Effects of Conformism on the Cultural Evolution of Social Behaviour

Lucas Molleman1*, Ido Pen2, Franz J. Weissing3

1 PhD student, Theoretical Biology Group, Centre for Ecological and Evolutionary Studies, University of Groningen, Groningen, The Netherlands, 2 Professor of Theoretical Biology, Theoretical Biology Group, Centre for Ecological and Evolutionary Studies, University of Groningen, Groningen, The Netherlands, 3 Professor of Theoretical Biology, Theoretical Biology Group, Centre for Ecological and Evolutionary Studies, University of Groningen, Groningen, The Netherlands

Abstract

Models of cultural evolution study how the distribution of cultural traits changes over time. The dynamics of cultural evolution strongly depend on the way these traits are transmitted between individuals by social learning. Two prominent forms of social learning are payoff-based learning (imitating others that have higher payoffs) and conformist learning (imitating locally common behaviours). How payoff-based and conformist learning affect the cultural evolution of cooperation is currently a matter of lively debate, but few studies systematically analyse the interplay of these forms of social learning. Here we perform such a study by investigating how the interaction of payoff-based and conformist learning affects the outcome of cultural evolution in three social contexts. First, we develop a simple argument that provides insights into how the outcome of cultural evolution will change when more and more conformist learning is added to payoff-based learning. In a social dilemma (e.g. a Prisoner’s Dilemma), conformism can turn cooperation into a stable equilibrium; in an evasion game (e.g. a Hawk-Dove game or a Snowdrift game) conformism tends to destabilize the polymorphic equilibrium; and in a coordination game (e.g. a Stag Hunt game), conformism changes the basin of attraction of the two equilibria. Second, we analyse a stochastic event-based model, revealing that conformism increases the speed of cultural evolution towards pure equilibria. Individual-based simulations as well as the analysis of the diffusion approximation of the stochastic model by and large confirm our findings. Third, we investigate the effect of an increasing degree of conformism on cultural group selection in a group-structured population. We conclude that, in contrast to statements in the literature, conformism hinders rather than promotes the evolution of cooperation.

Introduction

Social learning enables humans to survive in a broad array of different habitats across the planet. By learning from their peers, individuals can rapidly acquire adaptive information about which behaviour is optimal under a variety of environmental conditions. Models of cultural evolution use insights from theories of genetic evolution to study how cultural variants, such as ideas and beliefs, spread through populations of individuals by social learning. Social learning based on imitating the behaviour of successful individuals can lead to an evolutionary dynamic similar to the spread of alleles under natural selection, whereas learning by adopting behaviours from others more randomly leads to a process resembling genetic drift.

Models of cultural evolution have to be adapted to the specific mechanisms by which cultural traits transmit between individuals. Traits can be transmitted not only vertically from parents to offspring, but in a range of different ways. For instance, traditional hunters may learn from their parents a social norm to share hunting revenues, and may learn the optimal design of an arrow from their fellow hunters. How humans learn from each other is a topic of extensive theoretical and empirical research (for a recent overview see [1]), and various specific forms of social learning (termed ‘social learning strategies’ [2] or ‘learning biases’ [3]) have been studied as to how they affect the spread of cultural traits through populations. Two forms of social learning received particular attention: conformism and payoff-based learning.

When individuals can evaluate the payoffs of the behavioural strategies of others, the preferential imitation of high-payoff individuals can lead to the rapid spread of adaptive behaviours in a population [4]. However, such payoff-based learning is not always feasible. Getting insights in the payoffs received by others is not always straightforward, especially for newcomers in a population. In cases like this, imitating the majority (conformism) can be a good alternative form of social learning, in particular if the success of cultural traits strongly depends on the local circumstances [3,5].

The role of conformism in cultural evolution has recently become the matter of considerable debate. In the context of a social dilemma, payoff-based learning will tend to inhibit the spread of cooperation because defectors obtain higher payoffs by reaping the benefits of cooperation without paying the costs. Theory suggest that when payoff-based learning is complemented by other forms of social learning, the dynamics of cultural evolution can be strongly affected. For instance, adding random learning to payoff-based learning can facilitate the rapid solution of coordination problems [6], and conformism can stabilise...
cooperative equilibria under specific conditions [3,7–14]. Moreover, conformism can homogenise groups internally, thereby augmenting the relative amount of variation between groups [14]. This decreases the scope for selection within groups (i.e. payoff-based transmission disfavouring cooperation), and increases the potential role of ‘cultural group selection’. In group-structured populations, cooperation can spread when groups of cooperators have some advantage over groups of defectors. This advantage can manifest itself in a number of different ways; cooperative groups may send out more migrants, grow to larger sizes, or replace other groups (e.g. [8,15–17]). Selection at the group level may also occur when individuals occasionally learn from members of other groups that perform better[4,19–20]. Through such a process, cooperation can be promoted since individuals in cooperative groups have higher payoffs than individuals in groups of non-cooperators.

