

# Social learning strategies regulate the wisdom and madness of interactive crowds

Wataru Toyokawa<sup>1,2,3,\*</sup>, Andrew Whalen<sup>1</sup>, and Kevin N. Laland<sup>1</sup>

<sup>1</sup>*School of Biology, University of St Andrews, Harold Mitchel Building, St Andrews, Fife, KY16 9TH,  
Scotland*

<sup>2</sup>*Japan Society for the Promotion of Science, Kojimachi, Chiyoda-ku, Tokyo, 102-0083, Japan*

<sup>3</sup>*Department of Evolutionary Studies of Biosystems, School of Advanced Sciences, SOKENDAI (The  
Graduate University for Advanced Studies), Shonan Village, Hayama, Kanagawa 240-0193, Japan*

\* *corresponding author: wt25@st-andrews.ac.uk*

## Abstract

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14

Why groups of individuals sometimes exhibit collective ‘wisdom’ and other times maladaptive ‘herding’ is an enduring conundrum. Here we show that this apparent conflict is regulated by the social learning strategies deployed. We examined the patterns of human social learning through an interactive online experiment with 699 participants, varying both task uncertainty and group size, then used hierarchical Bayesian model-fitting to identify the individual learning strategies exhibited by participants. Challenging tasks elicit greater conformity amongst individuals, with rates of copying increasing with group size, leading to high probabilities of herding amongst large groups confronted with uncertainty. Conversely, the reduced social learning of small groups, and the greater probability that social information would be accurate for less-challenging tasks, generated ‘wisdom of the crowd’ effects in other circumstances. Our model-based approach provides evidence that the likelihood of collective intelligence versus herding can be predicted, resolving a longstanding puzzle in the literature.

15 Understanding the mechanisms that account for accurate collective decision-making amongst  
16 groups of animals – ‘collective intelligence’ – has been a central focus of animal behaviour re-  
17 search<sup>1-5</sup>. There are a large number of biological examples showing that collectives of poorly  
18 informed individuals can achieve a high performance in solving cognitive problems under un-  
19 certainty<sup>6-10</sup>. Although these findings suggest fundamental cognitive benefits of grouping<sup>11</sup>,  
20 there is also a long-standing recognition, especially for humans, that interacting individuals may  
21 sometimes be overwhelmed by the ‘*extraordinary popular delusions and madness of crowds*’<sup>12</sup>.  
22 Herd behaviour (i.e. an alignment of thoughts or behaviours of individuals in a group) occurs  
23 because individuals imitate each other<sup>13-15</sup>, even if each is a rational decision-maker<sup>16</sup>. Imita-  
24 tion is thought to be a cause of financial bubbles<sup>12;17</sup>, ‘groupthink’<sup>18</sup> and volatility in cultural  
25 markets<sup>19;20</sup>. More generally, interdependence between individual decisions may undermine the  
26 wisdom of crowds effect<sup>21</sup> (but see<sup>22</sup>), whilst potential disadvantages of information transfer are  
27 well-recognised in the biological literature<sup>23;24</sup>. It seems that information transmission among  
28 individuals, and making decisions collectively, is a double-edged sword: combining decisions  
29 may provide the benefits of collective intelligence, but at the same time, increase the risk of an  
30 informational cascade<sup>16</sup>. Collectively, an understanding of whether and, if so, how it is possible  
31 to prevent or reduce the risk of maladaptive herding, while concurrently keeping or enhancing  
32 collective intelligence, is largely lacking.

33 A balance between using individual and social information may play a key role in deter-  
34 mining the trade-off between collective wisdom and ‘madness’<sup>25</sup>. If individuals are too reliant  
35 on copying others’ behaviour, any idea, even a maladaptive one, can propagate in the social  
36 group through positive feedbacks<sup>2;26</sup>. For instance, disproportionately strong positive responses

37 to recruitment signals in social insects have been shown to trap the whole colony to exploit a  
38 suboptimal, out-dated resource<sup>24;27</sup>. Likewise, conformity-biased transmission in humans and  
39 other animals can potentially lead groups to converge on a maladaptive behaviour<sup>16;23;28;29</sup>. On  
40 the other hand, however, if individuals completely ignore social information so as to be indepen-  
41 dent, they will fail to exploit the benefits of aggregating information through social interactions.  
42 The extent to which individuals should use social information should fall between these two  
43 extremes. Evolutionary models predict that the balance between independence and interdepen-  
44 dence in collective decision-making may be changeable, contingent upon the individual-level  
45 flexibility and inter-individual variability associated with the social learning strategies deployed  
46 in diverse environmental states<sup>28;30;31</sup>.

47 Experimental studies report that animals (including humans) increase their use of social in-  
48 formation as the returns from asocial learning become more unreliable<sup>32-37</sup>, whilst theory and  
49 data suggest that the benefits to individuals of social learning increase with group size<sup>34;38-42</sup>.  
50 Selectivity in the predicted use of social information may impact on collective decision-making  
51 because slight differences in the parameter values of social information use are known to be able  
52 to alter qualitatively the collective behavioural dynamics<sup>1;2;5;43;44</sup>. Therefore, researchers should  
53 expect populations to exhibit a higher risk of being trapped with maladaptive behaviour with  
54 increasing group size and decreasing reliability of asocial learning (and concomitant increased  
55 reliance on social learning).

56 From the viewpoint of the classic wisdom of crowds theory, increasing group size may in-  
57 crease collective accuracy<sup>45-48</sup>. The relative advantage of the collective over solitary individuals  
58 may also be highlighted by increased task difficulty, because there would be more room for the

59 performance of difficult tasks to be improved compared to easier tasks in which high accuracy  
60 can be achieved by asocial learning only. To understand the collective decision performance of  
61 social learners fully requires fine-grained quantitative studies of social learning strategies and  
62 their relations to collective dynamics, linked to sophisticated computational analysis.

63 The aims of this study were twofold. First, we set to test the hypothesis that the circumstances  
64 under which collective decision making will generate ‘wisdom’ can be predicted with knowledge  
65 of the precise learning strategies individuals deploy, through a combination of experimentation  
66 and theoretical modelling. The choice of an abstract decision-making task allowed us to imple-  
67 ment a computational modelling approach, which has been increasingly deployed in quantitative  
68 studies of animal social learning strategies<sup>35;49-51</sup>. In particular, computational modelling al-  
69 lowed us to conduct a parametric description of different information-gathering processes and  
70 to estimate the parameter values at an individual-level resolution. This approach allows us to  
71 characterise the complex relationship between individual-level decision, learning strategies and  
72 collective-level behavioural dynamics.

73 Second, we added resolution to our analyses by manipulating both task uncertainty and group  
74 size in our web-based experiments with adult human subjects, predicting that these factors would  
75 induce heavier use of social information in humans, and thereby alter the balance between col-  
76 lective intelligence and the risk of inflexible herding. To do this, we focused on human groups  
77 exposed to a simple gambling task called a multi-player ‘multi-armed bandit’, where both asocial  
78 and social sources of information were available<sup>35;51;52</sup>. Through development of an interactive,  
79 web-based collective decision-making task, and use of hierarchical Bayesian statistical meth-  
80 ods in fitting our computational model to the experimental data, we identify the individual-level

81 learning strategies of participants as well as quantify variation in different learning parameters,  
82 allowing us to conduct an informed exploration of the population-level outcomes. The results  
83 provide clear evidence that the collective behavioural dynamics can be predicted with knowledge  
84 of human social learning strategies.

85 Below, we firstly deploy agent-based simulation to illustrate how the model parameters re-  
86 lating to social learning can in principle affect the collective-level behavioural dynamics. The  
87 simulation provides us with precise, quantitative predictions concerning the complex relation-  
88 ship between individual behaviour and group dynamics. Second, we present the findings of  
89 a multi-player web-based experiment with human participants that utilises the gambling task  
90 framework. Applying a hierarchical Bayesian statistical method, we estimated the model's pa-  
91 rameters for each of 699 different individuals, allowing us to (i) examine whether and, if so, how  
92 social information use is affected by different group size and task uncertainty, and (ii) whether  
93 and how social-information use affects both collective intelligence and the risk of maladaptive  
94 herding.

## 95 **1 Results**

### 96 **1.1 The relationship between social learning and the collective behaviour**

97 Figure 1 shows the relationship between the average decision accuracy and individual-level social  
98 information use obtained from our individual-based model simulations, highlighting the trade-  
99 off between accuracy and flexibility of collective decision-making. When the mean *conformity*  
100 *exponent* is small (i.e.  $\bar{\theta} = (\sum_i \theta_i)/individuals = 1$ ), large groups are able to recover the decision

101 accuracy quickly as do small groups after the location of the optimal option has been switched,  
102 whereas overall improvement by increasing group size in decision accuracy is subtle when the  
103 average *social learning weight* is also small (i.e.  $\bar{\sigma} = (\sum_i \sum_t \sigma_{i,t}) / (\text{individuals} \times \text{rounds}) = 0.3$ ;  
104 Figure 1A and 1C). On the other hand, when both the conformity exponent  $\bar{\theta}$  and the social  
105 learning weight  $\bar{\sigma}$  are large, average performance is no longer monotonically improving with in-  
106 creasing group size, and it is under these circumstances that the strong herding effect becomes  
107 prominent (Figure 1D). Although the high conformity bias with low social learning weight makes  
108 large groups more accurate before the environment changes, larger groups are less flexible in per-  
109 formance recovery (Figure 1B). The pattern is robust for other parameter regions (Supplementary  
110 Figure 2).

111 **Figure 2** indicates that when both  $\bar{\theta}$  and  $\bar{\sigma}$  are large the collective choices converged either on  
112 the good option or on one of the poor options almost randomly, regardless of the option's quality,  
113 and that once individuals start converging on an option the population gets stuck. As a result,  
114 the distribution of the groups' average performance over the replications becomes a bimodal  
115 'U-shape'. Interestingly, however, the maladaptive herding effect remains relatively weak in  
116 smaller groups (see Figure 1D; the dotted line). This is because the majority of individuals in  
117 smaller groups (i.e. two individuals out of three) are more likely to break the cultural inertia by  
118 simultaneously exploring another option by chance than are the majority in larger groups (e.g.  
119 six out of ten).

120 In summary, the model simulation suggests an interaction between social learning weight  
121  $\bar{\sigma}$  and conformity exponent  $\bar{\theta}$  on decision accuracy and the risk of inflexible herding. When  
122 the conformity exponent is not too large, increasing group size can increase decision accuracy

123 while concurrently retaining decision flexibility across a broad range of the mean social learning  
124 weights. When the conformity bias becomes large, however, the risk of inflexible herding arises,  
125 and, when both social learning parameters are large, collective intelligence is rare and inflexible  
126 herd behaviour dominates.

## 127 **1.2 Collective performance of human participants**

128 Figure 3A shows behavioural dynamics of human participants in different group sizes and dif-  
129 ferent task uncertainty conditions (see Supplementary Figure 3 for each group's behaviour). The  
130 average decision performance of collectives (i.e. group size  $\geq 2$ ) exceeded that of solitary in-  
131 dividuals (i.e. group size = 1) in the Moderate-uncertainty condition (i.e. the 95% Bayesian  
132 CI of  $\xi_t$  exceeds 0 at regions  $t \in [9, 40]$  and  $[67, 70]$ ; Figure 3B). In other uncertainty condi-  
133 tions, no global positive effect of grouping was observed, suggesting that collective intelligence  
134 was prominent only in the Moderate-uncertainty condition. However, the main effect of group  
135 size was positive in the post-change period of the Low-uncertainty condition (mean and the 95%  
136 Bayesian CI of  $\omega_2 = 0.67 [0.44, 0.91]$ ; Table 1), suggesting that the average performance of large  
137 groups (e.g.  $12 \leq \text{group size} \leq 16$ ) were better, and hence more flexible, than smaller groups  
138 and solitaries (Figure 3A). On the other hand, in the Moderate-uncertainty condition, the aver-  
139 age performance of the collectives dropped below that of the solitaries after the environmental  
140 change (i.e.  $\xi_t < 0$  at a region  $t \in [42, 45]$ ; Figure 3B). Also, the main effect of group size  
141 was negative in the post-change period (mean and the 95% Bayesian CI of  $\omega_2 = -0.26 [-0.44,$   
142  $-0.11]$ ; Table 1), suggesting that larger groups were more likely to get stuck in the out-dated op-  
143 tion in the Moderate-uncertainty condition. In the High-uncertainty condition, the main effect of

144 group size was positive in the prior-change period and negative post-change (mean and the 95%  
145 Bayesian CIs are  $\omega_1 = 0.07$  [0.00, 0.15] and  $\omega_2 = -0.10$  [-0.17, -0.02]; Table 1), although the  
146 effect size was too small to differentiate performances of different group sizes visually (Figure  
147 3A). Using monetary earnings as an outcome variable of decision performance did not change  
148 our conclusions qualitatively (supporting Supplementary Figure 4 and Supplementary Table 2).

149 Our phenomenological model regression established that manipulating both task uncertainty  
150 and group size indeed affected the collective decision dynamics. Below, we address whether or  
151 not the pattern could be explained with knowledge of human social learning strategies estimated  
152 through our learning and decision-making computational model.

