

1 Applying evolutionary models to the laboratory study of social
2 learning

3 Richard McElreath^{1,2,3,1} Mark Lubell^{2,4} Peter J. Richerson^{2,3,4}
4 Timothy M. Waring^{2,4} William Baum⁴ Edward Edsten³ Charles Efferson^{2,4}
5 Brian Paciotti^{2,4}

6 ¹*Department of Anthropology, UC Davis, Davis CA 95616.*

7 ²*Graduate Group in Ecology, UC Davis, Davis CA 95616.*

8 ³*Animal Behavior Graduate Group, UC Davis, Davis CA 95616.*

9 ⁴*Department of Environmental Science and Policy, UC Davis, Davis CA 95616.*

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¹Corresponding author. Email: mcelreath@ucdavis.edu. Phone: (530) 752-2660. Fax: (530) 752-8885.

Abstract

14 Cultural evolution is driven in part by the strategies individuals employ to acquire behavior from
15 others. These strategies themselves are partly products of natural selection, making the study of so-
16 cial learning an inherently Darwinian project. Formal models of the evolution of social learning sug-
17 gest that reliance on social learning should increase with task difficulty and decrease with the proba-
18 bility of environmental change. These models also make predictions about how individuals integrate
19 information from multiple peers. We present the results of micro-society experiments designed to
20 evaluate these predictions. The first experiment measures baseline individual learning strategy in a
21 two-armed bandit environment with variation in task difficulty and temporal fluctuation in the pay-
22 offs of the options. Our second experiment addresses how people in the same environment use mini-
23 mal social information from a single peer. Our third experiment expands on the second by allowing
24 access to the behavior of several other individuals, permitting frequency-dependent strategies like
25 conformity. In each of these experiments, we vary task difficulty and environmental fluctuation. We
26 present several candidate strategies and compute the expected payoffs to each in our experimental
27 environment. We then fit to the data the different models of the use of social information, and iden-
28 tify the best-fitting model via model comparison techniques. We find substantial evidence of both
29 conformist and non-conformist social learning and compare our results to theoretical expectations.

32 1 Introduction

33 Unlike most animals, humans acquire large and important parts of their behavioral repertoire via
34 imitation and other forms of social learning. Therefore, students of human behavior seek to un-
35 derstand how individuals acquire beliefs and behavior from their parents, peers and others. At
36 another level, social scientists attempt to fathom the resulting complex interactions that take place
37 at the level of the society. Whether one is interested in the emergence of political institutions,
38 languages, art, technologies, or moral traditions, these cultural elements all arose over long time
39 periods through the combined effects of many individual-level decisions. Understanding how people
40 use information available from the behavior of others is therefore important not only for under-
41 standing individual decisions, but also for comprehending patterns of change and variation among
42 human societies. And since the psychological mechanisms that make social learning possible are
43 partly products of natural selection, evolutionary models are necessary to fully understand their
44 design.

45 In this paper, we use micro-society experiments to investigate the psychological foundations of
46 social learning. Experimental micro-societies (Baum *et al.*, 2004) consist of human participants who
47 repeatedly interact in controlled ways within a laboratory. Over a series of rounds, the participants
48 make decisions that lead to real payoffs, receive feedback, and can access some information about
49 the decisions of their peers. Thus choices evolve over time, in response to both individual and
50 social learning. Our goal is to test and refine hypotheses, developed through formal models of the
51 evolution of cultural evolution, about how people regulate their reliance on individual and social
52 learning and the structural details of how people use social information. We are interested in (1)
53 how social learning changes in response to the difficulty of a task and (2) how it changes in response
54 to the probability of changes in the environment. We develop computational models for estimating

55 decision-making strategies and the strength of different components of these strategies. We find
56 considerable evidence of social learning, and the details of the strategies used in any particular
57 treatment tend to agree with numerical analyses of strategy efficacy. Nevertheless, individuals
58 sometimes imitate in ways that are not obviously profitable. Our results match some of the model
59 predictions concerning task difficulty and environmental change, but by no means all of them.

60 The first section of the paper reviews the theory that motivates our experiments and lays out
61 qualitative predictions for how people will respond to changes in different kinds of environmental
62 uncertainty. Then we present in detail our experimental choice environment and analyze how human
63 participants learn on their own within it. These estimates allow us to analyze the effectiveness of
64 different social learning strategies within our experimental environment. We present a quantitative
65 analysis of the payoffs to three different social learning strategies and the optimal reliance on each,
66 as functions of two types of environmental uncertainty. Then we present two experiments that
67 allow access to social information and estimate participant strategies in each. Finally, we relate the
68 findings to the predictions derived from the theoretical literature and our own analysis.

69 **2 The evolution of social learning**

70 In economics and political science, researchers seemingly discovered social learning as a “rational”
71 phenomenon in the early 1990s (Banerjee, 1992; Bikhchandani *et al.*, 1992). These models some-
72 times go by the labels of “herding” or “herd behavior” and other times as “informational cascades.”
73 A number of models have been developed that show how ignoring private information and choosing
74 based upon the behavior of others can be optimal. This result is surprising to many social scientists,
75 because the intuition dominating the study of judgment and decision-making has long been that
76 more objective information improves decisions (Gigerenzer *et al.*, 1999, demonstrate other ways in

77 which less information use can be optimal).

78 These models are very similar to models of social learning first derived in biology and anthro-
79 pology in the 1980s. Formal models by Boyd and Richerson and others derived conditions for
80 natural selection to favor various forms of imitation (Boyd & Richerson, 1985; Rogers, 1988). A
81 rich body of theory now exists arguing both that natural selection will often favor an extensive
82 reliance on imitation and that imitation can lead to unanticipated population-level effects (Henrich
83 & McElreath, 2003, review much of it). A robust result of these models is that social learning never
84 entirely replaces individual learning—no matter how difficult and costly—but that social learning
85 may be broadly adaptive even when it severely undermines a society’s ability to track changes in the
86 environment or leads to the spread of maladaptive behavior (Boyd & Richerson, 1985; Richerson
87 & Boyd, 2004).

88 A handful of empirical studies have addressed formal models of social learning, whether its
89 general properties (Kameda & Nakanishi, 2002; Anderson & Holt, 1997; Kameda & Nakanishi,
90 2003) or specific aspects of its design (Kameda & Nakanishi, 2002; Henrich, 2001; Schotter &
91 Sopher, 2003; McElreath, 2004; Apestegua *et al.*, 2003; Galef & Whiskin, 2004; Coultas, 2004;
92 Camerer & Ho, 1999). Social psychologists, most notably Albert Bandura (Bandura, 1977), of
93 course did a lot of work in 1970’s exploring the existence of various cues people use in social
94 learning. However, no formal models of social learning developed in psychology, and most of this
95 work was guided by intuition alone and lead to no enduring analytical work. Economists interested
96 in learning in games have developed a handful of candidate models that may honestly be called
97 “social learning models” (Camerer, 2003). These models are very descriptive and consider only
98 a tiny fraction of the social learning strategies specified in the evolutionary models developed by
99 Boyd and Richerson and others.

100 We are interested in addressing two variables that recur in many of the formal evolutionary
101 models: (1) the difficulty of learning on one's own and (2) the frequency of temporal fluctuations
102 in the payoffs of behavioral options. These can both be thought of as types of environmental
103 uncertainty, however existing models show that each leads to different qualitative effects on the
104 evolutionarily stable amount of social learning.

105 First, when it is difficult for individuals to determine the best behavior on their own, a greater
106 reliance on social learning arises at equilibrium (Boyd & Richerson, 1985; Rogers, 1988; Henrich &
107 Boyd, 1998). One way individual learning might be error prone is if the information available to
108 individuals is of poor quality. For example, if inter-annual variation in crop yields is large, learners
109 will have difficulty telling if some change in cultivation improves yield in the long-run. Crop yield
110 is uncertain, but there is a single best crop. A greater reliance on social learning evolves because
111 social learning can both reduce noise in estimates obtained individually as well as help one avoid
112 costly mistakes others have already endured.

113 Second, a principle problem with imitation is that changes in the environment may make past
114 behavior a poor guide to current payoffs. Environments are not perfectly stationary. If the climate
115 or pest populations change, it may no longer be a good idea to plant what one's father planted.
116 Thus when the frequency of such changes is high, less social learning exists at equilibrium (Boyd &
117 Richerson, 1985; Rogers, 1988; Henrich & Boyd, 1998). Essentially, environmental fluctuation can
118 render useless the adaptive knowledge stored in cultural systems. This fluctuation is another kind
119 of uncertainty, but it reduces rather than increases the amount of social learning at equilibrium.

120 On the basis of these models, as individual learning becomes more difficult, we expect more
121 social learning, and as the probability of environmental change increases, we expect less social
122 learning. While the evolutionary models do not contain enough psychological detail to say if

123 individuals should facultatively adjust reliance on social learning in different contexts, we think it
124 is reasonable to interpret the intuitions of these models in this way. Based upon cues of difficulty
125 of learning or fluctuation in the environment, people might adaptively regulate their attention to
126 the behavior of others (McElreath, 2004). People may have developed suites of adaptive strategies
127 from which they select, depending on different environmental cues.

