




Collective decision-making under changing social environments among agents adapted to sparse connectivity

Collective Intelligence
Volume 1:2: 1–16
© The Author(s) 2022
Article reuse guidelines:
sagepub.com/journals-permissions
DOI: 10.1177/26339137221121347
journals.sagepub.com/home/col


Richard P Mann¹ 

¹Department of Statistics, School of Mathematics, University of Leeds, Leeds, UK

Abstract

Humans and other animals often follow the decisions made by others because these are indicative of the quality of possible choices, resulting in ‘social response rules’, that is, observed relationships between the probability that an agent will make a specific choice and the decisions other individuals have made. The form of social responses can be understood by considering the behaviour of rational agents that seek to maximise their expected utility using both social and private information. Previous derivations of social responses assume that agents observe all others within a group, but real interaction networks are often characterised by sparse connectivity. Here, I analyse the observable behaviour of rational agents that attend to the decisions made by a subset of others in the group. This reveals an adaptive strategy in sparsely connected networks based on highly simplified social information, that is, the difference in the observed number of agents choosing each option. Where agents employ this strategy, collective outcomes and decision-making efficacy are controlled by the social connectivity at the time of the decision, rather than that to which the agents are accustomed, providing an important caveat for sociality observed in the laboratory and suggesting a basis for the social dynamics of highly connected online communities.

Keywords

collective behaviour, social decision-making, rationality, social connectivity

Significance statement

Many animal populations are undergoing rapid declines in density and diversity resulting from climate change and habitat destruction. At the same time, human societies have entered a new era of global connectivity through ubiquitous telecommunications and social media. As a result, the social environment for both human and animal communities is changing dramatically. Predicting the likely consequences of these social changes is vital for understanding the resilience of these communities in the future, and such predictions rely on understanding what individual animals and humans are seeking to achieve through their social interactions. In this paper, I show how the behaviour of agents adapted to succeed in a particular social world is likely to change when the number of available social connections increases or decreases. The results of this study suggest that the welfare of all social communities depends on careful stewardship of social connections. Animal populations must be maintained at densities high enough to effectively utilise social information, while human societies need to adapt intelligently to rapid increases in social connectivity.

Corresponding author:

Richard P Mann, Department of Statistics, University of Leeds, Woodhouse Lane, Leeds LS2 9JT, UK.

Email: r.p.mann@leeds.ac.uk



Creative Commons CC BY: This article is distributed under the terms of the Creative Commons Attribution 4.0 License (<https://creativecommons.org/licenses/by/4.0/>) which permits any use, reproduction and distribution of the work without further permission provided the original work is attributed as specified on the SAGE and Open Access pages (<https://us.sagepub.com/en-us/nam/open-access-at-sage>).

Introduction

Living in groups provides many advantages to social animals (Krause and Ruxton, 2002), including the opportunity to make use of ‘social information’ to improve the quality of decision-making (Ward et al., 2011; Wolf et al., 2013). Choices made by others are often a useful indicator of which option is best for oneself; if one trusts that others are acting rationally (Mann, 2018), and that they have similar preferences to oneself (Mann, 2020), then their decisions provide valuable clues about the quality of different options. However, when others are also attending to each other this can cause information cascades (Anderson and Holt, 1997; Bikhchandani et al., 1992), rendering all but a few decisions meaningless as sources of information. A crucial factor likely to determine how individuals learn from observing each other is therefore the social connectivity of the group, who each individual can observe and respond to. Previous research has investigated how rational agents make use of social information (Arganda et al., 2012; Mann, 2018, 2020; Karamched et al., 2020; Pérez-Escudero and De Polavieja, 2011) but these studies have focused on groups in which decisions constitute ‘common knowledge’ (Aumann, 1976), whereas in real groups and societies agents typically observe only a subset of other individuals. For example, visual occlusion by other individuals can restrict fish to observing a small subset of other group members (Poel et al., 2021; Rosenthal et al., 2015; Strandburg-Peshkin et al., 2013); social relationships in bird populations may form a sparse but connected network across broad geographical regions (Psorakis et al., 2015); and historically most human societies were characterised by sparse social connections relative to the total population (Apicella et al., 2012; Dunbar, 1998). As such, evolved social behaviour is likely to be explicable as an adaptive response to a social environment characterised by sparse connectivity.

Understanding how the structure of the social environment shapes the way that individuals interact is of critical importance due to rapid changes to this structure in both human and animal populations (Bak-Coleman et al., 2021; Fisher et al., 2021). One effect of large scale loss of animal abundance (Gerardo et al., 2020; Wagner et al., 2021) is to reduce the number and diversity interactions between con- and hetero-specifics. How animals individually and collectively respond to this social change will be a key determinant of their robustness to climate change and habitat destruction. Human societies have also been subjected to waves of dramatic change in the social landscape, the most recent of which is the rise of social media and ubiquitous, instant global connectivity. Each step change in communications technology has had profound social consequences (Eisenstein, 1980; McQuail, 1977) to understand the possible implications of the latest communications revolution

one must identify the factors that determine how people exchange social information and extrapolate these to determine the collective consequences of changing social connectivity (Fisher et al., 2021).

Learning on social networks is a well-studied phenomenon, with many theoretical treatments. The De Groot Model DeGroot (1974) and the Voter Model Holley and Liggett (1975) present the canonical examples, and the majority of subsequent treatments have followed these by assuming that agents update their opinions based on those of their social connections according to heuristic rules that are specified in advance, such as adopting the majority or average opinion of those in the social neighbourhood. While this permits precise analysis of the resulting opinion dynamics, the conclusions of such models hinges on the validity of the specified update rules, which must be justified either intuitively or empirically, rather than with respect to a theory of what the agents are seeking to achieve through their decisions.

An alternative to heuristic models is to imbue agents with goals, through utility functions, and the ability to make rational decisions through statistical inference. Sequential decision-making based on Bayesian rational decision theory is a paradigm for understanding the mechanisms of collective decision-making through such inference in both human and animal groups. Within social science (particularly economics), research has primarily focused on the efficacy of individual and collective decisions by rational agents under a variety of different scenarios (Bikhchandani et al., 2021). This research usually takes agents to be rational under the strong assumption that individuals are aware of the full details of the context in which they make the decision. Under this assumption, researchers often seek to determine properties of collective decision-making in large groups ($n \rightarrow \infty$), permitting formal mathematical answers to questions such as the conditions under which information cascades will occur (Anderson and Holt, 1997; Bikhchandani et al., 1992) and whether the majority of agents will decide correctly (Acemoglu et al., 2011; Bikhchandani et al., 1992; Mossel et al., 2015). Closely related to the current study, Acemoglu et al. (2011) investigated sequential decision making by individuals in social networks that determine which other agents they can observe and identified conditions under which agents either are or are not guaranteed to converge to the correct decision. In particular, they show that convergence to the correct decision is guaranteed when agents’ private beliefs are unbounded (an agent may receive arbitrarily strong private signals), but otherwise convergence depends on the network topology – a result that is further extended by Mossel et al. (2015). However, these theorems strictly pertain only to the limiting behaviour of very large social networks, and

depend on the assumption that agents hold correct beliefs regarding the structure of the social world and how other agents will respond to information, an assumption that may not hold under conditions of rapid social change or displacement to a new environment.

In animal biology, a greater focus has been placed on identifying the ‘interaction rules’ that specify the probability that a focal individual will make a specific decision conditioned on the choices made by other animals (typically con-specifics) within the group (Arganda et al., 2012; Mann, 2018, 2020, 2021; Pérez-Escudero and De Polavieja, 2011). Of particular interest is the goal of deriving theoretically motivated interaction rules that closely match empirical observations from collective decision-making in animal groups, whether in the laboratory Arganda et al. (2012); Kadak and Miller (2020); Miller et al. (2013); Pérez-Escudero and De Polavieja (2011) or the field Farine et al. (2014). As such, this research typically prioritises identifying what can be observed in social decision-making over establishing theoretical bounds on decision-making performance. In pursuit of this goal, past modelling has generally considered group sizes consistent with experimental research, often below 10 individuals and rarely above 100, rather than deriving results for the limiting case of very large group sizes. In this context, most models have made the assumption, often implicitly, that each individual within the group can observe all other group members and their decisions, including the order in which those decisions were made. Recent work, however, has shown how agents can respond optimally to highly specific forms of restricted social information, such as the aggregate number of previous agents selecting each option (Mann, 2021).

Of special interest from a biological perspective is the question of how variation in the observable behaviour of individuals and groups can be explained through variation in context such as the quality of environmental information (Mann, 2018), degree of alignment between individual preferences (Mann, 2020) or changes to the overall risk level of the choices available (Pérez-Escudero and De Polavieja, 2017). A distinguishing feature of the biological approach has been the consideration when investigating the effect of changing these model parameters that agents may remain adapted to a previously established context (Mann, 2018, 2020; Pérez-Escudero and De Polavieja, 2017), with particular consideration as to how animals are likely to behave in the laboratory as opposed to in the natural environment (Mann, 2018).