### 153 1.3 Estimation of human social information use

154 Using posterior estimation values obtained by the hierarchical Bayesian model fitting method  
155 (Table 2), we were able to categorise the participants as deploying one of three different learn-  
156 ing strategies based on their fitted conformity exponent values; namely, the ‘positive frequency-  
157 dependent copying’ strategy ( $\theta_i \gg 0$ ), the ‘negative-frequency dependent copying’ strategy  
158 ( $\theta_i \ll 0$ ) and the ‘random choice’ strategy ( $\theta_i \approx 0$ ). Note that we could not reliably detect  
159 the ‘weak positive’ frequency-dependent strategy ( $0 < \theta_i \leq 1$ ) due to the limitation of statisti-  
160 cal power (Supplementary Figure 5). Some individuals whose ‘true’ conformity exponent fell  
161 between zero and one would have been categorised as exhibiting a random choice strategy (Sup-  
162 plementary Figure 7). Individuals identified as exhibiting a positive frequency-dependent copiers  
163 were mainly those whose conformity exponent was larger than one ( $\theta_i > 1$ ).

164 Figure 4A show the estimated frequencies of different learning strategies. Generally speak-

165 ing, participants were more likely to utilize a positive frequency-dependent copying strategy  
166 than the other two strategies (the 95% Bayesian CI of the intercept of the GLMM predicting the  
167 probability to use the positive frequency-dependent copying strategy is above zero, [1.05, 2.50];  
168 Supplementary Table 4). We found that positive frequency-dependent copying decreased with  
169 increasing task uncertainty (the 95% Bayesian CI of task uncertainty effect is below zero, [-1.88,  
170 -0.25]; Supplementary Table 4). We found no clear effects of either the group size, age or gender  
171 on adoption of the positive frequency-dependent copying strategy, except for the negative inter-  
172 action effect between age and task uncertainty (the 95% Bayesian CI of the age  $\times$  uncertainty  
173 interaction = [-1.46, -0.15]; Supplementary Table 4).

174 We also investigated the effects of group size and task uncertainty on the fitted individual  
175 parameter values. We found that the individual mean social learning weight parameter (i.e.  $\bar{\sigma}_i =$   
176  $(\sum_t \sigma_{i,t})/(\text{total rounds})$ ) increased with group size (the 95% Bayesian CI = [0.15, 0.93]; Figure  
177 4B; Supplementary Table 5), and decreased with uncertainty (the 95% Bayesian CI = [-0.98,  
178 -0.22]), and age of subject (the 95% Bayesian CI = [-0.36, -0.02]). However, the negative effects  
179 of task uncertainty and age disappeared when we focused only on  $\bar{\sigma}_i$  of the positive frequency-  
180 dependent copying individuals, and only the positive effect of the group size was confirmed  
181 (Supplementary Table 6; Supplementary Figure 6). It is worth noting that the meaning of the  
182 social learning weight is different between these three different strategies: The social learning  
183 weight regulates positive reactions to the majorities' behaviour for positive frequency-dependent  
184 copiers, whereas it regulates avoidance of the majority for negative-frequency dependent copiers,  
185 and determines the probability of random decision-making for the random choice strategists.

186 The individual conformity exponent parameter  $\theta_i$  increased with task uncertainty (the 95%

187 Bayesian CI = [0.38, 1.41]), but we found no significant effects of group size, age, gender or  
188 interactions (Figure 4C; Supplementary Table 7). These results were qualitatively unchanged  
189 when we focused only on the positive frequency-dependent copying individuals (Supplementary  
190 Table 8; Supplementary Figure 6).

191 We observed extensive individual variation in social information use. The greater the task's  
192 uncertainty, the larger were individual variances in both the mean social learning weight and the  
193 conformity exponent (the 95% Bayesian CI of the GLMM's variation parameter for  $\bar{\sigma}_i$  was [1.11,  
194 1.62] (Supplementary Table 5) and for  $\theta_i$  was [1.07, 1.54] (Supplementary Table 7)). This was  
195 confirmed when focusing only on the positive frequency-dependent copying individuals: The  
196 Bayesian 95% CIs were [1.14, 1.80] (Supplementary Table 6) and [0.71, 1.10] (Supplementary  
197 Table 8), respectively.

198 The manner in which individual variation in social-information use of positive frequency-  
199 dependent copying individuals changes over time is visualised in Figure 5. The social learn-  
200 ing weights generally decreased with experimental round. However, some individuals in the  
201 Moderate- and the High-uncertain conditions accelerated rather than decreased their reliance on  
202 social learning over time. Interestingly, those accelerating individuals tended to have a larger  
203 conformity exponent (Supplementary Figure 5). In addition, the time-dependent  $\theta_{i,t}$  in our al-  
204 ternative model generally increased with experimental round in the Moderate- and the High-  
205 uncertainty conditions (Supplementary Figure 10), although the fitting of  $\theta_{i,t}$  in the alternative  
206 model was relatively unreliable (Supplementary Figure 9). These findings suggest that con-  
207 formists tended to use asocial learning at the outset (i.e. exploration asocially) but increasingly  
208 started to conform as the task proceeded (i.e. exploitation socially).

209 Extensive variation in the temporal dynamics of the social learning weight  $\sigma_{i,t}$  was also found  
210 for the negative-frequency dependent copying individuals but not found for random choice indi-  
211 viduals (Supplementary Figure 5). Individuals deploying a random choice strategy exhibited a  
212  $\sigma_{i,t}$  that approached to zero, indicating that their decision-making increasingly relied exclusively  
213 on the softmax choice rule, rather than unguided random choices, as the task proceeded.

214 No significant fixed effects were found in other asocial learning parameters such as the learn-  
215 ing rate  $\alpha_i$  and the mean inverse temperature  $\bar{\beta}_i = (\sum_t \beta_{i,t})/(\text{total rounds})$  (Supplementary Table  
216 9, Supplementary Table 10 and Supplementary Figure 6).

217 In summary, our experiments on adult humans revealed asymmetric influences of increasing  
218 task uncertainty and increasing group size on the social learning parameters. The conformity  
219 exponent increased with task uncertainty on average but the proportion of positive frequency-  
220 dependent copying individuals showed a corresponding decrease, due to the extensive individual  
221 variation emerging in the High-uncertain condition. Conversely, group size had a positive effect  
222 on the mean social learning weight, but did not affect conformity.

#### 223 **1.4 Social learning strategies explain the collective dynamics**

224 The post-hoc simulation provides statistical predictions on how likely it is, given the fitted learn-  
225 ing model parameters, that groups of individuals make accurate decisions and that they exhibit  
226 inflexible herding. Figure 3C shows the change over time in performance with different group  
227 sizes and different uncertainty conditions, generated by the post-hoc simulation (see also Sup-  
228 plementary Figure 3). The trajectories of the simulated dynamics recovered nicely the pattern  
229 observed in the experiment (Figure 3A and 3C), suggesting that the strategic changes in the

230 individual-level social information use (Figure 4) could explain the collective-level behavioural  
231 pattern.

232 Figure 3D shows that larger groups are more likely to make accurate decisions than are  
233 both small groups and solitaries in the period prior to change across all uncertainty conditions,  
234 suggesting collective intelligence was operating. In the post-change period, however, perfor-  
235 mance differed between the conditions. In the Low-uncertainty condition, where we found that  
236 the participants were most likely to have a relatively weak positive frequency-dependence (i.e.  
237  $\bar{\theta} = 1.65$ ), large groups performed better than did small groups over 59.5% of total 10,000  
238 repetitions. However, in the Moderate-uncertainty condition, where we found that participants  
239 were most likely to have strong positive frequency dependence ( $\bar{\theta} = 3.00$ , c.f. 1.65 in the Low-  
240 uncertainty condition), the large groups were more likely to get stuck on the suboptimal option,  
241 and hence the small groups performed better than did the large groups over 69.5% of repetitions  
242 (Figure 3D). The decision accuracy did not substantially differ with group size in the post-change  
243 period in the High-uncertainty condition although the large groups performed slightly better than  
244 did the small groups (50.8% of the repetitions).

245 Interestingly, although their relatively low conformity biases, there were some groups in the  
246 Low-uncertainty condition that seemed to exhibit herding (the ‘humped’ area at the lefthand side  
247 to the peak of the performance distribution in the post-change period; Figure 3D). This might be  
248 due to the lower softmax exploration rates among social learners in the Low-uncertainty condition  
249 (i.e. both  $\mu_{\beta_0^*}$  and  $\mu_\epsilon$  were large; Table 2): the whole population gets stuck because all individuals  
250 are very exploitative on their past experience.

## 251 **2 Discussion**

252 We investigated whether and how human social learning strategies regulate the trade-off between  
253 collective intelligence and inflexible herding behaviour using a collective learning-and-decision-  
254 making task combined with simulation and model fitting. We examined whether manipulat-  
255 ing the reliability of asocial learning and group size would affect the use of social information,  
256 and thereby alter the collective human decision dynamics, as suggested by our computational  
257 model simulation. Although a theoretical study has suggested that reliance on social learning  
258 and conformity bias would play a role in collective dynamics<sup>2:5:53</sup>, thus far no empirical studies  
259 have quantitatively investigated the population-level consequences of these two different social  
260 learning processes. Our high-resolution, model-based behavioural analysis using a hierarchi-  
261 cal Bayesian statistics enabled us to identify individual-level patterns and variation of different  
262 learning parameters and to explore their population-level outcomes. The results provide **quanti-**  
263 **tative** support for our hypothesis that the collective decision performance can be predicted with  
264 quantitative knowledge of social learning strategies.

265 Overall, our individual-based computational model recovered the behavioural pattern sug-  
266 gested by the phenomenological regression (Figure 3). Using the post-hoc simulation with  
267 individually-fit model parameters, we confirmed that in the Low-uncertainty condition, where  
268 individuals had weaker positive frequency bias (i.e.  $\bar{\theta} \approx 1.65$ ), larger groups **were able to be**  
269 **more accurate than smaller groups while retaining flexibility in their decision-making<sup>9</sup>**, although  
270 their low asocial exploration rates seemed to undermine the potential flexibility. However, in the  
271 Moderate- and the High-uncertain conditions where individuals had the higher conformity ex-

272 ponent parameters (i.e.  $\bar{\theta} \approx 3.0$  and  $2.7$ , respectively), larger groups performed better prior to  
273 environmental change but were vulnerable to getting stuck with an out-dated maladaptive option  
274 post change. Therefore, the changes in the level of conformity in human individuals<sup>34;41</sup> indeed  
275 incurred a trade-off between the collective intelligence effect and the risk of inflexible herding.

276 Although the social learning weight increased with increasing group size, the overall mean  
277 value was  $\bar{\sigma}_i \approx 0.3$  (Figure 4B; Supplementary Figure 5; Supplementary Figure 6) and it de-  
278 creased on average as the task proceeded (Figure 5). This implies a weaker social than asocial  
279 influence on decision-making as reported in several other experimental studies<sup>35;54-56</sup> although  
280 evolutionary models tend to predict heavier reliance on social learning than experimental studies  
281 report<sup>57;58</sup>. Thanks to this relatively weak reliance of social learning, the kind of extreme herding  
282 that would have blindly led a group to any option regardless of its quality, such as the ‘symmetry  
283 breaking’ known in trail-laying ant collective foraging systems<sup>2;5;26</sup>, did not occur (Figure 2).

284 Individual differences in rates of exploration might also help to mitigate potential herding.  
285 Although a majority of participants adopted a positive frequency-dependent copying strategy,  
286 some individuals exhibited negative frequency dependence or random decision-making (Figure  
287 4A). The random choice strategy was associated with more exploration than the other strate-  
288 gies, because it led to an almost random choice at a rate  $\sigma_i$ , irrespective of the options’ quality.  
289 Negative-frequency dependent copying individuals could also be highly exploratory. These indi-  
290 viduals tended to avoid choosing an option upon which other people had converged and would ex-  
291 plore the other two ‘unpopular’ options. Interestingly, in the High-uncertain condition the mean  
292 social learning weights of the negative-frequency dependent copying individuals ( $\bar{\sigma}_i \approx 0.5$ ) were  
293 larger than that of the other two strategies ( $\bar{\sigma}_i \approx 0.1$ , Supplementary Figure 5), indicating that

294 these individuals engaged in such majority-avoiding exploration relatively frequently. Such a  
295 high variety in social information use<sup>59-62</sup> and exploratory tendencies would prevent individuals  
296 from converging on a single option, leading to a mitigation of herding but concurrently dimin-  
297 ishing the decision accuracy in high-uncertainty circumstances (Figure 3).

298 A methodological advantage of using computational models to study social learning strate-  
299 gies is its explicitness of assumptions about the temporal dynamics of behaviour, which enabled  
300 us to distinguish different learning strategies<sup>63-65</sup>. For example, very exploitative asocial re-  
301 inforcement learners (i.e. for whom exploitation parameter  $\beta_{i,t}$  is large and the social learning  
302 weight  $\sigma_{i,t}$  is nearly zero, as seen in the Low-uncertainty condition) and conformity-biased social  
303 learners (where the conformity exponent  $\theta_i$  is large and  $\sigma_{i,t}$  is positive, as seen in the Moderate-  
304 uncertain condition) would eventually converge on the same option, resulting in the same final  
305 behavioural steady state. However, how they explored the environment, as well as how they re-  
306 acted to the other individuals in the same group, are significantly different and they could produce  
307 qualitatively different collective temporal dynamics.