128 Coincidental support for the prediction concerning task difficulty comes from a study of con-
129 formity in a perceptual task by Baron et al (1996), in which conformity appears stronger when the
130 task is made more difficult. Another study that indirectly supports these predictions is a study of
131 consumer choice (Fasolo, 2002), in which participants indicated they would be more likely to use
132 the opinions of their peers in a treatment in which there was no clear best option. McElreath (2004)
133 finds field evidence consistent with qualitative predictions about difficulty of learning. Galef and
134 Whiskin (2004) study the effects of environmental change on social learning in rats, and find results
135 that qualitatively support model predictions. Pingle (1995) constructed a production task in which
136 participants could see the choices of others and finds imitation when changes in the production
137 function (the underlying payoffs to options) were announced.

138 Most of the formal models addressing this problem have modeled imitation of members of a
139 previous generation—parents, elders, older siblings—rather than imitation of peers. In our experi-
140 ments, there are no naive individuals imitating experienced individuals, as in the models. Instead,
141 individuals of equal experience have the opportunity to imitate one another. The same predic-
142 tions hold in this purely-horizontal case, however, with some caveats. We demonstrate this in a
143 later section, in which we simulate the performance of different social learning strategies, combined
144 with estimates of how individuals learn individually in our experimental design. First, however,
145 we present the decision-making environment for our experiments and a pure individual learning

146 experiment.

147 **3 Experiment 1: Individual learning**

148 To correctly estimate the use of social learning, we have to take individual learning seriously. In the
149 first experiment, we introduce the task environment and explore patterns of individual learning,
150 before introducing the possibility of social learning in later experiments.

151 **3.1 Participants**

152 36 undergraduates at UC Davis participating in a psychology subject pool took part in this ex-
153 periment. They participated in groups of 6-10, but each made individual decisions and interacted
154 only with their computer terminal through the course of the experiment. Each participant re-
155 ceived course extra credit, in addition to their monetary earnings (see below), for completing the
156 experiment.

157 **3.2 Design**

158 The experiment was programmed using z-Tree (Fischbacher, 2002, the Zürich Toolbox for Ready-
159 made Economic Experiments) and administered via computer. The experimental task was framed
160 as a simulated agricultural choice. Each participant faced the decision of planting one of two al-
161 ternative crops (“wheat” or “potatoes”) each of 20 seasons on each of 6 sequentially encountered
162 farms (for a total of 120 decisions per participant). On each farm, the mean yield of one crop
163 was higher than the other, but which was higher was random on each farm. Each season, each
164 participant made a planting decision and was informed of his or her yield from this decision. Only
165 the most recent yield was ever displayed to the participant, and obviously no previous yield was

166 displayed in the first season on each farm.

167 On the first three farms, participants were told that the means were constant on each farm across
168 seasons, but potentially different across farms. In the last three farms, participants were told that
169 which crop was best could change in any given season, and that changes occurred randomly each
170 season, with a chance of 1/20 (communicated as a fraction).

171 The yields for crop i in each season were drawn from a normal distribution with mean μ_i and
172 variance σ^2 , $y_i \sim \mathcal{N}(\mu_i, \sigma^2)$. The variance was the same for both crops, while the mean of the more
173 profitable crop was 13 units and that of the less profitable 10 units. Participants were told at the
174 beginning of the experiment that they would be paid US\$.045 per 10 units of yield, for average
175 total winnings between \$4 and \$8. The stated goal was to maximize their yield by planting the
176 crop with the higher mean yield. We manipulated σ^2 to adjust the difficulty of learning on each
177 farm. When the variance in yield is large, it is harder to learn which of the two crops is best.
178 When the variance is small, the quality of information obtained from planting is much better, and
179 consequently individual learning more easily discovers the best crop. Each participant planted on
180 farms with three different unknown (but stable) variances in yield: 0.25, 4 and 16. The different
181 variances came in random order for each participant, although the sequence was the same for each
182 participant on the first three and last three farms.

183 In one extra session using 8 participants, we doubled the stakes to check for any large motivation
184 effects. The proportion of optimal planting decisions in this session was slightly lower than the other
185 sessions. We concluded that any motivational effect from the size of the stakes was quite minor
186 relative to the variation in behavior in the experiment.

187 3.3 Results

188 [Figure 1 about here.]

189 [Table 1 about here.]

190 This decision environment is a variant of the common two-arm bandit with a finite horizon.
191 There is a considerable literature on optimal strategies in such environments (Anderson, 2001;
192 Gittins, 1989), however it is usually very difficult or impossible to actually compute optimal choices
193 in practice. A smaller number of researchers have investigated how people actually make decisions
194 in these environments (Banks *et al.*, 1997; Gans *et al.*, 2003; Horowitz, 1973; Meyer & Shi, 1995),
195 and we know of only one serious study of a Gaussian bandit like our own (Anderson, 2001). There
196 are many possible models (Camerer, 2003), based upon several different views of learning. Our
197 goal here is not to improve upon this literature, but instead to find a robust individual learning
198 model (or models) that we can use as the basis of more complex models in later experiments. The
199 models we fit in this section are minimally-parametric generalizations of some popular candidates.

200 Both changing the variance of yields and the probability of fluctuation had noticeable effects
201 on the rate of optimal choices. The probabilities of optimal planting decisions under different
202 treatments is summarized in Table 1. Both variance and fluctuation in the means lowers the rate
203 of optimal choices. Under lower variance, individuals eventually reach much higher optimal choice
204 rates.

205 [Table 2 about here.]

206 [Table 3 about here.]

207 In order to address how participants were using yields to make choices, we fit three different
208 individual learning models to the 4320 planting decisions from this experiment. This allows us to

209 narrow the candidate individual-learning models to use in later fitting exercises. The three models
 210 are explained in Table 2. Each of the models uses a different rule to update the estimated mean
 211 yields of each crop i in season n , m_i^n , using the yields from each season $y_i^1 \dots y_i^n$. The Bayes 1
 212 model updates the estimate in a Bayesian fashion, assuming the individual knows the real long-
 213 run variance in yield, σ^2 . The Bayes 2 model relaxes this assumption, which means the sufficient
 214 statistic of the estimate of the mean is just the running average of the observed yields (Lehmann,
 215 1983). The final model, Memory Decay, is a generalization of the basic Bayesian model. Instead
 216 of the importance of recent information on the estimate being a function of the variance of the
 217 estimate and σ^2 , here it is a parameter γ to be fit from the data itself. When $\gamma = 0$, only the
 218 most recent information influences the estimate. As γ increases, past observed yields have a greater
 219 effect on an individual's estimate of the profitability of a crop.

Each model then uses the same functional form, a logit, to model how much each participant
 cares about differences in the estimated mean yields when choosing a crop to plant in season n .
 The probability of a participant planting crop i in season n is given by:

$$\Pr_n(i) = \frac{\exp(\beta m_i^n)}{\exp(\beta m_i^n) + \exp(\beta m_j^n)}.$$

220 The parameter β captures how much the difference between the estimated means influences choice.
 221 When $\beta = 0$, choice is random with respect to the estimates of the means. When $\beta = \infty$, the
 222 farmer always chooses the crop with the higher mean estimated yield.

223 The probability model above specifies a likelihood of observing each data point, and we fit each
 224 model to the data by finding the values of the parameters that maximize the joint likelihood of
 225 observing the data. It is possible to fit these models on an individual-by-individual basis, estimating

226 the strategy that best explains choice for each participant, or across individuals, assuming each
 227 individual is using the same strategy. Using all the data available for each individual, Memory
 228 Decay is the best fitting model for 32 of 36 individuals, with an average estimate for γ of 0.11
 229 (maximum 0.50, minimum 0, median 0.065). Bayes 1 is the best fitting model for 3 individuals,
 230 and Bayes 2 for only one individual.

There is too little data for each subject for estimates for each treatment to be reliable, but it is informative to lump together the individuals and fit the models within each treatment. The relative fits of each model may still indicate relative proportions of strategies within the participant pool. Table 3 shows the fits for the three models for the three different variances in yield and the two different fluctuation conditions. The parameter estimates in each case are shown below the Akaike Information Criteria (AIC), Δ value, and Akaike weight (w) of each model. AIC is twice the natural log of the likelihood of observing the data, given the model, plus twice the number of parameters in the model. Thus smaller AIC values indicate better fits. There is no threshold AIC value that is “good enough”. Fits must be judged relative to one another. The measure Δ is a goodness-of-fit measure analogous to the common R^2 for linear models. For a given model x with minus log-likelihood LL_x , $\Delta_x = 1 - LL_x/LL_{random}$, where LL_{random} is the fit of a model in which individuals simply guess at each decision (choose randomly). This measure varies from 0, when the fit of model x is the same of the random model, to 1, when the fit of model x is perfect. Δ therefore measures the absolute predictive power of a model, compared to a random choice model. Akaike weights (w), in contrast, measure relative fit among the set of considered models. These are computed from the AIC values. The Akaike weight w_i for a model i in a set of n models is:

$$w_i = \frac{\exp(-0.5(\text{AIC}_i - \text{AIC}_{min}))}{\sum_{j=1}^n \exp(-0.5(\text{AIC}_j - \text{AIC}_{min}))},$$

231 where AIC_{min} is the smallest AIC value in the set of models considered. Thus the best-fitting
232 model has the largest w value. One interpretation of Akaike weights is that each indicates the
233 probability that a given model is the correct one. See Burnham and Anderson (2002) for details
234 on these and other measures used to compare models.