In this paper, I take a primarily biological approach to the sequential decision-making problem, characterising the observable features of the interactions between agents who act rationally based on expectations originating from their adaptation to habitual conditions. I extend previous biologically motivated models (Mann, 2018, 2020, 2021) by

considering agents that live on a random network, such that each individual can only observe the decisions made by a limited number of other group members: that subset of the group to which it is connected. I identify the optimal decision-making rules for agents inhabiting such networks, and in particular, I focus on the form of those rules in the case where the social network is sparse, corresponding to the biologically relevant examples given above. In this context, I explore how the form of the resulting interaction rules depend on the habitual environmental and social parameters controlling the reliability of private information and the typical alignment of preferences to which individuals are adapted. I then investigate the collective consequences of those individual interactions in groups of varying sizes and show how the characteristics and efficacy of collective decisions change when the connectivity of the group is varied over timescales that do not allow for full adaptation, with a discussion of the implications of these results for both animal and human communities.

Model

I consider a group of n agents sequentially choosing between two options, A and B. The true utilities of A and B are unknown by the agents and may differ between agents; for agent i these are denoted as $U_{A,i}$ and $U_{B,i}$, and I define the difference between these utilities as $x_i \equiv U_{A,i} - U_{B,i}$ (Mann, 2018, 2020). Agents are assumed to be rational decision-makers who seek to maximise the expected utility of their decision Von Neumann and Morgenstern (2007) based on the total information I that they have.

$$P(i \rightarrow A) = \begin{cases} 1 & \text{if } \mathbb{E}(x_i | I) > 0 \\ 0 & \text{otherwise} \end{cases} \quad (1)$$

Agents estimate the value of x_i based on both private information received from the environment and social information in the form of observed decisions by other agents. Following previous work (Mann, 2018, 2020, 2021), agents start with normally distributed prior beliefs over x_i , with zero mean (no prior information in favour of A or B) and unit variance (setting the arbitrary scale of utility)

$$p(x_i) = \phi(x_i), \quad (2)$$

where $\phi(\cdot)$ is the standard normal probability density function. Alignment between agents’ preferences are represented by the covariance of utility differences, defined as $\text{cov}(x_k, x_l) = \rho$ for any two different agents k and l .

Each agent receives private information, Δ_i , from the environment that represents a noisy signal of the true value of x_i

$$p(\Delta_i | x_i) \propto \phi((x_i - \Delta_i)/\epsilon), \quad (3)$$

where ϵ controls the reliability of this private information. Given social information, \mathbf{s} , agent i forms a posterior belief about the value of x_i by combining its prior, its private information and the social information

$$p(x_i | \Delta_i, \mathbf{s}) \propto p(x_i)p(\Delta_i | x_i)P(\mathbf{s} | x_i), \quad (4)$$

where the private information and social information are independent conditional on the value of x_i . In the context of this model, social information \mathbf{s} constitutes an ordered sequence of previous decisions observed by the focal agent prior to making its own decision. The effect of social information \mathbf{s} on the decision of agent i is therefore determined by the detailed structure of $P(\mathbf{s}|x_i)$ – the probability that this social information would be observed conditioned on the value of x_i . If the observed social information is more probable conditioned on high values of x_i , agent i will accordingly adjust its belief in favour of option A and vice versa, weighing this against its private information Δ_i .

In judging the social information it observes, a rational agent must recognise that this information is potentially incomplete, since not all other individuals may be observable. That is, the social information the focal agent observes, \mathbf{s} , is a subset of the full sequence of past decisions, \mathbf{c} . Those that the focal agent can observe may in turn have been influenced by each other, or by further, non-observed individuals. In this paper, I assume that each agent observes any other with a fixed probability q , and this set of observation relations thus forms a random directed network that may range from fully connected ($q = 1$), where all agents observe each other, to sparse ($q \ll 1$), where each agent only observes a small subset of the overall group. I assume that in any given group, agents are habituated to a specific social connectivity q_{habitual} , whether by experience or evolutionary adaptation, under which their behaviour is optimised. That is, they act so as to maximise their expected utility, with that expectation being conditioned social connectivity remaining at this habitual value. This may differ from the true connectivity at the time behaviour is observed, q_{actual} , as a result of social or environmental changes that alter the connectivity more quickly than adaptation can occur, leading to potential changes in behaviour that may not be optimal for the new context.

The assumption above that agents interact with a fixed probability implies that the network of interactions forms an Erdős-Renyi random graph. This is a strong assumption, but necessary for the mathematical treatment here. It excludes explicit treatment of, for example, spatially organised groups or clustered social networks. In the discussion section, I elaborate on how these might be combined with results from fully connected networks to consider the likely behaviour in real groups.

An optimal decision-making rule within this model is one that provides a unique decision (A or B) for any combination of private information (Δ_i) and social

information (\mathbf{s}), such that the individual maximises the expected utility of their choice, given the conditional expectation of x_i

$$E(x_i | \Delta_i, \mathbf{s}) \propto \int_{-\infty}^{\infty} x_i p(x_i) p(\Delta_i | x_i) P(\mathbf{s} | x_i) dx_i. \quad (5)$$

Therefore, the appropriate decision rule is as follows:

$$P(i \rightarrow A) = \begin{cases} 1 & \text{if } \int_{-\infty}^{\infty} x_i p(x_i) p(\Delta_i | x_i) P(\mathbf{s} | x_i) dx_i > 0 \\ 0 & \text{otherwise} \end{cases} \quad (6)$$

Equation (6) implies that there is a critical threshold Δ_s^* , associated with any given social information (as observed by the agent), such that agent i will choose A if and only if $\Delta_i > \Delta_s^*$. From the perspective of a rational agent, decisions are fully determined by the value of their private information relative to the appropriate threshold specified by the social information they observe. However, when considering the response of an agent to social information as seen by an external observer, both the private and social information they observe is unknown. This results in uncertainty in the decision the agent will make conditioned on a given set of social information as seen by the external observer, \mathbf{c} , such that the response to this social information can be characterised by an apparently stochastic decision rule

$$P(i \rightarrow A | \mathbf{c}) = \sum_{s \in S} P(\Delta_i > \Delta_s^*) P(\mathbf{s} | \mathbf{c}), \quad (7)$$

where the summation is over the set, representing all possible full sequences of past decisions. When considering the response of an agent to specific social observations, we may choose to assume that we know which social information it has observed. In this case, we can collapse the summation to consider only one observed sequence, \mathbf{s} :

$$P(i \rightarrow A | \mathbf{s}) = P(\Delta_i > \Delta_s^*). \quad (8)$$

More details on the mathematical structure and derivation of optimal decision-making rules is given in the Methods section, and a full list of model parameters and definitions is given in Table 1.

Results

Effect of sparse connectivity on optimal social response

I first derived in full the optimal decision-making rule for agents in a group of 10 identical individuals ($\rho = 1$), sequentially choosing between two options A and B, with varying degrees of habitual connectivity, q_{habitual} .

Table 1. Definitions of model parameters.

Parameter	Definition
$U_{A,i}, U_{B,i}$	True utility of options A and B for agent i
x_i	Difference in utilities between the options A and B for agent i : $x_i = U_{A,i} - U_{B,i}$
z	Average utility difference across all agents
ρ	Correlation in utility differences for any pair of agents
Δ_i	Private signal received by agent i
ϵ	Scale of noise in private signals
C_k	The decision made by the k th agent in a sequence (1 for A, -1 for B)
\mathbf{c}	A full sequence of past decisions by other agents
\mathbf{s}	A partial sequence of past decisions as seen by a focal agent
Δ_s^*	The threshold applied to private signals by an agent that observes a sequence of past decisions \mathbf{s} , such that the agent chooses A if $\Delta_i > \Delta_s^*$
q_{habitual}	Habitual connectivity: The probability a focal agent can observe any other individual in the habitual environment
q_{actual}	Actual connectivity: The probability a focal agent can observe any other individual in the current environment
N_A, N_B	Actual number of agents that have chosen A and B
n_A, n_B	Number of agents that have chosen A and B as observed by a focal agent
Δ_w^*	The threshold applied to private signals by an agent that observes a difference $w = n_A - n_B$ in the number of previous choices for A and B, such that the agent chooses A if $\Delta_i > \Delta_w^*$

Specifically, I consider cases where agents observe 90% of other decisions ($q_{\text{habitual}} = 0.9$), 50% ($q_{\text{habitual}} = 0.5$) and 10% ($q_{\text{habitual}} = 0.1$) on average. To characterise the optimal behaviour, I mapped the probability that a focal agent will select option A conditioned on the number of previous individuals that it observes to have chosen A or B (n_A and n_B , respectively), assuming that (unknown to the decision-maker) the two options are in fact identical ($x = 0$). This probability is shown for the three cases in Figure 1 panels A ($q_{\text{habitual}} = 0.9$), B ($q_{\text{habitual}} = 0.5$) and C ($q_{\text{habitual}} = 0.1$). Panel A (high connectivity) shows a pattern of social response very similar to the previously derived case of full connectivity (Mann, 2018). In panel B (intermediate connectivity), the social response is stronger (greater probability to follow the majority) and in panel C (low connectivity) stronger still – agents adapt their strategy to attend more strongly to social information when it is limited. As the connectivity is decreased, the contour lines of equal probability also transform from a radial pattern (panel A), which maps to a proportional difference between n_A and n_B , to a parallel pattern (panel C) that more closely maps to an absolute difference.