308 However, our computational model could not fully capture other, potentially more sophisti-  
309 cated forms of social learning strategies that participants might deploy, which might be a reason  
310 for the seemingly low rate of social learning observed in the experiment compared to theory<sup>57,58</sup>.  
311 Indeed, the post-hoc simulation sometimes failed to recover the observed behavioural trajecto-  
312 ries. **In particular, experimental groups with  $n = 12$ ,  $n = 16$ , and one group in  $n = 9$ , in the**  
313 **Low-uncertainty condition performed very well, exceeding the 95% CIs of the post-hoc simu-**  
314 **lation after the environmental change (Supplementary Figure 3). This indicates that collective**  
315 **behaviour in these groups was more flexible than our model predicted. Further empirical studies**

316 that consider a wider range of possible social learning strategies, e.g. ‘copy-rapidly-increasing-  
317 option’ strategy<sup>66</sup> or Bayesian updating<sup>57;67</sup>, are needed to explore computational underpinnings  
318 of social learning and collective behaviour.

319 The Internet-based experimentation allowed us to conduct a real-time interactive behavioural  
320 task with larger subject pools than a conventional laboratory-based experiment. This enabled us  
321 not only to quantify the individual-level learning-and-decision processes<sup>68</sup> but also to map these  
322 individual-level processes on to the larger-scale collective behaviour<sup>5;15;20</sup>. Although there are  
323 always questions about the validity of participants’ behaviour when recruited via web-based tools,  
324 we believe that the computational modelling approach coupled with higher statistical power due  
325 to the large sample size, compensates for any drawbacks. The fact that our learning model could  
326 approximate the participants’ decision trajectories effectively suggest that most of the participants  
327 engaged seriously with solving the task. An increasing body of evidence supports the argument  
328 that web-based behavioural experiments are as reliable as results from the laboratory<sup>69;70</sup>.

329 The diverse effects of social influence on the collective wisdom of a group has been draw-  
330 ing substantial attention<sup>19;21;22;71;72</sup>. The bulk of this literature, including many jury models and  
331 election models<sup>45;73</sup>, has focused primarily on the static estimation problem, where the ‘truth’ is  
332 fixed from the outset. However, in reality, there are many situations under which the state of the  
333 true value is changing over time so that monitoring and tracking the pattern of change is a crucial  
334 determinant of decision performance<sup>74</sup>. In such temporally dynamic environments, decision-  
335 making and learning are coordinated to affect future behavioural outcomes recursively<sup>75</sup>. Our  
336 experimental task provides a simple vehicle for exploring collective intelligence in a dynamic  
337 situation, which encompasses this learning-and-decision-making feedback loop. Potentially, in-

338 tegrating the wisdom of crowds with social learning and collective dynamics research will facil-  
339 itate the more tractable use of collective intelligence in a temporary changing world.

340 In summary, a combination of experimentation and theoretical modelling sheds light on when  
341 groups of individuals will exhibit the wisdom of the crowds and when inflexible herding. Our  
342 analysis implies that herding is most likely amongst individuals in large groups exposed to chal-  
343 lenging tasks. That is because challenging tasks lead to greater uncertainty and thereby elicit  
344 greater conformist learning amongst individuals, whilst rates of copying increase with group  
345 size. Difficult tasks, by definition, render identification of the optimal behaviour harder, allow-  
346 ing groups sometimes to converge on maladaptive outcomes. Conversely, the reduced conformity  
347 levels of individuals in small groups, and the greater probability that social information would  
348 be accurate for less-challenging tasks, generated ‘wisdom of the crowd’ effects in most other cir-  
349 cumstances. Our findings provide clear evidence that the conflict between collective intelligence  
350 and maladaptive herding can be predicted with knowledge of human social learning strategies.

### 351 **3 Material and methods**

#### 352 **3.1 Participants**

353 The experimental procedure was approved by the Ethics Committee at the University of St An-  
354 drews (BL10808). A total of 755 subjects (354 females, 377 males, 2 others and 22 unspecified;  
355 mean age ( $1\ SD.$ ) = 34.33 (10.9)) participated through Amazon’s Mechanical Turk. All partic-  
356 ipants consented to participation through an online consent form at the beginning of the task.  
357 We excluded subjects who disconnected to the online task before completing at least the first 30

358 rounds from our computational-model fitting analysis due to unreliability of the model-parameter  
359 estimation, resulted in 699 subjects (573 subjects entered the group (i.e.  $N \geq 2$ ) and 126 entered  
360 the solitary (i.e.  $N = 1$ ) condition). The task was only available for individuals who had greater  
361 than or equal to 90% HIT approval rate and who accessed from the United States. Although  
362 no sample-size calculation was performed in advance, our parameter recovery test confirmed  
363 that the sample size was sufficient for estimation of individual parameters using a hierarchical  
364 Bayesian method.

### 365 **3.2 Design of the experimental manipulations**

366 The three uncertainty conditions were: Low-uncertainty condition (differences between mean  
367 payoffs were 1.264), Moderate-uncertainty condition (differences between mean payoffs were  
368 0.742) and High-uncertainty condition (differences between mean payoffs were 0.3). The mean  
369 payoff associated with the ‘excellent’ slot in all three conditions was fixed to 3.1 cents (Supple-  
370 mentary Figure 1). Each task uncertainty condition was randomly assigned for each different HIT  
371 session, and participants were allowed to participate in one HIT only. Sample size after the data  
372 exclusion for each uncertainty condition was:  $N = 113$  (Low-uncertainty condition),  $N = 132$   
373 (Moderate-Uncertain condition), and  $N = 454$  (High-uncertain condition). We assigned more  
374 sessions to the High-uncertainty condition compared to the other two because we expected that  
375 larger group sizes would be needed to generate the collective wisdom in noisier environments.

376 To manipulate the size of each group, we varied the capacity of the waiting room from 10 to  
377 30. Because the task was being advertised on the Worker website at AMT for approximately 2  
378 hours, some participants occasionally arrived after the earlier groups had already started. In that

379 case the participant entered the newly opened waiting room which was open for the next 5 min-  
380 utes. The number of participants arriving declined with time because newly posted alternative  
381 HITs were advertised on the top of the task list, which decreased our task's visibility. This meant  
382 that a later-starting session tended to begin before reaching maximum room capacity, resulting  
383 in the smaller group size. Therefore, the actual size differed between groups (Supplementary  
384 Figure 3, Supplementary Table 1). Data collection and analysis were not performed blind to the  
385 conditions of the experiments.

### 386 **3.3 The multi-player three-armed bandit task**

387 To study the relationship between social information use and collective behavioural dynamics,  
388 we focused on a well-established learning-and-decision problem called a 'multi-armed bandit'  
389 task, represented here as repeated choices between three slot machines (Supplementary Figure 1,  
390 Video 1, for detail see Supplementary Method). Participants played the task for 70 rounds. The  
391 slots paid off money noisily (in the US cents), varying around two different means during the  
392 first 40 rounds such that there was one 'good' slot and two other options giving poorer average  
393 returns. From the round 41st, however, one of the 'poor' slots abruptly increased its mean payoff  
394 to become 'excellent' (i.e. superior to 'good'). The purpose of this environmental change was  
395 to observe the effects of maladaptive herding by potentially trapping groups in the out-of-date  
396 suboptimal (good) slot, as individuals did not know whether or how an environmental change  
397 would occur. Through making choices and earning a reward from each choice, individuals could  
398 gradually learn which slot generated the highest rewards.

399 In addition to this asocial learning, we provided social information for each member of the

400 group specifying the frequency with which group members chose each slot. All group mem-  
 401 bers played the same task with the same conditions simultaneously, and all individuals had been  
 402 instructed that this was the case, and hence understood that the social information would be in-  
 403 formative.

404 Task uncertainty was experimentally manipulated by changing the difference between the  
 405 mean payoffs for the slot machines. In the task with the least uncertainty, the distribution of  
 406 payoffs barely overlapped, whilst in the task with the greatest uncertainty the distribution of  
 407 payoffs overlapped considerably (Supplementary Figure 1).

### 408 3.4 The computational learning-and-decision-making model

409 We modelled individual behavioural processes by assuming that individual  $i$  makes a choice for  
 410 option  $m$  at round  $t$ , in accordance with the choice-probability  $P_{i,t}(m)$  that is a weighted average  
 411 of social and asocial influences:

$$P_{i,t}(m) = \sigma_{i,t} \times \text{Social influence}_{i,m,t} + (1 - \sigma_{i,t}) \times \text{Asocial influence}_{i,m,t}, \quad (1)$$

412 where  $\sigma_{i,t}$  is the *social learning weight* ( $0 \leq \sigma_{i,t} \leq 1$ ).

413 For the social influence, we assumed a frequency-dependent copying strategy by which an  
 414 individual copies others' behaviour in accordance with the distribution of social frequency infor-  
 415 mation [49–51:55](#):

$$\text{Social influence}_{i,m,t} = \frac{\left(F_{t-1}(m) + 0.1\right)^{\theta_i}}{\sum_{k \in \text{options}} \left(F_{t-1}(k) + 0.1\right)^{\theta_i}}, \quad (2)$$

416 where  $F_{t-1}(m)$  is a number of choices made by other individuals (excluding her/his own choice)  
 417 for the option  $m$  in the preceding round  $t - 1$  ( $t \geq 2$ ).  $\theta_i$  is individual  $i$ 's *conformity exponent*,  
 418  $-\infty \leq \theta_i \leq +\infty$ . When this exponent is larger than zero, higher social influence is given  
 419 to an option which was chosen by more individuals (i.e. positive frequency bias). When this  
 420 exponent is larger than zero ( $\theta_i > 0$ ), higher social influence is afforded to an option chosen  
 421 by more individuals (i.e. positive frequency bias), with conformity bias arising when  $\theta_i > 1$ ,  
 422 such that disproportionately more social influence is given to the most common option<sup>28</sup>. When  
 423  $\theta_i < 0$ , on the other hand, higher social influence is afforded to the option that fewest individuals  
 424 chose in the preceding round  $t - 1$  (i.e. negative frequency bias). To implement the negative  
 425 frequency dependence, we added a small number 0.1 to  $F$  so that an option chosen by no one  
 426 (i.e.  $F_{t-1} = 0$ ) could provide the highest social influence when  $\theta_i < 0$ . Note, there is no social  
 427 influence when  $\theta_i = 0$  because in this case the 'social influence' favours an uniformly random  
 428 choice, i.e.,  $S_{i,t}(m) = f_m^0 / (f_1^0 + f_2^0 + f_3^0) = 1/3$ , independent of the social frequency distribution.  
 429 Note also that, in the first round  $t = 1$ , we assumed that the choice is only determined by the  
 430 asocial softmax function because there is no social information available.

431 For the asocial influence, we used a standard reinforcement learning with 'softmax' choice  
 432 rule<sup>75</sup>, widely applied in human social learning studies e.g.<sup>35:51;55</sup>. An individual  $i$  updates the  
 433 estimated average reward associated with an option  $m$  at round  $t$ , namely Q-value ( $Q_{i,t}(m)$ ), ac-  
 434 cording to the Rescorla-Wagner rule as follows:

$$Q_{i,t+1}(m) = Q_{i,t}(m) + \alpha_i \mathbb{1}(m, m_{i,t}) (r_{i,t}(m) - Q_{i,t}(m)), \quad (3)$$

435 where  $\alpha_i$  ( $0 \leq \alpha_i \leq 1$ ) is a *learning rate* parameter of individual  $i$  determining the weight given to

436 new experience and  $r_{i,t}(m)$  is the amount of monetary reward obtained from choosing the option  
 437  $m$  in round  $t$ .  $\mathbb{1}(m, m_{i,t})$  is the binary action-indicator function of individual  $i$ , given by

$$\mathbb{1}(m, m_{i,t}) = \begin{cases} 1, & \text{if } m_{i,t} = m \text{ or } t = 1, \\ 0, & \text{otherwise.} \end{cases} \quad (4)$$

438 Therefore,  $Q_{i,t}(m)$  is updated only when the option  $m$  was chosen; when the option  $m$  was not  
 439 chosen,  $Q_{i,t}(m)$  is not updated (i.e.  $Q_{i,t+1}(m) = Q_{i,t}(m)$ ). Note that, only in the first round  $t = 1$ ,  
 440 all Q-values are updated by using the chosen option's reward  $r_{i,1}(m)$ , so that the individual can  
 441 set a naive 'intuition' about the magnitude of reward values she/he would expect to earn from a  
 442 choice in the task; namely,  $Q_{i,t=2}(1) = Q_{i,t=2}(2) = Q_{i,t=2}(3) = \alpha_i r_{i,t=1}(m)$ . In practical terms,  
 443 this prevents the model from being overly sensitive to the first experience. Before the first choice,  
 444 individuals had no prior preference for either option (i.e.  $Q_{i,1}(1) = Q_{i,1}(2) = Q_{i,1}(3) = 0$ ).

