically equitable education, and (iii) taxonomically
24 diverse research experiences.

25

26 **Key-words:** charismatic species, knowledge imbalance, model organism, model system, research
27 bias, taxonomic chauvinism, taxonomic hyperfocus.

28

29 **Introduction**

30 One of the main goals of ecology and evolution is to comprehend biological processes and patterns
31 across Earth's vast diversity of life. This goal is hindered when certain taxa receive
32 disproportionately more attention than their diversity warrants, a pattern known as taxonomic bias
33 or taxonomic chauvinism (Shine and Bonnet 2000; Bonnet et al. 2002). Taxonomic bias leads to
34 the accumulation of knowledge on few organisms at the cost of the perpetuation of ignorance on
35 all others, ultimately precluding theoretical generality (Bonnet et al. 2002; Dochtermann et al.
36 2026) and innovation (Ng et al. 2021; Broeckhoven and du Plessis 2022; Penick et al. 2022; Stuart-
37 Fox et al. 2023; Snell-Rood and Smirnoff 2025). Unfortunately, taxonomic bias is widespread in
38 ecology and evolution (Bonnet et al. 2002; Clark and May 2002; Seddon et al. 2005; Leather 2009;
39 Stahlschmidt 2011; Zuk et al. 2014; Rosenthal et al. 2017; Titley et al. 2017; Troudet et al. 2017;
40 Mammola et al. 2020*b*; Pollo et al. 2024, 2025*a*; Guénard et al. 2025; Mizuno et al. 2025; Pollo
41 and Kasumovic 2026). For instance, even though birds and mammals comprise less than 1% of
42 animal biodiversity (Zhang 2013), these animals dominate both conservation efforts (Seddon et al.
43 2005; Mammola et al. 2020*b*; Guénard et al. 2025) and scientific articles in most subpockets of
44 ecology and evolution (Bonnet et al. 2002; Clark and May 2002; Leather 2009; Stahlschmidt 2011;
45 Rosenthal et al. 2017; Titley et al. 2017; Pollo et al. 2024, 2025*a*; Mizuno et al. 2025; Pollo and
46 Kasumovic 2026). Taxonomic bias also extends to finer taxonomic scales, with a handful of genera
47 or species concentrating much of the research attention given to entire clades (Zuk et al. 2014).
48 Investigations of the literature on specific groups of organisms (e.g. amphibians: Silva et al. 2020;
49 birds: Fischer et al., 2025; Murray et al., 2015; Yarwood et al., 2019; mammals: dos Santos et al.,
50 2020; Tam et al., 2022; parasites: Simoncini et al. 2025; plants: Adamo et al. 2021; primates:
51 Ellison et al. 2021; Chen et al. 2023; see also Mammola et al. 2023) have shown that organisms'

52 traits (e.g. size and colour), range, and location can influence the amount of attention they receive,
53 shedding some light on the drivers of taxonomic bias. However, such literature assessments present
54 only indirect and taxon-specific evidence regarding the origins of taxonomic bias, highlighting the
55 absence of a general framework spanning multiple taxonomic levels and empirical support for it.
56 In other words, why ecologists and evolutionary biologists choose the organisms they study largely
57 remains a conundrum. Understanding this question requires understanding the scientists that
58 perform this research.

59 Scientists are often expected to be entirely rational in their research choices (Fig. 1) but, in
60 reality, their emotions and context can play major roles in such decisions (Damasio 1994). For
61 instance, researchers may be drawn to particular organisms out of fondness for them (Lorimer
62 2007). Opinions about organisms, such as vertebrate animals being more valuable than
63 invertebrate ones (Miralles et al. 2019; Possidónio et al. 2019), often arise from societal
64 perceptions that develop from a young age. For example, children asked to draw natural landscapes
65 are much more likely to depict vertebrates, particularly mammals and birds, than invertebrates
66 (Strommen 1995; Snaddon et al. 2008). This skewed perception is then reinforced by the media
67 (e.g. documentaries, Wei et al. 2024) and educational materials (e.g. textbooks, Gangwani &
68 Landin, 2018), helping to cement the idea that certain organisms are more important or interesting
69 than others well into adulthood. It is thus plausible that the image that biologists have of nature,
70 which is subjected to cultural factors and constructed before even joining academia, can influence
71 their research choices. In fact, feelings of admiration and fascination towards certain taxonomic
72 groups potentially entice early-career biologists to work with these groups or may even encourage
73 people to pursue a career in biology in the first place. By contrast, organisms that produce anxiety,
74 fear, or disgust (i.e. biophobias, Simaika and Samways 2010; Soga et al. 2023) may suffer from

75 disinterest and avoidance from the general public and, consequently, have a lower research effort
76 directed at them.

77 While some emotional links to taxa arise from society at large, others can be a product of
78 habits and perceptions restricted to smaller social spheres. For instance, having hobbies that
79 involve particular organisms (e.g. hunting or bird watching) may help construct positive feelings
80 towards these organisms (Casola et al. 2026) and therefore skew research effort in their favour.
81 Similarly, preferences for certain work-related aspects, such as a strong inclination toward
82 fieldwork (a sentiment shared by many biologists and reflected in the popularity of field courses;
83 Fleischner et al. 2017), can influence researchers' taxonomic focus. This is because some
84 organisms (e.g. many mammals and birds) can only be studied in the field, potentially attracting
85 scientists that enjoy spending their time outdoors. Furthermore, although researchers may have
86 predilections for certain organisms, their choice of study species can also be driven by other
87 personal motivations. For example, researchers may be primarily motivated to investigate a
88 particular topic (e.g. sexual selection), making the selection of a particular study organism less
89 important to them. Understanding researchers' preferences and motivations then becomes pivotal
90 to grasp drivers of taxonomic bias.

91 Science is built on prior research, so the popularity of an organism in an academic context
92 may reinforce and amplify its own popularity, generating a feedback loop. This process partly
93 stems from pragmatism: as more resources become available for an organism, further research on
94 it becomes easier. However, there is also a subjective element to this process, as scientists may
95 come to view certain organisms as "ideal" study systems because of their established reputation
96 rather than their intrinsic characteristics. This is essentially how so-called model organisms arise
97 (Leonelli and Ankeny 2012, 2013). While the intense use of such organisms have allowed

98 important insights into biological mechanisms (Fields and Johnston 2005), ecologists and
99 evolutionary biologists not necessarily interested in mechanistic questions have also frequently
100 used model organisms in their research, leading to an unnecessary accumulation of knowledge on
101 a few species (Zuk et al. 2014). Moreover, scientists' individual context may also influence the
102 choice of research organisms. Researchers who continue working with the same taxa (often
103 introduced by their supervisors early in their careers) enjoy a logistical advantage over those who
104 attempt to study unfamiliar taxa. At the same time, fear of failure may also play a role in
105 maintaining this focus, particularly as careers progress and the practical advantage of studying a
106 familiar taxon over a new one diminishes. Ultimately, this form of taxonomic inheritance can
107 perpetuate or even exacerbate existing taxonomic biases.

108 Here, we explored potential overarching drivers of taxonomic bias in ecology and evolution
109 by directly surveying researchers (specifically ethologists and behavioural ecologists) about their
110 experiences, preferences, and perceptions regarding different taxa. We hypothesise that multiple
111 rational, contextual, and emotional factors are associated with the selection of research organisms
112 (Fig. 1), leading to a variety of predictions (Table S1; preregistered in Pollo and Kasumovic 2025).
113 Based on our findings, we then suggest interventions to reduce taxonomic bias in the field of
114 ecology and evolution.

115

116 **Material and methods**

117 *Deviations from the preregistration*

118 Our methodology was described in our preregistration (Pollo and Kasumovic 2025). Although we
119 adhered to it as much as possible, we included several post-hoc analyses in our study. We also note
120 that part of our original plan was used to generate a separate study assessing existing patterns of

121 taxonomic bias in ecology and evolution (Pollo and Kasumovic 2026). Aside from this, two main
122 deviations from our pre-registration occurred. First, although some of our predictions were pre-
123 registered (Table S1), many relationships we tested were not explicitly made in our pre-
124 registration. Second, we initially planned to assess relationships between variables using a 7-point
125 Likert scale with polychoric multivariate analyses. However, because applying this method on
126 several pairs of variables from our dataset often violated statistical assumptions related to this
127 method, we opted to use Spearman's correlations instead.

128

129 *Survey with ethologists and behavioural ecologists*

130 PP and MK formulated a survey with three sections: (1) demographics, (2) preferences,
131 experiences, and feelings, and (3) perceptions regarding species used by participants for
132 behavioural research (Appendix 1). For example, we asked participants from which organisms
133 they collected behavioural data (questions 12 to 15, Appendix 1), as well as how they felt towards
134 distinct taxonomic groups on a spectrum ranging from negative (e.g. fear, anxiety) to positive (e.g.
135 cuteness, fascination) feelings, with ambivalent feelings in between (question 11, Appendix 1).
136 This survey was available online from 27 May 2025 to 21 August 2025 to any person that desired
137 to participate, was at least 18 years old, and had directly collected behavioural data from at least
138 one non-human organism that led to at least one research output of which they were an author (e.g.
139 conference poster or presentation, preprint, peer-reviewed publication, dissertation, thesis, book).
140 No financial incentives were provided to participants, and anonymity was maintained by collecting
141 only their survey responses. Ethical approval for this study was obtained from the University of
142 New South Wales' Human Research Ethics Committee (reference number iRECS8512).

143 PP recruited participants by sharing the link to the survey on the social media site *Bluesky*,
144 through email chains, and via a targeted email list. This email list was obtained by extracting the
145 emails associated with articles published in specific behaviour journals since 2015 and in certain
146 general ecology and evolution journals since 2020 (see Table S2). This resulted in a list containing
147 20,818 unique email addresses, albeit 2,879 were invalid (i.e. undeliverable) and another 708
148 generated automatic replies indicating that their users were unavailable (e.g. out of the office) at
149 the time of delivery. After accounting for these, 17,231 emails were considered to have most likely
150 reached their intended recipients.

151 The survey was completed by 946 individuals, representing approximately 5.49% of the
152 recruitment emails sent that likely reached recipients. However, an exact estimate of the
153 recruitment success for our survey could not be calculated as the identity of participants was not
154 collected and the total number of individuals that were reached and were eligible to participate in
155 the survey was unknown. For instance, emails associated with studies published in ecology and
156 evolution journals may have reached researchers that had never collected behavioural data.

157 PP carefully examined the responses obtained and excluded 77 of them from the dataset:
158 31 from participants that incorrectly answered the attention question in the survey (i.e. selected
159 other options than agree for the question “please select the option agree”); seven responses with
160 dubious veracity (a 99 years old participant, a participant that was 18 years old when they
161 completed their PhD, and five participants with editorial experience despite not having a PhD); 37
162 contradictory responses (nine participants that worked with a greater number of taxonomic groups
163 than of species, 28 participants that collected behavioural data of a species from a taxon they
164 reported having no experience with); and three visibly duplicated responses (identical age, gender,

165 country, year of PhD, species mentioned). As a result of these exclusions, 868 responses (91.75%
166 of all responses obtained) were used for most analyses in the present study.

167 Participants mentioned up to two species for which they collected behavioural data in the
168 survey (question 14, Appendix 1). PP only considered these specific responses as valid for analyses
169 involving them when they represented a genus or a species (common names were transformed to
170 latin names when possible). Consequently, PP excluded 86 answers out of the 1,666 obtained
171 (5.16%) that were too vague (e.g. “insect” instead of a specific insect species or genus), contained
172 multiple species, or were repeated by a participant (i.e. most and least recent species mentioned
173 were the same).

174

175 *Statistical analyses*

176 First, PP fitted two similar generalised linear models (GLMs): one with a negative binomial error
177 distribution and the other with a Poisson error distribution. The response variable in the first GLM
178 was the number of species that participants experienced, while the response variable in the second
179 GLM was the number of taxonomic groups experienced (out of 11 options). Participant age
180 (continuous), gender (men *vs.* women *vs.* other or unknown), editorial experience (binary: with *vs.*
181 without), preference for fieldwork, and motivation from taxa served as predictor variables (the last
182 two were treated as continuous variables even though participants provided responses in discrete
183 increments of 10, ranging from 0 to 100). Additionally, the first GLM included the number of taxa
184 experienced as a predictor variable, and the second GLM included the number of species
185 experienced as a predictor variable.

186 Second, PP fitted cumulative linked models (CLMs) with a logit link on participants’
187 feelings for each of the nine animal groups included in the survey (“other invertebrates” and “other

188 organisms” were not included in question 11, Appendix 1). In addition to the variables used in
189 GLMs aforementioned (i.e. age, gender, editorial experience, preference for fieldwork, motivation
190 from taxa), these CLMs also included experience with the taxon in question as a predictor variable
191 (binary: with *vs.* without). Furthermore, to compare fondness for distinct taxonomic groups across
192 participants, PP fitted a cumulative linked mixed model (CLMM) on all nine responses of how
193 participants felt towards distinct taxa together, using participant ID as random factor and
194 taxonomic group as a predictor variable in these CLMMs.

195 Third, PP fitted a separate GLM for each of the 11 taxonomic groups included in the survey
196 (amphibians, arachnids, birds, crustaceans, fish, insects, mammals, molluscs, other invertebrates,
197 other organisms, and reptiles), all using a binomial error distribution. The response variable in
198 each of these GLMs indicated whether a participant had experience with the taxon in question. All
199 of these GLMs included the same predictors aforementioned (age, gender, editorial experience,
200 preference for fieldwork, and motivation from taxa) and, except for “other invertebrates” and
201 “other organisms”, participants’ feelings for the taxon in question (factor with up to 7 levels).

202 Fourth, PP fitted a generalised linear mixed model (GLMM) using a binomial error
203 distribution on whether species mentioned by participants were recommended to them for research
204 purposes by their supervisor *vs.* others (i.e. other people or themselves). Predictor variables in this
205 GLMM included whether the species mentioned was the most or least recently experienced by
206 them, and when participants first worked with the species they mentioned (factor with three levels:
207 during/before their PhD, during postdoc, or after obtaining a permanent position). Participant ID
208 was then used as a random factor.

209 Fifth, PP fitted a distinct CLMM for each participants’ agreement (7-point Likert scale) to
210 a statement made in our survey related to the species they most recently experienced (questions

211 19-31, Appendix 1) as well as to the species they least recently experienced (only questions 27 and
212 28, Appendix 1). In these CLMMs, the taxonomic group of species mentioned by participants
213 served as the only predictor variable. The genus of the species mentioned was used as a random
214 factor in all CLMMs, while participant ID also served as a random factor for the CLMMs related
215 to questions 27 and 28 (as there were up to two responses from each participant).

216 Sixth, PP fitted another set of CLMMs, again on participants' agreement to several
217 statements, but particularly those involving easiness to work with the species mentioned (questions
218 19-23, Appendix 1). Two predictor variables were used in these CLMMs: (1) the number of
219 participants mentioning the genus that the species belonged to, and (2) participants' preference for
220 fieldwork. In addition, the genus of the species mentioned was used as a random factor in CLMMs.

221 In all models described above, continuous predictor variables were scaled by subtracting
222 the mean from each value and then dividing the result by the standard deviation. PP identified
223 predictor variables associated with response variables tested using model selection. He selected
224 models whose Akaike Information Criterion corrected for small sample sizes (AICc) was within
225 two units of the model with the lowest AICc. He then fitted simplified versions of these models
226 by including only predictor variables that appeared in all selected models, i.e. he assumed that only
227 these variables were associated with response variables. Lastly, PP used Spearman's correlations
228 to verify pairwise relationships between variables that were on a 7-point Likert scale. Unless
229 otherwise stated, means were presented with standard error.