235 The reasons for analyzing data in this way, rather than using common null-hypothesis tests,
236 has been covered many times (Cohen, 1994; Anderson *et al.*, 2000; Gigerenzer *et al.*, 2004, for
237 example). Model comparison allows an arbitrary number of competing hypotheses, each of which
238 competes on equal footing. Different specific non-linear quantitative predictions thus compete to
239 explain the observed data, rather than predictions from only a model we know a priori to be wrong.

240 In every case, Memory Decay is the best fitting model, and only when $\sigma = 0.5$ and there is no
241 fluctuation do Bayes 1 and Bayes 2 even approach Memory Decay's fit. The estimates of β show
242 that choice becomes more random with respect to observed payoffs as both variance in yields and
243 the probability of fluctuation increase. The estimates of γ are small in most cases, and below 0.25
244 in every case. Both increasing variance and the probability of fluctuation reduce the estimates of
245 γ . In some sense, this result is quasi-Bayesian: a Bayesian pays less attention to older data when
246 the long-run variance in the data is smaller. New data is informative when the data is not highly
247 variable. Similarly, when the variance in yields here is smaller, γ , the weight given to previous
248 estimates, is smaller. When the means may change each season, previous estimates may become
249 unreliable, and therefore new data about yield has a stronger influence on the estimate.

250 3.4 Discussion

251 Our purpose in the first experiment was to understand individual learning in this decision environ-
252 ment, so we can seriously model individual learning in the later experiments that also allow social

253 learning. The Memory Decay model, even accounting for its extra parameter, fits much better
254 than the two Bayesian models. We found that the degree to which individuals are influenced by
255 differences in yield trends downwards (as indicated by lower β estimates) when the variances in
256 yield are high or the environment is not stochastic. We also mapped out the difficulty of learning
257 the optimal crop, as a function of variance in yield. Given the large difference in rate of learning
258 between $\sigma = 0.5$ and $\sigma = 4$, we chose these two standard deviations as easy and difficult treat-
259 ments, respectively, for the next two experiments. Given the clear advantage of Memory Decay in
260 predicting choice in these data, it forms the basis of individual learning in the following analyses.

261 4 Analysis of strategies

262 In this section we use simulations to analyze the performance of three alternative social learning
263 strategies in the experimental environment introduced in Experiment 1. This analysis allows us to
264 make specific predictions about which social learning strategies we expect in each experiment to
265 follow, as well as how much we expect subjects to use them. The conditions of our analysis exactly
266 mimic those of the Experiments we present afterwards.

267 Unlike the environment in most of the models we discussed earlier in the paper, our experiments
268 allow only peer-to-peer cultural transmission. The simulations we present here allow us to see how
269 well the predictions about the effects of difficulty of learning and fluctuations in payoffs hold in our
270 modified case. They also allow us to make immediately relevant comparisons of the effectiveness
271 of different social learning strategies.

272 Many social learning strategies are available to people in natural environments. Our experiments
273 restrict people to strategies that rely upon the frequencies of different alternative behaviors. We
274 outline three different imitation strategies of this type.

275 **Linear Imitation.** When individuals choose a target individual at random and copy their ob-
 276 served behavior, we refer to this as *Linear Imitation*. The imitation is linear with respect to each
 277 behavior’s frequency in the population of potential targets. For example, if two alternative behav-
 278 iors are present with frequencies 0.6 and 0.4, then linear imitation has a chance 0.6 of copying the
 279 first and 0.4 of copying the second. Across iterations of social learning, linear imitation does not
 280 change the expected frequencies of behaviors in the population.

We model Linear Imitation in a nested model with the individual learning model fit in the
 previous section. Let L_i^n be the probability of choosing behavior i in round n from the Memory
 Decay model. Then the probability of choosing behavior i in round n when using Linear Imitation
 is:

$$\Pr_n(i) = (1 - \alpha)L_i^n + \alpha \frac{x_i^{n-1}}{N}, \quad (1)$$

281 where x_i^n is the number of observable target individuals who choose option i in round n and N
 282 is the total number of observable targets. The parameter α specifies the strength of reliance on
 283 imitation versus individual learning. When $\alpha = 0$, the model reduces to the pure Memory Decay
 284 model. When $\alpha = 1$, the model reduces to pure Linear Imitation.

285 **Confirmation.** Another way to use the behavior of a single target individual, without simply
 286 copying his behavior, is to practice *Confirmation*. By Confirmation, we mean keeping one’s previous
 287 behavior, when a randomly chosen target individual also previously chose the same behavior, and
 288 relying upon individual judgment otherwise.

We model Confirmation in a nested model, as we do with Linear Imitation. Assuming only
 two behavioral options, the probability of adopting behavior i in round n , given access to N target

individuals, x_i^{n-1} of whom practiced behavior i in round $n - 1$, is:

$$\begin{aligned} \Pr_n(i) &= (1 - \alpha)L_i^n \\ &+ \alpha \left(\frac{x_i^{n-1}}{N} \begin{cases} c^{n-1} = i, & 1 \\ c^{n-1} \neq i, & L_i^n \end{cases} \right. \\ &\left. + \frac{N - x_i^{n-1}}{N} \begin{cases} c^{n-1} = i, & L_i^n \\ c^{n-1} \neq i, & 0 \end{cases} \right), \end{aligned} \tag{2}$$

289 where c^{n-1} is the individual's behavior (choice) in round $n - 1$.

290 **Conformity.** When at least three target individuals are observable, one can do better by using
 291 information from each of them. We define *Conformity* as adopting the majority behavior among
 292 a group of targets. When there is no clear majority among the targets, we assume individuals fall
 293 back on individual judgment.

In our two-alternative choice environment, the probability of choosing behavior i in round n is:

$$\begin{aligned} \Pr_n(i) &= (1 - \alpha)L_i^n \\ &+ \alpha \begin{cases} x_i^{n-1} > N/2, & 1 \\ x_i^{n-1} < N/2, & 0 \\ x_i^{n-1} = N/2, & L_i^n \end{cases} . \end{aligned} \tag{3}$$

294 4.1 Comparison of strategies

295 [Figure 2 about here.]

296 Which of these strategies is best in our experimental setting, and what is the optimal amount

297 of reliance on each? The theory we mentioned in section 2 (Henrich & Boyd, 1998, especially)
298 suggests that Conformity is broadly adaptive and likely to perform better than either alternative
299 we have nominated. However, Confirmation has not yet been analyzed in the thorough way that
300 Conformity has. Section 2 also suggests that reliance on any social learning strategy should increase
301 with increasing difficulty of the task and decrease with increasing fluctuation in payoffs. To compare
302 the three strategies above, therefore, we conducted simulations to compute the expected payoffs to
303 each strategy, under different values of the variance in yields and the probability of fluctuation in
304 the means, as well as across the range of reliance on each (varying α from 0 to 1). The simulations
305 use the exact experimental design described in Experiment 1, except the virtual participants are
306 in groups of five and can freely observe the previous choices, but not payoffs, of each other group
307 member, each planting round. We conducted 100,000 simulations at each parameter combination,
308 where each simulation modeled decisions in 20 rounds of planting. Performance was measured by the
309 mean payoff over all 100,000 simulations. The values of the individual learning parameters β and γ
310 used in the simulation were taken from the maximum-likelihood estimates from the Memory Decay
311 model in Experiment 1, which are the best guess as to how participants are learning individually.
312 Of course, individual learning may change when social information is introduced. However, as
313 we will demonstrate later, the estimates of these parameters change very little in our subsequent
314 experiments.

315 We found no situation in which Conformity, at its optimal value of α , does not out-perform
316 both Linear Imitation and Confirmation. Figure 2 shows the expected payoffs to Linear Imita-
317 tion, Confirmation, and Conformity, as functions of the reliance on social learning (the value of
318 the parameter α in the models above). Linear Imitation is never useful, in this environment. The
319 expected payoff to Linear Imitation is always highest when the reliance on social learning is zero.

320 Confirmation and Conformity both lead to gains over both pure individual learning (when $\alpha = 0$)
321 and Linear Imitation, but Conformity out-performs both other strategies, provided individual use
322 the optimal value of α . These results are typical of other experimental settings. Conformity lever-
323 ages the extra information available from multiple target individuals, while neither other strategy
324 does so. Increasing the size of social groups would increase the advantage Conformity holds.

325 However, when information from only one target is available, Confirmation is better than Linear
326 Imitation. Simulations with two-person groups, in which only one other individual is observable,
327 confirm that Confirmation's effectiveness generalizes to these smallest possible social groups. Con-
328 formity is not possible in these groups, however.

329 4.2 Optimal amounts of social learning

330 Confirmation and Conformity both lead to gains in payoff, but the optimal reliance on social
331 learning, measured in the parameter α will vary as a function of the experimental variables. We
332 demonstrate here how variance in yields and fluctuation in the means leads to the predicted effects
333 we summarized in section 2.

334 Figures 3 and 4 plot the relative performance of Confirmation and Conformity under different
335 experimental conditions. In both cases, increases in the variance in crop yields makes an increased
336 reliance on social learning optimal, while an increase in the probability of fluctuation in the means
337 of the yields makes a decrease in reliance on social learning optimal. These computations verify
338 the relevance of the general predictions from the theory reviewed in section 2. However, in some
339 cases, this difference is quite small. The results of our next two experiments allow us to address to
340 qualitative nature of these predictions, as well as measure how calibrated participants are to the
341 decision environment. We do not expect participants to select their strategies optimally, but we do

342 expect detectable shifts in the direction of optimal strategy.