445 The Q-value is then translated into the asocial influence through the softmax (or logit choice)  
 446 function:

$$A_{i,t}(m) = \frac{\exp(\beta_{i,t} Q_{i,t}(m))}{\sum_{k \in \text{options}} \exp(\beta_{i,t} Q_{i,t}(k))}, \quad (5)$$

447 where  $\beta_{i,t}$ , called *inverse temperature*, manipulates individual  $i$ 's sensitivity to the Q-values (in  
 448 other words, controlling the proneness to explore). As  $\beta_{i,t}$  goes to zero, asocial influence ap-  
 449 proximates to a random choice (i.e. highly explorative). Conversely, if  $\beta_{i,t} \rightarrow +\infty$ , the aso-  
 450 cial influence leads to a deterministic choice in favour of the option with the highest Q-value  
 451 (i.e. highly exploitative). For intermediate values of  $\beta_{i,t}$ , individual  $i$  exhibits a balance be-  
 452 tween exploration and exploitation<sup>35:68</sup>. We allowed for the possibility that the balance between

453 exploration-exploitation could change as the task proceeds. To depict such time dependence in  
 454 exploration, we used the equation:  $\beta_{i,t} = \beta_{i,0}^* + \epsilon_i t/70$ . If the slope  $\epsilon_i$  is positive (negative), aso-  
 455 cial influence  $A_{i,t}$  becomes more and more exploitative (explorative) as round  $t$  increases. For a  
 456 model fitting purpose, the time-dependent term  $\epsilon_i t$  is scaled by the total round number 70.

457 We allowed that the social learning weight  $\sigma_{i,t}$  could also change over time as assumed in  
 458 the inverse temperature  $\beta_{i,t}$ . To let  $\sigma_{i,t}$  satisfy the constraint  $0 \leq \sigma_{i,t} \leq 1$ , we used the following  
 459 sigmoidal function:

$$\sigma_{i,t} = \frac{1}{1 + \exp(-(\sigma_{i,0}^* + \delta_i t/70))}. \quad (6)$$

460 If the slope  $\delta_i$  is positive (negative), the social influence increases (decreases) over time. We  
 461 set the social learning weight equal to zero when group size is one (i.e. when an individual  
 462 participated in the task alone and/or when  $\sum_{k \in \text{options}} F_{t-1}(k) = 0$ ).

463 We modelled both the inverse temperature  $\beta_{i,t}$  and the social learning weight  $\sigma_{i,t}$  as a time  
 464 function since otherwise it would be challenging to distinguish different patterns of learning in  
 465 this social learning task<sup>63</sup>. The parameter recovery test confirmed that we were able to differenti-  
 466 ate such processes under these assumptions (Supplementary Figure 7, Supplementary Figure 8).  
 467 While we also considered the possibility of the conformity exponent being time-dependent (i.e.  
 468  $\theta_{i,t} = \theta_{i,0}^* + \gamma_i t/70$ ), the parameter recovery test suggested that the individual slope parameter  
 469  $\gamma_i$  was not reliably recovered (Supplementary Figure 9), and hence we concentrated our anal-  
 470 ysis on the time-independent  $\theta_i$  model. We confirmed that instead using the alternative model  
 471 where both social learning parameters were time-dependent (i.e.  $\sigma_{i,t}$  and  $\theta_{i,t}$ ) did not qualitatively  
 472 change our results (Supplementary Figure 10).

473 One concern might be the asymmetry between the asocial softmax influence which takes  
474 many prior experiences into account (depending upon a learning rate) and the social influence  
475 referring only the most recent frequency information  $F_{t-1}$ . The choice frequency appeared at  
476 round  $t$  is the most reliable social information, compared to the past frequencies, because it could  
477 be the most ‘updated’ information as long as the other individuals have made informed decisions  
478 in their best knowledge. In contrast, option’s reward just obtained at  $t - 1$ , which was inde-  
479 pendently and randomly drawn from a probability distribution, is less reliable than accumulated  
480 Q-values taking past rewards into account. Although many other formulations for asocial and  
481 social learning processes were possible, we believe that our current choice – time-depth asocial  
482 reinforcement learning with the most-updated-frequency-dependent copying was a reasonable  
483 first step.

484 In summary, the model has six free parameters that were estimated for each individual human  
485 participant; namely,  $\alpha_i$ ,  $\beta_{i,0}^*$ ,  $\epsilon_i$ ,  $\sigma_{i,0}^*$ ,  $\delta_i$ , and  $\theta_i$ . To fit the model, we used a hierarchical Bayesian  
486 method (HBM), estimating the global means ( $\mu_\alpha$ ,  $\mu_{\beta_0^*}$ ,  $\mu_\epsilon$ ,  $\mu_{\sigma_0^*}$ ,  $\mu_\delta$ , and  $\mu_\theta$ ) and the global vari-  
487 ations ( $\nu_\alpha$ ,  $\nu_{\beta_0^*}$ ,  $\nu_\epsilon$ ,  $\nu_{\sigma_0^*}$ ,  $\nu_\delta$ , and  $\nu_\theta$ ) for each of the three experimental conditions (i.e. the Low-,  
488 Moderate- and High-uncertain condition), which govern overall distributions of individual pa-  
489 rameter values. It has become recognised that the HBM can provide more robust and reliable  
490 parameter estimation than conventional maximum likelihood point estimation in complex cog-  
491 nitive models<sup>76</sup>, a conclusion with which our parameter recovery test agreed (Supplementary  
492 Figure 7, Supplementary Figure 8).

### 493 **3.5 Agent-based model simulation**

494 We ran a series of individual-based model simulations assuming that a group of individuals play  
495 our three-armed bandit task for 90 rounds (under the Moderate-uncertainty condition) and that in-  
496 dividuals behave in accordance with the computational learning-and-decision model. We varied  
497 the group size ( $n \in \{3, 10, 30\}$ ), the mean social learning weight ( $\bar{\sigma} \in \{0.01, 0.1, 0.2, 0.3, \dots, 0.9\}$ )  
498 and the mean conformity exponent ( $\bar{\theta} \in \{0.5, 1, 3, 6\}$ ), running 10,000 replications for each of  
499 the possible parameter  $\times$  group size combinations. As for the other parameter values (e.g. the  
500 asocial reinforcement learning parameters;  $\alpha, \beta_0^*, \epsilon$ ), here we used the experimentally fitted global  
501 means (Table 2 and Supplementary Table 3). Relaxation of this assumption (i.e. using a different  
502 set of asocial learning parameters) does not qualitatively change our story (Supplementary Fig-  
503 ure 2). Note that each individual's parameter values were randomly drawn from the distributions  
504 centred by the global mean parameter values fixed to each simulation run. Therefore, the actual  
505 composition of individual parameter values were different between individuals even within the  
506 same social group.

### 507 **3.6 Generalised linear mixed models**

508 To directly analyse the effects of group size and task uncertainty on the time evolution of decision  
509 performance, we conducted a statistical analysis using a phenomenological model, namely, a  
510 hidden Markov process logistic regression without assuming any specific learning-and-decision-  
511 making processes. The dependent variable was whether the participant chose the best option (1)  
512 or not (0). The model includes fixed effects of grouping  $\xi$ , standardised group size  $\omega$ , and an  
513 intercept with a random effect of individuals  $\mu + \rho_i$ . We assumed that the intercept and the effect

514 of grouping change from round to round, as a random walk process. For the effect of group size  
515 we considered the effect of the 1st environment  $1 \leq t \leq 40$  and that of the 2nd environment,  
516 namely,  $\omega_1$  and  $\omega_2$ , separately.

517 To examine whether increasing group size and increasing task uncertainty affected individ-  
518 ual use of the positive frequency-dependent copying strategy, we used a hierarchical Bayesian  
519 logistic regression model with a random effect of groups. The dependent variable was whether  
520 the participant used the positive frequency-dependent copying (1) or not (0). The model includes  
521 fixed effects of group size (standardised), task uncertainty (0: Low, 0.5: Moderate, 1: High), age  
522 (standardised), gender (0: male, 1: female, NA: others or unspecified), and possible two-way  
523 interactions between these fixed effects.

524 We also investigated the effects of both group size and the task's uncertainty on the fitted  
525 values of the learning parameters. We used a hierarchical Bayesian gaussian regression model  
526 predicting the individual fitted parameter values. The model includes effects of group size (stan-  
527 dardised), task uncertainty (0: Low, 0.5: Moderate, 1: High), age (standardised), gender (0:  
528 male, 1: female, NA: others or unspecified), and two-way interactions between these fixed ef-  
529 fects. We assumed that the variance of the individual parameter values might be contingent upon  
530 task uncertainty because we had found in the computational model-fitting result that the fitted  
531 global variance parameters (i.e.  $v_{\sigma_0^*}$ ,  $v_\delta$  and  $v_\theta$ ) were larger in more uncertain conditions (Sup-  
532 plementary Table 2).

## 533 **3.7 Statistical analysis**

534 We used a hierarchical Bayesian method (HBM) to estimate the free parameters of our sta-  
535 tistical models, including both the phenomenological regression model and the computational  
536 learning-and-decision-making model. The HBM allows us to estimate individual differences,  
537 while ensures these individual variations are bounded by the group-level global parameters. The  
538 HBM was performed under Stan 2.16.2 (<http://mc-stan.org>) in R 3.4.1 ([https://www.  
539 r-project.org](https://www.r-project.org)) software. The models contained at least 4 parallel chains and we confirmed  
540 convergence of the MCMC using both the Gelman-Rubin statistics and the effective sample sizes.  
541 Full details of the model fitting procedure and prior assumptions are shown in the appendix.

### 542 **3.7.1 Parameter recovery test**

543 To check the validity of our model-fitting method, we conducted a ‘parameter recovery test’  
544 so as to examine how well our model fitting procedure had been able to reveal true individual  
545 parameter values. To do this, we generated synthetic data by running a simulation with the  
546 empirically fitted global parameter values, and then re-fitted the model with this synthetic data  
547 using the same procedure. The parameter recovery test showed that the all true global parameter  
548 values were fallen into the 95% Bayesian credible interval (Supplementary Figure 7), and at least  
549 93% of the true individual parameter values were correctly recovered (i.e. 96% of  $\alpha_i$ , 93% of  $\beta_{i,0}^*$ ,  
550 95% of  $\epsilon_i$ , 97% of  $\sigma_{i,0}^*$ , 96% of  $\delta_i$  and 97% of  $\theta_i$  values were fallen into the 95% Bayesian CI.  
551 Supplementary Figure 7).

### 552 **3.7.2 Categorisation of individual learning strategies**

553 Based on the 50% CI of the individual *conformity exponent* parameter values  $\theta_i$ , we divided  
554 the participants into the following three different social learning strategies. If her/his 50% CI  
555 of  $\theta_i$  fell above zero ( $\theta_{lower} > 0$ ), below zero ( $\theta_{upper} < 0$ ) or including zero ( $\theta_{lower} \leq 0 \leq$   
556  $\theta_{upper}$ ), she/he was categorised as a ‘positive frequency-dependent copier’, a ‘negative frequency-  
557 dependent copier’, or a ‘random choice individual’, respectively. We used the 50% Bayesian CI  
558 to conduct this categorisation instead of using the more conservative 95% CI because the latter  
559 would cause much higher rates of ‘false negatives’, by which an individual who applied either a  
560 positive frequency-dependent copying or a negative-frequency dependent copying strategy was  
561 falsely labelled as an asocial random choice individual (Supplementary Figure 7). Four hundred  
562 agents out of 572 ( $\approx 70\%$ ) were falsely categorised as a random choice learner in the recovery  
563 test when we used the 95% criterion (Supplementary Figure 7). On the other hand, the 50%  
564 CI criterion seemed to be much better in terms of the false negative rate which was only 18.5%  
565 (i.e. 106 agents), although it might be slightly worse in terms of ‘false positives’: Thirty-seven  
566 agents (6.5%) were falsely labelled as either a positive frequency-dependent copier or a negative-  
567 frequency dependent copier by the 50% CI, whereas the false positive rate of the 95% CI was  
568 only 0.2% (Supplementary Figure 7). To balance the risk of false positives and false negatives,  
569 we decided to use the 50% CI which seemed to have more strategy detecting power.

### 570 **3.7.3 The post-hoc model simulation**

571 So as to evaluate how accurately our model can generate observed decision pattern in our task  
572 setting, we ran a series of individual-based model simulation using the fitted individual param-

573 eter values (i.e. means of the individual posterior distributions) for each group size for each  
574 uncertainty condition. At the first step of the simulation, we assigned a set of fitted parameters  
575 of a randomly-chosen experimental subject from the same group size and the same uncertain  
576 condition to an simulated agent, until the number of agents reaches the simulated group size. We  
577 allowed duplicate choice of experimental subject in this parameter assignment. At the second  
578 step, we let this synthetic group of agents play the bandit task for 90 rounds. We repeated these  
579 steps 10,000 times for each group size, task uncertainty.

### 580 **3.8 Data availability**

581 Both experimental and simulation data are available on an online repository (<https://github.com/WataruToyokawa/ToyokawaWhalenLaland2018>).