230 PP conducted all analyses using R (R Core Team 2025) v. 4.5.1. GLMMs were fitted using
231 the package *glmmTMB* (Brooks, Mollie et al. 2017; McGillicuddy et al. 2025) v. 1.1.12 and
232 ordinal regressions (CLMs and CLMMs) were conducted using the package *ordinal* (Christensen
233 2023) v. 2023.12.4.1. PP verified GLMMs' assumptions using the package *DHARMA* (Hartig

234 2016) v. 0.4.7. and computed model predictions using the package *emmeans* (Lenth 2025) v.
235 1.11.2.8. PP used the package *MuMIn* (Bartoń 2023) v. 1.48.11 to obtain and compare all possible
236 candidate models from complex models.

237

238 **Results**

239 *Participant profile*

240 Participants in our survey ($N = 868$) were, on average, 42.89 ± 11.56 years old (mean \pm SD).
241 52.76% of them identified as men, 44.82% as women, 1.73% as another gender identity (e.g.
242 gender-fluid, non-binary), and 0.69% did not disclose their gender identity. We asked each
243 participant to name the country where they spent most of their life before and after reaching
244 adulthood (18 years old), to which we obtained 68 unique countries before age 18 and 59 after age
245 18 (Fig. S1), with 80.65% of participants selecting the same country for both periods. Most
246 participants (89.97%) held a PhD degree and more than a third of all participants (36.52%) had
247 editorial experience (question 8, Appendix 1).

248

249 *Preferences and overall experience collecting behavioural data*

250 On a scale from 0% to 100%, we asked participants to determine how they would split their time
251 collecting behavioural data between the field and the lab based purely on personal enjoyment
252 (hereby *preference for fieldwork*; question 9, Appendix 1), and how much their desire to work with
253 certain taxa drove their behavioural research interests compared with other motivations (hereby
254 *motivation from taxa*; question 10, Appendix 1). On average, preference for fieldwork was 59.27%
255 $\pm 1.02\%$ and motivation from taxa was $50.76\% \pm 0.96\%$, but responses to these questions varied
256 across participants (Fig. S2A, B).

257 Participants experienced (i.e. directly collected behavioural data that led to at least one
258 research output in which they were an author; question 13, Appendix 1), on average, 9.48 ± 0.45
259 species (Fig. S2C). However, this estimate would be higher if we had not (unintentionally) set a
260 maximum of 80 in the survey platform for this question (16 participants answered this maximum
261 number; Fig. S2C). From the 11 taxonomic groups listed in our survey as options (amphibians,
262 arachnids, birds, crustaceans, fish, insects, mammals, molluscs, other invertebrates, other
263 organisms, and reptiles), participants experienced, on average, 2.1 ± 0.04 of them (Fig. S2D).
264 Participants that experienced more species, on average, also experienced more taxonomic groups
265 (Fig. 2; Table S3). Age and motivation from taxa were associated with both the number of species
266 and the number of taxonomic groups experienced, while gender and editorial experience were only
267 related to the number of species (Fig. 2A, C; Table S3, S4). By contrast, field preference was
268 unrelated to the number of species or taxonomic groups experienced (Fig. 2; Table S3, S4).

269

270 *Feelings towards animal groups*

271 All animal groups mostly elicited positive feelings from participants, although some more than
272 others (Fig. S3). On average, participants declared more favourable perceptions of taxa they
273 experienced than of other taxa ($\beta = 2.52 \pm 0.07$, $z = 33.91$, $p < 0.001$; Fig. 3A; Fig. S3B, C; Table
274 S5). Multiple other factors were also associated with how participants felt about certain taxa (Fig.
275 3A; Table S5).

276

277 *Experiences with taxonomic groups*

278 Birds and mammals were the taxa that participants most frequently experienced (each by 42.58%
279 of participants), followed by insects (35.21%) and fish (24.05%; Fig. S4). Unsurprisingly,

280 participants that experienced more taxonomic groups were more likely to experience any given
281 taxon (Fig. 3B; Table S6). We also found associations between experience with certain taxa and
282 other factors, especially preference for fieldwork (Fig. 3B; Table S6).

283

284 *Experiences with specific species*

285 We asked participants to name the species they most and least recently experienced (question 14,
286 Appendix 1), for which we obtained 839 and 741 valid responses, respectively. We then asked
287 participants several questions related to the species they mentioned, beginning with when they first
288 worked with these species (question 17, Appendix 1) and who advised them to do so (question 18,
289 Appendix 1). Regardless if the species named was the most or least recently experienced by the
290 participant, it was suggested by supervisors as a research system in most cases, except when
291 participants already held a permanent position (Fig. 4; Table S7).

292 The taxonomic group of species mentioned by participants in our survey (Fig. S5A)
293 followed a similar pattern to the one shown for participants' overall experience (Fig. S4). However,
294 we noted that many researchers mentioned the same species and genera. For example, *Drosophila*
295 *melanogaster*, *Rattus norvegicus*, and *Parus major* were mentioned by 22, 21, and 17 participants,
296 respectively. In fact, only 65.29% and 49.68% of all responses (for organisms experienced by
297 participants) represented unique species and genera, respectively. The most popular taxonomic
298 groups, especially mammals, were the ones with the relatively fewest unique species (Fig. S5B)
299 and genera (Fig. S5C).

300 We additionally asked participants to rate their agreement with 13 statements related to the
301 organism they most recently experienced on a 7-point Likert scale (from “completely disagree” to
302 “completely agree”; two particular statements were also rated for the organism least recently

303 experienced; questions 19-31, Appendix 1). The first five statements tackled the ease of accessing
304 and working with the organisms mentioned, in which we found distinct average scores across
305 taxonomic groups (Fig. 5). Furthermore, after controlling for the genus and the taxonomic group
306 of the species mentioned by participants, a stronger preference for fieldwork was associated with
307 perceiving the species experienced as easier to observe in the field (though not necessarily easier
308 to find there), and harder to buy, rear, or observe in the laboratory (Tables S8, S9). Lastly, within
309 taxonomic groups, the number of times a genus was mentioned by distinct participants was
310 positively related to all aspects aforementioned regarding how easy they are to work with (Tables
311 S8, S9).

312 No universal taxonomic patterns emerged from participants' agreement with other
313 statements regarding the organisms they most and least recently worked with (Fig. 6). We also
314 found that participants' perception that the species they most recently experienced is a model
315 organism was positively related to multiple other attitudes about the same species with varying
316 magnitudes: (1) strongly related to the impression that the behavioural literature on the species is
317 extensive ($r_s = 0.643$); (2) moderately related to the impression that the species is easy to buy (r_s
318 = 0.456), rear ($r_s = 0.404$), and observe in the laboratory ($r_s = 0.401$); (3) weakly related to the
319 impression that the species is easy to find ($r_s = 0.184$) and observe in the field ($r_s = 0.119$).
320 Furthermore, we also noted that participants' agreement that the research they conducted (with the
321 most recently experienced species mentioned) was applied was moderately linked to their
322 perception that the species is economically important ($r_s = 0.331$) and weakly associated with their
323 perception that the species is endangered or threatened in the wild ($r_s = 0.241$).

324

325 **Discussion**

326 In the present study, we hypothesised that several factors (stemming from reason, context, or
327 emotion, Fig. 1) are related to the selection of research organisms and thus drivers of taxonomic
328 bias. The survey we conducted with ethologists and behavioural ecologists provides evidence in
329 favour of this framework, although, in some cases, only at certain taxonomic levels. Below we
330 discuss each of our main findings in detail, but see Appendix 2 for the discussion of our secondary
331 results.

332 The taxonomic experiences of participants in our survey mirrored patterns of taxonomic
333 bias present in the animal behaviour literature (Rosenthal et al. 2017; Pollo and Kasumovic 2026).
334 For instance, birds and mammals were the taxonomic groups participants in our survey were most
335 likely to have worked with (Fig. S4; Fig. S5A). Conversely, with the exception of insects,
336 participants rarely used invertebrates as research organisms (Fig. S4; Fig. S5A) despite their
337 immense diversity (Zhang 2013; Eisenhauer and Hines 2021). Moreover, we observed that certain
338 organisms (at the species or genus level) concentrated much of the attention given to their
339 taxonomic group (Fig. S5B, C), reiterating that taxonomic bias also occurs at fine taxonomic levels
340 (Zuk et al. 2014). Note, however, that the popularity of some organisms may be a spillover from
341 the popularity of the organisms they are associated with, such as parasites of charismatic mammals
342 (Simoncini et al. 2025).

343 The popularity of taxonomic groups among participants in our survey was rarely associated
344 with how easy they are to access and work with, showing that organisms' logistic aspects do not
345 appear to be associated with their research use at high taxonomic levels. For example, with some
346 exceptions, birds and (especially) mammals were considered as equally or more difficult to find,
347 observe, rear, or buy than other taxonomic groups (Fig. 5). As a counterpoint, molluscs, which

348 were rarely used by surveyed scientists for research purposes (only 5.88% of participants worked
349 with them; Fig. S4), were largely perceived as easy to work with both in the field and in the lab
350 (Fig. 5). These results are not surprising as the necessary fieldwork duration to collect data from
351 overlooked taxa (beetles and moths) is usually shorter than from popular taxa (mammals and birds)
352 (Pawar 2003). Nonetheless, within taxonomic groups, genera mentioned more frequently by
353 participants tended to be perceived as easier to work with. This suggests that organisms with
354 logistic advantages may be selected over other closely related taxa, a result that is congruent with
355 studies investigating literature patterns related to specific taxonomic groups (Yarwood et al. 2019;
356 dos Santos et al. 2020; Silva et al. 2020; Ellison et al. 2021; Tam et al. 2022; Chen et al. 2023;
357 Fischer et al. 2025).

358 We hypothesised that researchers also consider the possible applications of organisms to
359 society, such as their economic relevance or conservation status, when deciding which research to
360 pursue. We found that insects and mammals were deemed more economically important than
361 arachnids and birds (mammals were also considered more economically important than reptiles).
362 By contrast, insects were less frequently deemed as threatened or endangered in the wild than most
363 taxonomic groups (including mammals, which were also more frequently deemed as endangered
364 than arachnids). This finding is worrisome as invertebrates are under severe threat (Cardoso et al.
365 2020) yet remain underrepresented in biodiversity databases (e.g. IUCN's red list; Cardoso et al.
366 2012) as most biodiversity assessments focus on birds and mammals, making the decline and
367 extinction of invertebrates unnoticed (Régnier et al. 2015). This represents a striking example of
368 the taxonomic bias vortex, in which lack of data generates the impression that no problem exists,
369 leading to even less attention to the issue. Participants were also more likely to state that they had
370 conducted applied research with species they deemed more economically important or more

371 endangered in the wild. Strangely, the likelihood of participants conducting applied research was
372 greater for mammals than for arachnids and insects (Fig. 6G) despite the latter being perceived as
373 economically important as the former. Within taxa, the number of participants working with a
374 genus was positively related to its perception as economically relevant, but not with its perception
375 as endangered in the wild. Our results indicate that organisms' aspects connected to their explicit
376 usefulness to society can be involved in researchers' taxonomic decisions, but this influence seems
377 to be weak, at least among ethologists and/or behavioural ecologists.

378 Participants appear to let their emotions lead their choice of research organisms as stronger
379 affection for a taxonomic group corresponded with a higher likelihood of working with it among
380 participants in our survey (Fig. 3B; Fig. S3). Given that the most popular taxonomic groups (e.g.
381 birds and mammals) overwhelmingly elicited positive feelings from researchers (including those
382 that never worked with them; Fig. S3), we argue that positive views of a taxon make researchers
383 more likely to work with that taxon. Importantly, the patterns of fondness for distinct taxonomic
384 groups we observed from surveyed researchers echoes values present in society, in which
385 vertebrates are generally more well perceived than invertebrates (Miralles et al. 2019; Possidónio
386 et al. 2019). We also found that participants' experiences with an organism stimulated more
387 positive perceptions of its taxonomic group (Fig. 6B), whereas this effect was stronger for
388 arachnids than for birds and mammals, and also stronger for fish than for mammals. The
389 relationship between fondness and experience appears to be reciprocal: appreciation of a
390 taxonomic group increases the likelihood of working with it, while experience with a taxonomic
391 group enhances appreciation of it. This self-reinforcing cycle resembles a positive counterpart of
392 the conceptual framework for biophobia, in which exposure to negative information about
393 organisms causes fear and avoidance, leading to disconnection from nature, fortifying the cycle by

394 strengthening the belief in and impact from negative perceptions of feared organisms (Soga et al.
395 2023). Outreach programs thus remain crucial to promote awareness and appreciation of neglected
396 taxonomic groups among the general public, countering fear-based narratives commonly
397 perpetuated by the media that only aggravate this dire situation (Mammola et al. 2020a; Soga et
398 al. 2023). Moreover, endorsing diverse taxonomic experiences in teaching and supervision within
399 academia may also allow researchers to evaluate their options with less interference from
400 damaging and alienating taxonomic perceptions.

401 Specific activities and preferences also appear to be connected to researchers' taxonomic
402 choices. For example, participants were more likely to have engaged in (the equivalent of)
403 birdwatching prior to working with a bird species than of having engaged in a similar hobby with
404 an invertebrate taxon before having an experience with that taxon (Fig. 6A). This bolsters the idea
405 that positive exposure to a taxonomic group elicits individuals to work with that group,
406 highlighting that encouraging activities involving neglected taxonomic groups (e.g. bugwatching;
407 Eaton 2025) can generate research interest in the organisms involved. Furthermore, we found that
408 surveyed researchers were diverse in their preference for fieldwork over lab work, ranging from
409 complete to no preference (Fig. S2A). Greater preference for fieldwork was not associated with
410 how many species or taxonomic groups a surveyed researcher worked with (Fig. 2), but it was
411 positively related to greater appreciation of arachnids and vertebrates other than fish (Fig. 3A).
412 Yet, greater preference for fieldwork was only positively related to the likelihood of working with
413 birds and mammals, but not with amphibians and reptiles, and negatively related to the likelihood
414 of working with fish and invertebrates other than molluscs (Fig. 3B). These findings suggest that
415 professional preferences may influence scientists' fondness for certain taxa and their choices of
416 research organisms (sometimes in conflicting ways).

417 Researchers also appear to vary in what motivates them, as some participants in our survey
418 were exclusively driven by specific taxa while others were fully motivated by other factors (e.g.
419 particular topics, professional prospects) (Fig. S2B). The more participants were motivated by
420 taxa, the more species but the fewer taxonomic groups they worked with (Fig. 2). This suggests
421 that researchers primarily motivated by taxa are open to expanding their experience towards other
422 (preferably neglected) species from the same taxonomic group, while their counterparts can be
423 more easily inspired to diversify their experiences at a higher taxonomic level. Additionally,
424 although motivation from taxa was not associated with fondness for any taxonomic group (Fig.
425 3A), the fact that it was positively related to the likelihood of working with amphibians and
426 arachnids (Fig. 3B) indicates that these two taxonomic groups are more likely to receive attention
427 from researchers specifically interested in studying only them or few other taxa.

428 In our survey, participants frequently deemed several species as model organisms (e.g. barn
429 swallows, crab-eating macaques) despite them not being officially listed as such (by USA's
430 National Institute of Health; NIH 2025). Participants' agreement that an organism is a model
431 system was positively related to the number of distinct participants mentioning the genus of that
432 organism, indicating that this term is loosely applied to commonly studied species in a certain
433 context (in our case, behavioural research). Species considered model organisms by participants
434 in our survey were also more likely to be deemed easy to work with in the lab than in the field,
435 confirming the connection of this term with experimental and laboratory work (Leonelli and
436 Ankeny 2012, 2013; Dietrich et al. 2014; Matthews and Vosshall 2020). Despite this, the only
437 difference across taxonomic groups we found was between fish and arachnids (Fig. 6C), which is
438 surprising given that they are perceived as similarly accessible to work in the laboratory (Fig. 5C,
439 D).