More insight into the form of the social response is gained by conditioning on those cases in which the most recent observable decision-maker selected option A, shown in panels D–F. Previous work has shown that the most recent decision can be more important than all other social cues when agents can observe all previous decisions (Mann, 2018, 2020). In the case of high connectivity (panel D), this conditioning indeed makes a large difference to the probability for the focal agent to choose A. As connectivity is decreased, the effect of this most recent decision diminishes,

and in the case of sparse connectivity is negligible (panel F), showing that the agents in this case respond only to aggregate quantities (n_A and n_B) and effectively ignore the ordering of previous decisions.

Panels G–I show the social response projected onto a single dimension: the absolute difference $n_A - n_B$. Grey points indicate the probability conditioned on each potential sequence of past decisions, while red points indicate the average of these (weighted by the probability of the sequence arising) for each possible value of the difference. This shows that the initially sequence-dependent response in panel G transforms to depend only on the quantity $n_A - n_B$ when the connectivity is lowered (panels H–I).

The collective consequences of these decision rules can be quantified by the total number of agents that ultimately select option A, N_A (upper-case N will in general denote actual numbers of agents making a decision, while lower case n denotes the number observed by an agent). Here, one can distinguish between two possible scenarios. The first is a collective decision made under the conditions of social connectivity the agents are optimised for ($q_{\text{actual}} = q_{\text{habitual}}$). This is shown in panels J–L, with a tendency towards consensus ($N_A = 10$ or $N_A = 0$) in the case of high connectivity (panel J), which decreases in line with decreasing connectivity. Although agents optimised for low social connectivity have a stronger response to the other agents they can observe (see above), this does not compensate for the overall reduction in social information, so agents operating with low connectivity are much less likely to all select the same option (panels K–L). This pattern reverses, however, if the decision is made under a different regime,

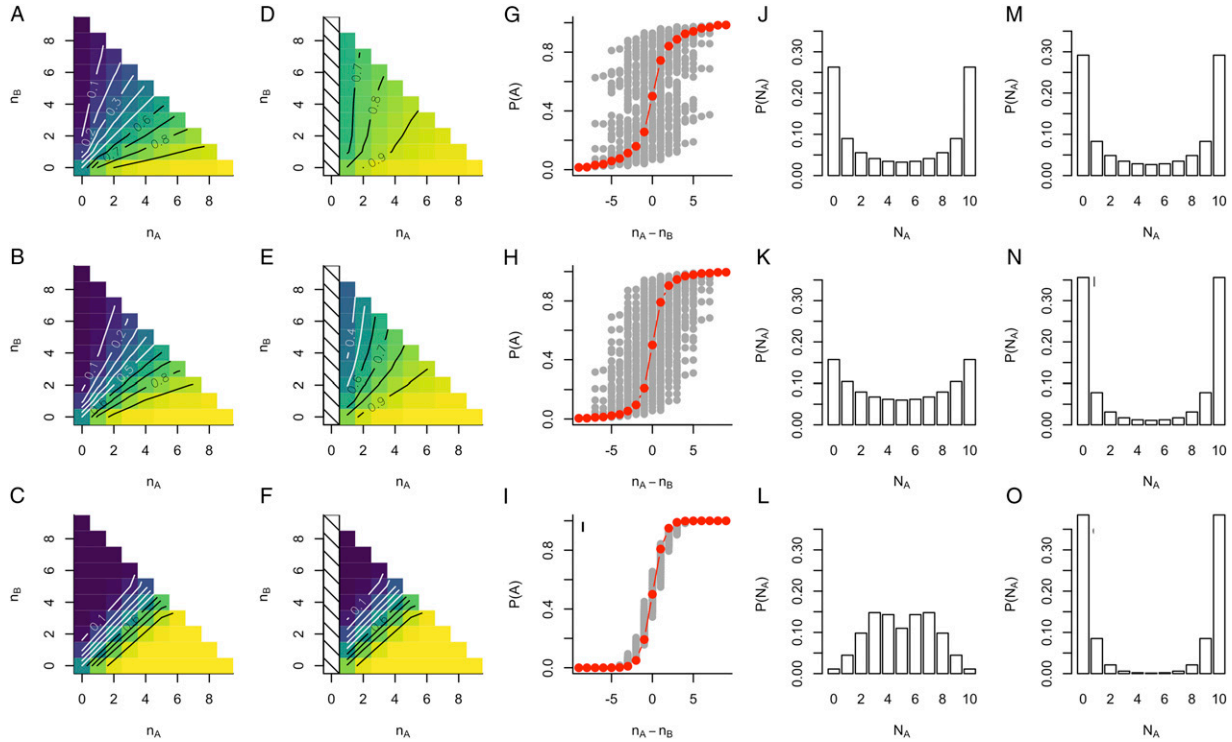


Figure 1. Characterising the optimal social information strategy for differing values of social connectivity in a group of 10 agents (other group sizes give similar results). Each row shows differing perspectives on the social response by agents habituated to a specific level of social connectivity: $q_{\text{habitual}} = 0.9$ (top row), $q_{\text{habitual}} = 0.5$ (middle row) or $q_{\text{habitual}} = 0.1$ (bottom row). Panels A–C show the probability a focal agent will choose option A, conditioned on the number of previous decisions for A and B observed by that agent. Panels D–F show the same, but further conditioned on the most recently observed decision being for A (thus also ruling out instances where $n_A = 0$). Panels G–I show the social response projected onto one dimension: the difference $n_A - n_B$; grey points indicate probabilities arising from specific sequences, while red points show the mean probability for each value of $n_A - n_B$. Panels J–L show the consequence of these decision rules in terms of the distribution of collective outcomes (total number of decisions for A) under habitual conditions ($q_{\text{actual}} = q_{\text{habitual}}$, where social connectivity remains at the level the agents are adapted to), while panels M–O show the distribution of collective outcomes under conditions of full connectivity ($q_{\text{actual}} = 1$) if agents continue to apply the behavioural rules optimised for their habitual conditions.

where all agents can observe each other ($q_{\text{actual}} = 1$), but continue to apply the behavioural rules that are optimised for their habitual conditions. As shown in panels M–O, under this treatment, agents optimised for conditions of low connectivity are more social, resulting in a greater degree of group consensus. Such a change in social connectivity might result from, for example, translating animals into a laboratory environment that lacks the usual sources of sensory occlusion that the wild habitat contains, or by placing human subjects in a computer-mediated environment that reveals the decisions of all group members in a way that would not occur in more naturalistic settings.

Behaviour of large, sparsely connected groups

To analyse the behaviour of larger group sizes that characterise many animal and human societies, one can utilise the emergent property of sparsely connected groups revealed above: the dependence of the social response solely

on the absolute difference $n_A - n_B$. This property means that an optimal decision rule for agents in a group of size n can be characterised by responses to $2n + 1$ distinct possible values of $n_A - n_B$, and this can be further reduced by applying the symmetry in response to absolute differences of equal magnitude but differing sign. Using a Monte Carlo method (see Methods), I derived the optimal decision-making rule for agents in a group of 101 individuals, with a habitual connectivity of either $q_{\text{habitual}} = 0.01$, $q_{\text{habitual}} = 0.05$ or $q_{\text{habitual}} = 0.1$ (equivalent to a mean degree of 1, 5 or 10). This optimal strategy is illustrated in Figure 2(a), which shows that the probability for a focal agent to select option A, conditioned on the value of $n_A - n_B$ it observes, is almost identical across the differing levels of habitual connectivity. I then calculated the probability that an agent will select option A, conditioned on the difference $N_A - N_B$, where N_A and N_B refer to the actual number of previous decision-makers selecting either A or B, not only those that the focal agent can observe. That is, this probability