### 583 **3.9 Code availability**

584 The browser based online task was built by Node.js (<https://nodejs.org/en/>) and socket.io  
585 (<https://socket.io>), and the code are available on a GitHub repository (<https://github.com/WataruToyokawa/MultiPlayerThreeArmedBanditGame>). Analyses were conducted in  
586 R (<https://www.r-project.org>) and simulations of individual based models were conducted  
587 in Mathematica (<https://www.wolfram.com>), both are available on an online repository (<https://github.com/WataruToyokawa/ToyokawaWhalenLaland2018>).

## 590 **4 Competing interest**

591 The authors declare no competing interests.

## 592 **5 Authors' contributions**

593 WT, AW and KNL planned the study and built the computational model. WT ran simulations.  
594 WT and AW made the experimental material, ran the web-base experiment, and collected the  
595 experimental data. WT, AW and KNL analysed the data and wrote the manuscript.

## 596 **6 Acknowledgements**

597 The experiment was supported by The John Templeton Foundation (KNL; 40128) and Suntory  
598 Foundation Research Support (WT; 2015-311). The computer simulations and computational  
599 model analyses were supported by JSPS Overseas Research Fellowships (WT; H27-11). The  
600 phenomenological model analyses were supported by JSPS KAKENHI Grant Number 17J01559.  
601 The funders had no role in study design, data collection and analysis, decision to publish or  
602 preparation of the manuscript.

## 603 **7 Figure captions and tables**

**Figure 1: Findings of the individual-based model showing the effects of social information use on the average decision accuracy over replications.** The x-axis gives the round and y-axis gives the proportion of individuals expected to choose the optimal slot (i.e. decision accuracy) averaged over all replications. The vertical dashed line indicates the timing of environmental (i.e. payoff) change (at  $t = 41$ ). Different group sizes are shown by different styles (black (dotted):  $n = 3$ , orange (dashed):  $n = 10$ , red (solid):  $n = 30$ ). We set the average slopes for the *social learning weight* to be equal to zero for the sake of simplicity; namely,  $\mu_\delta = 0$ . Other free parameter values (i.e.  $\mu_\alpha$ ,  $\mu_{\beta_0^*}$ ,  $\mu_\epsilon$ ,  $\nu_\alpha$ ,  $\nu_{\beta_0^*}$ ,  $\nu_\epsilon$ ,  $\nu_\sigma$ ,  $\nu_\delta$  and  $\nu_\theta$ ) are best approximates to the experimental fitted values (see Table 2 and Supplementary Table 1).

**Figure 2: Results from the individual-based model simulations showing the distribution of each group's mean accuracy before environmental change ( $t \leq 40$ ).** The x-axis gives the mean decision accuracy over the first 40 rounds (i.e. the environment 1) for each replication. Different group sizes are shown by different styles (black (dotted):  $n = 3$ , orange (dashed):  $n = 10$ , red (solid):  $n = 30$ ). The other free parameter values are the same as in Figure 1.

**Table 1:** The mean and the 95% Bayesian credible intervals of the posterior for the group size effect at the phenomenological logistic model

	Low Uncertainty		Moderate Uncertainty		High Uncertainty	
$\omega_1$	0.08	[-0.15, 0.33]	0.10	[-0.06, 0.26]	0.07	[0.00, 0.15]
$\omega_2$	0.67	[0.44, 0.91]	-0.26	[-0.44, -0.11]	-0.10	[-0.17, -0.02]

Note: All  $\hat{R}$  values are 1.0 and the effective sample sizes are larger than 837.

**Figure 3: Time evolutions and distributions of decision performance for each condition.** **A:** The average decision accuracies of the experimental participants (red: large groups, orange: small groups, dark grey: lone individuals). All individual performances were averaged within the same size category (solid lines). The light-shaded areas, dark-shaded areas, and dashed curves show the 95%, 50%, and median Bayesian credible intervals of the phenomenological, time-series logistic regression. **Sample sizes for large, small, and lone groups are:  $N = 43$ ,  $N = 44$  and  $N = 38$  for the Low-uncertainty condition;  $N = 52$ ,  $N = 56$  and  $N = 37$  for the Moderate-uncertainty condition; and  $N = 259$ ,  $N = 168$  and  $N = 58$  for the High-uncertainty condition, respectively.** **B:** Change in the main effect of the dummy variable of grouping on the decision accuracy at the phenomenological regression model. The shaded areas are the Bayesian CIs and solid curves are the median. **C, D:** Change and distribution in average decision accuracy of the individual-based post-hoc simulations of the learning process model using the experimentally fit parameter values. **C:** All replications were averaged within the same size category (solid lines). The shaded areas give the 50% quantiles. The experimental horizon (i.e.  $t = 70$ ) is indicated by the vertical line. **D:** Performance was averaged within prior- and post-change periods for each replication for each group sizes category.

**Figure 4: Model fitting for the three different task's uncertain conditions (the Low-, Moderate- and High-uncertainty) and the different group size.** Three different learning strategies are shown in different styles (red-triangle: positive frequency-dependent learning, blue-circle: negative frequency-dependent learning; grey-circle: nearly random choice strategy). (A) Frequencies of three different learning strategies. Note that a sum of the frequencies of these three strategies in the same group size does not necessarily equal to 1, because there are a small number of individuals eliminated from this analysis due to insufficient data. (B) Estimated social learning weight, and (C) estimated conformity exponent, for each individual shown for each learning strategy. The 50% Bayesian CIs of the fitted GLMMs are shown by dashed lines and shaded areas. The horizontal lines in (C) show a region  $-1 < \theta_i < 1$ . **Sample sizes for Negative Frequency Dependent, Positive Frequency Dependent, and Random Choice strategies are:  $N = 2$ ,  $N = 61$  and  $N = 14$  for the Low-uncertainty condition;  $N = 3$ ,  $N = 80$  and  $N = 15$  for the Moderate-uncertainty condition; and  $N = 32$ ,  $N = 260$  and  $N = 106$  for the High-uncertainty condition, respectively.**

**Figure 5: Change in fitted values (i.e. median of the Bayesian posterior distribution) of the social learning weight  $\sigma_{i,t}$  with time for each **Positive Frequency Dependent** individual, for each level of task uncertainty.**

Thick dashed lines are the median values of  $\sigma_{i,t}$  across the subjects for each uncertainty condition. Individual conformity exponent values  $\theta_i$  are shown in different colours (higher  $\theta_i$  is darker). **Sample size for each task uncertainty condition is:  $N = 61$  (Low-uncertainty),  $N = 80$  (Moderate-uncertainty) and  $N = 260$  (High-uncertainty).**

**Table 2: The mean and the 95% Bayesian credible intervals of the posterior global means for the parameter values.**

The number of participants ( $N$ ) for each experimental condition are also shown.

Uncertainty:	Groups			Solitary individuals		
	Low	Moderate	High	Low	Moderate	High
$\mu_{\alpha^*}$ (learning rate)	0.99 [0.34, 1.73]	0.90 [0.43, 1.44]	0.61 [0.21, 1.03]	0.85 [-0.07, 1.95]	-0.17 [-1.27, 0.89]	0.46 [-0.39, 1.36]
$\mu_{\beta_0^*}$ (inv. temp.)	1.84 [1.15, 2.70]	1.68 [1.25, 2.18]	1.38 [1.16, 1.62]	1.10 [0.69, 1.54]	1.44 [0.80, 2.07]	0.85 [0.46, 1.22]
$\mu_{\epsilon}$ (inv. temp.)	3.70 [1.98, 5.71]	3.01 [1.88, 4.27]	2.97 [2.37, 3.60]	2.39 [1.46, 3.53]	2.81 [1.64, 4.07]	2.27 [1.40, 3.31]
$\mu_{\sigma_0^*}$ (soc. wight)	-1.55 [-2.71, -0.71]	-2.37 [-4.12, -1.01]	-2.16 [-2.81, -1.63]	– –	– –	– –
$\mu_{\delta}$ (soc. wight)	-1.39 [-2.66, -0.03]	-1.55 [-4.29, 0.91]	-1.87 [-3.04, -0.81]	– –	– –	– –
$\mu_{\theta}$ (conformity coeff.)	1.65 [0.83, 2.82]	3.00 [1.57, 4.85]	2.67 [1.80, 3.73]	– –	– –	– –
$N$	77	98	398	36	34	56

604 **References**

- 605 [1] Eric Bonabeau, Marco Dorigo, and Guy Theraulaz. *Swarm Intelligence: From Natural to*  
606 *Artificial Systems*. Oxford University Press, New York, NY, 1999. ISBN 0-19-513159-2.
- 607 [2] Scott Camazine, Jean-Louis Deneubourg, Nigel R. Franks, James Sneyd, Guy Theraulaz,  
608 and Eric Bonabeau. *Self-Organization in Biological Systems*. Princeton University Press,  
609 Princeton, NJ, 2001. ISBN 9780691116242.
- 610 [3] Jens Krause, Graeme D Ruxton, and Stefan Krause. Swarm intelligence in animals and  
611 humans. *Trends in ecology & evolution*, 25(1):28–34, jan 2010. ISSN 0169-5347. doi:  
612 10.1016/j.tree.2009.06.016. URL <http://www.ncbi.nlm.nih.gov/pubmed/19735961>.
- 613 [4] Thomas D. Seeley. *The Wisdom of the Hive*. Harvard University Press, Cambridge, MA,  
614 1995. ISBN 9780674953765.
- 615 [5] David J. T. Sumpter. *Collective Animal Behavior*. Princeton University Press, Princeton,  
616 NJ, 2010. ISBN 9781400837106.
- 617 [6] Andrew J. King and Cédric Sueur. Where Next? Group Coordination and Collective  
618 Decision Making by Primates. *International Journal of Primatology*, 32(6):1245–1267,  
619 jun 2011. ISSN 0164-0291. doi: 10.1007/s10764-011-9526-7. URL [http://www.  
620 springerlink.com/index/10.1007/s10764-011-9526-7](http://www.springerlink.com/index/10.1007/s10764-011-9526-7).
- 621 [7] J. Morand-Ferron and J. L. Quinn. Larger groups of passerines are more efficient problem  
622 solvers in the wild. *Proceedings of the National Academy of Sciences*, 108(38):15898–  
623 15903, 2011. ISSN 0027-8424. doi: 10.1073/pnas.1111560108.

- 624 [8] Takao Sasaki and Dora Biro. Cumulative culture can emerge from collective intelligence  
625 in animal groups. *Nature Communications*, 8:1–6, 2017. ISSN 20411723. doi: 10.1038/  
626 ncomms15049. URL <http://dx.doi.org/10.1038/ncomms15049>.
- 627 [9] Zachary Shaffer, Takao Sasaki, and Stephen C. Pratt. Linear recruitment leads to alloca-  
628 tion and flexibility in collective foraging by ants. *Animal Behaviour*, 86(5):967–75, sep  
629 2013. ISSN 00033472. doi: 10.1016/j.anbehav.2013.08.014. URL [http://linkinghub.  
630 elsevier.com/retrieve/pii/S0003347213003825](http://linkinghub.elsevier.com/retrieve/pii/S0003347213003825).
- 631 [10] Chris R. Reid and Tanya Latty. Collective behaviour and swarm intelligence in slime  
632 moulds. *FEMS Microbiology Reviews*, 40(6):798–806, 2016. ISSN 15746976. doi:  
633 10.1093/femsre/fuw033.
- 634 [11] Jens Krause and Graeme D. Ruxton. *Living in groups*. Oxford University Press, Oxford,  
635 New York, 2002. ISBN 978-0-19-850818-2.
- 636 [12] Charles Mackay. *Extraordinary Popular Delusions and the Madness of Crowds*. Richard  
637 Bentley, London, UK, 1841.
- 638 [13] Tatsuya Kameda and Reid Hastie. Herd Behavior. In *Emerging Trends in the Social and  
639 Behavioral Sciences: An Interdisciplinary, Searchable, and Linkable Resource*, pages 1–  
640 14. Wiley Online Library, 2015. ISBN 9781118900772.
- 641 [14] Gustav Le Bon. *The crowd: study of the popular mind*. Unwin, London, UK, 4th edition,  
642 1896.
- 643 [15] Ramsey M Raafat, Nick Chater, and Chris Frith. Herding in humans. *Trends in cognitive*

- 644 *sciences*, 13(10):420–8, oct 2009. ISSN 1364-6613. doi: 10.1016/j.tics.2009.08.002. URL  
645 <http://www.ncbi.nlm.nih.gov/pubmed/19748818>.
- 646 [16] Sushil Bikhchandani, David Hirshleifer, and Ivo Welch. A Theory of Fads, Fashion, Custom,  
647 and Cultural Change as Informational Cascades. *Journal of Political Economy*, 100  
648 (5):992, 1992. ISSN 0022-3808. doi: 10.1086/261849.
- 649 [17] V. V. Chari and Patrick J. Kehoe. Financial crises as herds: Overturning the critiques.  
650 *Journal of Economic Theory*, 119(1 SPEC. ISS.):128–150, 2004. ISSN 00220531. doi:  
651 10.1016/S0022-0531(03)00225-4.
- 652 [18] Irving L. Janis. *Victims of groupthink: A psychological study of foreign policy*. Houghton  
653 Mifflin Company, Boston, NY, 1972.
- 654 [19] Lev Muchnik, Sinan Aral, and Sean J Taylor. Social influence bias: a randomized exper-  
655 iment. *Science (New York, N.Y.)*, 341(6146):647–51, aug 2013. ISSN 1095-9203. doi:  
656 10.1126/science.1240466. URL <http://www.ncbi.nlm.nih.gov/pubmed/23929980>.
- 657 [20] Matthew J Salganik, Peter Sheridan Dodds, and Duncan J Watts. Experimental Study of  
658 Inequality and Cultural Market. *Science*, 311(February), 2006.
- 659 [21] Jan Lorenz, Heiko Rauhut, Frank Schweitzer, and Dirk Helbing. How social influence can  
660 undermine the wisdom of crowd effect. *Proceedings of the National Academy of Sciences of  
661 the United States of America*, 108(22):9020–5, may 2011. ISSN 1091-6490. doi: 10.1073/  
662 pnas.1008636108. URL [http://www.pubmedcentral.nih.gov/articlerender.  
663 fcgi?artid=3107299&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=3107299&tool=pmcentrez&rendertype=abstract).