440 Given that the model organism label propulses taxonomic bias by rewarding and promoting
441 species that have acquired an arbitrary status among researchers, we argue that this term should be
442 abolished in most spheres of ecology and evolution. Moreover, even though species considered
443 model organisms have generated breakthroughs in science (Fields and Johnston 2005), they cannot
444 replace the hidden gems that biodiversity has to offer (e.g. bioinspired designs and materials; Ortiz
445 and Boyce 2008). We should thus facilitate research with different species, both in the field and in
446 the laboratory, instead of re-using the same ones as a cheap trick to bypass a broken funding
447 system. Recent technologies (e.g. drones, minute tracking devices; Farine et al. 2024) and data
448 sources (e.g. citizen science; Niemiller et al. 2021; Mason et al. 2025) can certainly help with this,
449 allowing ecology and evolution to reach its full potential.

450 Precedence effects can also occur at the individual level. Most participants in our survey
451 mentioned working, in multiple stages of their career, with species that were suggested by their
452 supervisors, albeit this pattern was weakened after participants obtained a permanent position (Fig.
453 4). This means that the taxonomic experiences of early career researchers (ECRs) are largely
454 influenced by late-career researchers (LCRs) they work with. ECRs often interact with few LCRs
455 who can act as their mentors and/or supervisors (e.g. those employed by the university in which
456 they study), representing a major bottleneck for ECRs' possible experiences, especially if LCRs
457 only accept supervising research projects with a specific taxon. Because few LCRs work with
458 neglected taxonomic groups, ECRs desiring to work with such groups are less likely to find
459 supervisors who can support their interests. ECRs that manage to work with their predilected
460 taxonomic group or those that do not have taxonomic preferences are then mostly bound to the
461 specific organism suggested by the LCR supervising them. These processes can then dramatically
462 reinforce existing taxonomic biases if researchers continue to work with the same taxonomic group

463 or species throughout their career. Our findings suggest that researchers commonly take this route,
464 stagnating or even amplifying taxonomic biases at multiple taxonomic levels. Fortunately, there
465 are several tools that can help researchers to diversify their taxonomic experiences. For example,
466 citizen science data (e.g. iNaturalist) can show species that are abundant in a given researcher's
467 area, and enthusiasts (e.g. photographers, social media influencers) and other researchers (e.g.
468 taxonomists) may be happy to suggest species to study and ways of investigating them in the field
469 and in the lab. To ameliorate this precedence effect, we recommend that (1) institutions consider
470 the diversity of taxonomic experiences of LCRs when recruiting them, (2) LCRs adopt a flexible
471 approach, supporting and motivating ECRs to engage with diverse taxa, (3) ECRs search for
472 distinct sources of information to expand their experiences.

473 Taxonomic bias generates a skewed perspective of natural processes and patterns, but it is
474 only one of many biases affecting the study of ecology and evolution (Winder et al. 2025). For
475 example, biases in our perceptions of the sexes likely produce stereotypic portrayals of them (Pollo
476 and Kasumovic 2022) (e.g. coy females and eager males), which are common in the literature
477 (Green and Madjidian 2011; Ah-King 2022a, 2022b; Spaulding and Fuselier 2023).
478 Misrepresentations of taxa, sexes, and other biological aspects curb the power of studies attempting
479 to draw broad conclusions about nature. On one hand, gaps in the available data, if properly
480 detected, only allow part of the puzzle to be revealed. For example, lack of data on sexual signals
481 expressed by females and animals other than birds precludes a holistic understanding of the
482 evolution of these traits (Pollo et al. 2025a). Studies pointing out gaps in the literature can then
483 direct researchers to collect data that is urgently needed (Pollo et al. 2025b). However, researchers
484 may infer erroneous conclusions when they fail to recognise that the data they analyse are biased.
485 This is apparently a frequent issue in meta-analyses related to animal behaviour, in which the

486 underrepresentation of invertebrates leads to improper generalisations (Dochtermann et al. 2026).
487 Awareness of biases in science is essential for their detection and mitigation (Winder et al. 2025)
488 and thus an important first step. However, we argue that we need to go beyond theoretical
489 discussions by implementing actions that minimise the existing biases in ecology and evolution.
490 Similar to pleas made by researchers combating biases in how sexes are represented in science
491 (Ah-King 2013; Hughes 2022; Zemenick et al. 2022), we advocate for a more taxonomically
492 equitable education (as early as childcare) and data collection. We hope that our recommendations
493 throughout the present study can then influence ecologists and evolutionary biologists to strive for
494 such endeavour (but see also Pollo and Kasumovic 2026).

495

496 **Code and data accessibility**

497 All data and code used in this study are available at <https://zenodo.org/records/18719247>.

498

499 **Acknowledgements**

500 We thank all survey participants for generously contributing their time, especially those that helped
501 us to recruit more participants or reached out to us with feedback. We are also grateful to
502 Jigmidmaa Boldbaatar, Russell Bonduriansky, Caitlin Nicole Creak, Kyle Morrison, Daniela
503 Wilner, and Wai Chi Vicky Ying for helping us test our survey. We thank Catharina Vendl for
504 feedback on the manuscript.

505

506 **Author contributions**

507 PP: conceptualisation, methodology, formal analysis, investigation, data curation, writing -
508 original draft, writing - review & editing, visualisation, project administration.

509 MK: methodology, writing - review & editing.

510

511 **Competing interests**

512 The authors declare no conflicts of interest.

513

514 **Declaration of AI use**

515 The authors declare that they occasionally used GPT-5 (OpenAI) to improve the clarity and
516 readability of this work. After using these tools, the authors reviewed and edited the content as
517 needed and took full responsibility for the content of the publication.

518

519 **References**

520 Adamo, M., M. Chialva, J. Calevo, F. Bertoni, K. Dixon, and S. Mammola. 2021. Plant scientists'
521 research attention is skewed towards colourful, conspicuous and broadly distributed flowers.
522 *Nature Plants* 7:574–578.

523 Ah-King, M. 2013. Queering animal sexual behavior in biology textbooks. *Confero Essays on*
524 *Education Philosophy and Politics* 1:46–89.

525 ———. 2022*a*. The history of sexual selection research provides insights as to why females are
526 still understudied. *Nature Communications* 13:6976.

527 ———. 2022*b*. *The female turn*. Springer Nature Singapore, Singapore.

528 Bartoń, K. 2023. MuMIn: multi-model inference.

529 Bonnet, X., R. Shine, and O. Lourdais. 2002. Taxonomic chauvinism. *Trends in Ecology &*
530 *Evolution* 17:2000–2002.

531 Broeckhoven, C., and A. du Plessis. 2022. Escaping the labyrinth of bioinspiration: biodiversity
532 as key to successful product innovation. *Advanced Functional Materials* 32:1–8.

533 Brooks, Mollie, E., K. Kristensen, V. Benthem, Koen, J., A. Magnusson, W. Berg, Casper, A.
534 Nielsen, J. Skaug, Hans, et al. 2017. glmmTMB balances speed and flexibility among
535 packages for zero-inflated generalized linear mixed modeling. *The R Journal* 9:378.

536 Cardoso, P., P. S. Barton, K. Birkhofer, F. Chichorro, C. Deacon, T. Fartmann, C. S. Fukushima,
537 et al. 2020. Scientists’ warning to humanity on insect extinctions. *Biological Conservation*
538 242.

539 Cardoso, P., P. A. V. Borges, K. A. Triantis, M. A. Ferrández, and J. L. Martín. 2012. The
540 underrepresentation and misrepresentation of invertebrates in the IUCN Red List. *Biological*
541 *Conservation* 149:147–148.

542 Casad, B. J., J. E. Franks, C. E. Garasky, M. M. Kittleman, A. C. Roesler, D. Y. Hall, and Z. W.
543 Petzel. 2021. Gender inequality in academia: problems and solutions for women faculty in
544 STEM. *Journal of Neuroscience Research* 99:13–23.

545 Casola, W., M. N. Peterson, D. F. Lawson, S. B. Jackson, K. T. Stevenson, and R. X. Valdez.
546 2026. Evaluating household dynamics of wildlife preferences using toys. *Human*
547 *Dimensions of Wildlife* 31:155–160.

548 Chen, T., P. A. Garber, L. Zhang, L. Yang, and P. Fan. 2023. The pattern and drivers of taxonomic
549 bias in global primate research. *Global Ecology and Conservation* 46:e02599.

550 Chen, Z., R. T. Corlett, X. Jiao, S.-J. Liu, T. Charles-Dominique, S. Zhang, H. Li, et al. 2018.
551 Prolonged milk provisioning in a jumping spider. *Science* 362:1052–1055.

552 Christensen, R. H. B. 2023. ordinal - regression models for ordinal data.

553 Clark, J. A., and R. M. May. 2002. Taxonomic bias in conservation research. *Science* 297:191–
554 192.

555 Clutton-Brock, T. H. 1991. *The evolution of parental care*. Princeton University Press, Princeton,
556 New Jersey.

557 Damasio, A. 1994. *Descartes' error: emotion, reason, and the human brain*. Avon Books, New
558 York, NY.

559 Dietrich, M. R., R. A. Ankeny, and P. M. Chen. 2014. Publication trends in model organism
560 research. *Genetics* 198:787–794.

561 Dochtermann, N., M. Sekhar, and S. Nakagawa. 2026. Ungeneralizable generalizations? A meta-
562 meta-analysis of the influence of taxonomic bias on the study of behaviour. *Animal*
563 *Behaviour* 236:123542.

564 dos Santos, J. W., R. A. Correia, A. C. M. Malhado, J. V. Campos-Silva, D. Teles, P. Jepson, and
565 R. J. Ladle. 2020. Drivers of taxonomic bias in conservation research: a global analysis of
566 terrestrial mammals. *Animal Conservation* 23:679–688.

567 Eaton, E. R. 2025. *Bugwatching: the art, joy, and importance of observing insects*. Princeton
568 University Press, Princeton, NJ.

569 Eisenhauer, N., and J. Hines. 2021. Invertebrate biodiversity and conservation. *Current Biology*
570 31:R1214–R1218.

571 Ellison, G., M. Jones, B. Cain, and C. M. Bettridge. 2021. Taxonomic and geographic bias in 50
572 years of research on the behaviour and ecology of galagids. *PLOS One* 16:1–22.

573 Farine, D. R., J. Penndorf, S. Bolcato, B. Nyaguthii, and L. M. Aplin. 2024. Low-cost animal
574 tracking using Bluetooth low energy beacons on a crowd-sourced network. *Methods in*
575 *Ecology and Evolution* 15:2247–2261.

576 Fields, S., and M. Johnston. 2005. Whither model organism research? *Science* 307:1885–1886.

577 Fischer, S. E., J. G. Otten, A. M. Lindsay, D. Miles, and H. Streby. 2025. Six-decade research bias
578 towards fancy and familiar bird species. *Proceedings of the Royal Society B: Biological*
579 *Sciences* 292.

580 Fleischner, T. L., R. E. Espinoza, G. A. Gerrish, H. W. Greene, R. W. Kimmerer, E. A. Lacey, S.
581 Pace, et al. 2017. Teaching biology in the field: importance, challenges, and solutions.
582 *BioScience* 67:558–567.

583 Fox, C. W., J. Meyer, and E. Aimé. 2023. Double-blind peer review affects reviewer ratings and
584 editor decisions at an ecology journal. *Functional Ecology* 37:1144–1157.

585 Fredrikson, M., P. Annas, H. Fischer, and G. Wik. 1996. Gender and age differences in the
586 prevalence of specific fears and phobias. *Behaviour Research and Therapy* 34:33–39.

587 Gangwani, K., and J. Landin. 2018. The decline of insect representation in biology textbooks over
588 time. *American Entomologist* 64:252–257.

589 Green, K. K., and J. A. Madjidian. 2011. Active males, reactive females: stereotypic sex roles in
590 sexual conflict research? *Animal Behaviour* 81:901–907.

591 Guénard, B., A. C. Hughes, C. Lainé, S. Cannicci, B. D. Russell, and G. A. Williams. 2025.
592 Limited and biased global conservation funding means most threatened species remain
593 unsupported. *Proceedings of the National Academy of Sciences of the United States of*
594 *America* 122:e2412479122.

595 Hartig, F. 2016. DHARMA: residual diagnostics for hierarchical (multi-level/mixed) regression
596 models.

597 Hughes, M. 2022. Student preconceptions of sex differences in animal behaviour classes:
598 observations, recommendations and need for research. *Animal Behaviour* 192:95–100.

599 Leather, S. R. 2009. Taxonomic chauvinism threatens the future of entomology. *Biologist* 56:10–
600 13.

601 Lenth, R. V. 2025. emmeans: estimated marginal means, aka least-squares means.

602 Leonelli, S., and R. A. Ankeny. 2012. Re-thinking organisms: the impact of databases on model
603 organism biology. *Studies in History and Philosophy of Biological and Biomedical Sciences*
604 43:29–36.

605 ———. 2013. What makes a model organism? *Endeavour* 37:209–212.

606 Lorimer, J. 2007. Nonhuman charisma. *Environment and Planning D: Society and Space* 25:911–
607 932.

608 Mailho-Fontana, P. L., M. M. Antoniazzi, G. R. Coelho, D. C. Pimenta, L. P. Fernandes, A.
609 Kupfer, E. D. Brodie, et al. 2024. Milk provisioning in oviparous caecilian amphibians.
610 *Science* 383:1092–1095.

611 Mammola, S., M. Adamo, D. Antić, J. Calevo, T. Cancellario, P. Cardoso, D. Chamberlain, et al.
612 2023. Drivers of species knowledge across the tree of life. *eLife* 12:1–23.

613 Mammola, S., V. Nanni, P. Pantini, and M. Isaia. 2020*a*. Media framing of spiders may exacerbate
614 arachnophobic sentiments. *People and Nature* 2:1145–1157.

615 Mammola, S., N. Riccardi, V. Prié, R. Correia, P. Cardoso, M. Lopes-Lima, and R. Sousa. 2020*b*.
616 Towards a taxonomically unbiased European Union biodiversity strategy for 2030.
617 *Proceedings of the Royal Society B: Biological Sciences* 287:20202166.

618 Mann, M. D. 2015. Choosing the right species in research. *Lab Animal* 44:274–278.

619 Mason, B. M., T. Mesaglio, J. Barratt Heitmann, M. Chandler, S. Chowdhury, S. B. Z. Gorta, F.
620 Grattarola, et al. 2025. iNaturalist accelerates biodiversity research. *BioScience* 75:953–965.

621 Matthews, B. J., and L. B. Vosshall. 2020. How to turn an organism into a model organism in 10
622 “easy” steps. *Journal of Experimental Biology* 223.

623 McGillicuddy, M., D. I. Warton, G. Popovic, and B. M. Bolker. 2025. Parsimoniously fitting large
624 multivariate random effects in glmmTMB. *Journal of Statistical Software* 112.

625 Miralles, A., M. Raymond, and G. Lecointre. 2019. Empathy and compassion toward other species
626 decrease with evolutionary divergence time. *Scientific Reports* 9:1–8.

627 Mizuno, A., M. Lagisz, P. Pollo, L. Guillette, M. Soma, and S. Nakagawa. 2025. Systematic
628 mapping and bibliometric analysis of meta-analyses on animal cognition. *Neuroscience and*
629 *Biobehavioral Reviews* 177:106342.

630 Murray, H. J., E. J. Green, D. R. Williams, I. J. Burfield, and M. L. de Brooke. 2015. Is research
631 effort associated with the conservation status of European bird species? *Endangered Species*
632 *Research* 27:193–206.