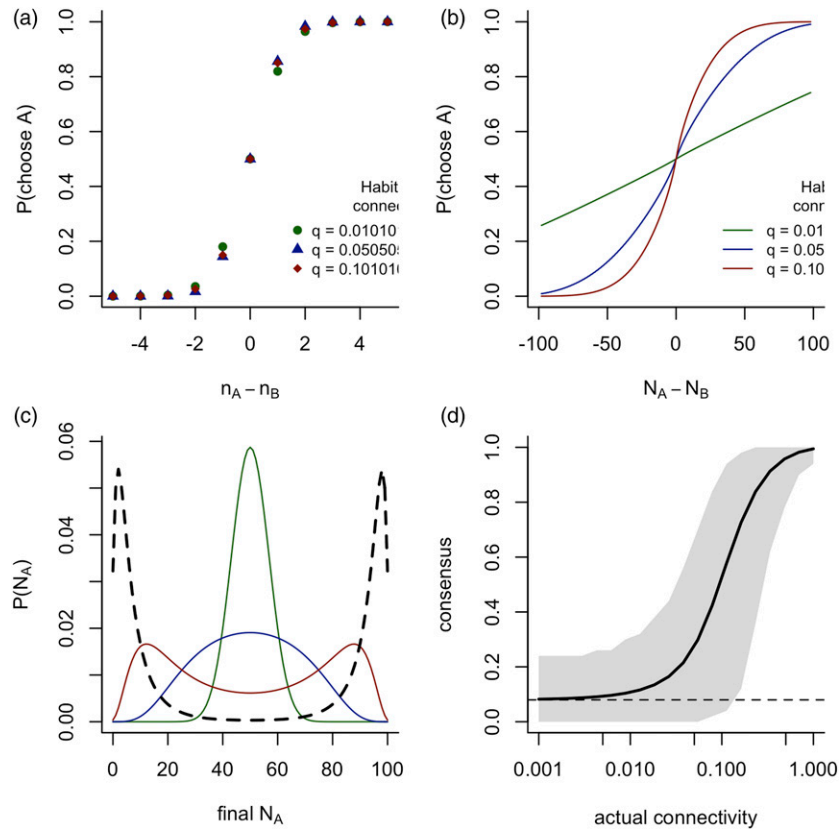


Figure 2. Characterising the optimal social information strategy in a large ($n = 100$), sparsely connected network, for agents habituated to differing connectivity ($q_{\text{habitual}} = \{0.01, 0.05, 0.1\}$). (a) Agents adopt a decision rule that depends on the observed difference in the number of prior decisions between options ($n_A - n_B$). This rule is essentially independent of the connectivity to which they are habituated. (b) When this rule is applied under the habitual conditions, the observable response to the true difference ($N_A - N_B$) is weaker in conditions of lower connectivity. (c) More sparsely connected groups are less cohesive under natural conditions (solid lines), as measured by the total number of agents choosing one option. Increasing network connectivity to $q_{\text{actual}} = 0.2$ and applying the same decision rule results in a substantially more cohesive collective outcome that is independent of the habitual connectivity (dashed line). (d) The degree of consensus ($|N_A - N_B|/(N_A + N_B)$) varies monotonically with the true connectivity of the network and is independent of habitual connectivity. The solid line indicates the expected consensus for varying actual connectivity; the grey region indicates the 95% probability region. The dashed lines indicate the expected consensus when agents choose independently.

represents that which would be inferred by making a simple empirical analysis of the proportion of agents choosing option A based on the number of previous decisions for A and B seen by an external observer who can observe all decisions but who does not have access to the network defining which agents can observe each other. This probability, shown in Figure 2(b), varies strongly with habitual connectivity. Taken together, the results in panels A and B indicate that agents in more densely connected (but still sparse) networks exhibit more sociality by virtue of increased social connections, rather than as a result of any difference in how they use that social information. To see how this is reflected in collective outcomes, I calculated the probability that a given proportion of agents will ultimately select option A under each level of habitual connectivity, under habitual conditions ($q_{\text{actual}} = q_{\text{habitual}}$). The results in Figure 2(c) show that agents in the more

connected network ($q_{\text{habitual}} = 0.1$, red line) exhibit a tendency to consensus, with the peaks of the distribution near 0 and 100, while those in the less connected networks show a distribution that is peaked at 50, indicating a tendency for the group to split evenly between the options; when $q_{\text{habitual}} = 0.01$ (green line) this is barely distinguishable from a binomial distribution in which each agent chooses independently. However, the same three sets of agents, transferred to a new condition with increased connectivity of $q_{\text{actual}} = 0.2$ (mean degree = 20), all exhibit identical collective behaviour, with a strong bias towards choosing the same option (dashed black line). Finally, I repeated this analysis for actual connectivity levels from $q_{\text{actual}} = 0.01$ (mean degree = 1) to $q_{\text{actual}} = 1$ (mean degree = 100) and measured the resulting consensus, defined as $|N_A - N_B|/(N_A + N_B)$, once all decisions have been made. As shown in Figure 2(d),

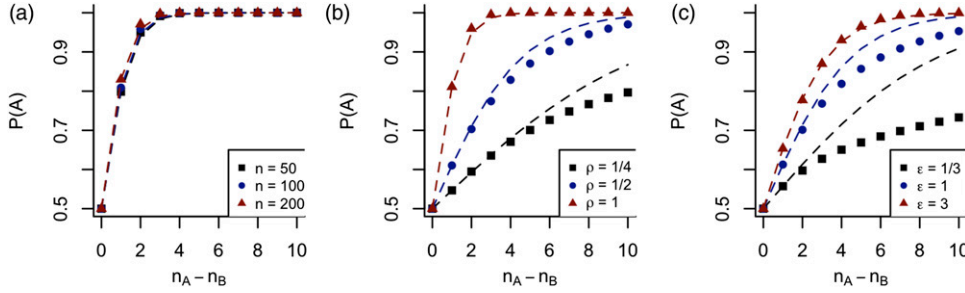


Figure 3. Variation of optimal social responses with differing group size n , preference alignment ρ and private information variance ϵ . (a) Variable n with fixed $\rho = 1$ and $\epsilon = 1$, showing that the social response does not depend on group size; (b) variable ρ , with fixed $\epsilon = 1$, $n = 100$, showing an increasing social response as preference correlation increases and (c) variable ϵ with fixed $\rho = 0.5$ and $n = 100$, showing an increasing social response with increasing environmental noise. The probability given by equation (9) is also plotted in each case (dashed lines) (dashed lines), with sociality parameter S given by equation (10), which provides a close fit between the two when agents share identical preferences ($\rho = 1$), but otherwise consistently over-estimates the probability the follow the majority of previous decisions, especially for high values of ϵ .

group consensus varies strongly and monotonically with q_{actual} .

Effect of differing preferences

In Figure 3, I show the probability that a focal agent will choose option A in a symmetric setup when considering agents, as a function of the value of $n_A - n_B$ observed by the agent, while varying group size (n), agents' preference correlation (ρ) and environmental noise (ϵ). These results show that an agent's predicted response to the observed social information is essentially invariant to group size (panel A), but depends strongly on the degree of preference correlation (panel B) and the environmental noise (panel C). For any given value of $n_A - n_B$, the focal agent is more likely to follow the majority of previous decisions if ρ or ϵ is increased. This follows similar results obtained in the case of a fully connected social network (Mann, 2020).

The binary decision problem studied here was also considered in Pérez-Escudero and De Polavieja (2011). There the authors present a derivation for a decision rule based on an approximation under which other agents are assumed to have made their choices independently. In the case of a symmetric choice, they arrive at a decision rule of the form

$$P(i \rightarrow A | n_A, n_B) = \frac{1}{1 + S^{-(n_A - n_B)}}, \quad (9)$$

where S is a sociality parameter that represents the relative likelihood that another individual will make a 'correct' or 'incorrect' choice. Since the effect of social information in this rule also depends on the value of $n_A - n_B$ it is natural to compare how that study relates to this one. In the case of a symmetric choice ($x_i = 0 \forall i$), rearranging equation (9) suggests an effective value of the sociality parameter

$$S = (1/P(i \rightarrow A | n_A - n_B = 1) - 1)^{-1} \quad (10)$$

where $P(i \rightarrow A | n_A - n_B = 1)$ is the probability that agent i will choose option A, having observed social information $n_A - n_B = 1$ (see Methods).

Figure 3 shows that equation (9) provides a highly accurate approximation to the predicted social response when agents share identical preferences ($\rho = 1$). However, substantial departures from equation (9) are observed when agent preferences are non-identical ($\rho < 1$). In these cases, the probability that an agent will follow the current majority is always lower than that suggested by equation (9) (with a value of S given by equation (10)), and this difference is larger for larger values of $|n_A - n_B|$. Deviations from equation (9) are particularly acute when $\rho < 1$ and ϵ is small, representing situations where an agent has highly trustworthy private information and cannot assume that others share the same preferences as itself. Thus, although the model considered in this paper supports a social response that depends on $n_A - n_B$, as in Pérez-Escudero and De Polavieja (2011), this must be treated with caution. First, this approximation is only valid in the case where interactions between agents are sparse. Second, within the set of functions of $n_A - n_B$, the particular form of equation (9) is only a valid approximation when $\rho \approx 1$; that is, when agents can effectively be considered to be identical. Thirdly, this model highlights that the n_A and n_B that the focal agent responds to are the number of decisions for A and B that the agent itself perceives, which may be different to those recorded by, say, an experimental observer (cf. Figures 2(a) and (b)). Thus, statistical efforts to fit equation (9) to experimental or observational data (see e.g. Eguíluz et al. (2015); Farine et al. (2014) as well as examples in Pérez-Escudero and De Polavieja (2011)) are valid when the agents under observation habitually inhabit a sparse interaction network, when those agents have closely aligned preferences, and insofar as the recorded social information

accurately represents the individuals the focal agent actually attended to.