- 664 [22] Bertrand Jayles, Hye-rin Kim, Ramón Escobedo, Stéphane Cezera, Adrien Blanchet, Tat-  
665 suya Kameda, Clément Sire, and Guy Theraulaz. How social information can improve  
666 estimation accuracy in human groups. *Proceedings of the National Academy of Sciences*,  
667 114(47):201703695, 2017. ISSN 0027-8424. doi: 10.1073/pnas.1703695114. URL  
668 <http://www.pnas.org/lookup/doi/10.1073/pnas.1703695114>.
- 669 [23] Luc-Alain Giraldeau, Thomas J Valone, and Jennifer J Templeton. Potential dis-  
670 advantages of using socially acquired information. *Philosophical transactions*  
671 *of the Royal Society of London. Series B, Biological sciences*, 357(1427):1559–  
672 66, nov 2002. ISSN 0962-8436. doi: 10.1098/rstb.2002.1065. URL [http:](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1693065&tool=pmcentrez&rendertype=abstract)  
673 [//www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1693065&tool=](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1693065&tool=pmcentrez&rendertype=abstract)  
674 [pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1693065&tool=pmcentrez&rendertype=abstract).
- 675 [24] Claire Detrain and Jean Louis Deneubourg. *Collective Decision-Making and Foraging Pat-*  
676 *terns in Ants and Honeybees*, volume 35. Elsevier, 2008. ISBN 9780123743299. doi: 10.  
677 1016/S0065-2806(08)00002-7. URL [http://dx.doi.org/10.1016/S0065-2806\(08\)](http://dx.doi.org/10.1016/S0065-2806(08)00002-7)  
678 [00002-7](http://dx.doi.org/10.1016/S0065-2806(08)00002-7).
- 679 [25] Christian List, Christian Elsholtz, and Thomas D Seeley. Independence and interdepen-  
680 dence in collective decision making: an agent-based model of nest-site choice by honeybee  
681 swarms. *Philosophical transactions of the Royal Society of London. Series B, Biological*  
682 *sciences*, 364(1518):755–762, 2009. ISSN 0962-8436. doi: 10.1098/rstb.2008.0277.
- 683 [26] J. L. Deneubourg, S. Aron, S. Goss, and J. M. Pasteels. The self-organizing exploratory pat-

- 684 tern of the argentine ant. *Journal of Insect Behavior*, 3(2):159–168, 1990. ISSN 08927553.  
685 doi: 10.1007/BF01417909.
- 686 [27] R Beckers, J L D Deneubourg, S Goss, and J M Pasteels. Collective decision making  
687 through food recruitment. *Insectes Sociaux*, 37:258–267, 1990.
- 688 [28] Robert Boyd and Peter J. Richerson. *Culture and the Evolutionary Process*. University of  
689 Chicago Press, Chicago, IL, 1985.
- 690 [29] Peter J. Richerson and Robert Boyd. *Not By Genes Alone*. University of Chicago Press,  
691 Chicago, IL, 2005.
- 692 [30] Marcus W. Feldman, Kenichi Aoki, and Jochen Kumm. Individual Versus Social Learn-  
693 ing: Evolutionary Analysis in a Fluctuating Environment. *Anthropological Science*,  
694 104(3):209–231, 1996. ISSN 0918-7960. doi: 10.1537/ase.104.209. URL [http:  
695 //www.santafe.edu/media/workingpapers/96-05-030.pdf](http://www.santafe.edu/media/workingpapers/96-05-030.pdf)  
696 [//publication/uuid/2408B672-F4CE-4EE0-A7B0-AFAC56879407](http://publication.uuid/uuid/2408B672-F4CE-4EE0-A7B0-AFAC56879407).
- 697 [31] Kevin N. Laland. Social learning strategies. *Animal Learning & Behavior*, 32(1):4–14,  
698 2004. ISSN 0090-4996. doi: 10.3758/BF03196002.
- 699 [32] Tatsuya Kameda and Daisuke Nakanishi. Cost–benefit analysis of social/cultural learn-  
700 ing in a nonstationary uncertain environment. *Evolution and Human Behavior*, 23(5):  
701 373–393, 2002. ISSN 10905138. doi: 10.1016/S1090-5138(02)00101-0. URL [http:  
702 //www.sciencedirect.com/science/article/pii/S1090513802001010](http://www.sciencedirect.com/science/article/pii/S1090513802001010).
- 703 [33] Rachel L. Kendal, Isabelle Coolen, and Kevin N. Laland. The role of conformity in foraging

- 704 when personal and social information conflict. *Behavioral Ecology*, 15(2):269–277, 2004.  
705 ISSN 10452249. doi: 10.1093/beheco/arh008.
- 706 [34] T J H Morgan, L E Rendell, M Ehn, W Hoppitt, and K N Laland. The evolutionary basis of  
707 human social learning. *Proceedings. Biological sciences / The Royal Society*, 279(1729):  
708 653–62, feb 2012. ISSN 1471-2954. doi: 10.1098/rspb.2011.1172. URL [http://www.  
709 ncbi.nlm.nih.gov/pubmed/21795267](http://www.ncbi.nlm.nih.gov/pubmed/21795267).
- 710 [35] Wataru Toyokawa, Yoshimatsu Saito, and Tatsuya Kameda. Individual differences in learn-  
711 ing behaviours in humans: asocial exploration tendency does not predict reliance on so-  
712 cial learning. *Evolution and Human Behavior*, 38(3):325–333, 2017. ISSN 10905138.  
713 doi: 10.1016/j.evolhumbehav.2016.11.001. URL [http://linkinghub.elsevier.com/  
714 retrieve/pii/S1090513816301180](http://linkinghub.elsevier.com/retrieve/pii/S1090513816301180).
- 715 [36] M.M Webster and K.N Laland. Social learning strategies and predation risk: minnows copy  
716 only when using private information would be costly. *Proceedings of the Royal Society B:  
717 Biological Sciences*, 275(1653):2869–2876, 2008. ISSN 0962-8452. doi: 10.1098/rspb.  
718 2008.0817. URL [http://rspb.royalsocietypublishing.org/cgi/doi/10.1098/  
719 rspb.2008.0817](http://rspb.royalsocietypublishing.org/cgi/doi/10.1098/rspb.2008.0817).
- 720 [37] M M Webster and K N Laland. Reproductive state affects reliance on public information  
721 in sticklebacks. *Proceedings. Biological sciences / The Royal Society*, 278(1705):619–627,  
722 2011. ISSN 0962-8452. doi: 10.1098/rspb.2010.1562.
- 723 [38] Robert Boyd and Peter J. Richerson. Social learning as an adaptation. *Lectures on Mathe-  
724 matics in the Life Sciences*, 20:1–26, 1989.

- 725 [39] R. Bond. Group Size and Conformity. *Group Processes & Intergroup Relations*, 8(4):  
726 331–354, oct 2005. ISSN 1368-4302. doi: 10.1177/1368430205056464. URL [http:  
727 //gpi.sagepub.com/cgi/doi/10.1177/1368430205056464](http://gpi.sagepub.com/cgi/doi/10.1177/1368430205056464).
- 728 [40] M. A. Kline and R. Boyd. Population size predicts technological complexity in  
729 Oceania. *Proceedings of the Royal Society B: Biological Sciences*, 277(1693):2559–  
730 2564, 2010. ISSN 0962-8452. doi: 10.1098/rspb.2010.0452. URL [http://rspb.  
731 royalsocietypublishing.org/cgi/doi/10.1098/rspb.2010.0452](http://rspb.royalsocietypublishing.org/cgi/doi/10.1098/rspb.2010.0452).
- 732 [41] Michael Muthukrishna, Ben W Shulman, Vlad Vasilescu, and Joseph Henrich. Sociality  
733 influences cultural complexity. *Proceedings of Royal Society B*, 281(November 2013):  
734 20132511, 2014.
- 735 [42] Sally E Street, Ana F Navarrete, Simon M Reader, and Kevin N Laland. Coevolution of  
736 cultural intelligence , extended life history , sociality , and brain size in primates. *Pro-  
737 ceedings of the National Academy of Sciences*, 114(30):1–7, 2017. ISSN 0027-8424. doi:  
738 10.1073/pnas.1620734114.
- 739 [43] Sc Nicolis and JI Deneubourg. Emerging patterns and food recruitment in ants: an analytical  
740 study. *Journal of theoretical biology*, 198(4):575–92, jul 1999. ISSN 1095-8541. doi:  
741 10.1006/jtbi.1999.0934. URL <http://www.ncbi.nlm.nih.gov/pubmed/10373356>.
- 742 [44] Stephen C Pratt and David J T Sumpter. A tunable algorithm for collective decision-  
743 making. *Proceedings of the National Academy of Sciences of the United States  
744 of America*, 103(43):15906–10, oct 2006. ISSN 0027-8424. doi: 10.1073/pnas.

- 745 0604801103. URL [http://www.pubmedcentral.nih.gov/articlerender.fcgi?](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1635101&tool=pmcentrez&rendertype=abstract)  
746 [artid=1635101&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1635101&tool=pmcentrez&rendertype=abstract).
- 747 [45] Christian List. Democracy in animal groups: a political science perspective. *Trends in*  
748 *Ecology and Evolution*, 19(4):166–168, 2004. ISSN 01695347. doi: 10.1016/j.tree.2004.  
749 02.001.
- 750 [46] Andrew J King and Guy Cowlshaw. When to use social information: the advantage of  
751 large group size in individual decision making. *Biology letters*, 3(2):137–139, 2007. ISSN  
752 1744-9561. doi: 10.1098/rsbl.2007.0017.
- 753 [47] Max Wolf, Ralf H J M Kurvers, Ashley J W Ward, Stefan Krause, and Jens Krause.  
754 Accurate decisions in an uncertain world: collective cognition increases true positives  
755 while decreasing false positives. *Proceedings. Biological sciences / The Royal Society*,  
756 280(1756):20122777, apr 2013. ISSN 1471-2954. doi: 10.1098/rspb.2012.2777. URL  
757 <http://www.ncbi.nlm.nih.gov/pubmed/23407830>.
- 758 [48] Andres Laan, Gabriel Madirolas, and Gonzalo G De Polavieja. Rescuing collective wisdom  
759 when the average group opinion is wrong. *Frontiers in Robotics and AI*, 4(November):1–28,  
760 2017. ISSN 2296-9144. doi: 10.3389/frobt.2017.00056.
- 761 [49] Lucy M. Aplin, Ben C. Sheldon, and Richard McElreath. Conformity does not per-  
762 petuate suboptimal traditions in a wild population of songbirds. *Proceedings of the*  
763 *National Academy of Sciences*, page 201621067, 2017. ISSN 0027-8424. doi: 10.  
764 1073/PNAS.1621067114. URL [http://www.pnas.org/content/early/2017/07/18/](http://www.pnas.org/content/early/2017/07/18/1621067114.full)  
765 [1621067114.full](http://www.pnas.org/content/early/2017/07/18/1621067114.full).

- 766 [50] Brendan J Barrett, Richard L McElreath, Susan E Perry, and Brendan J Barrett. Pay-off-  
767 biased social learning underlies the diffusion of novel extractive foraging traditions in a  
768 wild primate. *Proceedings of Royal Society B*, 284, 2017.
- 769 [51] Richard McElreath, Adrian V Bell, Charles Efferson, Mark Lubell, Peter J Rich-  
770 erson, and Timothy Waring. Beyond existence and aiming outside the labora-  
771 tory: estimating frequency-dependent and pay-off-biased social learning strategies.  
772 *Philosophical transactions of the Royal Society of London. Series B, Biological sci-*  
773 *ences*, 363(1509):3515–28, nov 2008. ISSN 1471-2970. doi: 10.1098/rstb.2008.  
774 0131. URL [http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2607339&tool=pmcentrez&rendertype=abstract)  
775 [2607339&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2607339&tool=pmcentrez&rendertype=abstract).
- 776 [52] Wataru Toyokawa, Hye-Rin Kim, and Tatsuya Kameda. Human Collective  
777 Intelligence under Dual Exploration-Exploitation Dilemmas. *PloS one*, 9(4):  
778 e95789, jan 2014. ISSN 1932-6203. doi: 10.1371/journal.pone.0095789.  
779 URL [http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=3995913&tool=pmcentrez&rendertype=abstract)  
780 [3995913&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=3995913&tool=pmcentrez&rendertype=abstract).
- 781 [53] Anne Kandler and Kevin N Laland. Tradeoffs between the strength of conformity and  
782 number of conformists in variable environments. *Journal of theoretical biology*, 332:191–  
783 202, sep 2013. ISSN 1095-8541. doi: 10.1016/j.jtbi.2013.04.023. URL [http://www.](http://www.ncbi.nlm.nih.gov/pubmed/23643630)  
784 [ncbi.nlm.nih.gov/pubmed/23643630](http://www.ncbi.nlm.nih.gov/pubmed/23643630).
- 785 [54] Charles Efferson, Rafael Lalive, Peter J. Richerson, Richard McElreath, and Mark Lubell.  
786 Conformists and mavericks: the empirics of frequency-dependent cultural transmission.