343 [Figure 3 about here.]

344 [Figure 4 about here.]

345 4.3 Predictions for Experiments 2 and 3

346 Our next two experiments allow access to different amounts of social information. Using the analysis
347 above, we outline a set of predictions for how our estimates of participant strategy will respond
348 to changes in experiment parameters. In Experiment 2, we modify Experiment 1 to allow access
349 to the behavior of a single peer. In Experiment 3, participants have access to the behavior of all
350 group members.

351 **Choice of social learning strategy:** We expect participants to rely on Conformity when possi-
352 ble, but to rely on Confirmation in Experiment 2, where information from only one peer is available.
353 Linear Imitation is not useful in either experiment, and so we predict participants will not use it,
354 provided the cues provided in the experiment lead them to select appropriate strategies.

355 **Response to variance:** We expect participants to rely on social learning more (as indicated by
356 increased estimates of α) when variance in yield increases, regardless of which strategy they select.

357 **Response to fluctuation:** We expect participants to rely on social learning less when fluctuation
358 in the mean yields increases, regardless of which strategy they select.

359 **5 Experiment 2: One social target**

360 In the second experiment, we added simple one-model social learning in order to estimate partic-
361 ipants tendencies to access this information and how they use it. Recall that our reading of the
362 theory suggests that linear social learning in this purely peer-to-peer laboratory culture is of little
363 value. Instead we expect to see Confirmation and an increase in reliance upon it when the variance
364 in yield increase and a decrease in reliance upon it when fluctuation in the means is possible.

365 **5.1 Participants**

366 55 undergraduates from a UC Davis psychology subject pool participated in this experiment. None
367 of them had participated in the previous experiment. They participated in sessions of size 6-10,
368 which were divided into anonymous groups of 4-6 individuals, depending only upon the contin-
369 gencies of daily attendance. There were 12 groups total: 8 of size 4, 3 of size 5, and 1 of size 6.
370 Participants always knew the actual size of their group, but they never knew the identities of the
371 other people in their group. Participants received course extra credit, in addition to their monetary
372 payments.

373 **5.2 Design**

374 This experiment builds upon experiment 1 by providing one additional piece of information each
375 season, prior to planting. Again, the experiment was programmed using z-Tree (Fischbacher, 2002).
376 The software assigned participants to groups at random and passed decisions among the clients
377 in response to participant behavior. After the first season at each farm, each participant had
378 the option of clicking a button to view the most recent planting decision (but not yield) of one
379 randomly chosen, anonymous member of their own group. Participants were told that members of

380 the same group always experienced the same environment: the means and variances of the crops
381 were the same for all members of a group, at all times, even when the means occasionally switched
382 in the last three farms. Environmental fluctuations occurred simultaneously within groups, and
383 the participants knew this as well.

384 **5.3 Results**

385 [Figure 5 about here.]

386 We collected information on the rates at which participants accessed the decisions of other
387 members of their groups (“social information”). We use this data, together with individual planting
388 decisions, to model social learning strategy.

389 There is impressive variation among participants in the rate they access social information.
390 Figure 5(a) plots the distribution of individual click frequencies. 20 participants never or very rarely
391 accessed social information. The remainder are spread over the entire range of click frequencies.
392 These data alone suggest considerable variation in participant strategy. Figure 5(b) plots the
393 frequency of clicks for social information averaged across participants but by season (experimental
394 round). The frequency peaks at 0.5 in the second season, the first season social information is
395 available, and declines to just above 0.2 by the final season.

396 Table 4 shows the probabilities of participants’ clicking to access social information each round
397 of the experiment. Access to social information increases with increasing variance and decreases
398 with increasing probability of fluctuation in the means. The effect of increasing variance is large
399 in the absence of fluctuation—the confidence intervals in this case do not overlap. The effect of
400 fluctuation is large in the case of high variance—again the confidence intervals do not overlap.

401 [Table 4 about here.]

[Table 5 about here.]

402

403 While the descriptive presentation of the frequencies of access to social information give hints
404 that social information is of interest to individuals and that participants vary in their interest,
405 they do not tell us much about how participants might be using the information they acquire from
406 other group members. To address this question, we fit the 6360 decisions from this experiment to
407 three candidate models. The first model is the pure individual learning model Memory Decay from
408 the previous experiment. The second model is the Linear Imitation model we presented in section
409 4 (Equation 1), which models social learning by introducing one new parameter to the Memory
410 Decay model.

411 The third model we fit the data from Experiment 2 is Confirmation (Equation 2), also introduced
412 in section 4, which models participants using social information in a way distinct from copying. For
413 this strategy, we assume the individual checks another participant in order to see that someone else
414 is doing the same thing as themselves. If the other individual planted the same crop last round as
415 the focal individual, then the focal individual keeps their previous behavior. Otherwise, she relies
416 upon individual learning.

417 Since participants could see the behavior of only one other individual, Conformity as modeled
418 in Equation 3 is not possible in this experiment.

419 Table 5 summarizes the fit and parameter estimates of each of three models, across individuals.
420 In the easy standard deviation (0.5), the Linear Imitation model fits the best ($w = .78$) when
421 there is no fluctuation is possible, and the Confirmation model is superior when there is fluctuation
422 ($w = .99$). In the hard standard deviation (4), there is much less dominance of social learning. Each
423 of three models earns good support, although Confirmation consistently does somewhat better than
424 Linear Imitation.

425 5.4 Discussion

426 In this experiment, we added only the option of seeing the most recent planting decision of a single
427 anonymous member of one’s own group, who planted under the same conditions. We find that
428 participants choose to view social information slightly more often when the variance in yields is
429 high and when there is no fluctuation in the means through time. We also found substantial evidence
430 of social learning from the model fitting exercise. However, while the social models fit considerably
431 better than the individual learning model for the low-variance farms, this should not be interpreted
432 to mean that all participants were using social information. The social models only differ from the
433 individual learning model when an individual in fact viewed social information. Some individuals
434 rarely did so. By the end of an experimental farm, only about 20 percent of participants choose to
435 view social information. Thus the better fit of the social model applies only in those cases, which
436 are overall the minority. When participants did view social information, the evidence indicates it
437 had a detectable effect on their choices, as indicated by the model fits. Many individuals never or
438 almost never viewed social information, and so these individuals must be described as individual
439 learners, despite the better fit of Linear Imitation or Confirmation.

440 In the general discussion we interpret the rates of access to social information together with the
441 model fits to evaluate the results of the experiments with respect to the expectations we developed
442 at the beginning of the paper.

443 6 Experiment 3: Conformity

444 While experiment 2 addresses the most simple kind of social learning possible in our experimental
445 design, rarely do people find themselves in a situation in which they can observe the behavior

446 of only one peer at a time. Experiment 3 was designed to address how participants use social
447 information from more than one individual. Our reading of the formal literature suggests that
448 the use of a majority rule, adopting the most common behavior among models, is more valuable
449 here than simply imitating in the linear or confirmation fashion that was possible in the previous
450 experiment.

451 **6.1 Participants**

452 49 undergraduates from a UC Davis, recruited from classrooms, participated in this experiment.
453 None of them had participated in the previous experiments. They participated in sessions of
454 size 6-10, which were divided into anonymous groups of 4-7 individuals, depending only upon the
455 contingencies of daily attendance. There were 9 groups total: 2 of size 4, 3 of size 5, 2 of size 6,
456 and 2 of size 7. Participants always knew the actual size of their group.

457 **6.2 Design**

458 This experiment builds upon experiment 2 by allowing participants to click a button each season
459 after the first, in order to view the most recent planting decisions of all other group members. As
460 in experiment 2, participants viewing social information could not identify individuals by name
461 or number or any other identifying information, nor could they view the payoffs these individuals
462 received. They simply saw a randomized vector of crop planting choices (*wheat, wheat, potatoes,*
463 *wheat*).

464 Again, participants were told that members of the same group always experienced the same
465 environment: the means and variances of the crops were the same for all members of a group,
466 at all times, even when the means occasionally switched in the last three farms. Environmental

467 fluctuations occurred simultaneously within groups, and the participants knew this.

468 **6.3 Results**

469 [Table 6 about here.]

470 [Figure 6 about here.]

471 [Table 7 about here.]

472 As in experiment 2, we collected information about how often participants chose to view the
473 decisions of their peers. Overall, the pattern of clicks is similar to that in experiment 2: the fre-
474 quency is highest at the beginning of each farm and stabilizes above zero before season 20. Table 6
475 summarizes the frequencies of clicks for social information. There is more access to social informa-
476 tion in the high variance farms, especially when fluctuation in the means is absent. There is less
477 access when fluctuation in the means is possible. Figure 6 demonstrates that there is considerable
478 variation in participant strategy, as in experiment 2. Some individuals never or only very rarely
479 click to access social information. Other access social information nearly every experimental round.
480 Overall, the frequencies of access are higher than in experiment 2.

481 While it makes sense to hypothesize that individuals might click more often in larger groups,
482 which contain more information, there is no discernible relationship between group size and fre-
483 quency of clicks: $N = 4$, 0.29; $N = 5$, 0.45; $N = 6$, 0.36; $N = 7$, 0.25.