633 Ng, L., M. A. Elgar, and D. Stuart-Fox. 2021. From bioinspired to bioinformed: benefits of greater
634 engagement from biologists. *Frontiers in Ecology and Evolution* 9:1–9.

635 Niemiller, K. D. K., M. A. Davis, and M. L. Niemiller. 2021. Addressing ‘biodiversity naivety’
636 through project-based learning using iNaturalist. *Journal for Nature Conservation* 64.

637 NIH. 2025. Model organism sharing policy.

638 Ortiz, C., and M. C. Boyce. 2008. Bioinspired structural materials. *Science* 319:1053–1054.

639 Pawar, S. 2003. Taxonomic chauvinism and the methodologically challenged. *BioScience* 53:861–
640 864.

641 Penick, C. A., G. Cope, S. Morankar, Y. Mistry, A. Grishin, N. Chawla, and D. Bhate. 2022. The
642 comparative approach to bio-inspired design: integrating biodiversity and biologists into the
643 design process. *Integrative and Comparative Biology* 62:1153–1163.

644 Pollo, P., and M. M. Kasumovic. 2022. Let's talk about sex roles: what affects perceptions of sex
645 differences in animal behaviour? *Animal Behaviour* 183:1–12.

646 Pollo, P., and M. M. Kasumovic. 2025. Taxonomic bias in ethology and behavioural ecology:
647 patterns, origins, and solutions, a protocol.

648 ———. 2026. Taxonomic bias: a persistent issue in ecology and evolution. *EcoEvoRxiv*.

649 Pollo, P., M. Lagisz, R. C. Macedo-Rego, A. Mizuno, Y. Yang, and S. Nakagawa. 2025*a*.
650 Synthesis of nature's extravaganza: an augmented meta-meta-analysis on (putative) sexual
651 signals. *Ecology Letters* 28:e70215.

652 Pollo, P., M. Lagisz, Y. Yang, A. Culina, and S. Nakagawa. 2024. Synthesis of sexual selection: a
653 systematic map of meta-analyses with bibliometric analysis. *Biological Reviews* 99:2134–
654 2175.

655 Pollo, P., A. R. Martinig, A. Mizuno, K. Morrison, P. Pottier, L. Ricolfi, J. Tam, et al. 2025*b*.
656 Harnessing meta-analyses' insights in ecology and evolution research. *Royal Society Open*
657 *Science* 12:250759.

658 Possidónio, C., J. Graça, J. Piazza, and M. Prada. 2019. Animal images database: validation of 120
659 images for human-animal studies. *Animals* 9:475.

660 R Core Team. 2025. R: a language and environment for statistical computing. R foundation for
661 statistical computing, Vienna, Austria.

662 Régnier, C., G. Achaz, A. Lambert, R. H. Cowie, P. Bouchet, and B. Fontaine. 2015. Mass
663 extinction in poorly known taxa. *Proceedings of the National Academy of Sciences of the*
664 *United States of America* 112:7761–7766.

665 Rosenthal, M. F., M. Gertler, A. D. Hamilton, S. Prasad, and M. C. B. Andrade. 2017. Taxonomic
666 bias in animal behaviour publications. *Animal Behaviour* 127:83–89.

667 Seddon, P. J., P. S. Soorae, and F. Launay. 2005. Taxonomic bias in reintroduction projects.
668 *Animal Conservation* 8:51–58.

669 Shine, R., and X. Bonnet. 2000. Snakes: a new ‘model organism’ in ecological research? Trends
670 in Ecology & Evolution 15:221–222.

671 Silva, A. F. da, A. C. M. Malhado, R. A. Correia, R. J. Ladle, M. V. C. Vital, and T. Mott. 2020.
672 Taxonomic bias in amphibian research: are researchers responding to conservation need?
673 *Journal for Nature Conservation* 56:125829.

674 Simaika, J. P., and M. J. Samways. 2010. Biophilia as a universal ethic for conserving biodiversity.
675 *Conservation Biology* 24:903–906.

676 Simoncini, A., G. F. Ficetola, and M. Lattanzi. 2025. Taxonomic bias towards charismatic and
677 easy-to-find mammals shapes knowledge of parasites. *Biodiversity and Conservation*
678 34:4035–4048.

679 Snaddon, J. L., E. C. Turner, and W. A. Foster. 2008. Children’s perceptions of rainforest
680 biodiversity: which animals have the lion’s share of environmental awareness? *PLOS One*
681 3:1–5.

682 Snell-Rood, E. C., and D. Smirnov. 2025. Biology for biomimetics: II. A tutorial on how to
683 diversify a list of biological models. *Bioinspiration and Biomimetics* 20:62001.

684 Soga, M., K. J. Gaston, Y. Fukano, and M. J. Evans. 2023. The vicious cycle of biophobia. Trends
685 in Ecology and Evolution 38:512–520.

686 Spaulding, S., and L. Fuselier. 2023. Images of nonhuman animals in animal behaviour textbooks
687 communicate an androcentric view of reproductive-related behaviours. *Animal Behaviour*
688 205:117–129.

689 Spoon, K., N. LaBerge, K. H. Wapman, S. Zhang, A. C. Morgan, M. Galesic, B. K. Fosdick, et al.
690 2023. Gender and retention patterns among U.S. faculty. *Science Advances* 9:1–12.

691 Srivastava, D. S., J. Bernardino, A. T. Marques, A. Proença-Ferreira, A. F. Filipe, L. Borda-de-
692 Água, and J. Gameiro. 2024. Editors are biased too: an extension of Fox et al. (2023)'s
693 analysis makes the case for triple-blind review. *Functional Ecology* 38:278–283.

694 Stahlschmidt, Z. R. 2011. Taxonomic chauvinism revisited: insight from parental care research.
695 *PLOS One* 6:e24192.

696 Stay, B., and A. C. Coop. 1974. “Milk” secretion for embryogenesis in a viviparous cockroach.
697 *Tissue and Cell* 6:669–693.

698 Strommen, E. 1995. Lions and tigers and bears, oh my! Children’s conceptions of forests and their
699 inhabitants. *Journal of Research in Science Teaching* 32:683–698.

700 Stuart-Fox, D., L. Ng, L. Barner, A. T. D. Bennett, S. J. Blamires, M. A. Elgar, A. R. Evans, et al.
701 2023. Challenges and opportunities for innovation in bioinformed sustainable materials.
702 *Communications Materials* 4:1–12.

703 Tam, J., M. Lagisz, W. Cornwell, and S. Nakagawa. 2022. Quantifying research interests in 7,521
704 mammalian species with h-index: a case study. *GigaScience* 11:1–11.

705 Titley, M. A., J. L. Snaddon, and E. C. Turner. 2017. Scientific research on animal biodiversity is
706 systematically biased towards vertebrates and temperate regions. *PLOS One* 12:e0189577.

707 Troudet, J., P. Grandcolas, A. Blin, R. Vignes-Lebbe, and F. Legendre. 2017. Taxonomic bias in
708 biodiversity data and societal preferences. *Scientific Reports* 7:9132.

709 Wei, H., V. Berdejo-Espinola, Y. Ma, and T. Amano. 2024. Content analysis of nature
710 documentaries in China: challenges and opportunities to raise public conservation
711 awareness. *Biological Conservation* 292:110522.

712 Winder, L. A., E. Brignall, F. S. E. Dawson Pell, M. Germain, C. Halliwell, J. A. Hibberd, F.
713 Morland, et al. 2025. Known and unknown biases: a framework for contextualising and
714 identifying bias in animal behaviour research. *Ethology* 1–8.

715 Yarwood, M. R., M. A. Weston, and M. R. E. Symonds. 2019. Biological determinants of research
716 effort on Australian birds: a comparative analysis. *Emu* 119:38–44.

717 Zemenick, A. T., S. Turney, A. J. Webster, S. C. Jones, and M. G. Weber. 2022. Six principles for
718 embracing gender and sexual diversity in postsecondary biology classrooms. *BioScience*
719 72:481–492.

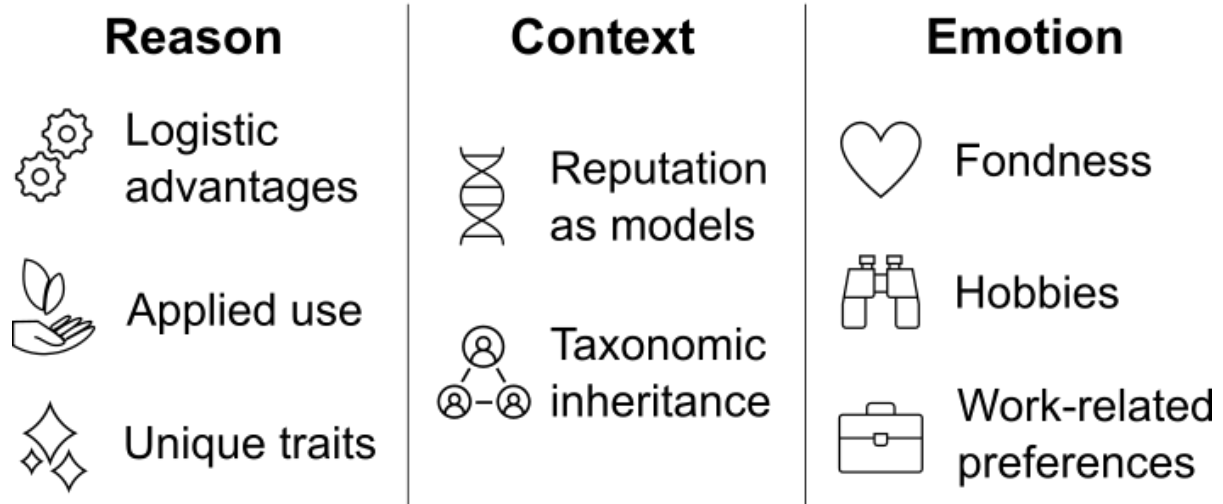
720 Zhang, Z.-Q. 2013. Animal biodiversity: an update of classification and diversity in 2013. *Zootaxa*
721 3703:5–11.

722 Zuk, M., F. Garcia-Gonzalez, M. E. Herberstein, and L. W. Simmons. 2014. Model systems,
723 taxonomic bias, and sexual selection: beyond *Drosophila*. *Annual Review of Entomology*
724 59:321–338.

725

726

Why are organisms selected for research?

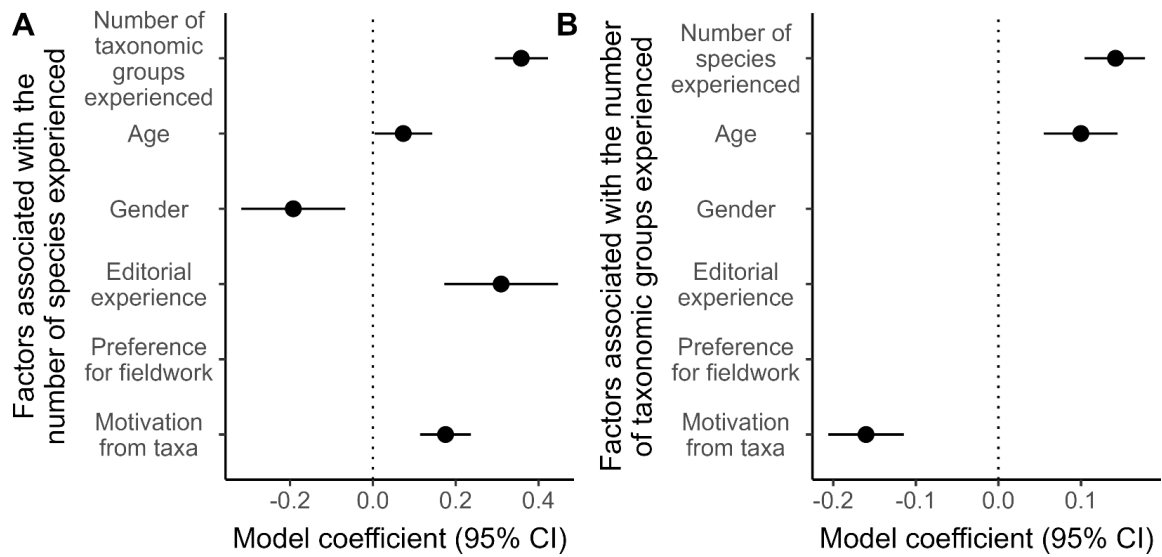


729

730 *Figure 1.*

731 Factors hypothesised to be associated with the selection of research organisms in ecology and

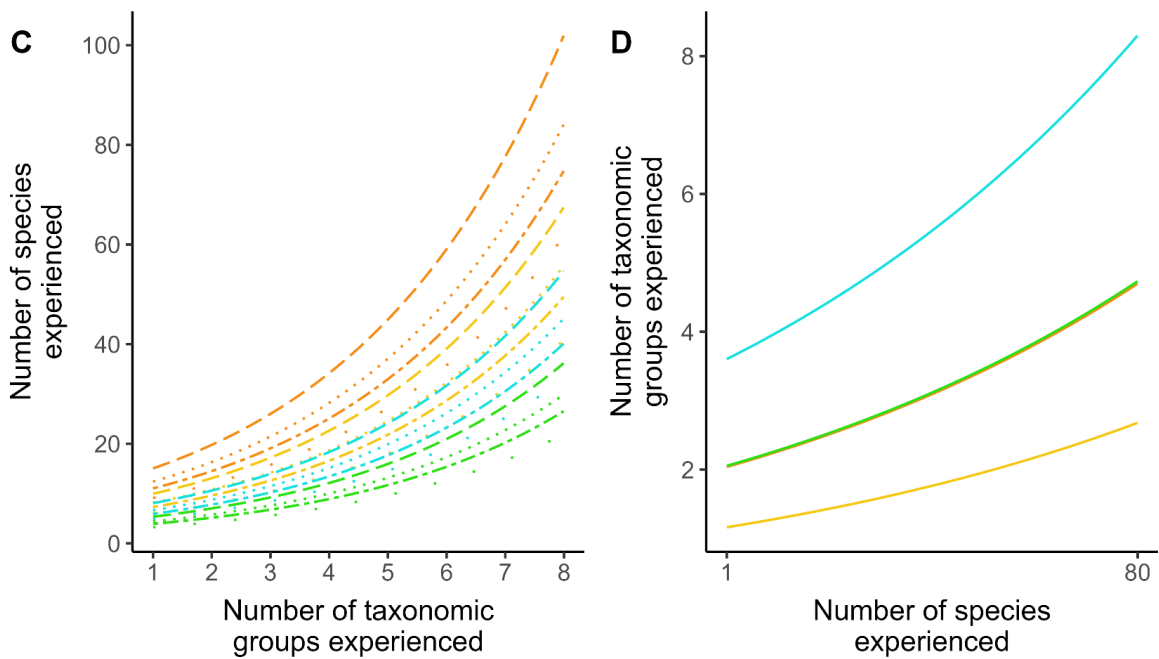
732 evolution.