787 *Evolution and Human Behavior*, 29:56–64, 2008. ISSN 10905138. doi: 10.1016/j.  
788 evolhumbehav.2007.08.003.

789 [55] Richard McElreath, Mark Lubell, Peter J. Richerson, Timothy M. Waring, William Baum,  
790 Edward Edsten, Charles Efferson, and Brian Paciotti. Applying evolutionary models to  
791 the laboratory study of social learning. *Evolution and Human Behavior*, 26(6):483–508,  
792 nov 2005. ISSN 10905138. doi: 10.1016/j.evolhumbehav.2005.04.003. URL [http://](http://linkinghub.elsevier.com/retrieve/pii/S1090513805000231)  
793 [linkinghub.elsevier.com/retrieve/pii/S1090513805000231](http://linkinghub.elsevier.com/retrieve/pii/S1090513805000231).

794 [56] Alex Mesoudi. An experimental comparison of human social learning strategies: payoff-  
795 biased social learning is adaptive but underused. *Evolution and Human Behavior*, 32(5):  
796 334–342, sep 2011. ISSN 10905138. doi: 10.1016/j.evolhumbehav.2010.12.001. URL  
797 <http://linkinghub.elsevier.com/retrieve/pii/S1090513810001443>.

798 [57] Charles Perreault, Cristina Moya, and Robert Boyd. A Bayesian approach to the evo-  
799 lution of social learning. *Evolution and Human Behavior*, 33(5):449–459, sep 2012.  
800 ISSN 10905138. doi: 10.1016/j.evolhumbehav.2011.12.007. URL [http://linkinghub.](http://linkinghub.elsevier.com/retrieve/pii/S1090513811001437)  
801 [elsevier.com/retrieve/pii/S1090513811001437](http://linkinghub.elsevier.com/retrieve/pii/S1090513811001437).

802 [58] L Rendell, R Boyd, D Cownden, M Enquist, K Eriksson, M W Feldman, L Fog-  
803 arty, S Ghirlanda, T Lillicrap, and K N Laland. Why copy others? Insights  
804 from the social learning strategies tournament. *Science (New York, N.Y.)*, 328  
805 (5975):208–13, apr 2010. ISSN 1095-9203. doi: 10.1126/science.1184719.  
806 URL [http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2989663&tool=pmcentrez&rendertype=abstract)  
807 [2989663&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2989663&tool=pmcentrez&rendertype=abstract).

- 808 [59] Jolle W Jolles, Kate L Laskowski, Neeltje J Boogert, and Andrea Manica. Repeatable group  
809 differences in the collective behaviour of stickleback shoals across ecological contexts. *Pro-*  
810 *ceedings of Royal Society B*, pages 13–16, 2018.
- 811 [60] Pablo Michelena, Raphaël Jeanson, Jean-Louis Deneubourg, and Angela M Sibbald. Per-  
812 sonality and collective decision-making in foraging herbivores. *Proceedings. Biological*  
813 *sciences / The Royal Society*, 277(1684):1093–9, apr 2010. ISSN 1471-2954. doi: 10.  
814 1098/rspb.2009.1926. URL [http://www.pubmedcentral.nih.gov/articlerender.](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2842771&tool=pmcentrez&rendertype=abstract)  
815 [fcgi?artid=2842771&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2842771&tool=pmcentrez&rendertype=abstract).
- 816 [61] Isaac Planas-sitja, Jean-louis Deneubourg, Celine Gibon, and Gregory Sempo. Group per-  
817 sonality during collective decision-making : a multi-level approach. *Proceedings of the*  
818 *Royal Society B: Biological Sciences*, 282:20142515, 2015.
- 819 [62] Alex Mesoudi, Lei Chang, Sasha R.X. Dall, and Alex Thornton. The Evolution of Indi-  
820 vidual and Cultural Variation in Social Learning. *Trends in Ecology & Evolution*, xx:1–11,  
821 2016. ISSN 01695347. doi: 10.1016/j.tree.2015.12.012. URL [http://linkinghub.](http://linkinghub.elsevier.com/retrieve/pii/S0169534715003237)  
822 [elsevier.com/retrieve/pii/S0169534715003237](http://linkinghub.elsevier.com/retrieve/pii/S0169534715003237).
- 823 [63] Brendan J Barrett. Equifinality in empirical studies of cultural transmission. *Behavioural*  
824 *Processes*, 2018. ISSN 03766357. doi: 10.1016/j.beproc.2018.01.011. URL [http://dx.](http://dx.doi.org/10.1016/j.beproc.2018.01.011)  
825 [doi.org/10.1016/j.beproc.2018.01.011](http://dx.doi.org/10.1016/j.beproc.2018.01.011).
- 826 [64] Dora Biro, Takao Sasaki, and Steven J. Portugal. Bringing a Time-Depth Perspective to  
827 Collective Animal Behaviour. *Trends in Ecology and Evolution*, 31(7):550–562, 2016.

828 ISSN 01695347. doi: 10.1016/j.tree.2016.03.018. URL <http://dx.doi.org/10.1016/>  
829 [j.tree.2016.03.018](http://dx.doi.org/10.1016/j.tree.2016.03.018).

830 [65] William Hoppitt and Kevin N. Laland. *Social Learning: An Introduction to Mecha-*  
831 *nisms, Methods, and Models*. Princeton University Press, 2013. ISBN 0691150702,  
832 9780691150703.

833 [66] Ulf Toelch, Matthew J. Bruce, Marius T.H. Meeus, and Simon M. Reader. Humans copy  
834 rapidly increasing choices in a multiarmed bandit problem. *Evolution and Human Behavior*,  
835 31(5):326–333, sep 2010. ISSN 10905138. doi: 10.1016/j.evolhumbehav.2010.03.002.  
836 URL <http://linkinghub.elsevier.com/retrieve/pii/S1090513810000292>.

837 [67] Bahador Bahrami, Karsten Olsen, Peter E. Latham, Andreas Roepstorff, Geraint Rees, and  
838 Chris D. Frith. Optimally interacting minds. *Science*, 329:1081–1085, 2010. doi: 10.5061/  
839 dryad.5t110.Supplementary.

840 [68] Nathaniel D Daw, John P O’Doherty, Peter Dayan, Ben Seymour, and Ray-  
841 mond J Dolan. Cortical substrates for exploratory decisions in humans. *Na-*  
842 *ture*, 441(7095):876–9, jun 2006. ISSN 1476-4687. doi: 10.1038/nature04766.  
843 URL [http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2635947&tool=pmcentrez&rendertype=abstract)  
844 [2635947&tool=pmcentrez&rendertype=abstract](http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2635947&tool=pmcentrez&rendertype=abstract).

845 [69] Frédéric Dandurand, Thomas R. ShultzEmail, and Kristine H. Onishi. Comparing online  
846 and lab methods in a problem-solving experiment. *Behavior Research Methods*, 40(2):  
847 428–434, 2008. doi: 10.3758/BRM.40.2.428.

- 848 [70] Jérôme Hergueux and Nicolas Jacquemet. Social preferences in the online laboratory: a  
849 randomized experiment. *Experimental Economics*, 18(2):251–283, 2015. ISSN 13864157.  
850 doi: 10.1007/s10683-014-9400-5.
- 851 [71] Joshua Becker, Devon Brackbill, and Damon Centola. Network dynamics of social in-  
852 fluence in the wisdom of crowds. *Proceedings of the National Academy of Sciences*,  
853 page 201615978, 2017. ISSN 0027-8424. doi: 10.1073/pnas.1615978114. URL [http:  
854 //www.pnas.org/lookup/doi/10.1073/pnas.1615978114](http://www.pnas.org/lookup/doi/10.1073/pnas.1615978114).
- 855 [72] I. Lorge, D. Fox, J. Davitz, and M. Brenner. A survey of studies contrasting the quality of  
856 group performance and individual performance, 1920-1957. *Psychological Bulletin*, 55(6):  
857 337–372, 1958. ISSN 0033-2909. doi: 10.1037/h0042344.
- 858 [73] Reid Hastie and Tatsuya Kameda. The robust beauty of majority rules in group deci-  
859 sions. *Psychological review*, 112(2):494–508, apr 2005. ISSN 0033-295X. doi: 10.1037/  
860 0033-295X.112.2.494. URL <http://www.ncbi.nlm.nih.gov/pubmed/15783295>.
- 861 [74] Elise Payzan-Lenestour and Peter Bossaerts. Risk, unexpected uncertainty, and estimation  
862 uncertainty: Bayesian learning in unstable settings. *PLoS Computational Biology*, 7(1),  
863 2011. ISSN 1553734X. doi: 10.1371/journal.pcbi.1001048.
- 864 [75] R S Sutton and a G Barto. *Reinforcement learning: an introduction*. MIT Press, Cambridge,  
865 MA, 1998. ISBN 0262193981. doi: 10.1109/TNN.1998.712192.
- 866 [76] Woo Young Ahn, Georgi Vasilev, Sung Ha Lee, Jerome R. Busemeyer, John K. Kruschke,  
867 Antoine Bechara, and Jasmin Vassileva. Decision-making in stimulant and opiate addicts in

868 protracted abstinence: Evidence from computational modeling with pure users. *Frontiers*  
869 *in Psychology*, 5(AUG):1–15, 2014. ISSN 16641078. doi: 10.3389/fpsyg.2014.00849.