484 As with the previous experiments, we analyzed the data from experiment 3 to determine the
485 ability of different choice models to predict participant planting decisions. The first model we
486 applied to these data is the Memory Decay model from experiment 1. This is the baseline individual
487 learning model. The second model we fit to the data is the Linear Imitation model presented in

488 section 4 and analyzed already in Experiment 2 (Equation 1). The third model we fit to these data
489 is Confirmation (Equation 2), and the fourth Conformity, introduced in our analysis in setion 3
490 (Equation 3).

491 We fit these three models to the 5880 decisions made in experiment 3. Table 7 summarizes
492 the overall model fits. As in previous tables of this kind, we show the overall fit using the Akaike
493 Information Criteria (AIC), Akaike weights (w), and Δ values. Each model fit is followed by its
494 maximum likelihood parameter estimates. Overall, Linear Imitation and Conformity fit the data
495 much better than either the pure individual learning model or Confirmation. Confirmation does
496 a comparatively poor job everywhere, even compared to pure individual learning. In the absence
497 of environmental fluctuation, Linear Imitation better predicts choice, compared to Conformity,
498 whether variance in yields is high or low. When environmental fluctuation is present, Conformity
499 better predicts choice, when the variance in yields is low ($\sigma = 0.5$). When the variance is high
500 ($\sigma = 4$), Linear Imitation and Conformity are essentially tied in fit to the data.

501 Looking at the parameter estimates of α , the reliance on social learning relative to individual
502 learning, the proportions of social learning are overall much higher in this experiment than in
503 experiment 2. Moving from small variance in yield to large, the estimates of reliance on social
504 learning are relatively much smaller.

505 Environmental fluctuation—the probability of change in the mean yield of each crop—seems
506 to favor Conformity. In the low-variance farms, Conformity is a much better predictor of choice
507 than Linear Imitation, provided the probability of fluctuation is above zero. In the high-variance
508 farms, Conformity ties with Linear Imitation when fluctuation is possible but is far inferior to
509 it when fluctuation is not possible. However, looking at the fits to the individual-by-individual
510 data, when $\sigma = 4$ and the probability of fluctuation is 0.05, Conformity fits the data better than

511 Linear Imitation. Of 49 participants, the choices of 38 are best predicted by Conformity, while the
512 choices of 11 are best predicted by Linear Imitation. The evidence suggests participants are likely
513 to use some strategy approximating Conformity, provided there is the possibility of fluctuation in
514 the means. Otherwise, there is little evidence of the use of a strategy that integrates the social
515 information in a positive frequency-dependent way.

516 **6.4 Discussion**

517 This experiment allowed participants to see behavior from all other members of their group, and
518 we expected this to lead to conformist crop planting decisions. This expectation was partly upheld,
519 however not in the absence of environmental fluctuation. When there is no chance of fluctuation
520 in the means of the crops, participants appear to learn socially, but the majority of them are not
521 conformist. We did find, in agreement with the analyses in section 4, that participants are likely
522 not using a Confirmation strategy.

523 **7 General Discussion**

524 The most obvious and least surprising result of our experiments is that many participants used the
525 choices of their peers in making their own choices. It is more surprising that we found evidence of
526 simple Linear Imitation in experiment 2, where it is little use. Of course, imitation in the one-peer
527 experiment is not much worse than learning on one's own; the payoff difference between imitating
528 a random peer and learning on one's own in this case is not very large. Nevertheless, the sizable
529 proportion (although not the majority) of participants who seem to have used simple imitation
530 deserves an explanation. We imagine two reasons individuals may imitate when there is no struc-
531 tural feature of the environment or strategy that makes it profitable. First, some individuals learn

532 better than others. Because some individuals are more likely to arrive at optimal behavior faster,
533 for those who imagine themselves slower than average, even Linear Imitation can be profitable.
534 Second, people may be carrying over strategies which are broadly useful in their daily lives into
535 the experiment. It is unreasonable to expect that participants approach experiments as naive yet
536 rational agents. Patterns of imitation behavior in normal life may encourage people to imitate in
537 these experiments, even when there is no apparent advantage to such a strategy.

538 It is also unclear why participants use Linear Imitation, instead of Conformity, in the no-
539 fluctuation treatments of Experiment 3. On those farms, individuals would have done better had
540 they used Conformity, yet we found little evidence of Conformity there, even though we found con-
541 siderable evidence of it when the environment could fluctuate. Participants are clearly responding
542 to the experimental treatments, and further work will be needed to understand how to cues the
543 experiment provides activate existing strategies designed for learning in natural settings.

544 [Table 8 about here.]

545 The predictions from theory suggested that individuals would rely more on social learning when
546 (1) the variance in yields was larger and (2) there was no possibility of environmental fluctuation.
547 The clicks to access social information agree with these predictions. Only in experiment 2, when
548 the standard deviation of yields was small, is there little noticeable effect on the frequency of clicks
549 to access social information (although the measured effect is in the right direction even then). In
550 all other cases, rates of clicks increase in treatments with higher standard deviations and decrease
551 in those with fluctuation in the means.

However, estimates of reliance on social learning do not generally agree with the predictions.
In order to make the process data (clicks) and model estimates comparable, we need to multiply

frequencies of access to social information by the estimated reliances on social learning. We compute the estimated reliance on social learning in each treatment by computing the model-averaged estimate of α in each case, using Akaike weights (w) for weighting the different estimates from different models. The model-averaged estimate of reliance on social learning for a set of n models is:

$$\bar{\alpha} = \sum_i^n w_i \alpha_i^*,$$

552 where α_i^* is the maximum-likelihood estimate of α for model i . For Memory Decay, we set α^* to zero
553 (no reliance on social learning). We compute the total rate of social learning then by multiplying
554 each rate of information access by the model-weighted reliance on that information.

555 Table 8 summarizes the total estimated frequencies computed in this way. In experiment 3,
556 frequencies of social learning decrease, as predicted, when we introduce fluctuation in the mean
557 yields (probability of fluctuation 0.05). However, in experiment 2, fluctuation appears to have had
558 the opposite effect: social learning increases in that case. The effect of increasing the variance
559 in yields is contrary to prediction in every case. In both experiments 2 and 3, total estimated
560 frequencies of social learning decrease when the variance in yield increases.

561 One possible explanation of this counter-theoretical result is that participants are interpreting
562 large variance in the mean yields as environmental unpredictability of the sort introduced by fluctuation.
563 We cannot address this possibility with our data, but new experiments using a different
564 mechanism for manipulating the difficulty of individual learning would help to deal with it. Another
565 other possibility is that we simply have the wrong models. All models are simplifications. If the
566 models of social learning we have considered are missing some structurally important feature of

567 individuals' imitation behavior, then our model estimates will simply tell the wrong story. Notice
568 that the process data, the clicks to access social information, agree with the theory. These process
569 data have the virtue of not being constructed through an intervening model: they are plainly mea-
570 sured. Skeptics of our model estimates may therefore take comfort in the less uncertain process
571 data. Either way, whether because of the details of the difficulty manipulation or the structural
572 inaccuracy of our models, there is a problem to be solved, in order to reconcile theory with our
573 experimental results.

574 We found good evidence of individual variation in strategy in all three experiments. Yet we
575 have made little effort yet to explain this variation. Variation in individual learning ability/skill
576 may explain some of the estimated variation in strategy. It is tempting also to hypothesize about
577 covariance with other individual characteristics. We hope to address the data in this way in later
578 work.

579 **8 Conclusion**

580 By way of conclusion, we offer several cautions. First, these experiments obviously explore only a
581 tiny fraction of the universe of meaningful learning environments and potential strategies available
582 to people. Progress in understanding the design of social learning will come from a body of detailed
583 work fully exploring a number of decision environments and transmission schemes (information
584 structures) while iteratively revising the quantitative models that motivate them. We find work
585 by Tatsuya Kameda and his colleagues (Kameda & Nakanishi, 2003; Kameda & Nakanishi, 2002)
586 inspiring in this regard. It is not enough to simply nominate the existence of a collection of "effects"
587 and test for their existence. Mature predictive models of some depth come about by iteratively
588 building complexity into a research design and the models it is meant to address.

589 Second, the depth of this kind of work needs to be balanced by breadth. Replication, of both
590 parameter estimates and general results, both cross-culturally and across cultural domains, is es-
591 sential. We do not imagine social learning strategies, which themselves can be learned, are invariant
592 human universals. The strength of conformity, in particular, likely varies cross-culturally and situ-
593 ationally. Students in Western societies are repeatedly admonished to “think for themselves.” It is
594 also important to notice that students, the favorite subjects of psychologists and economists alike,
595 are an odd population to study in order to understand how people learn. Students in university
596 are trained to learn in particular ways that are unlikely to be representative of most adults. Con-
597 structing theories of human nature based on student data is always hazardous, but particularly so
598 in this case.

599 Even when considering members of a single study population, parameter and strategy estimates
600 from any one sample are notoriously prone to overfitting. Using the estimates we have developed
601 here to predict the choices of new subjects would go a long way to estimating the narrow-sense
602 robustness of our results.