Gender and editorial experience Motivation from taxa and age

-- Man, with -.- Man, without — Maximum, oldest — Maximum, youngest

···· Woman, with ··· Woman, without — Minimum, oldest — Minimum, youngest



733

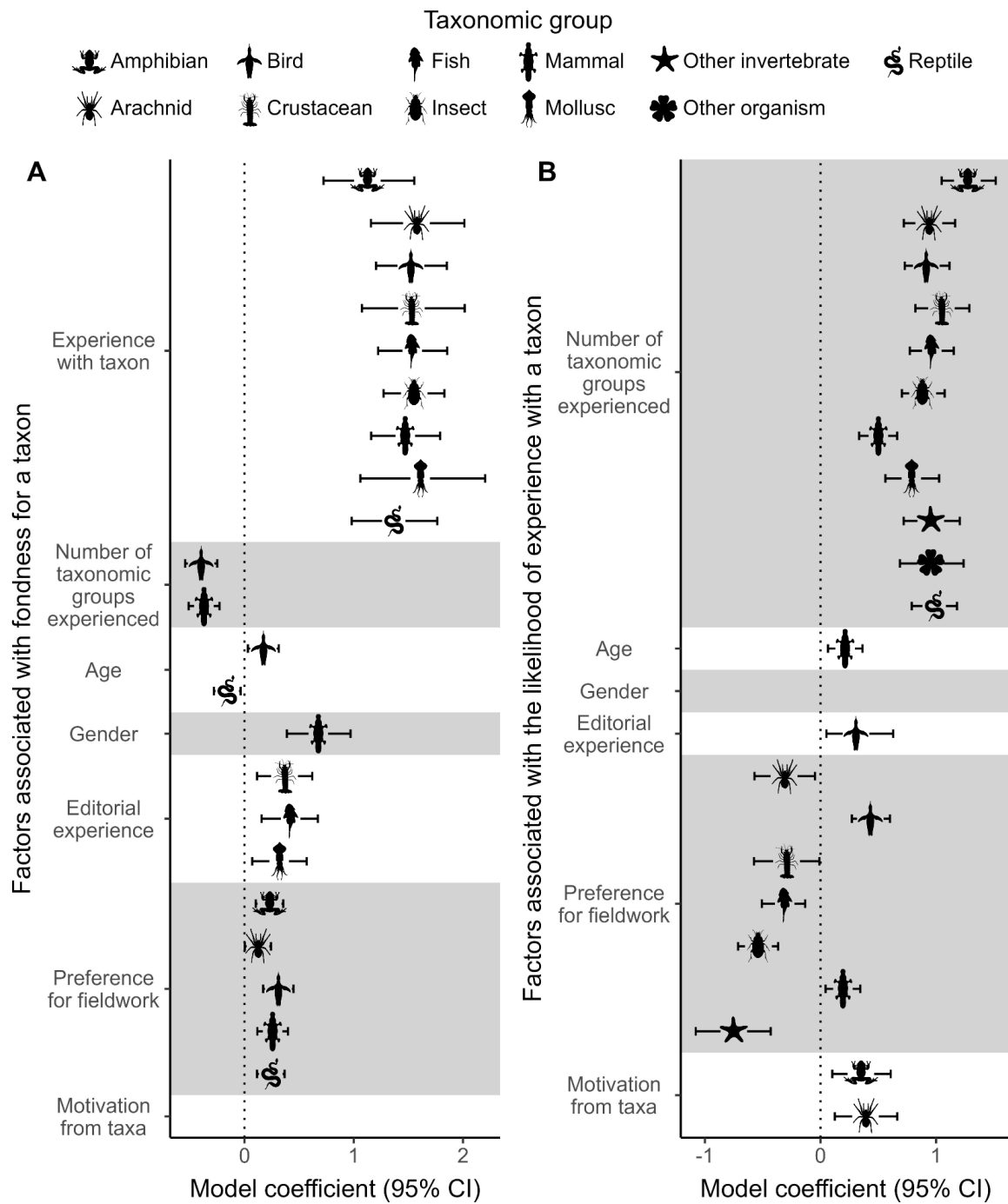
734 *Figure 2.*

735 Factors associated with participants' experiences. Top panels show coefficient values for factors

736 associated with the number of species (A) and taxonomic groups (B) that participants worked with.

737 Gender refers to the comparison of women vs. men (comparisons with other or unknown gender
738 identities are not shown). Note that only coefficients from factors included in all selected models
739 are shown. Bottom panels show the number of species (C) and taxonomic groups (D) that
740 participants worked with predicted by selected models. Maximum and minimum motivation from
741 taxa refer to 100% and 0% of this variable (respectively), while oldest and youngest represent 82
742 and 21 years old individuals (respectively).

743



744

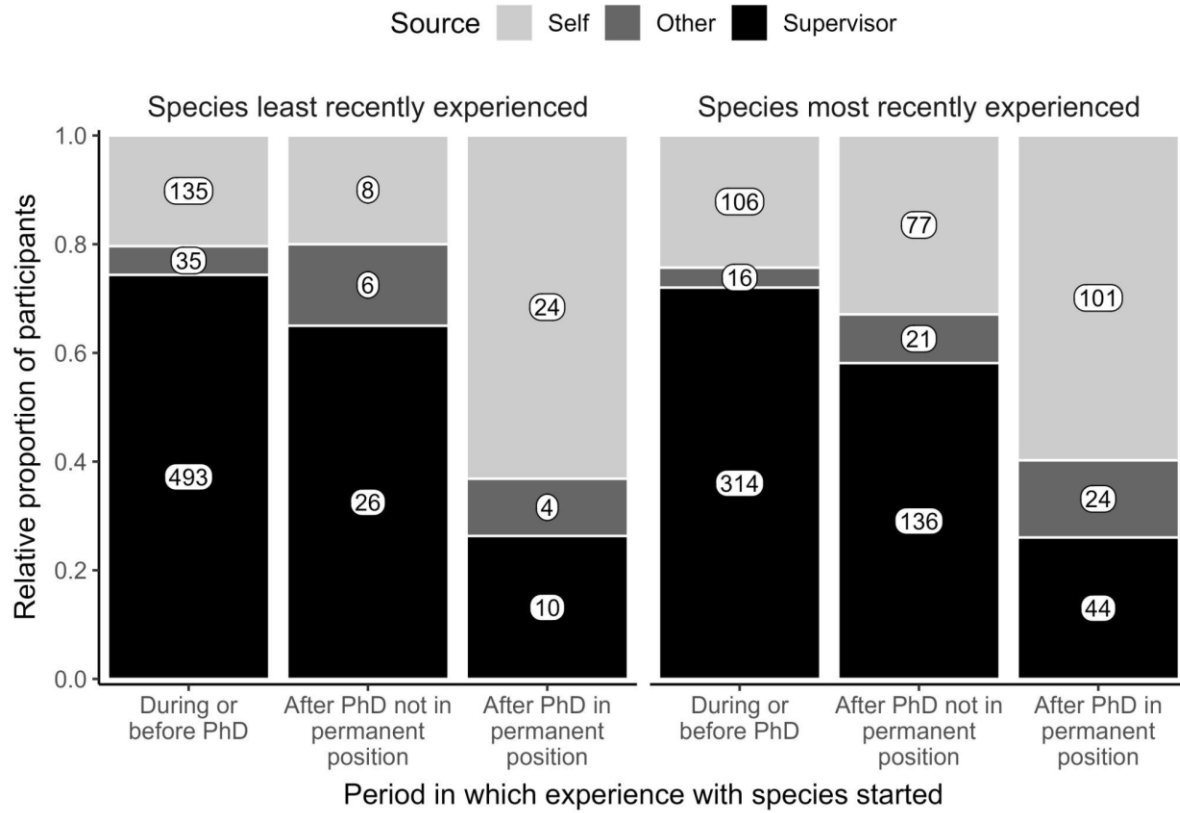
745 *Figure 3.*

746 Factors associated with fondness for (A) and experience with (B) a given taxonomic group. Gender

747 refers to the comparison of women vs. men (comparisons with other or unknown gender identities

748 are not shown). Fondness for a taxon was also included as a predictor in models exploring the
749 factors associated with the likelihood of experience with that taxon (B), but coefficients are not
750 shown. Background colours highlight coefficients associated with different factors.

751



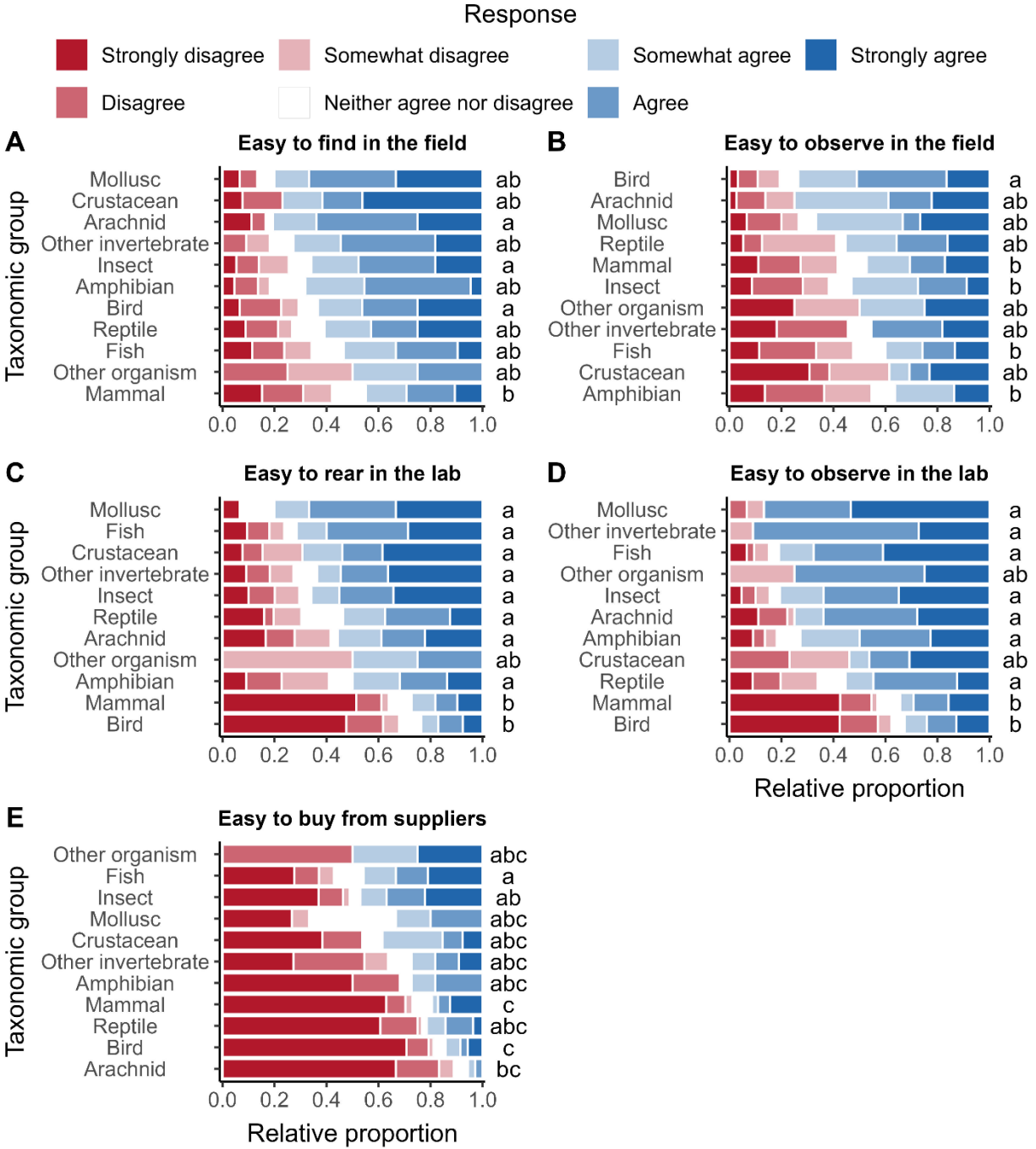
752

753 *Figure 4.*

754 How and when participants started working with certain species. Numbers within bars represent

755 the number of participants within that category.

756



757

758 *Figure 5.*

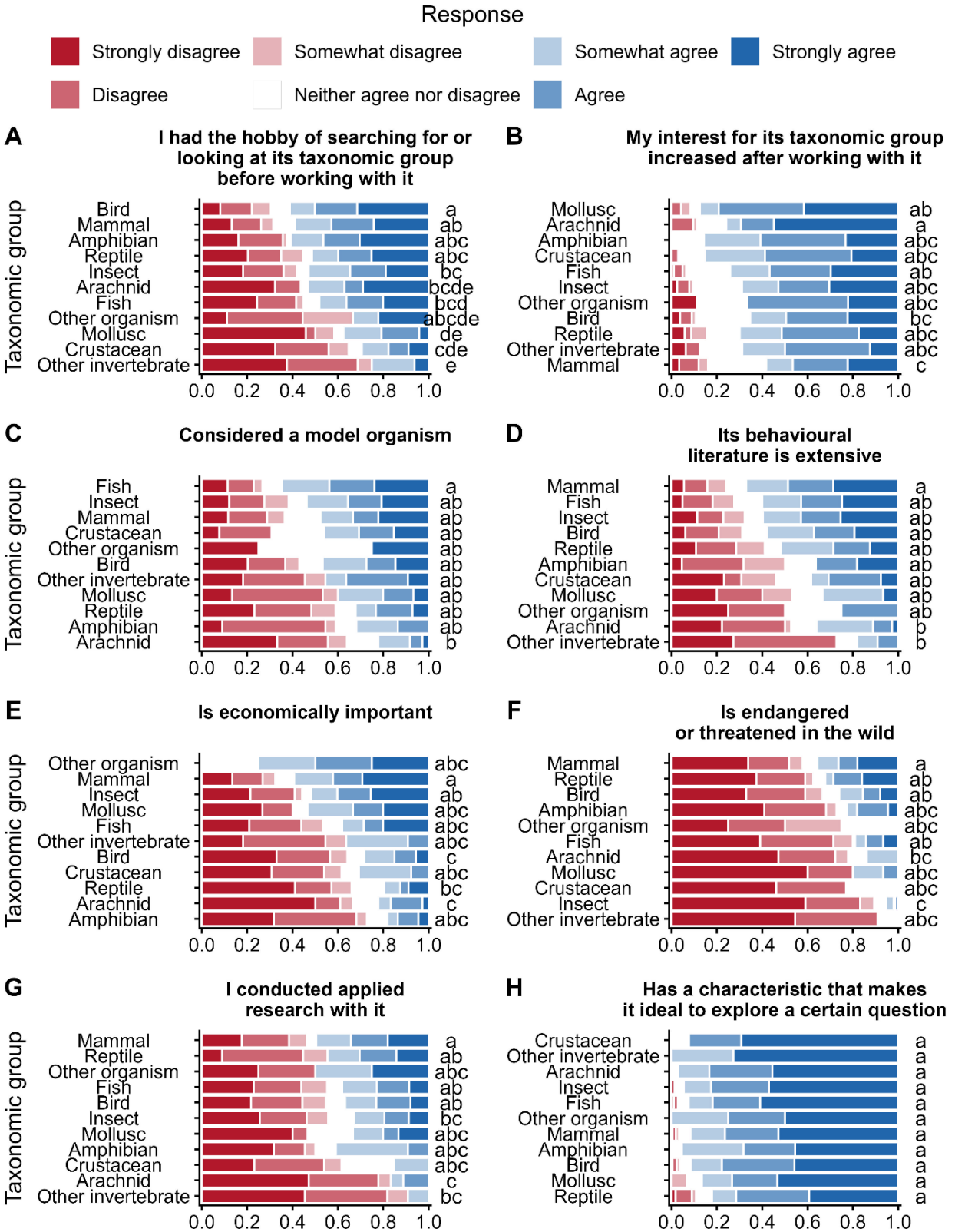
759 Ease of working with certain organisms. Regarding the organism they most recently worked with,

760 participants' agreement with statements related to the ease to locate it in the field (A), observe it

761 in the field (B), rear it in the lab (C), observe it in the lab (D), buy it from suppliers (E). Distinct

762 letters on the right-side of bars represent statistical differences between taxonomic groups within
763 that panel (z-values with $p < 0.05$ for all pairwise comparisons).