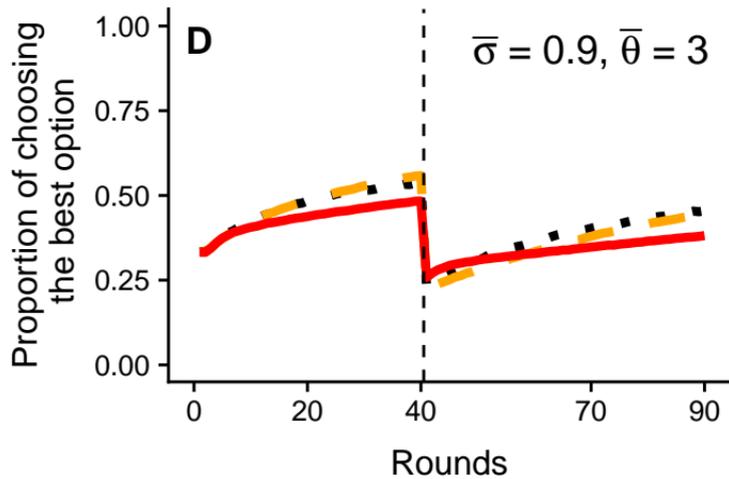
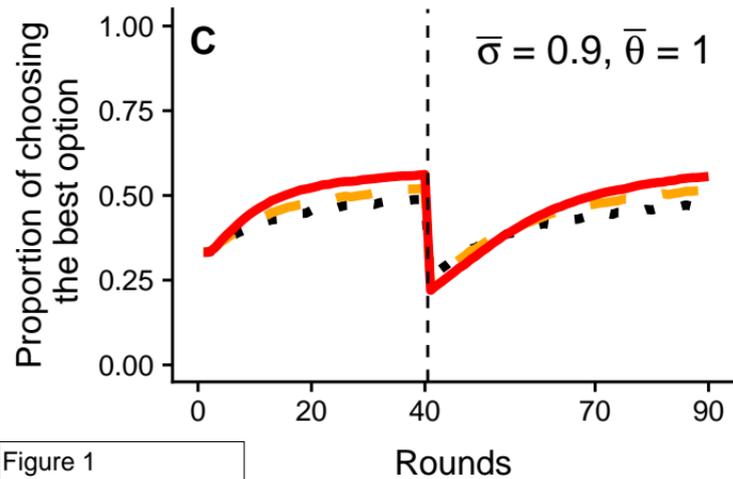
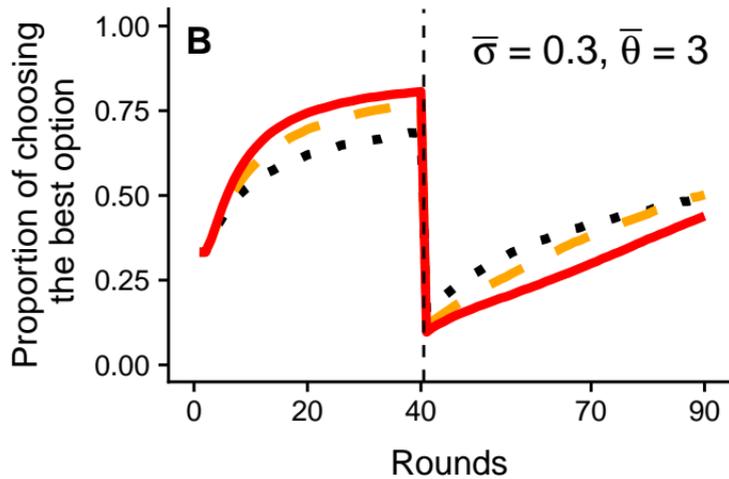
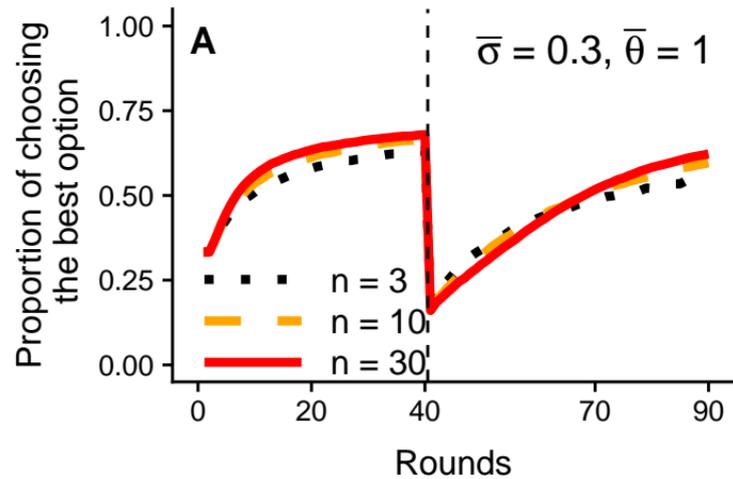
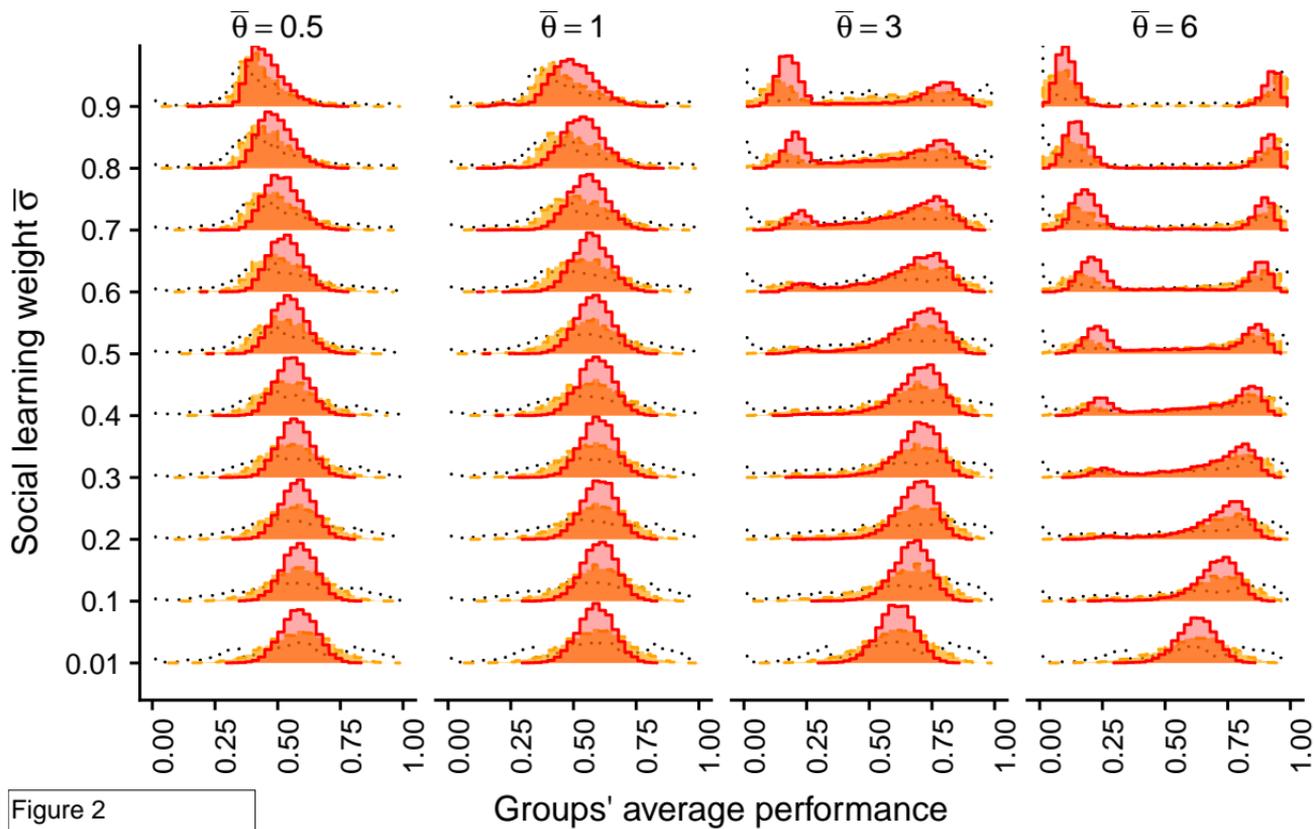


Figure 1

Group size:  $\dots$   $n = 3$ ;  $\square$   $n = 10$ ;  $\square$   $n = 30$



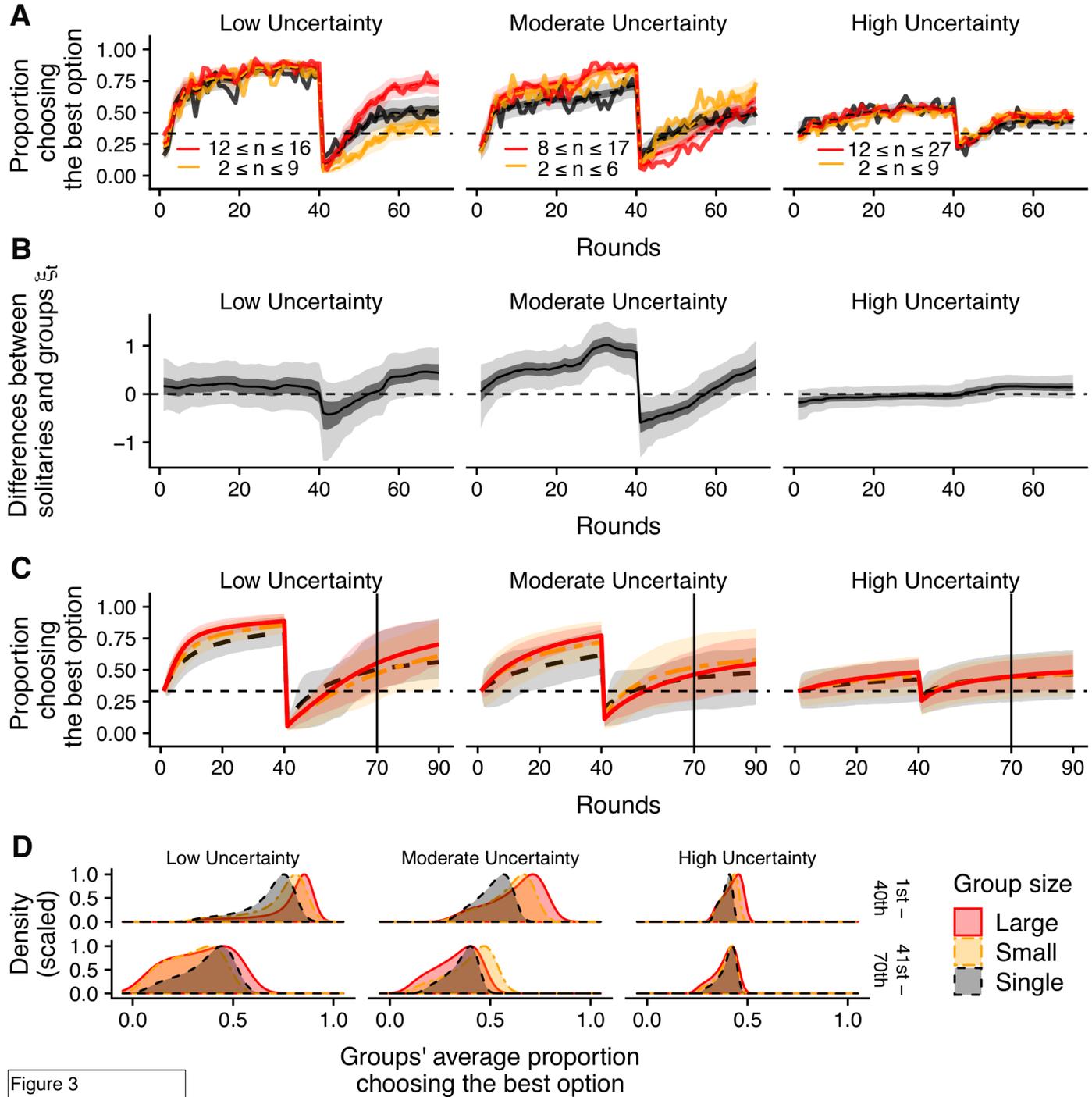


Figure 3

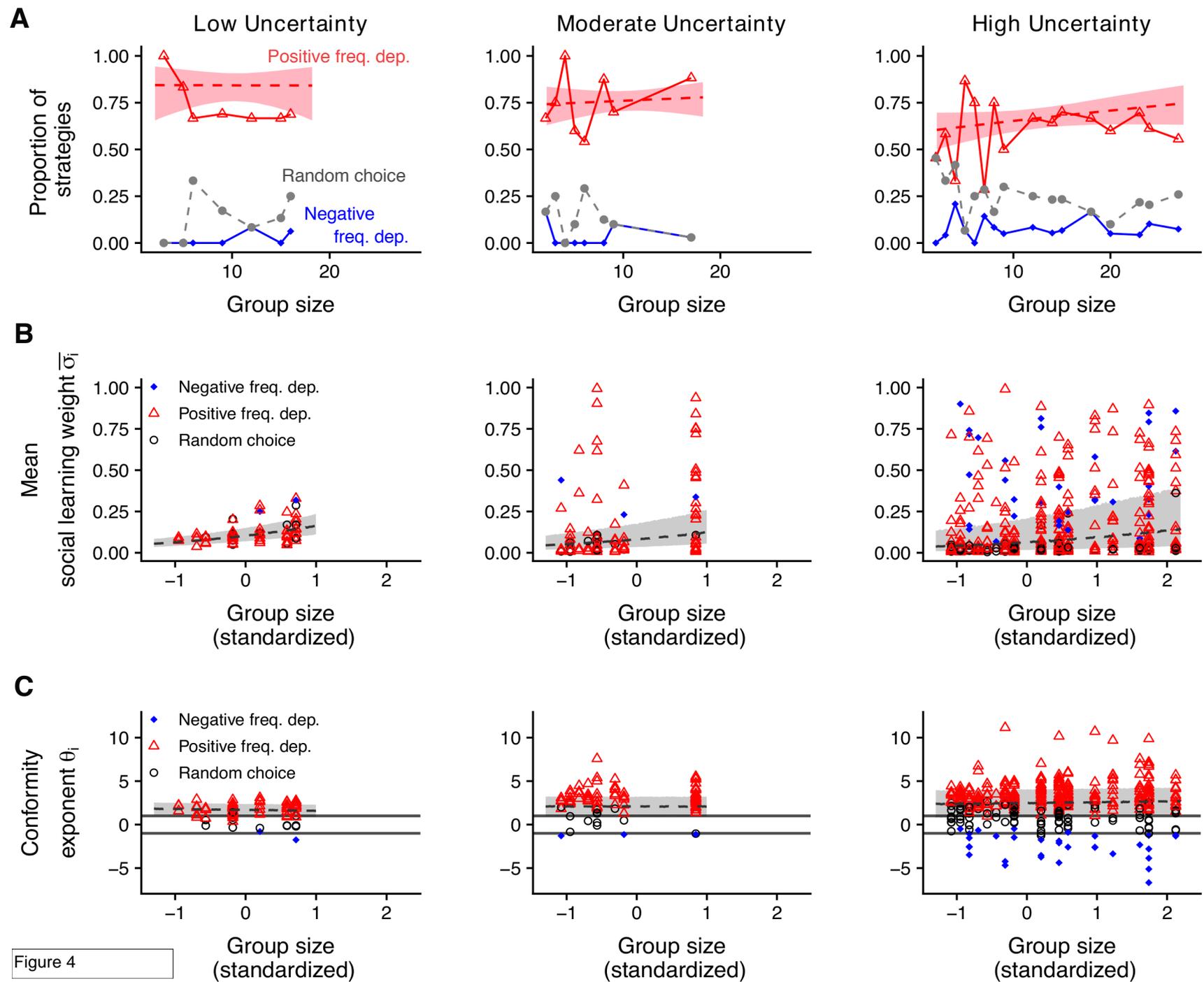


Figure 4