Impact of changing connectivity on individual rewards

The results above show that agents adapted to sparse social connections will exhibit stronger social responses when connectivity is increased. What impact will this have on the rewards the agents are able to extract from the environment? Naively one might expect that any deviation from the condition they agents are adapted to will be detrimental, but that expectation may be deceptive in this case; social connectivity increases the total social information available to each agent, and the benefit of this additional information may outweigh the costs of being adapted to a different social environment.

I calculated the expected payoff per decision for identical agents ($\rho = 1$) adapted to sparse connectivity (habitual mean degree = 5) in groups of varying size between $n = 50$ and $n = 200$ under a range of experimental social connectivities between $q_{\text{actual}} = 0.01$ and $q_{\text{actual}} = 1$. This expected payoff is plotted in Figure 4 as a function of mean degree $(n - 1)q_{\text{actual}}$ (panel A) and connectivity q_{actual} (panel B). These results show that for any group size there is an optimal connectivity, at which the expected payoff is maximised. This optimal connectivity varies consistently with group size, such that the optimal mean degree varies proportionally to the square root of the group size (panel C), and equivalently the optimal value of q_{actual} varies in inverse proportion to \sqrt{n} (panel D). That is, in larger groups, the optimal absolute number of social connections per agent increases, but less than linearly, such that the optimal proportion of possible social connections that are actually realised decreases with group size. Above this optimal value, further increases in connectivity reduce expected payoffs.

Discussion

In this paper, I derived the optimal use of social information arising from incomplete observations of the decisions of other agents. I showed that when such information is sufficiently sparse, rational agents develop a strategy that attends only to the absolute difference between the number of other individuals that have selected each option, ignoring both the total number of previous decisions and the order in which they were made. Exploiting this property to analyse large groups revealed that the optimal social response is essentially independent of the degree of social connectivity the agents are habituated to and behaviour is instead controlled by the connectivity at the time the decision is made. In particular, greater connectivity leads to increasing consensus among group members. Finally, I showed that

increases in social connectivity are beneficial to sparsely connected groups up to a critical value of q , increasing the average payoff of the options individuals choose. However, sufficiently large increases in connectivity result in extremely high social consensus that is associated with lower average rewards.

This model predicts that the number of social connections individuals experience at the time they make a decision determines the degree of group consensus. This has practical consequences for the scientific study of collective behaviour. For example, an animal that lives in conditions of high sensory occlusion such as muddy water may respond weakly to local conspecifics simply because it does not perceive many of them; if translated to an environment with less occlusion such animals may exhibit increased apparent social responses without fundamentally changing how they respond to the other individuals that they can perceive. While in experiments one might expect animal or human subjects to adjust somewhat to the new context (and indeed animal subjects are often given time in the new environment to acclimatise), this finding provides an important context for laboratory-based research, especially since the effect operates in the same direction as earlier findings that decreased noise in environmental signals could also increase observed sociality in laboratory conditions (Mann, 2018).

Where rational individuals experience sparse social connectivity they are able to treat the decisions of others as being effectively independent. As long as this assumption remains valid, more social information, in the form of additional social connections, is beneficial to a decision-maker, resulting in a greater average payoff from their decisions. However, if social connectivity exceeds a critical value this independence assumption begins to breakdown, and if agents do not recognise and adjust for this they will make worse decisions. In this model, this critical value of social connectivity was found to depend on the square root of the total group size, such that $q_{\text{optimal}} \propto 1/\sqrt{n}$, giving a critical mean degree proportional to \sqrt{n} . This scaling resembles that found in previous studies of optimal network structure for collective decision-making Kao and Couzin (2019), suggesting a common basis in agents applying decision rules that assume independence between those others who they observe.

The relationship between connectivity and decision-making efficacy has profound implications for the welfare and sustainability of animal and human communities. Under pressure from climate change and habitat destruction, many animal populations have experienced precipitous declines. Many of the affected populations are social animals, and these may depend on social information, for instance to locate food sources or navigate to seasonal habitats. Reduction in group size while holding constant the probability of social interaction between any pair of

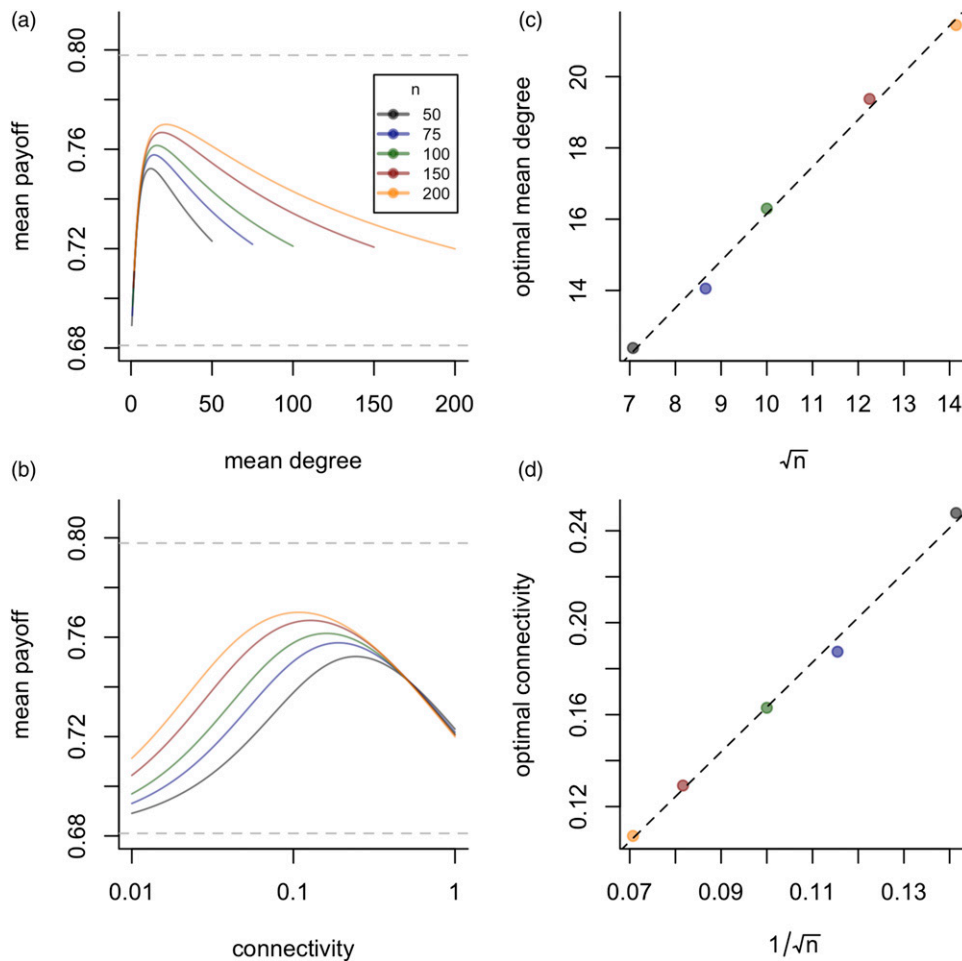


Figure 4. Average payoff obtained by agents in sparse networks of varying sizes as a function of actual connectivity and mean degree. Agents in each network are habituated to a connectivity $q_{\text{habitual}} = 5/(n - 1)$ implying a mean degree of 5, though the results are not sensitive to this factor as long as habitual connectivity is sparse. (a) Mean payoffs increase as mean degree increases in all networks up to an optimal value, and then fall, with a differing optimal mean degree for different group sizes. The dashed grey lines indicate the average reward for a single agent without any social information (lower line) and the average reward for an agent that always chooses the option with greater utility (upper line); (b) Mean payoffs increase with connectivity up to an optimal value and then fall, with a differing optimal connectivity for different group sizes; (c) The optimal mean degree generating maximum payoffs varies proportionally to the square root of network size and (d) The optimal connectivity varies inversely with the square root of network size.

individuals is likely to lead to individuals making poorer decisions on average (see Figure 4(c)), potentially creating additional difficulties for populations already under pressure. There is evidence that in some cases animals may seek to compensate for lost social bonds by creating new connections (Firth et al., 2017), but this may depend on whether populations are adapted to significant changes in population size over time. These results suggest therefore that, where populations are in decline, maintaining the physical cohesion and density of animal groups, so that social connectivity remains at appropriate levels, should be an important conservation goal alongside wider efforts to promote landscape connectivity (Doherty and Driscoll, 2018; Rudnick et al., 2012).