603 Ultimately, results from laboratory studies like this one need to be validated in naturalistic or
604 quasi-naturalistic settings. Accurate models of individual-level processes can be scaled up to predict
605 large-scale dynamics, much as evolutionary biologists use micro-evolutionary models of events in
606 the lives of organisms to understand long-term macro-evolutionary trends. Studies like Prentice
607 and Miller (1993), Nisbett and Cohen (1996), Edgerton (1971), and Henrich et al. (2004) remind us
608 of the phenomena we ultimately intend to understand and provide significant constraints on theory
609 development. More direct applications of micro-level theory to macro-level problems, like Henrich’s
610 (2001) application of social learning models to data on the spread of technological innovations,
611 demonstrate the relevance of experimental studies to the cultural transformations we witness in

612 daily life. We think researchers should not be shy about extrapolating findings in both directions,
613 from the laboratory to the field and visa versa.

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619 **References**

- 620 Anderson, C. M. 2001. *Behavioral Models of Strategies in Multi-Armed Bandit Problems*. Ph.D.
621 thesis, California Institute of Technology.
- 622 Anderson, D. R., Burnham, K. P., & Thompson, W. L. 2000. Null hypothesis testing: Problems,
623 prevalence, and an alternative. *Journal of Wildlife Management*, **64**, 912–923.
- 624 Anderson, L. R., & Holt, C. A. 1997. Information Cascades in the Laboratory. *American Economic*
625 *Review*, **87**(2), 847–862.
- 626 Apesteguia, Jose, Huck, Steffen, & Oechssler, Jörg. 2003. *Imitation, Theory and Experimental*
627 *Evidence*.
- 628 Bandura, Albert. 1977. *Social learning theory*. New York: Prentice Hall. Book English.
- 629 Banerjee, Abhijit V. 1992. A Simple Model of Herd Behavior. *The Quarterly Journal of Economics*,
630 **107**(3), 797–817.

- 631 Banks, J., Olson, M., & Porter, D. 1997. An Experimental Analysis of the Bandit Problem.
632 *Economic Theory*, **10**, 55–77.
- 633 Baron, R. S., Vandello, J. A., & Brunsman, B. 1996. The forgotten variable in conformity research:
634 Impact of task importance on social influence. *Journal of Personality and Social Psychology*,
635 **71**(5), 915–927.
- 636 Baum, W. M., Richerson, P. J., Efferson, C. M., & Paciotti, B. M. 2004. Cultural evolution in
637 laboratory microsocieties including traditions of rule giving and rule following. *Evolution and*
638 *Human Behavior*, **25**, 305–326.
- 639 Bikhchandani, Suhsil, Hirschleifer, David, & Welch, Ivo. 1992. A Theory of Fads, Fashion, Custom,
640 and Cultural Change as Informational Cascades. *The Journal of Political Economy*, **100**(5),
641 992–1026.
- 642 Boyd, R., & Richerson, P. J. 1985. *Culture and the evolutionary process*. Chicago: University of
643 Chicago Press.
- 644 Burnham, Kenneth P., & Anderson, David. 2002. *Model Selection and Multi-Model Inference*.
645 Springer-Verlag Telos.
- 646 Camerer, C. F. 2003. *Behavioral Game Theory: Experiments in strategic interaction*. Princeton:
647 Princeton University Press.
- 648 Camerer, C. F., & Ho, T. 1999. Experience-Weighted Attraction Learning in Normal Form Games.
649 *Econometrica*, **67**(4), 827–874.
- 650 Cohen, J. 1994. The earth is round ($p < .05$). *American Psychologist*, **49**, 997–1000.

- 651 Coultas, Julie C. 2004. When in Rome... An Evolutionary Perspective on Conformity. *Group*
652 *Processes & Intergroup Relations*, **70**(4), 317–331.
- 653 Edgerton, R. B. 1971. *The individual in cultural adaptation: A study of four East African peoples*.
654 Berkeley: University of California Press.
- 655 Fasolo, Barbara. 2002. *Multi-Attribute Decisions Online: How People Handle Conflict Among*
656 *Attributes*. Ph.D. thesis, University of Colorado at Boulder.
- 657 Fischbacher, Urs. 2002. *z-Tree: Experimenter's Manual*. Institute for Empirical Research in Eco-
658 nomics.
- 659 Galef, B. G. Jr., & Whiskin, E. E. 2004. Effects of environmental stability and demonstrator age
660 on social learning of food preferences by young Norway rats. *Animal Behaviour*, **68**, 897–902.
- 661 Gans, Noah, Croson, Rachel, & Knox, George. 2003. *Customer Learning and Switching: A Two-*
662 *Armed Bandit Experiment*.
- 663 Gigerenzer, G., Todd, P. M., & the ABC Group. 1999. *Simple heuristics that make us smart*. New
664 York: Oxford University Press.
- 665 Gigerenzer, Gerd, Krauss, Stefan, & Vitouch, Oliver. 2004. The Null Ritual: What you always
666 wanted to know about significance testing but were afraid to ask. In: Kaplan, D. (ed), *Sage*
667 *Handbook of Quantitative Methods for the Social Sciences*. Sage.
- 668 Gittins, J. C. 1989. *Multi-Armed Bandit Allocation Indices*. John Wiley & Sons, Inc.
- 669 Henrich, J., & Boyd, R. 1998. The Evolution of Conformist Transmission and the Emergence of
670 Between-Group Differences. *Evolution and Human Behavior*, **19**, 215–242.

- 671 Henrich, J., Boyd, R., Bowles, S., Camerer, C.F., Fehr, E., Gintis, H., & McElreath, R. 2004.
672 Overview and synthesis. *Pages 8–54 of:* Henrich, J., Boyd, R., Bowles, S., Camerer, C., Fehr,
673 E., & Gintis, H. (eds), *Foundations of Human Sociality: Ethnography and Experiments in 15*
674 *Small-scale Societies*. Oxford: Oxford University Press.
- 675 Henrich, Joseph. 2001. Cultural transmission and the diffusion of innovations: Adoption dynam-
676 ics indicate that biased cultural transmission is the predominate force in behavioral change.
677 *American Anthropologist*, **103**(4), 992–1013.
- 678 Henrich, Joseph, & McElreath, Richard. 2003. The evolution of cultural evolution. *Evolutionary*
679 *Anthropology*, **12**, 123–135.
- 680 Horowitz, A. D. 1973. *Experimental Study of the Two-Armed Bandit Problem*. Ph.D. thesis,
681 University of North Carolina, Chapel Hill.
- 682 Kameda, T., & Nakanishi, D. 2002. Cost-benefit analysis of social/cultural learning in a non-
683 stationary uncertain environment: An evolutionary simulation and an experiment with human
684 subjects. *Evolution and Human Behavior*, **23**(5), 373–393.
- 685 Kameda, T., & Nakanishi, D. 2003. Does social/cultural learning increase human adaptability?
686 Rogers’ question revisited. *Evolution and Human Behavior*, **24**, 242–260.
- 687 Lehmann, E. L. 1983. *Theory of Point Estimation*. New York: John Wiley & Sons, Inc.
- 688 McElreath, Richard. 2004. Social learning and the maintenance of cultural variation: An evolu-
689 tionary model and data from East Africa. *American Anthropologist*, **106**(2), 308–321.
- 690 Meyer, R. J., & Shi, Y. 1995. Sequential Choice Under Ambiguity: Intuitive Solutions to the
691 Armed-Bandit Problem. *Management Science*, **415**, 817–834.

- 692 Nisbett, Richard E., & Cohen, Dov. 1996. *Culture of Honor: The Psychology of Violence in the*
693 *South*. Westview Press.
- 694 Pingle, M. 1995. Imitation vs. rationality: An experimental perspective on decision-making. *Journal*
695 *of Socio-Economics*, **24**, 281–315.
- 696 Prentice, Deborah A., & Miller, Dale T. 1993. Pluralistic ignorance and alcohol use on campus:
697 Some consequences of misperceiving the social norm. *Journal of Personality & Social Psychol-*
698 *ogy*, **64**(2), 243–256. Article English.
- 699 Richerson, P. J., & Boyd, R. 2004. *Not by genes alone: How culture transformed human biology*.
700 Chicago: University of Chicago Press.
- 701 Rogers, Alan R. 1988. Does biology constrain culture? *American Anthropologist*, **90**(4), 819–831.
- 702 Schotter, Andrew, & Sopher, Barry. 2003. Social Learning and Coordination Conventions in Inter-
703 generational Games: An Experimental Study. *Journal of Political Economy*, **111**(3), 498–529.