764



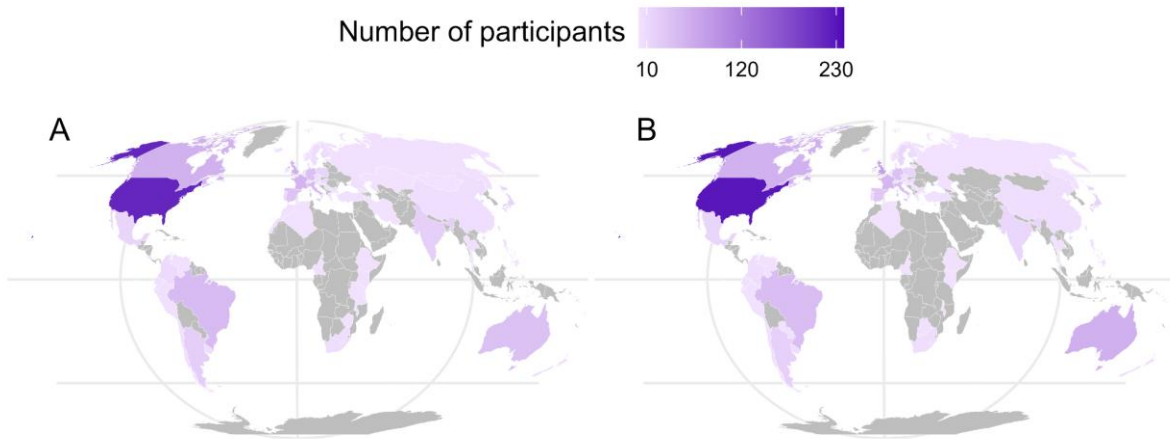
766 *Figure 6.*

767 Participants' agreement with various statements related to one or two organisms they worked with.

768 Distinct letters on the right-side of bars represent statistical differences between taxonomic groups

769 within that panel (z-values with $p < 0.05$ for all pairwise comparisons)

770 **Supplementary figures**

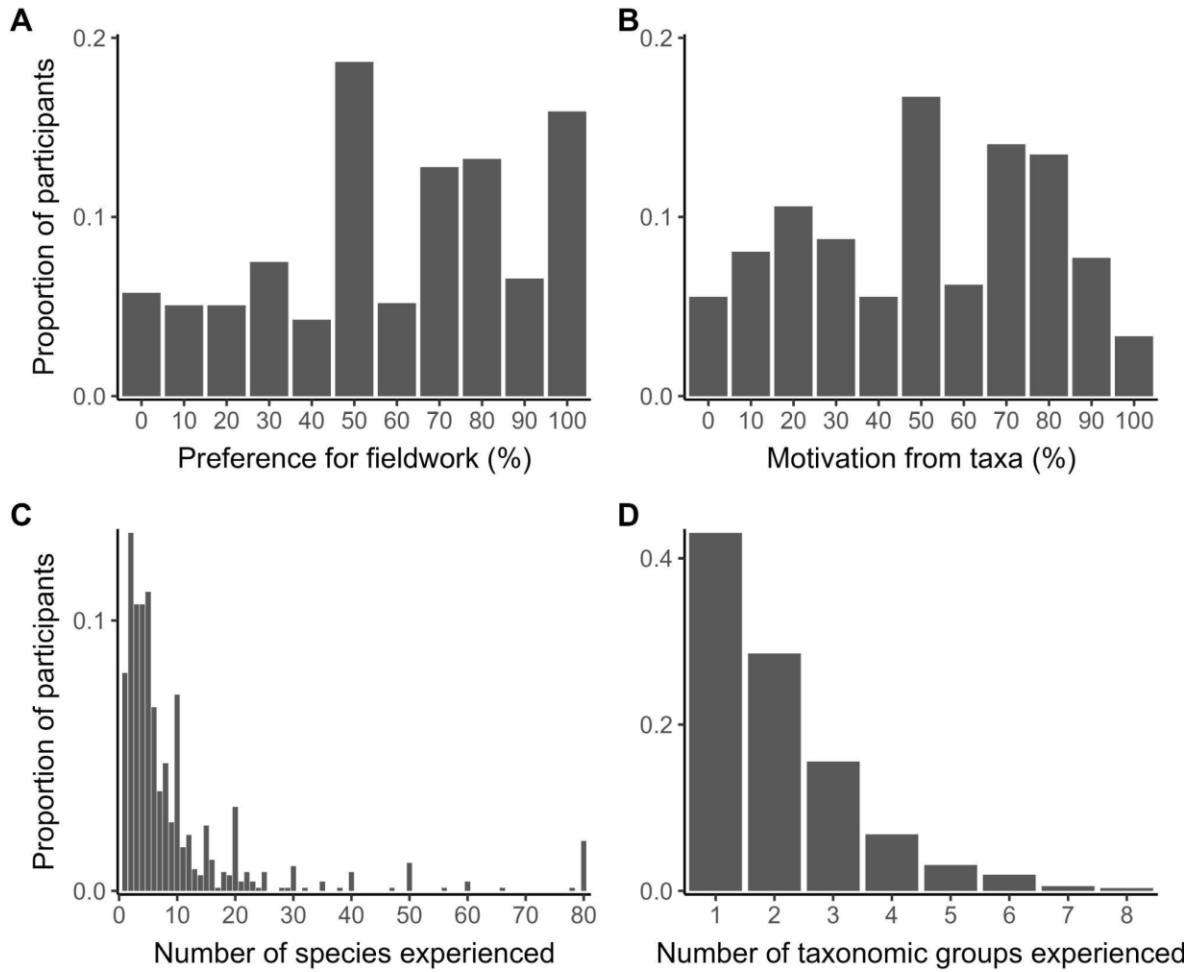


771

772 *Figure S1.*

773 Participants' country. Country in which participants spent most of their life before (A) and after

774 (B) being 18 years old. No participants mentioned countries or regions in grey.



775

776 *Figure S2.*

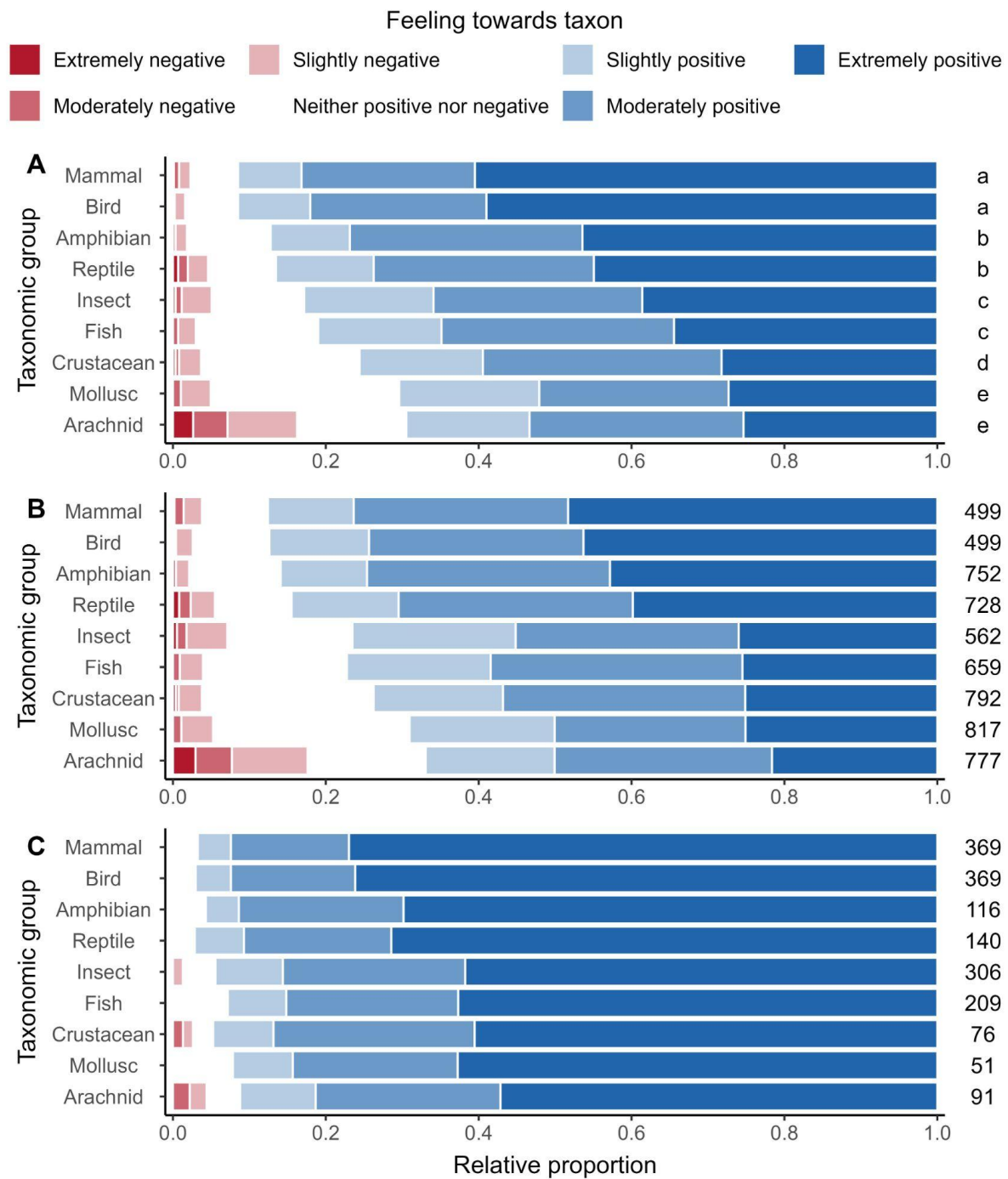
777 Participants' preferences, motivations, and overall experiences. Proportion of participants

778 expressing their preference for fieldwork over lab work (A) and their motivation from taxa over

779 other factors (B). Proportion of participants that experienced (i.e. collected behavioural data from)

780 distinct numbers of species (C), and of taxonomic groups (D).

781



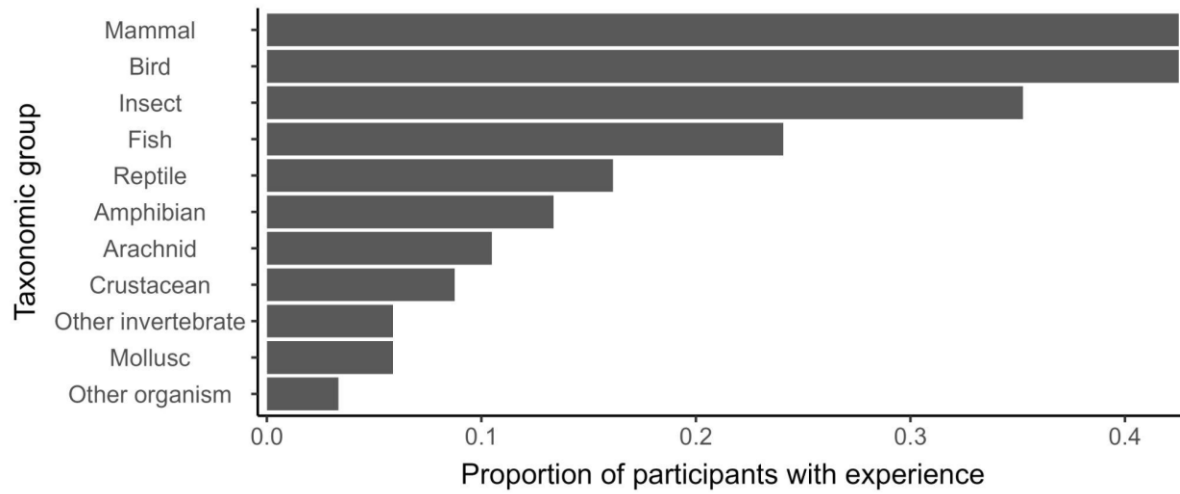
782

783 *Figure S3.*

784 Participants' feelings towards distinct animal groups. Responses from all participants together (A)

785 or by experience with the taxon in question (B: without, C: with). In panel A, distinct letters on

786 the right-side of bars represent statistical differences between taxonomic groups within that panel
787 (z-values with $p < 0.05$ for all pairwise comparisons). In panels B and C, numbers on the right-
788 side of bars represent the number of participants for each taxon in the subset presented, with a total
789 of 868 for each taxon.
790



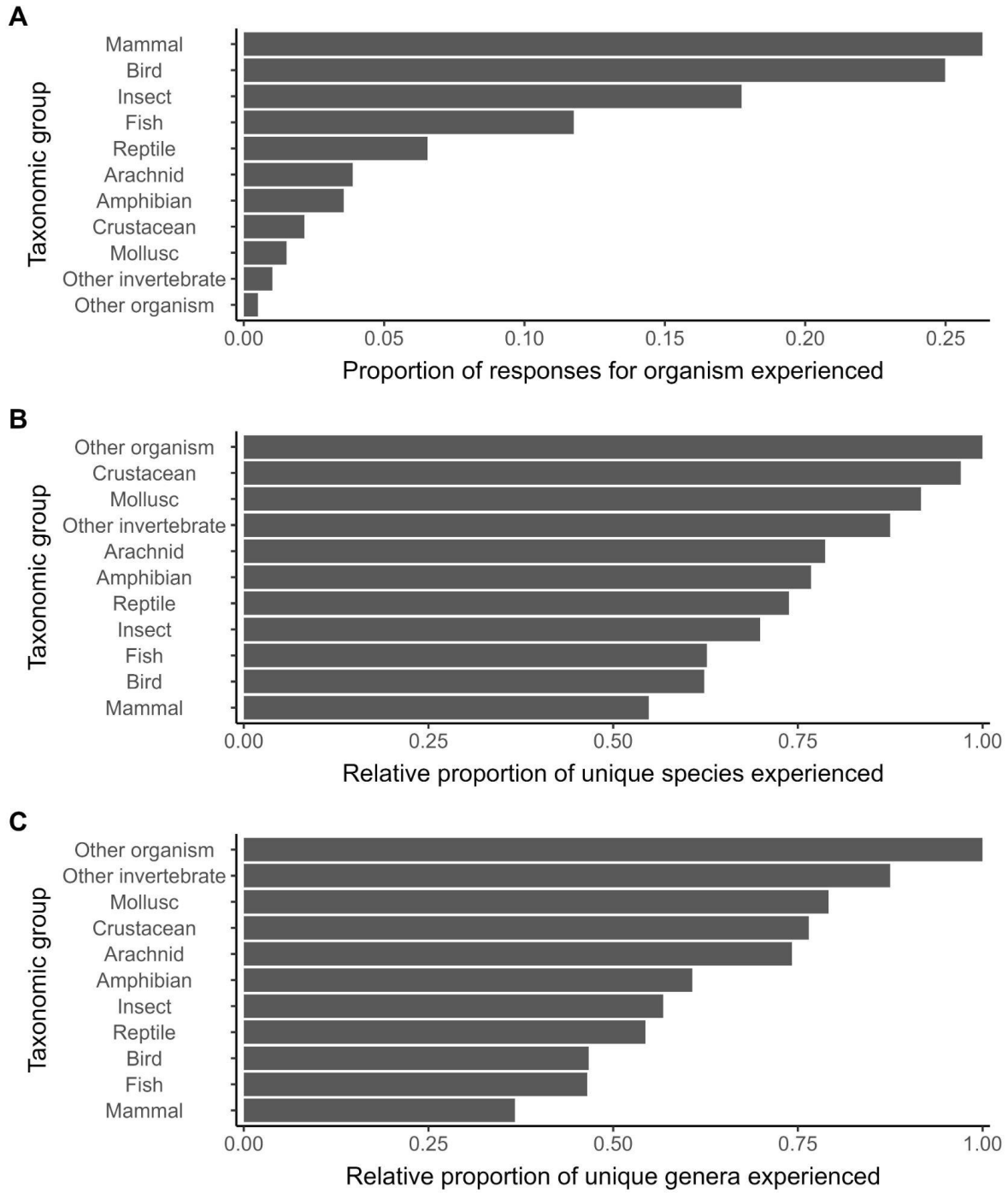
791

792 *Figure S4.*

793 Taxonomic groups with which participants had experience with. Note that each participant could

794 have experience with multiple taxa.