In human societies, a more salient concern is the efficacy of decision making by individuals exposed to recent increases in social connectivity (Bak-Coleman et al., 2021), most obviously through social media (Kietzmann et al., 2011), but also through the ubiquity on online recommendation and reputation systems (Dellarocas, 2003). These factors serve to increase the number of choices, decisions and opinions individuals can observe others making or expressing. The rapidity with which this social environment has changed for many people, and the time required to adapt to novel social contexts (Burton-Chellew and West, 2021), makes it highly plausible that they will continue to operate with heuristics that are still adapted to lower levels of connectivity. While some increase in

connectivity may be beneficial, large increases have the potential to reduce the quality of decision making by all participants by introducing a high degree of social feedback that swamps the true quality signals from the environment rather than amplifying them, such that collective decisions become highly contingent on variable early decision-makers (Muchnik et al., 2013; Salganik et al., 2006). A number of experimental studies have revealed reductions in collective wisdom when individuals are exposed to the complete or aggregated opinions of other group members (Frey and Armut 2020; Lorenz et al., 2011), but others have shown improvements under treatments where connectivity is limited (Becker et al., 2017; Jayles et al., 2021; Navajas et al., 2018; Kao and Couzin, 2014; Kao and Couzin, 2019). These apparently contrary findings may be reconciled by recognising that some additional communication between agents is beneficial, but that too much can be harmful when the opinions or decisions of agents become highly correlated in a manner not recognised or accounted for by another agent observing them.

The potential endogenous reduction in decision-making efficacy identified above has political implications in an age of ubiquitous online connectivity, but should also be a source of concern in an increasingly-connected scientific community (Mann and Woolley-Meza, 2017). Societies that have recently experienced large increases in connectivity may also be more vulnerable to exogenous misinformation (e.g. Spaiser et al., 2017), as false information can be more easily amplified throughout the population. In social networks that are naturally highly connected, such as small friendship groups individuals may intuit the correlations arising from those connections and apply appropriate decision rules (Mann, 2018, 2020, 2021). However, such correlations may now be induced in wider networks where they were previously absent as a result of increased connectivity, with the information nonetheless appearing to come from independent sources to an agent that has not adjusted to the new social context. It is these cases where habituation and expectations deviate from reality that are most ripe for the spread of ‘viral’ opinions. This should motivate efforts not only to identify and remove obvious misinformation online, but also to educate citizens in how to respond to social information in a highly connected world (Badrinathan, 2021; Bak-Coleman et al., 2022).

Most real-world networks are neither fully connected nor accurately described by an Erdős-Renyi network (one in which all possible social connections occur fixed probability). Animal groups on the move are generally considered to be characterised by short-range interactions between spatial neighbours (Couzin et al., 2002), while social networks in both humans and animals typically exhibit substantial structure such as clusters (a high probability that people share friends) and hierarchy (such as in a pecking order). As such, the behaviour of real social groups may

interpolate between that of the sparsely connected random network considered here, and the behaviour predicted in fully connected networks (Mann, 2018, 2020). For example, individuals may treat their local network (such as close friends or spatial neighbours) with social behaviour appropriate to a fully connected network, while treating more dispersed social connections, such as passing conversations with strangers, or online comments, in a manner appropriate to an Erdős-Renyi network. As indicated above, it is this second case where the effect of increasing social connectivity is likely to be observed. It is also plausible that specific network structures, such as small world networks, may give rise to behaviour that is not predicted by either of these scenarios. Future work should therefore consider how to extend the analysis in this paper to a wider class of social interaction networks.

Methods

Correlation between agent preferences

The decisions of other individuals are useful information insofar as they are assumed to share preferences with the focal agent. Following Mann (2020), I encode the degree to which individuals’ preferences align via a joint multivariate normal distribution $N(\cdot, \cdot, \cdot)$

$$p(x_1, x_2, \dots, x_n) = N(\mathbf{x}, \mathbf{0}, \Sigma). \quad (11)$$

where $\mathbf{X} = [x_1, \dots, x_n]^\top$, $\mathbf{0}$ is vector of n zeroes, and

$$\Sigma = \begin{bmatrix} 1 & \rho & \dots & \rho \\ \rho & \ddots & & \vdots \\ \vdots & & & \rho \\ \rho & \dots & \rho & 1 \end{bmatrix},$$

is the covariance matrix, with ρ being the correlation between the utilities of any pair of agents. Combined with equation (3), this further specifies a joint multivariate normal distribution over the true values of x_1, \dots, x_n and the observed private information of each individual $\Delta_1, \dots, \Delta_n$, where

$$\begin{aligned} \text{cov}(x_k, x_l) &= \text{cov}(\Delta_k, x_l) = \rho + \delta_{k,l}(1 - \rho) \\ \text{cov}(\Delta_k, \Delta_l) &= \rho + \delta_{k,l}(1 + \epsilon^2 - \rho), \end{aligned} \quad (12)$$

where $\delta_{k,l}$ is the Kronecker delta function. Throughout this paper, I assume that agents know or are habituated to the true value of ρ , and that it is this value that they base their inferences on.

At this point it is useful to introduce a new variable z , which represents the mean value of x for all agents. The reason for doing so is that the various values of x_i for all agents can be described as being distributed independently, conditioned on the value of z , which thus contains all the

correlations between agent preferences. One can then re-write equation (11) as

$$p(x_i | z) \propto \phi\left(\frac{x_i - z}{\sqrt{1 - \rho}}\right) \quad (13)$$

$$p(z) \propto \phi\left(\frac{z}{\sqrt{\rho}}\right). \quad (14)$$

Bayes' rule then provides an expression for the distribution of z , conditioned on x_i

$$p(z | x_i) \propto \phi\left(\frac{z - \rho x_i}{\sqrt{\rho - \rho^2}}\right) \quad (15)$$

Using this formulation, one can then write down the joint probability for a set of agents $j \neq i$ to make private observations $\{\Delta_j\}$, conditioned on the value of x_i

$$\begin{aligned} p(\{\Delta_j\} | x_i) &\propto \int_{-\infty}^{\infty} \prod_{j \neq i} \phi\left(\frac{z - \Delta_j}{\sqrt{1 + \epsilon^2 - \rho}}\right) p(z | x_i) dz \\ &\propto \int_{-\infty}^{\infty} \prod_{j \neq i} \phi\left(\frac{z - \Delta_j}{\sqrt{1 + \epsilon^2 - \rho}}\right) \phi\left(\frac{z - \rho x_i}{\sqrt{\rho - \rho^2}}\right) dz, \end{aligned} \quad (16)$$

where the product form inside the integral relies on the independence of the agents' private information, conditioned on z . Note that for the special case of $\rho = 1$, $x_i = z \forall i$ and the above expression simplifies to

$$p(\{\Delta_j\} | x_i) \propto \prod_{j \neq i} \phi\left(\frac{x_i - \Delta_j}{\epsilon}\right). \quad (17)$$

Form of social information strategy

Based on the methodological developments in previous studies (Mann, 2018, 2020, 2021), a social information strategy associates a real number, Δ_s^* with each distinct observable state of social information \mathbf{s} . This number represents a threshold, such that if an agent receives private information $\Delta > \Delta_s^*$ the agent will choose option A, otherwise it will choose option B.

In an optimal strategy, the thresholds Δ_s^* are identified as those that cause the agent to make rational decisions, that is, to choose the option that maximises its expected utility, conditioned on both the private and social information it has available. This implies that when an agent observes private information exactly equal to the threshold value, this makes the expected utility difference between the two options (x_i) equal to zero

$$\begin{aligned} 0 &= \mathbb{E}(x_i | \mathbf{s}, \Delta_i = \Delta_s^*) \\ &\propto \int_{-\infty}^{\infty} x_i p(x_i) p(\Delta_i = \Delta_s^* | x_i) p(\mathbf{s} | x_i) dx_i \end{aligned} \quad (18)$$

In this paper, the distinct observable states of social information are observed sequences of decisions, $\mathbf{s} = K_1, \dots, K_k$, which are themselves random subsets of the true full sequence of decisions $\mathbf{c} = C_1, C_2, \dots, C_m$, such that $k \leq m$. Given an observed decision sequence \mathbf{s} , equation (18) can be expanded as

$$\begin{aligned} 0 &= E(x_i | \mathbf{s}, \Delta_i = \Delta_s^*) \\ &\propto \int_{-\infty}^{\infty} x_i p(x_i) p(\Delta_i = \Delta_s^* | x_i) \sum_{\mathbf{c} \in \mathbf{S}} P(\mathbf{s} | \mathbf{c}) P(\mathbf{c} | x_i) dx_i \end{aligned} \quad (19)$$

where the summation is over the set \mathbf{S} of all possible sequences of $n - 1$ decisions that could have occurred prior to the focal agents decision.

Observation probability of a decision sequence

Given a true full sequence of decisions, $\mathbf{c} = C_1, C_2, \dots, C_m$ and a connectivity parameter q (the probability to observe any given prior decision), the probability that the focal agent will observe the sequence $\mathbf{s} = K_1, \dots, K_k$ is given by

$$P(\mathbf{s} | \mathbf{c}) = q^k (1 - q)^{m-k} C(\mathbf{s}, \mathbf{c}) \quad (20)$$

where $C(\mathbf{s}, \mathbf{c})$ is the number of distinct decimations of \mathbf{c} that can result in \mathbf{s} . To give an example, if the original sequence of decisions is $\mathbf{c} = 1, 1, -1, 1, -1$, the observed sequence $\mathbf{s} = 1, 1$ can be generated from this in three distinct ways: by selecting elements 1 and 2, elements 1 and 4 or elements 2 and 4. In contrast the observed sequence $\mathbf{s}' = -1, -1$ can only be generated in one way, by selecting elements 3 and 5. Hence, in this example, $C(\mathbf{s}, \mathbf{c}) = 3$ and $C(\mathbf{s}', \mathbf{c}) = 1$. By extension then, $P(\mathbf{s} | \mathbf{c}) = 3q^2(1 - q)^3$ and $P(\mathbf{s}' | \mathbf{c}) = q^2(1 - q)^3$.