704 **List of Tables**

705 1 Estimates of probability of optimal crop choice, categorized by variance in yield
706 and the probability of environmental fluctuation. 95% confidence intervals shown
707 in parentheses. Optimal planting choices decrease as variance increases and with
708 fluctuation in the means. 36

709 2 Models fit to individual learning data and their updating rules for computing the
710 estimate of the mean yield of crop i in season n . The first model, Bayes 1, updates
711 an estimate of the mean m_i and the variance in this estimate, v_i . The second model,
712 Bayes 2, uses the number of samples from crop i , N_i , to compute the running mean.
713 The third model is a parameterized generalization of Bayes 1. In each case, these
714 rules apply when crop i is chosen in round $n - 1$. When another crop was chosen in
715 the previous round, each rule is $m_i^n = m_i^{n-1}$ 37

716 3 Akaike Information Criteria (AIC), fit relative to a random model (Δ), Akaike
717 weights (w), and parameter estimates for the three individual learning models, by
718 experimental farm standard deviation and probability of fluctuation of means. The
719 measure Δ for a model x is defined as $\Delta_x = 1 - LL_x/LL_{\text{Random}}$. It gives the relative
720 improvement in fit of the model x , compared to the accuracy of a random model.
721 The weights (w) give the relative fit (1 best) of each model, adjusted for number of
722 parameters, to the other models in the analysis. 38

723 4 Probabilities of accessing social information each experimental round, by standard
724 deviation and probability of environmental fluctuation. 95% confidence intervals
725 shown in parentheses. Access to social information increases with increasing variance
726 and decreases with increasing probability of fluctuation in the means. 39

727 5 Akaike Information Criteria (AIC), fit relative to a random model (Δ), Akaike
728 weights (w), and parameter estimates for the three candidate models fit to the data
729 from experiment 2. 40

730 6 Frequencies of clicks to view social information in experiment 3. Ranges in paren-
731 theses are 95% confidence intervals. There is more access to social information in
732 the high variance farms, especially when fluctuation in the means is absent. There
733 is less access when fluctuation in the means is possible. 41

734 7 Akaike Information Criteria (AIC), fit relative to a random model (Δ), Akaike
735 weights (w), and parameter estimates for the three candidate models fit to the data
736 from experiment 3. 42

737 8 Total estimated frequencies of social learning in each experiment and treatment.
738 These frequencies come from multiplying the frequency of clicks to view social infor-
739 mation by the Akaike-weighted estimated influence of social learning in each treatment. 43

Table 1: Estimates of probability of optimal crop choice, categorized by variance in yield and the probability of environmental fluctuation. 95% confidence intervals shown in parentheses. Optimal planting choices decrease as variance increases and with fluctuation in the means.

Fluctuation	Variance		
	0.25	4	16
0	0.79 (0.75-0.82)	0.76 (0.72-0.79)	0.68 (0.64-0.72)
0.05	0.69 (0.66-0.73)	0.68 (0.64-0.71)	0.61 (0.56-0.65)

Table 2: Models fit to individual learning data and their updating rules for computing the estimate of the mean yield of crop i in season n . The first model, Bayes 1, updates an estimate of the mean m_i and the variance in this estimate, v_i . The second model, Bayes 2, uses the number of samples from crop i , N_i , to compute the running mean. The third model is a parameterized generalization of Bayes 1. In each case, these rules apply when crop i is chosen in round $n - 1$. When another crop was chosen in the previous round, each rule is $m_i^n = m_i^{n-1}$.

Model	Updating rule	Free parameters
Bayes 1	$m_i^n = am_i^{n-1} + (1 - a)y_i^{n-1}$ $v_i^n = av_i^{n-1}$ $a = \frac{\sigma^2}{\sigma^2 + v_i^{n-1}}$	β (see main text)
Bayes 2	$m_i^n = \frac{N_i^{n-1}m_i^{n-1} + y_i^{n-1}}{N_i^n}$	β
Memory Decay	$m_i^n = \gamma m_i^{n-1} + (1 - \gamma)y_i^{n-1}$	β, γ

Table 3: Akaike Information Criteria (AIC), fit relative to a random model (Δ), Akaike weights (w), and parameter estimates for the three individual learning models, by experimental farm standard deviation and probability of fluctuation of means. The measure Δ for a model x is defined as $\Delta_x = 1 - LL_x/LL_{\text{Random}}$. It gives the relative improvement in fit of the model x , compared to the accuracy of a random model. The weights (w) give the relative fit (1 best) of each model, adjusted for number of parameters, to the other models in the analysis.

Standard Deviation Fluctuation	0.5		2		4	
	0	0.05	0	0.05	0	0.05
Bayes 1: AIC	710.84	779.33	781.79	862.04	874.61	932.83
Δ	0.29	0.22	0.22	0.14	0.13	0.07
w	0.14	9.64E-31	3.72E-10	1.56E-40	4.02E-12	2.88E-36
β	0.53	0.63	0.40	0.49	0.28	0.21
Bayes 2: AIC	710.93	779.37	783.66	863.60	879.52	933.72
Δ	0.29	0.22	0.22	0.14	0.12	0.07
w	0.14	9.43E-31	1.46E-10	7.14E-41	3.44E-13	1.85E-36
β	0.53	0.63	0.39	0.47	0.22	0.17
Memory Decay: AIC	707.59	641.10	738.37	678.72	822.12	769.16
Δ	0.30	0.36	0.26	0.32	0.18	0.23
w	0.72	1.00E+00	1.00E+00	1.00E+00	1.00E+00	1.00E+00
β	0.53	0.71	0.42	0.54	0.21	0.25
γ	0.05	0.01	0.17	0.05	0.21	0.13

Table 4: Probabilities of accessing social information each experimental round, by standard deviation and probability of environmental fluctuation. 95% confidence intervals shown in parentheses. Access to social information increases with increasing variance and decreases with increasing probability of fluctuation in the means.

Fluctuation	Standard Deviation	
	0.5	4
0	0.2448 (0.22–0.27)	0.3213 (0.30–0.34)
0.05	0.2329 (0.21–0.25)	0.2591 (0.24–0.28)

Table 5: Akaike Information Criteria (AIC), fit relative to a random model (Δ), Akaike weights (w), and parameter estimates for the three candidate models fit to the data from experiment 2.

Standard Deviation Fluctuation	0.5		4	
	0	0.05	0	0.05
Memory Decay: AIC	1302.06	1353.61	1852.76	1490.37
Δ	0.39	0.41	0.19	0.30
w	0.06	1.97E-05	0.35	0.27
β	0.66	0.79	0.23	0.32
γ	0.01	0.00	0.19	0.00
Simple Social: AIC	1296.88	1345.37	1853.30	1491.17
Δ	0.39	0.41	0.19	0.30
w	0.78	1.21E-03	0.27	0.18
β	0.66	0.79	0.24	0.33
γ	0.02	0.00	0.19	0.00
α	0.14	0.18	0.04	0.05
Confirmation: AIC	1301.02	1331.94	1852.80	1489.04
Δ	0.39	0.42	0.19	0.30
w	0.10	9.99E-01	0.34	0.52
β	0.65	0.76	0.23	0.32
γ	0.00	0.00	0.20	0.00
α	0.17	0.42	0.09	0.17

Table 6: Frequencies of clicks to view social information in experiment 3. Ranges in parentheses are 95% confidence intervals. There is more access to social information in the high variance farms, especially when fluctuation in the means is absent. There is less access when fluctuation in the means is possible.

Fluctuation	Standard Deviation	
	0.5	4
0	0.3521 (0.33-0.38)	0.4178 (0.39-0.45)
0.05	0.2621 (0.24-0.29)	0.2879 (0.26-0.31)

Table 7: Akaike Information Criteria (AIC), fit relative to a random model (Δ), Akaike weights (w), and parameter estimates for the three candidate models fit to the data from experiment 3.

Standard Deviation Fluctuation	0.5		4	
	0	0.05	0	0.05
Memory Decay: AIC	1753.49	1321.05	1460.98	1518.32
Δ	0.24	0.33	0.18	0.28
w	7.23E-11	1.91E-10	2.54E-03	3.97E-03
β	0.50	0.64	0.21	0.28
γ	0.00	0.03	0.13	0.00
Linear Imitation: AIC	1706.80	1293.83	1449.17	1508.64
Δ	0.26	0.34	0.19	0.29
w	0.99	1.55E-04	0.93	0.50
β	0.49	0.60	0.23	0.31
γ	0.00	0.00	0.11	0.00
α	0.48	0.60	0.23	0.15
Confirmation: AIC	1755.49	1323.20	1462.96	1519.17
Δ	0.24	0.33	0.18	0.28
w	2.66E-11	6.51E-11	9.42E-04	2.60E-03
β	0.50	0.65	0.21	0.29
γ	0.00	0.06	0.11	0.00
α	0.00	0.00	0.00	0.05
Conformity: AIC	1716.61	1276.29	1454.40	1508.68
Δ	0.26	0.35	0.18	0.29
w	7.38E-03	1.00	0.07	0.49
β	0.47	0.58	0.21	0.30
γ	0.00	0.00	0.11	0.00
α	0.33	0.54	0.12	0.13

Table 8: Total estimated frequencies of social learning in each experiment and treatment. These frequencies come from multiplying the frequency of clicks to view social information by the Akaike-weighted estimated influence of social learning in each treatment.