795



796

797 *Figure S5.*

798 Taxonomic groups of specific species that participants had experience with.

799 **Supplementary tables**

800 *Table S1.*

801 Pre-registered predictions (see Pollo & Kasumovic 2025).

Prediction	Supported?
Participants that work with vertebrate animals, especially birds and mammals, disclose that their interest is more strongly driven by taxon (opposed to other motivations) than other participants	No
Vertebrate animals, especially birds and mammals, elicit more positive feelings from participants than other taxa	Yes
Taxa that participants worked with elicit more positive feelings from them compared with other taxa	Yes
An employer, supervisor, or collaborator is responsible for selecting research species in most cases, except when the participant holds a permanent research position mentioning their most recently studied species.	Yes
Vertebrate animals, especially birds and mammals, are perceived as harder to rear and maintain in the lab, and easier to observe or record in the field, compared with other taxa	Mixed
Participants working with birds are more likely to have engaged in cultural activities involving the observation of their study organism before conducting research on it than participants working with other organisms	Mixed
Participants from anglophone countries are more likely to have engaged in cultural activities involving bird observation (i.e. birdwatching) before conducting research on birds than participants from non-anglophone countries.	No
A participant's preference for data collection in the field opposed to in the lab is positively related to their opinion of the species they mention being easy to be located/attracted and recorded/observed in the field, and negatively related to their opinion of the species being easy to be maintained and reared in the lab.	Yes
A participant's opinion that a species is considered a model organism by the scientific community is positively associated with their opinion of how extensive the behavioural literature on that species is.	Yes
A participant's opinion that a species is considered a model organism by the scientific community is positively associated with their opinion of how easy and accessible it is to obtain, maintain, and observe that species in the lab or in the field, but rarely both	Mixed
Organisms perceived as economically relevant are more frequently used in applied research than those perceived as economically irrelevant	Yes

802

803

804 *Table S2.*

805 Journals specialised in ethology and/or behavioural ecology (i.e. behaviour journals) or general
 806 ecology and evolution (i.e. E&E journals) used in our study to retrieve researchers' emails. ISSN
 807 stands for International Standard Serial Number.

Set	Journal	Publisher	Supporting society or institution	ISSN(s)
Behaviour	Acta Ethologica	Springer Nature	Portuguese Ethological Society	0873-9749, 1437-9546
	Animal Behaviour	Elsevier	Association for the Study of Animal Behaviour, Animal Behavior Society	1095-8282, 0003-3472
	Animal Cognition	Springer Nature	-	1435-9448, 1435-9456
	Behavioral Ecology	Oxford University Press	International Society for Behavioral Ecology	1045-2249, 1465-7279
	Behavioral Ecology and Sociobiology	Springer Nature	-	0340-5443, 1432-0762
	Behaviour	Brill	-	0005-7959, 1568-539X
	Behavioural Processes	Elsevier	-	0376-6357, 1872-8308
	Ethology	Wiley	Ethological Society	0179-1613, 1439-0310
	Ethology Ecology and Evolution	Taylor & Francis	-	0394-9370, 1120-6705, 1828-7131,
	Journal of Comparative Psychology	American Psychological Association	-	0735-7036, 1939-2087
	Journal of Ethology	Springer Nature	Japan Ethological Society	0289-0771, 1439-5444
	E&E	BMC Ecology and Evolution	BMC	-
Current Zoology		Oxford University Press	China Zoological Society	1674-5507, 2396-9814
Ecology and Evolution		Wiley	British Ecological Society	2045-7758
Ecology Letters		Wiley	Centre National de la Recherche Scientifique	1461-023X, 1461-0248
Evolution		Oxford University Press	Society for the Study of Evolution	0014-3820, 1558-5646
Evolution Letters		Oxford University Press	Society for the Study of Evolution	2056-3744
Functional Ecology		Wiley	British Ecological Society	0269-8463, 1365-2435
Journal of Animal Ecology		Wiley	British Ecological Society	0021-8790, 1365-2656

Journal of Evolutionary Biology	Oxford University Press	European Society for Evolutionary Biology	1010-061X, 1420-9101
Nature Ecology and Evolution	Springer Nature	-	2397-334X
The American Naturalist	The University of Chicago Press	American Society of Naturalists	0003-0147, 1537-5323

809 *Table S3.*

810 Model comparison regarding the number of species experienced by participants in our survey.

811 Several possible predictor variables were tested (see main text for details). “+” represents that the

812 predictor variable was included in models shown. *AICc* and *W* stand for models’ Akaike

813 information criterion for small sample sizes and their weight, respectively. Only selected models

814 (i.e. $\Delta AICc < 2$) are shown.

Rank	Predictor variables						<i>AICc</i>	$\Delta AICc$	<i>W</i>
	Number of taxonomic groups experienced	Age	Gender	Editorial experience	Preference for fieldwork	Motivation from taxa			
1	+	+	+	+		+	5510.72	0.00	0.50
2	+	+	+	+	+	+	5512.45	1.73	0.21

815

816

817 *Table S4.*

818 Model comparison regarding the number of taxonomic groups experienced by participants in our
819 survey. Several possible predictor variables were tested (see main text for details). “+” represents
820 that the predictor variable was included in models shown. *AICc* and *W* stand for models’ Akaike
821 information criterion for small sample sizes and their weight, respectively. Only selected models
822 (i.e. $\Delta AICc < 2$) are shown.

Rank	Predictor variables						<i>AICc</i>	$\Delta AICc$	<i>W</i>
	Number of species experienced	Age	Gender	Editorial experience	Preference for fieldwork	Motivation from taxa			
1	+	+		+		+	2681.64	0.00	0.25
2	+	+				+	2682.02	0.39	0.21
3	+	+		+	+	+	2683.16	1.53	0.12
4	+	+	+	+		+	2683.16	1.53	0.12
5	+	+	+			+	2683.49	1.86	0.10
6	+	+			+	+	2683.59	1.96	0.09

823

824 Table S5.

825 Model comparison regarding the fondness by participants in our survey for each taxonomic group.
 826 Several possible predictor variables were tested (see main text for details). “+” represents that the
 827 predictor variable was included in in models shown. *AICc* and *W* stand for models’ Akaike
 828 information criterion for small sample sizes and their weight, respectively. Only selected models
 829 (i.e. $\Delta AICc < 2$) are shown.

Taxon	Rank	Predictor variables							<i>AICc</i>	$\Delta AICc$	<i>W</i>
		Experience	Number of taxonomic groups experienced	Age	Gender	Editorial experience	Preference for fieldwork	Motivation from taxa			
Amphibian	1	+				+	+	2186.14	0.00	0.09	
Amphibian	2	+				+	+	2186.35	0.21	0.08	
Amphibian	3	+					+	2186.40	0.26	0.08	
Amphibian	4	+		+		+	+	2186.58	0.44	0.07	
Amphibian	5	+	+		+	+	+	2187.30	1.16	0.05	
Amphibian	6	+			+		+	2187.38	1.24	0.05	
Amphibian	7	+	+				+	2187.44	1.30	0.05	
Amphibian	8	+	+			+	+	2187.49	1.35	0.04	
Amphibian	9	+	+	+		+	+	2187.51	1.37	0.04	
Amphibian	10	+		+	+	+	+	2187.55	1.41	0.04	
Amphibian	11	+	+		+		+	2187.94	1.81	0.03	
Amphibian	12	+				+	+	2188.10	1.97	0.03	
Arachnid	1	+		+			+	2958.94	0.00	0.08	
Arachnid	2	+		+			+	2959.60	0.66	0.06	
Arachnid	3	+					+	2960.18	1.24	0.04	
Arachnid	4	+				+	+	2960.51	1.57	0.04	
Arachnid	5	+		+		+	+	2960.61	1.67	0.04	
Arachnid	6	+	+	+			+	2960.67	1.74	0.03	
Arachnid	7	+	+	+			+	2960.74	1.8	0.03	
Arachnid	8	+					+	2960.78	1.84	0.03	
Arachnid	9	+		+	+		+	2960.78	1.85	0.03	
Bird	1	+	+	+			+	1846.31	0.00	0.26	
Bird	2	+	+	+	+		+	1846.94	0.63	0.19	
Bird	3	+	+	+		+	+	1847.96	1.65	0.11	
Crustacean	1	+				+		2544.91	0	0.14	
Crustacean	2	+				+		2545.31	0.4	0.12	
Crustacean	3	+	+			+		2545.65	0.74	0.10	
Crustacean	4	+	+			+	+	2546.59	1.68	0.06	
Crustacean	5	+		+		+		2546.85	1.94	0.05	
Fish	1	+				+		2399.70	0.00	0.10	
Fish	2	+				+	+	2399.97	0.28	0.09	
Fish	3	+		+		+		2400.10	0.40	0.08	
Fish	4	+		+		+	+	2400.50	0.80	0.07	
Fish	5	+				+	+	2400.97	1.27	0.05	
Fish	6	+				+	+	2401.11	1.41	0.05	
Fish	7	+	+			+		2401.39	1.69	0.04	
Fish	8	+		+		+	+	2401.47	1.77	0.04	
Fish	9	+		+		+	+	2401.48	1.78	0.04	
Fish	10	+	+	+		+		2401.51	1.82	0.04	
Fish	11	+	+			+	+	2401.62	1.92	0.04	
Insect	1	+				+		2425.11	0.00	0.07	
Insect	2	+						2425.51	0.40	0.05	
Insect	3	+	+			+		2425.64	0.52	0.05	
Insect	4	+				+	+	2426.12	1.01	0.04	
Insect	5	+					+	2426.49	1.38	0.03	
Insect	6	+	+			+	+	2426.50	1.39	0.03	
Insect	7	+	+					2426.55	1.43	0.03	
Insect	8	+		+				2426.79	1.67	0.03	
Insect	9	+			+			2426.88	1.77	0.03	
Insect	10	+			+	+		2426.96	1.85	0.03	
Insect	11	+				+	+	2427.04	1.93	0.03	

Insect	12	+		+		+		2427.08	1.96	0.02
Mammal	1	+	+		+		+	1812.87	0.00	0.29
Mammal	2	+	+		+		+	1813.39	0.51	0.22
Mammal	3	+	+	+	+		+	1814.46	1.58	0.13
Mollusc	1	+				+		2629.47	0.00	0.09
Mollusc	2	+			+			2630.14	0.67	0.06
Mollusc	3	+				+	+	2630.27	0.80	0.06
Mollusc	4	+	+			+		2630.31	0.84	0.06
Mollusc	5	+			+		+	2630.70	1.23	0.05
Mollusc	6	+				+	+	2630.88	1.41	0.04
Mollusc	7	+	+		+			2631.02	1.54	0.04
Mollusc	8	+				+	+	2631.36	1.89	0.03
Reptile	1	+		+			+	2343.39	0.00	0.22
Reptile	2	+		+		+	+	2344.82	1.43	0.11
Reptile	3	+		+	+		+	2345.21	1.82	0.09
Reptile	4	+		+			+	2345.34	1.95	0.08

830

831

832 Table S6.

833 Model comparison regarding whether participants in our survey had experience with each
 834 taxonomic group. Several possible predictor variables were tested (see main text for details). “+”
 835 represents that the predictor variable was included in in models shown. *AICc* and *W* stand for
 836 models’ Akaike information criterion for small sample sizes and their weight, respectively. Only
 837 selected models (i.e. $\Delta AICc < 2$) are shown.

Taxon	Rank	Predictor variables						<i>AICc</i>	$\Delta AICc$	<i>W</i>
		Fondness	Number of taxonomic groups experienced	Age	Gender	Editorial experience	Preference for fieldwork			
Amphibian	1	+	+	+				504.54	0.00	0.28
Amphibian	2	+	+	+		+		506.13	1.59	0.12
Amphibian	3	+	+					506.37	1.83	0.11
Amphibian	4	+	+	+			+	506.43	1.89	0.11
Arachnid	1	+	+	+	+		+	463.99	0.00	0.17
Arachnid	2	+	+	+			+	464.48	0.49	0.13
Arachnid	3	+	+	+	+	+	+	464.81	0.82	0.11
Arachnid	4	+	+			+	+	464.99	1.00	0.10
Arachnid	5	+	+		+	+	+	465.30	1.31	0.09
Arachnid	6	+	+	+		+	+	465.39	1.40	0.08
Arachnid	7	+	+				+	465.56	1.58	0.08
Bird	1	+	+	+		+	+	969.83	0.00	0.28
Bird	2	+	+			+	+	970.79	0.97	0.17
Bird	3	+	+	+		+	+	971.43	1.60	0.13
Crustacean	1	+	+			+	+	385.54	0.00	0.19
Crustacean	2	+	+				+	386.71	1.17	0.11
Crustacean	3	+	+			+	+	387.00	1.47	0.09
Crustacean	4	+	+	+		+	+	387.39	1.86	0.08
Fish	1	+	+	+			+	732.05	0.00	0.15
Fish	2	+	+	+	+		+	732.49	0.44	0.12
Fish	3	+	+	+		+	+	732.79	0.73	0.10
Fish	4	+	+		+	+	+	732.8	0.74	0.10
Fish	5	+	+	+	+	+	+	733.27	1.21	0.08
Fish	6	+	+			+	+	733.52	1.47	0.07
Fish	7	+	+		+		+	733.64	1.59	0.07
Fish	8	+	+	+			+	733.94	1.88	0.06
Insect	1	+	+				+	868.92	0.00	0.19
Insect	2	+	+				+	869.39	0.47	0.15
Insect	3	+	+	+			+	869.64	0.72	0.13
Insect	4	+	+			+	+	870.03	1.12	0.11
Insect	5	+	+	+			+	870.28	1.37	0.09
Insect	6	+	+			+	+	870.56	1.64	0.08
Mammal	1	+	+	+	+		+	1057.46	0.00	0.2
Mammal	2	+	+	+			+	1058.16	0.7	0.14
Mammal	3	+	+	+	+		+	1058.31	0.85	0.13
Mammal	4	+	+	+	+	+	+	1058.61	1.15	0.11
Mammal	5	+	+	+			+	1058.76	1.3	0.1
Mammal	6	+	+	+		+	+	1059.36	1.90	0.08
Mollusc	1	+	+				+	318.33	0.00	0.12
Mollusc	2	+	+	+		+	+	318.68	0.36	0.10
Mollusc	3	+	+			+	+	318.74	0.41	0.10
Mollusc	4	+	+	+			+	319.48	1.16	0.07
Mollusc	5	+	+					319.68	1.35	0.06
Mollusc	6	+	+				+	319.81	1.48	0.06
Mollusc	7	+	+	+		+		320.06	1.73	0.05
Mollusc	8	+	+			+		320.10	1.77	0.05
Reptile	1	+	+					607.85	0.00	0.14
Reptile	2	+	+				+	608.20	0.35	0.12
Reptile	3	+	+			+		609.31	1.46	0.07
Reptile	4	+	+				+	609.31	1.46	0.07
Reptile	5	+	+			+	+	609.75	1.90	0.06

Other invertebrate	1	+			+	314.41	0.00	0.21
Other invertebrate	2	+			+	314.65	0.23	0.19
Other invertebrate	3	+	+		+	316.20	1.79	0.09
Other invertebrate	4	+			+	316.37	1.96	0.08
Other organism	1	+				211.88	0.00	0.13
Other organism	2	+	+			212.14	0.25	0.11
Other organism	3	+		+		212.76	0.88	0.08
Other organism	4	+	+	+		213.75	1.87	0.05
Other organism	5	+			+	213.77	1.89	0.05
Other organism	6	+		+		213.79	1.91	0.05
Other organism	7	+			+	213.82	1.93	0.05

838

839

840 *Table S7.*

841 Model comparison regarding whether the species mentioned by participants in our survey were
842 suggested by a supervisor (*vs.* other people or themselves). Two possible predictor variables were
843 tested (see main text for details). “+” represents that the predictor variable was included in models
844 shown. *AICc* and *W* stand for models’ Akaike information criterion for small sample sizes and
845 their weight, respectively.