Generating probability of a decision sequence

The summation in equation (19) requires calculating the probability $P(\mathbf{c} | x_i)$ to generate any possible sequence of decisions, conditioned on a specific true value of x_i . This quantity can be evaluated by considering the various contiguous sub-sequences of \mathbf{c} : $\mathbf{c}_k = C_1, \dots, C_k$. Since the form of the social information strategy specified in equation (19) specifies a threshold for each of these sub-sequences, the probability of the full sequence is recursively defined by product of the probability for each decision within that sequence, conditioned on those thresholds and the preceding sequence of decisions

$$P(\mathbf{c} | x_i) = \prod_{k=1}^{|\mathbf{c}|} \sum_{\mathbf{s} \in S} P(C_k | x_i, \mathbf{s}, \mathbf{c}_{k-1}) P(\mathbf{s} | \mathbf{c}_{k-1}), \quad (21)$$

where the summation is over all the sequences $\mathbf{s} \in S$ that the agent may have observed as a result of the true sequence \mathbf{c} , and $|\mathbf{c}|$ is the length (number of decisions) of the sequence \mathbf{c} . The above expression can be further expanded by expressing the dependence on x_i through the average utility difference z , as in equation (16)

$$P(\mathbf{c} | x_i) = \int_{-\infty}^{\infty} \prod_{k=1}^{|\mathbf{c}|} \sum_{\mathbf{s} \in S} P(C_k | z, \mathbf{s}, \mathbf{c}_{k-1}) P(\mathbf{s} | \mathbf{c}_{k-1}) \times p(z | x_i) dz, \quad (22)$$

where $p(z | x_i) \propto \phi(z - \rho x_i / \sqrt{\rho - \rho^2})$. This expansion is useful because the values of the private information observed by each agent are independent conditioned on z . As such, $P(C_k | z, \mathbf{s}, \mathbf{c}_{k-1}) = P(C_k | z, \mathbf{s})$ and, with reference to equation (16), equation (22) can be rewritten as

$$P(\mathbf{c} | x_i) = \int_{-\infty}^{\infty} \prod_{k=1}^{|\mathbf{c}|} \sum_{\mathbf{s} \in S} \Phi\left(\frac{C_k(z - \Delta_s^*)}{\sqrt{1 + \epsilon^2 - \rho}}\right) P(\mathbf{s} | \mathbf{c}_{k-1}) \times p(z | x_i) dz, \quad (23)$$

where $\Phi(\cdot)$ is the cumulative normal distribution function. In this form the product now involves terms that depend only on the sequence \mathbf{s} observed by the agent and not on the full previous sequence \mathbf{c}_{k-1} .

Iterative calculation of thresholds

The formulation above poses a problem, since determining the value of the threshold Δ_s^* in equation (19) requires knowing the thresholds associated with all sequences in S , including Δ_s^* itself. This suggests an iterative approach. First, all thresholds are set to an initial value of zero: $\Delta_s^* = 0$ for all possible social information \mathbf{s} . I then used these initial values to calculate the sequence-generation probabilities $P(\mathbf{c} | x_i)$ (for any given agent identity i), by applying equation (23). Each threshold is then updated by solving equation (19) for each Δ_s^* , based on these sequence probabilities. These steps are then repeated, iterating until the thresholds reach a stable value that provides a self-consistent solution to equation (19) for all sequences.

Monte Carlo methods for large groups

On the basis of the results from small groups, I assume that in sparsely connected groups agents employ a social information strategy that depends on the difference between the number of agents they observe to have chosen

A and B. Thus, this strategy associates a threshold Δ_w^* with every value $w = n_A - n_B$ that the agent may observe as the difference between the observed number of decisions for A (n_A) and the observed number of decisions for B (n_B)

Given a true sequence of decisions \mathbf{s} that contains N_A agents choosing option A and N_B choosing B, the probability that the focal agent observes n_A and n_B decisions for each option is given by two independent binomial distributions

$$\begin{aligned} P(n_A | N_A) &= \binom{N_A}{n_A} q^{n_A} (1-q)^{(N_A-n_A)} \\ P(n_B | N_B) &= \binom{N_B}{n_B} q^{n_B} (1-q)^{(N_B-n_B)}. \end{aligned} \quad (24)$$

The resulting distribution for the difference $w = n_A - n_B$ observed by the focal agent has no closed form, but can be evaluated numerically by summation over the values of n_A and n_B

$$P(w | N_A, N_B) = \sum_{n_A=1}^{N_A} \sum_{n_B=1}^{N_B} P(n_A | N_A) P(n_B | N_B) \delta_{(n_A-n_B), w} \quad (25)$$

where $\delta_{(n_A-n_B), w}$ is the Kronecker delta function.

In large groups, exhaustively calculating the probability to generate every possible sequence of decisions is infeasible. Instead, I employ a Monte Carlo methodology. As in the case of small groups, I initialise all thresholds to zero and based on these thresholds I simulate a number Λ of candidate decision sequences, with random lengths drawn uniformly between 0 and $n - 1$ (since the focal agent, observing only the difference w , does not attend to how many previous decisions have been made). The average utility difference for all agents, z , is drawn from the distribution given in equation (14), and the actual utility differences for both the focal agent (x_i) and all other agents in the sequence are then drawn from the distribution specified in equation (13). Extracting the values of N_A and N_B from the generated sequences, and retaining the samples of x_i used to generate them, these sequences permit the following Monte Carlo approximation to the expected value of x_i , conditioned on the observable social information w

$$\begin{aligned} \mathbb{E}(x_i | \Delta_i, w) &\propto \int_{-\infty}^{\infty} x_i p(x_i) p(\Delta_i | x_i) P(w | x_i) dx_i \\ &\simeq (1/\Lambda) \sum_{\lambda=1}^{\Lambda} x_{i\lambda} \phi((x_{i\lambda} - \Delta_i)/\epsilon) P(w | N_{A,\lambda}, N_{B,\lambda}), \end{aligned} \quad (26)$$

where λ indexes the Λ Monte Carlo samples, and $P(w|N_{A,\lambda}, N_{B,\lambda})$ is given by equation (25). Using this approximation, one can calculate a value for the threshold $\Delta_w^{\text{proposal}}$ such that

$$\sum_{\lambda=1}^{\Lambda} x_{\lambda} \phi((x_{\lambda} - \Delta_w^{\text{proposal}}) / \epsilon) P(w | N_{A,\lambda}, N_{B,\lambda}) = 0. \quad (27)$$

This value of the threshold, which represents the rational strategy conditioned on all other agents continuing to apply the previously initialised threshold values, acts as a proposed value for the update step. To improve convergence I introduce a learning rate α , such that the updated threshold ($\Delta_{w,i+1}^*$) is a weighted average of the previous value ($\Delta_{w,i}^*$) and the proposal defined above

$$\Delta_{w,i+1}^* = \alpha \Delta_w^{\text{proposal}} + (1 - \alpha) \Delta_{w,i}^*. \quad (28)$$

In my analyses, I use $\Lambda = 1000$ Monte Carlo samples, and $\alpha = 0.1$ with 100 iterations to reliably ensure convergence to a stable set of threshold values.

Expected payoff

I calculated the expected payoff for an individual as follows: for a given value of utility difference (x – equal for all agents when $\rho = 1$) I evaluated the probability that a given number of agents would choose the option with greater utility, based on equation (23) for the probability of a given sequence of decisions and averaging over possible sequences. Agents choosing the higher reward option are assigned a payoff of x , such that the expected reward for a randomly-chosen agent is x multiplied by the expected proportion of agents selecting the better option. I then integrated over possible values of x according to the assumed distribution of rewards in the environment, which corresponds to the agents' prior belief about x given in equation (2).

Acknowledgements

I would like to thank Joseph Bak-Coleman and Albert Kao for helpful discussions and advice on preparing the manuscript.

Declaration of conflicting interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship and/or publication of this article.

Funding

The author(s) disclosed receipt of the following financial support for the research, authorship and/or publication of this article: This work was supported by UK Research and Innovation Future Leaders Fellowship MR/S032525/1.