Experiment	Standard Deviation	Prob of Fluctuation	Frequency of clicks	Weighted α	Total freq of social learning
2	0.5	0	0.24	0.13	0.032
	0.5	0.05	0.23	0.42	0.099
	4	0	0.32	0.04	0.014
	4	0.05	0.26	0.10	0.025
3	0.5	0	0.35	0.48	0.170
	0.5	0.05	0.26	0.54	0.140
	4	0	0.42	0.22	0.091
	4	0.05	0.29	0.14	0.040

List of Figures

740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763

1	Effects of standard deviation on task difficulty as shown by the proportion of optimal planting decisions in each season, for standard deviations 0.5 (easiest) and 4 (hardest). Standard deviation 2 (not shown) is intermediate between these two trends.	45
2	Relative performance of linear (unbiased) imitation, confirmation, and conformity. When possible, conformity outperforms confirmation. Both conformity and confirmation outperform linear imitation. Results shown for $\sigma = 4$ and $f = 0$. Ordering of performance same for other values of these parameters. Each point in the graph is the average from 100,000 simulations.	46
3	Relative performance of confirmation under different experimental conditions. (a) When the variance in yields increases, more Confirmation-based social learning is optimal. (b) When the environment fluctuates more, less Confirmation-based social learning is optimal.	47
4	Relative performance of Conformity-based social learning under different experimental conditions. (a) When the variance in yields increases, more Conformity-based social learning is optimal. (b) When the environment fluctuates more, less Conformity is optimal.	48
5	(a) Distribution of frequency of clicking to access social information, by individual. (b) Frequencies of clicks to access social information, averaged across individuals, by season. Access peaks in the second season and declines steadily until the final season of each farm.	49
6	Distribution of individual frequencies of viewing the decisions of other group members.	50

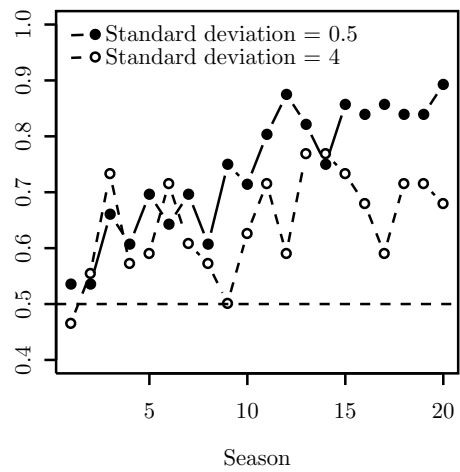


Figure 1: Effects of standard deviation on task difficulty as shown by the proportion of optimal planting decisions in each season, for standard deviations 0.5 (easiest) and 4 (hardest). Standard deviation 2 (not shown) is intermediate between these two trends.

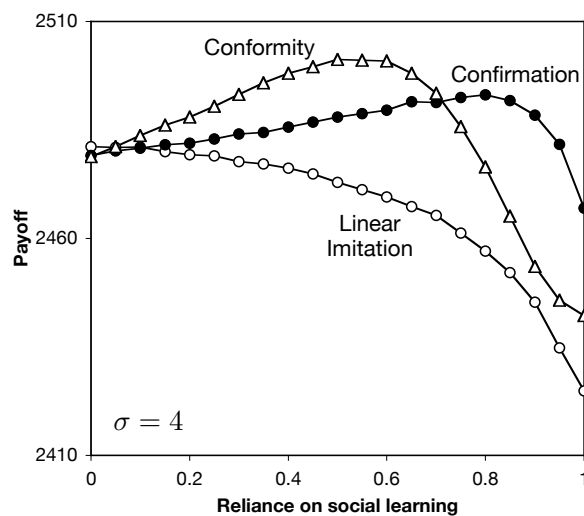
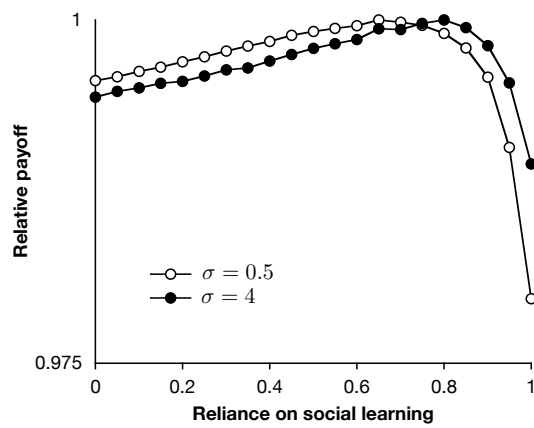
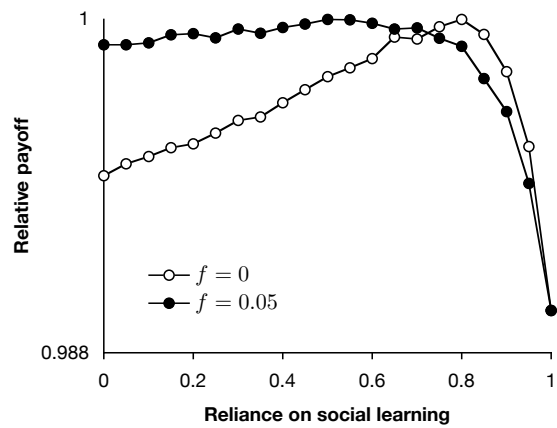


Figure 2: Relative performance of linear (unbiased) imitation, confirmation, and conformity. When possible, conformity outperforms confirmation. Both conformity and confirmation outperform linear imitation. Results shown for $\sigma = 4$ and $f = 0$. Ordering of performance same for other values of these parameters. Each point in the graph is the average from 100,000 simulations.

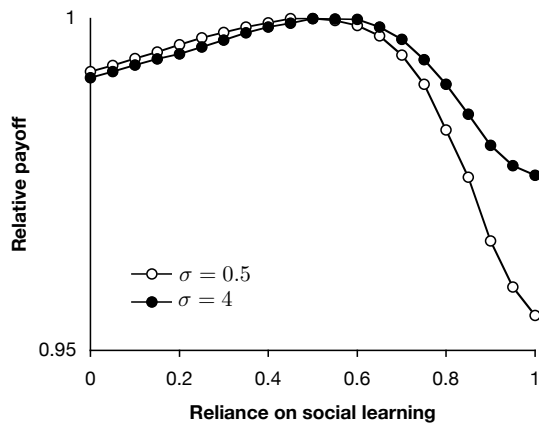


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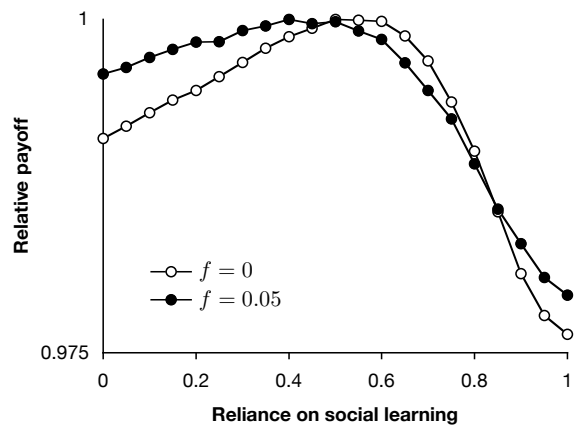


(b)

Figure 3: Relative performance of confirmation under different experimental conditions. (a) When the variance in yields increases, more Confirmation-based social learning is optimal. (b) When the environment fluctuates more, less Confirmation-based social learning is optimal.

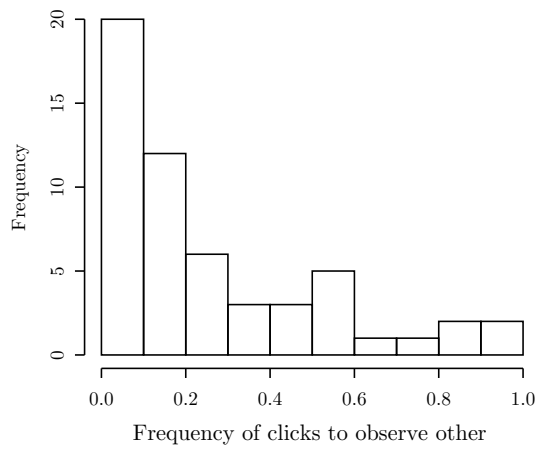


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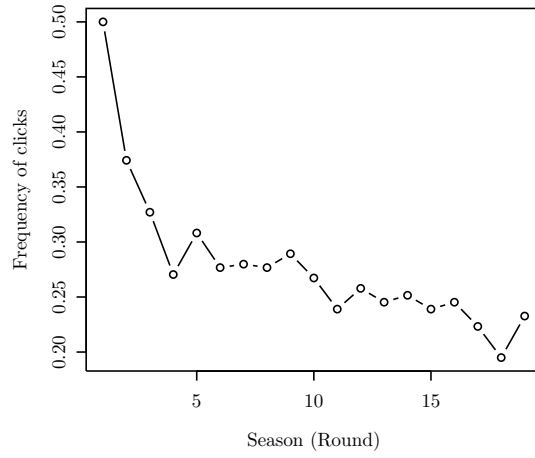


(b)

Figure 4: Relative performance of Conformity-based social learning under different experimental conditions. (a) When the variance in yields increases, more Conformity-based social learning is optimal. (b) When the environment fluctuates more, less Conformity is optimal.



(a)



(b)

Figure 5: (a) Distribution of frequency of clicking to access social information, by individual. (b) Frequencies of clicks to access social information, averaged across individuals, by season. Access peaks in the second season and declines steadily until the final season of each farm.

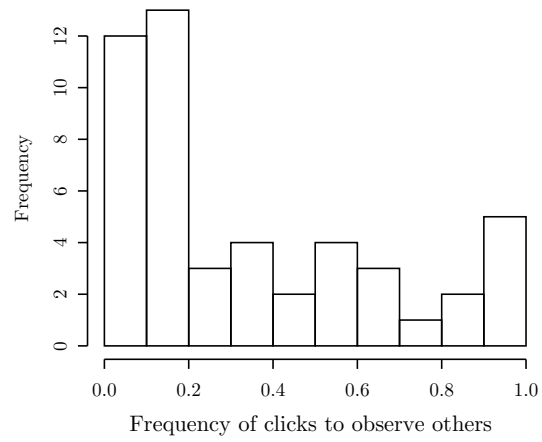


Figure 6: Distribution of individual frequencies of viewing the decisions of other group members.