Rank	Predictor variables		<i>AICc</i>	$\Delta AICc$	<i>W</i>
	Period	Version			
1	+		1871.87	0.00	0.61
2	+	+	1872.73	0.86	0.39
3		+	2013.53	141.66	0.00
4			2042.80	170.93	0.00

846

847

848 *Table S8.*

849 Model comparison regarding five statements (response variables) about the easiness to work with
 850 the most recent species experienced by participants in our survey. “+” represents that the predictor
 851 variable was included in models shown. *AICc* and *W* stand for models’ Akaike information
 852 criterion for small sample sizes and their weight, respectively.

Response variable	Rank	Predictor variables		<i>AICc</i>	$\Delta AICc$	<i>W</i>
		Genus mentions	Preference for fieldwork			
Easy to buy	1	+	+	2182.53	0.00	1.00
Easy to buy	2		+	2193.97	11.43	0.00
Easy to buy	3	+		2219.78	37.25	0.00
Easy to buy	4			2234.16	51.63	0.00
Easy to find in the field	1		+	3112.52	0.00	0.70
Easy to find in the field	2	+	+	3114.45	1.92	0.27
Easy to find in the field	3			3119.07	6.55	0.03
Easy to find in the field	4	+		3120.6	8.08	0.01
Easy to observe in the field	1	+	+	3139.08	0.00	0.93
Easy to observe in the field	2	+		3144.53	5.44	0.06
Easy to observe in the field	3		+	3148.09	9.00	0.01
Easy to observe in the field	4			3151.57	12.49	0.00
Easy to observe in the lab	1	+	+	2631.58	0.00	1.00
Easy to observe in the lab	2	+		2674.9	43.32	0.00
Easy to observe in the lab	3		+	2675.63	44.05	0.00
Easy to observe in the lab	4			2721.03	89.45	0.00
Easy to rear in the lab	1	+	+	2675.73	0.00	1.00
Easy to rear in the lab	2	+		2709.14	33.41	0.00
Easy to rear in the lab	3		+	2717.62	41.89	0.00

853

854

855 *Table S9.*

856 Coefficients from models selected regarding five statements (response variables) about the
857 easiness to work with the most recent species experienced by participants in our survey (see also
858 Table S8).

Response variable	Predictor variable	Coefficient	SE	z-score	p-value
Easy to buy	Genus mentions	0.45	0.07	6.02	<0.001
Easy to buy	Preference for fieldwork	-0.35	0.10	-3.68	<0.001
Easy to find in the field	Genus mentions	0.14	0.05	2.90	<0.001
Easy to observe in the field	Genus mentions	0.11	0.04	2.68	0.01
Easy to observe in the field	Preference for fieldwork	0.24	0.07	3.33	<0.001
Easy to observe in the lab	Genus mentions	0.39	0.06	6.64	<0.001
Easy to observe in the lab	Preference for fieldwork	-0.58	0.09	-6.76	<0.001
Easy to rear in the lab	Genus mentions	0.35	0.06	5.92	<0.001
Easy to rear in the lab	Preference for fieldwork	-0.57	0.09	-6.63	<0.001

859

860 **Appendix 1 – Survey**

861

862 [Participant Information Sheet and Consent was applied before the survey]

863

864 *PART 1: Eligibility*

865

866 **1. Have you ever directly collected behavioural data from non-human organisms that led to**
867 **at least one research output* of which you were an author?**

868 *Research outputs include conference posters or presentations, preprints, peer-reviewed
869 publications, dissertations, theses, or books.

870 Yes [proceed]

871 No [terminate the survey]

872

873 *PART 2: Demographics*

874

875 **2. How old are you?**

876 [fill]

877

878 **3. How do you identify?**

879 Agender

880 Gender fluid

881 Man

882 Non binary

883 Woman

884 Other (please specify) [fill]

885

886 **4. In which country have you spent most of your life UNTIL you were 18 years old? (please**
887 **elect only one country if multiple countries apply)**

888 [fill]
889

890 **5. In which country have you spent most of your life AFTER turning 18 years old? (please**
891 **elect only one country if multiple countries apply)**

892 [fill]
893

894 **6. What is your current highest level of education? (i.e. obtained degree; do NOT consider**
895 **current studies)**

- 896 High school/Technical school
- 897 Undergraduate/Honours degree
- 898 MSc/Master's degree
- 899 PhD/Doctoral degree
- 900

901 **7. [if Q6 is PhD/Doctoral degree] In what year did you complete your PhD?**

902 [fill]
903

904 *PART 3: General preferences and experiences*

905

906 **8. Are you or have you ever been an editor of any scientific journals that at least occasionally**
907 **publish ethological or behavioural research?**

- 908 Yes
- 909 No

910

911 **9. Based purely on personal enjoyment, how would you split your time collecting behavioural**
912 **data between the field and the lab?**

- 913 ○ 100% in the field, 0% in the lab
- 914 ○ 90% in the field, 10% in the lab
- 915 ○ 80% in the field, 20% in the lab
- 916 ○ 70% in the field, 30% in the lab
- 917 ○ 60% in the field, 40% in the lab
- 918 ○ 50% in the field, 50% in the lab
- 919 ○ 40% in the field, 60% in the lab
- 920 ○ 30% in the field, 70% in the lab
- 921 ○ 20% in the field, 80% in the lab
- 922 ○ 10% in the field, 90% in the lab
- 923 ○ 0% in the field, 100% in the lab
- 924

925 **10. How much does your desire to work with certain taxonomic groups drive your**
 926 **behavioural research interests compared with other possible motivations (e.g. exploration of**
 927 **specific topics pertinent to most organisms, professional prospects, etc)?**

- 928 ○ 100% driven by certain taxonomic group(s), 0% driven by other motivations
- 929 ○ 90% driven by certain taxonomic group(s), 10% driven by other motivations
- 930 ○ 80% driven by certain taxonomic group(s), 20% driven by other motivations
- 931 ○ 70% driven by certain taxonomic group(s), 30% driven by other motivations
- 932 ○ 60% driven by certain taxonomic group(s), 40% driven by other motivations
- 933 ○ 50% driven by certain taxonomic group(s), 50% driven by other motivations
- 934 ○ 40% driven by certain taxonomic group(s), 60% driven by other motivations
- 935 ○ 30% driven by certain taxonomic group(s), 70% driven by other motivations
- 936 ○ 20% driven by certain taxonomic group(s), 80% driven by other motivations
- 937 ○ 10% driven by certain taxonomic group(s), 90% driven by other motivations
- 938 ○ 0% driven by certain taxonomic group(s), 100% driven by other motivations
- 939

940 **11. How would you rate your feelings towards the following animals on a spectrum ranging**
 941 **from negative (e.g., anxiety, disgust, fear) to positive (e.g., admiration, cuteness, fascination),**
 942 **with ambivalent feelings in between?**

Taxonomic group	Extremely negative	Moderately negative	Slightly negative	Neither positive nor negative	Slightly positive	Moderately positive	Extremely positive
Amphibians							
Arachnids							
Birds							

Crustaceans							
Fish							
Insects							
Mammals							
Molluscs							
Reptiles							

943

944 **12. From how many different non-human organisms (at the species level) have you directly**
945 **collected behavioural data that led to at least one research output* for which you were an**
946 **author?**

947 *Research outputs include conference posters or presentations, preprints, peer-reviewed
948 publications, dissertations, theses, or books.

949 [fill]

950

951 **13. Select all taxonomic groups from which you directly collected behavioural data that led**
952 **to at least one research output* of which you were an author.**

953 *Research outputs include conference posters or presentations, preprints, peer-reviewed
954 publications, dissertations, theses, or books.

- 955 ▪ Amphibian
- 956 ▪ Arachnid
- 957 ▪ Bird
- 958 ▪ Crustacean
- 959 ▪ Fish
- 960 ▪ Insect
- 961 ▪ Invertebrate not on this list
- 962 ▪ Mammal
- 963 ▪ Mollusc
- 964 ▪ Non-animal organism
- 965 ▪ Reptile

966

967 *PART 4: Experiences and opinions regarding specific studied organisms*

968

969 If q12 answer is equal or greater than 2, some of the following questions (q15-18 and q27-28) will
970 be asked twice. However, repeated questions will be related to the species that the participant
971 worked with least recently (see q14).

972

973 **14. Specify the scientific name (genus and epithet, e.g. *Homo sapiens*) of the [most/least]**
974 **recent non-human species from which you directly collected behavioural data that led to at**
975 **least one research output* of which you were an author (elect one species if multiple apply).**

976 *Research outputs include conference posters or presentations, preprints, peer-reviewed
977 publications, dissertations, theses, or books.

978 [fill]

979

980 **15. [q14 answer] is a/an**

- 981 Amphibian
982 Arachnid
983 Bird
984 Crustacean
985 Fish
986 Insect
987 Invertebrate not on this list
988 Mammal
989 Mollusc
990 Non-animal organism
991 Reptile
992

993 **16. Approximately how many peer-reviewed articles have you published involving**
994 **behavioural data that you directly collected from [q14 answer]?**

- 995 0
996 1
997 2

- 998 ○ 3
- 999 ○ 4 or more

1000

1001 **17. When did you first directly collect behavioural data from [q14 answer]?**

- 1002 ○ During or before my PhD
- 1003 ○ After my PhD, but not while holding a permanent research position (e.g. unemployed,
1004 postdoc)
- 1005 ○ After my PhD, while holding a permanent research position (e.g. lecturer, professor)

1006

1007 **18. Which of these situations best describes how you INITIALLY decided to work with [q14
1008 answer]?**

- 1009 ○ My employer, supervisor, or collaborator used this species for research and/or suggested
1010 me to use this species in my research
- 1011 ○ Someone, but NOT my employer, supervisor, or collaborator, showed or told me about
1012 this species and/or suggested that I used it in my research
- 1013 ○ I had the idea to use this species in my research before anyone else directly suggested it
1014 or influenced me to do it

1015

1016 Questions below are answered in a 7-point Likert scale (i.e. strongly disagree to strongly agree):

1017

1018 **19. Reaching the place where [q14 answer] individuals are expected to be, and then locating
1019 or attracting them in the field is easy/accessible.**

1020

1021 **20. Once [q14 answer] individuals are located or attracted, observing or recording their
1022 behaviours in the field is easy/accessible.**

1023

1024 **21. Obtaining [q14 answer] individuals from suppliers (e.g. other research teams, farms,
1025 fisheries, pet stores) is easy/accessible.**

1026

1027 **22. Rearing and maintaining [q14 answer] individuals in the lab is easy/accessible.**

1028

1029 **23. Observing or recording [q14 answer] individuals in the lab is easy/accessible.**

1030

1031 **24. [q14 answer] is considered a model organism by the scientific community.**

1032

1033 **25. The behavioural literature on [q14 answer] is extensive.**

1034

1035 **26. [q14 answer] possesses a specific characteristic that makes it an ideal species to answer a**
1036 **question I am or was interested in.**

1037

1038 **27. I already had the hobby of going out specifically to search for or look at [q15 answer]s**
1039 **before I started working with [q14 answer].**

1040

1041 **28. My interest for [q15 answer]s increased after working with [q14 answer].**

1042

1043 **29. [q14 answer] is an economically important species (e.g. used commercially, provides**
1044 **ecological services, considered a pest or a disease vector).**

1045

1046 **30. [q14 answer] is endangered/threatened in the wild.**

1047

1048 **31. The behavioural research I have conducted with [q14 answer] can be considered applied**
1049 **(e.g. pest control, conservation).**

1050

1051 **32. Please select the option “agree”. [attention question]**

1052 **Appendix 2 – Additional discussion**

1053 Contradictory results we found, such that preference for fieldwork was positively related
1054 to fondness for arachnids but negatively related to the likelihood of working with them, may
1055 indicate that researchers are unaware that they can use neglected taxa to pursue their preferences
1056 (e.g. work with arachnids in the field), indicating that a diverse taxonomic education is key to
1057 addressing taxonomic bias in ecology and evolution. At finer taxonomic levels, participants with
1058 a stronger preference for fieldwork were more likely to report that the species they most recently
1059 worked with is easy to observe in the field, but harder to work with in the laboratory. This suggests
1060 that researchers select specific organisms to match their professional preferences also within a
1061 taxonomic group.

1062 Justifying the choice of study species has become a staple in research manuscripts (Mann
1063 2015), which may explain why most surveyed researchers agreed that the species they most
1064 recently worked with possesses a specific characteristic that makes it ideal to explore a certain
1065 topic. However, it is possible that other species could be equally useful for addressing the same
1066 questions, as similar traits often occur across diverse taxa, including (apparently) unique
1067 characteristics. For example, biologists would intuitively use mammals to understand the evolution
1068 of milk-like secretions as a form of parental care (Clutton-Brock 1991), but species of other
1069 taxonomic groups can also express this behaviour (e.g. cecilians: Mailho-Fontana et al. 2024;
1070 cockroaches: Stay and Coop 1974; spiders: Chen et al. 2018). We thus argue that we could discover
1071 more species with astonishing features by exploring the existent biodiversity, which would grant
1072 us new ideal organisms to study, instead of simply marketing a few species as ideal for a topic
1073 (especially those that have been investigated for decades by several people).

1074 We also investigated the relationship between age and gender on researchers' taxonomic
1075 preferences and experiences, even though we had no hypotheses on how these variables relate to
1076 taxonomic bias. We found that, on average, older participants in our survey worked with more
1077 species and taxonomic groups than their younger counterparts, which may simply reflect that they
1078 had more time to do so. However, it is also possible that researchers have been recently using
1079 fewer taxa in their research, making their experiences less taxonomically diverse compared with
1080 researchers in previous decades. We also observed that increasing age was related to greater
1081 fondness for birds, lower fondness for amphibians, and greater likelihood of having worked with
1082 mammals, which might show that distinct generations of researchers have different perspectives.
1083 Women that participated in our survey worked with fewer species and taxonomic groups than men,
1084 possibly reflecting the difficulties women face in academia (Casad et al. 2021; Spoon et al. 2023)
1085 if these obstacles limit their opportunities for taxonomic diversification. Although the literature
1086 suggests that women are more likely to have animal phobias than men (Fredrikson et al. 1996), we
1087 found no gender differences in the perception of different taxonomic groups, except for mammals,
1088 which women viewed more favourably than men (Fig. 3A).

1089 In our survey, we also verified the relationship between participants' editorial experience
1090 and their taxonomic focus. Given that editors are the main gatekeepers in the publishing system,
1091 they have more power than other individuals to change patterns of taxonomic bias in the literature.
1092 Participants with editorial experience worked with more species but not with more taxonomic
1093 groups than other participants (Fig. 2), showing that these individuals are not specifically selected
1094 because of the diversity of their experiences at a high taxonomic level. Despite participants with
1095 editorial experience having a more favourable perception of some animal groups (crustaceans,
1096 fish, molluscs) than others, they were more likely to work with birds than their counterparts. If

1097 editors are more likely to accept research manuscripts that match their personal experiences for
1098 publication, then taxonomic bias in the literature can worsen. This is certainly possible as biases
1099 in the publication process within ecology and evolution have been documented (Fox et al. 2023;
1100 Srivastava et al. 2024), highlighting the need for greater transparency in editorial decisions to
1101 assess whether individual preferences and experiences influence publication patterns.