ORCID iD

Richard P. Mann  <https://orcid.org/0000-0003-0701-1274>

References

- Acemoglu D, Dahleh MA, Lobel I, et al. (2011) Bayesian learning in social networks. *The Review of Economic Studies* 78: 1201–1236.
- Anderson LR and Holt CA (1997) Information Cascades in the Laboratory. *The American Economic Review* 847–862.
- Apicella CL, Marlowe FW, Fowler JH, et al. (2012) Social networks and cooperation in hunter-gatherers. *Nature* 481: 7382–7501.
- Arganda S, Pérez-Escudero A and De Polavieja GG (2012) A common rule for decision-making in animal collectives across species. *Proceedings of the National Academy of Sciences* 109: 20508–20513.
- Aumann RJ (1976) Agreeing to disagree. *The Annals of Statistics* 4: 1236–1239.
- Badrinathan S (2021) Educative Interventions to Combat Misinformation: Evidence from a Field Experiment in India. *American Political Science Review* 115(4), 1325–1341.
- Bak-Coleman JB, Alfano M, Barfuss W, et al. (2021) Stewardship of global collective behavior. *Proceedings of the National Academy of Sciences* 118: e2025764118.
- Bak-Coleman JB, Kennedy I, Morgan W, et al. (2022) Combining Interventions to Reduce the Spread of Viral Misinformation. *Nature Human Behaviour*.
- Becker J, Brackbill D and Centola D (2017) Network dynamics of social influence in the wisdom of crowds. *Proceedings of the National Academy of Sciences* 114: E5070–E5076.
- Bikhchandani S, Hirshleifer D and Welch I (1992) A theory of fads, fashion, custom, and cultural change as informational cascades. *Journal of Political Economy* 100(5): 992–1026.
- Bikhchandani S, Hirshleifer D, Tamuz O, et al. (2021) *Information Cascades and Social Learning*. National Bureau of Economic Research. Technical Report.
- Burton-Chellew MN and West SA (2021) Payoff-based learning best explains the rate of decline in cooperation across 237 public-goods games. *Nature Human Behaviour* 5: 1330–1338.
- Couzin ID, Krause J, James R, et al. (2002) Collective memory and spatial sorting in animal groups. *Journal of Theoretical Biology* 218: 1–11.
- DeGroot MH (1974) Reaching a Consensus. *Journal of the American Statistical Association*, 69: 118–121.
- Dellarocas C (2003) The digitization of word of mouth: Promise and challenges of online feedback mechanisms. *Management Science* 49: 1407–1424.
- Doherty TS and Driscoll DA (2018) Coupling movement and landscape ecology for animal conservation in production landscapes. *Proceedings of the Royal Society B: Biological Sciences* 285: 20172272.
- Dunbar RIM (1998) The social brain hypothesis. *Evolutionary Anthropology: Issues, News, and Reviews: Issues, News, and Reviews* 6: 178–190.

- Eguiluz M, Masuda N and Fernández-Gracia J (2015) Bayesian decision making in human collectives with binary choices. *PLoS ONE* 10: e0121332.
- Eisenstein EL (1980) *The printing press as an agent of change*. vol. 1. Cambridge, UK: Cambridge University Press.
- Farine DR, Aplin LM, Garroway CJ, et al. (2014) Collective decision making and social interaction rules in mixed-species flocks of songbirds. *Animal Behaviour*. 2014 95: 173–182.
- Firth JA, Voelkl B, Crates RA, et al. (2017) Wild birds respond to flockmate loss by increasing their social network associations to others. *Proceedings of the Royal Society B: Biological Sciences* 284: 20170299.
- Fisher DN, Kilgour RJ, R Siracusa E, et al. (2021) Anticipated effects of abiotic environmental change on intraspecific social interactions. *Biological Reviews* 96: 2661–2693.
- Frey V and Arnout VDR (2020) Social influence undermines the wisdom of the crowd in sequential decision making. *Management Science* 67: 4273–4286.
- Gerardo C, Ehrlich PR and Raven PH (2020) Vertebrates on the brink as indicators of biological annihilation and the sixth mass extinction. *Proceedings of the National Academy of Sciences* 117: 13596–13602.
- Holley RA and Liggett TM (1975) Ergodic theorems for weakly interacting infinite systems and the voter model. *The Annals of Probability* 3: 643–663.
- Jayles B, Sire C and Kurvers RHJM (2021) Impact of sharing full versus averaged social information on social influence and estimation accuracy. *Journal of the Royal Society Interface* 18: 20210231.
- Kadak K and Miller N (2020) *Follow the Straggler: Zebrafish Use a Simple Heuristic for Collective Decision-Making*. *Proceedings of the Royal Society B: Biological Sciences*.
- Kao AB and Couzin ID (2014) Decision accuracy in complex environments is often maximized by small group sizes. *Proceedings of the Royal Society B: Biological Sciences* 281: 20133305.
- Kao AB and Couzin ID (2019) Modular structure within groups causes information loss but can improve decision accuracy. *Philosophical Transactions of the Royal Society B* 374: 20180378.
- Karamched B, Stickler M, Ott W, et al. (2020) Heterogeneity improves speed and accuracy in social networks. *Physical Review Letters* 125: 218302.
- Kietzmann JH, Hermkens K, McCarthy IP, et al. (2011) Social media? Get serious! Understanding the functional building blocks of social media. *Business Horizons* 54(3): 241–251.
- Krause J and Ruxton GD (2002) *Living in Groups*. Oxford, UK: Oxford University Press.
- Lorenz J, Rauhut H, Schweitzer F, et al. (2011) How social influence can undermine the wisdom of crowd effect. *Proceedings of the National Academy of Sciences* 108: 9020–9025.
- Mann RP and Woolley-Meza O (2017) Maintaining intellectual diversity in data science. *Data Science* 1: 85–94.
- Mann RP (2018) Collective decision making by rational individuals. *Proceedings of the National Academy of Sciences* 115: E10387–E10396.
- Mann RP (2020) Collective decision-making by rational agents with differing preferences. *Proceedings of the National Academy of Sciences* 117: 10388–10396.
- Mann RP (2021) Optimal use of simplified social information in sequential decision-making. *Journal of the Royal Society Interface* 18: 20210082.
- McQuail D (1977) *The influence and effects of the mass media*, in Curran J, Gurevitch M and Woollacott J (eds) *Mass Communication and Society*, Edward Arnold.
- Miller N, Garnier S, Hartnett AT, et al. (2013) Both information and social cohesion determine collective decisions in animal groups. *Proceedings of the National Academy of Sciences* 110: 5263–5268.
- Mossel E, Allan S and Tamuz O (2015) Strategic learning and the topology of social networks. *Econometrica* 83: 1755–1794.
- Muchnik L, Aral S and Taylor SJ (2013) Social influence bias: A randomized experiment. *Science* 341: 6146–6651.
- Navajas J, Niella T, Garbulsky G, et al. (2018) Aggregated knowledge from a small number of debates outperforms the wisdom of large crowds. *Nature Human Behaviour* 2: 126–132.
- Neumann JV and Morgenstern O (2007) *Theory of Games and Economic Behavior*. Princeton, NJ: Princeton University Press.
- Pérez-Escudero A and De Polavieja GG (2011) Collective Animal Behavior from Bayesian Estimation and Probability Matching. *Plos Computational Biology* 7: e1002282.
- Pérez-Escudero A and de Polavieja GG (2017) Adversity magnifies the importance of social information in decision-making. *Journal of The Royal Society Interface* 14: 20170748.
- Poel W, Winklmayr C and Romanczuk P (2021) Spatial structure and information transfer in visual networks. *Frontiers in Physics* 623.
- Psorakis I, Voelkl B, J Garroway C, et al. (2015) Inferring social structure from temporal data. *Behavioral Ecology and Sociobiology* 69: 857–866.
- Rosenthal SB, Twomey CR, Hartnett AT, et al. (2015) Revealing the hidden networks of interaction in mobile animal groups allows prediction of complex behavioral contagion. *Proceedings of the National Academy of Sciences* 112: 4690–4695.
- Rudnick D, Ryan SJ, Paul B, et al. (2012) The role of landscape connectivity in planning and implementing conservation and restoration priorities. *Issues in Ecology* 16: 1–20.
- Salganik MJ, Dodds PS and Watts DJ (2006) Experimental study of inequality and unpredictability in an artificial cultural market. *Science* 311: 576–856.
- Spaiser V, Chadeaux T, Donnay K, et al. (2017) *Communication Power Struggles on Social Media: A Case Study of the 2011-12*

- Russian Protests. Journal of Information Technology & Politics* 14: 132–153.
- Strandburg-Peshkin A, Twomey CR, Bode NWF, et al. (2013) Visual sensory networks and effective information transfer in animal groups. *Current Biology* 23: R709–R711.
- Wagner DL, Grames EM, Forister ML, et al. (2021) Insect decline in the Anthropocene: Death by a thousand cuts. *Proceedings of the National Academy of Sciences* 118: e2023989118.
- Ward AJW, Herbert-Read JE, Sumpter DJT, et al. (2011) Fast and accurate decisions through collective vigilance in fish shoals. *Proceedings of the National Academy of Sciences* 108: 2312–2315.
- Wolf M, Kurvers RHJM, Ward AJW, et al. (2013) Accurate decisions in an uncertain world: collective cognition increases true positives while decreasing false positives. *Proceedings of the Royal Society B: Biological Sciences* 280: 20122777.