Experiments from psychology and behavioural economics suggest that humans indeed use both conformist and payoff-based learning in determining their behaviour. When individuals are allowed to use both conformist and payoff-based learning, experimental evidence suggests that cultural traits can spread through a mixture of these two forms of social learning [23,25,26]. This raises the question of how the interplay of conformist and payoff-based learning affects the spread of cultural traits through a population.

First, we develop a simple argument to delineate how the direction of cultural change is affected by the relative rate of conformist and payoff-based learning. This will give us an intuitive insight in the effects of conformism in contexts of social interaction. Second, we construct stochastic models that allow us to follow the spread of culturally transmitted behaviours in the course of time in a finite population. These models allow us to quantify how the relative degree of conformism (as opposed to payoff-based learning) affects the success of social strategies in reaching fixation. Third, we examine how cooperation can spread in a group-structured population by means of cultural group selection. With this model, we investigate whether conformism tends to promote – as often claimed in the literature – or hamper the spread of cooperation in populations that are structured into groups of finite size.

**Analysis and Results**

1. **Model Structure**

   We consider a population in which individuals are involved in social interactions. Individuals have a culturally acquired strategy that determines their behaviour in these interactions. We consider two variants of this behaviour (A or B). An individual has the inclination of playing either A or B, but this inclination can change over the course of time due to social learning. Learning is either based on payoffs (individuals tend to imitate successful individuals) or on conformism (individuals tend to imitate the majority of the population). The relative frequency γ of these two forms of social learning is the key parameter of interest. The value of γ ranges from 0 to 1. If γ = 0, all learning is payoff-based; if γ = 1, all learning is based on conformism. If 0 < γ < 1, individuals use a mixture of these two forms of social learning. We assume that all individuals use the same mixture of conformist and payoff-based learning.

   Individuals acquire payoffs by social interaction with others in their group. Payoffs depend linearly on the frequency p of A-individuals and 1−p of B-individuals, and are calculated using the payoff matrix \( G = \begin{pmatrix} a & b \\ c & d \end{pmatrix} \). The payoff of A-individuals is \( \pi_A = p\cdot a + (1-p)\cdot b \)

and the payoff of B-individuals is \( \pi_B = p\cdot c + (1-p)\cdot d. \)

There are three strategically different classes of games with two pure strategies, and we consider the evolutionary dynamics in each of these ‘interaction contexts’. In the first class of games, one of the pure strategies (say B) is dominant over the other: \( a < c \) and \( b < d \). In the special case where \( a < d < c < b \), this is a *social dilemma*. Collective interests are opposed to individual interests: when all individuals exhibit behaviour A (‘cooperate’), payoffs are higher than when all individuals exhibit behaviour B (‘defect’). Individually, however, B yields higher payoffs than A, irrespective of what others are doing. Second, we consider the class of *coordination games*, which are characterized by \( a > c \) and \( b < d \). In this class, the payoff of a pure strategy increases with the number of individuals using this strategy. In a coordination game, both pure strategies are Nash equilibrium strategies. In addition, there is a (dynamically unstable) mixed-strategy equilibrium at

\[
 p^* = \frac{d-b}{a-b-c+d}. \tag{2}
\]

Third, we consider the class of *evasion games*, where \( a < c \) and \( b > d \). Now the relative payoff of each pure strategy decreases with the frequency of this strategy in the population. In an evasion game, none of the pure strategies is a Nash equilibrium; instead an evasion game has a unique mixed-strategy equilibrium, which is given by eq. (2). The Hawk-Dove game and the Snowdrift game are prominent examples of evasion games.

2. **Representation of Conformism by a Coordination Game**

   Before studying the dynamics of cultural evolution in finite populations, we aim to get some intuition on how conformism might change the direction of cultural evolution. To this end, we represent conformism-based learning by payoff-based learning in the coordination game with payoff matrix \( K = \begin{pmatrix} s & 0 \\ 0 & s \end{pmatrix} \). In this coordination game, the payoff to each pure strategy is proportional to its frequency in the population: \( \pi_A = p\cdot s \) and \( \pi_B = (1-p)\cdot s \), where s is positive. Accordingly, the pure strategy with highest frequency is favoured by payoff-based learning, just as it is in case of conformism-based learning. Based on these considerations, a mixture of payoff-based learning (characterised by matrix G) and conformist learning (characterised by matrix K) can be described by the combined matrix

\[
 M(\gamma) = (1-\gamma) G + \gamma K = \begin{pmatrix} (1-\gamma)a + \gamma s & (1-\gamma)b \\ (1-\gamma)c & (1-\gamma)d + \gamma s \end{pmatrix} \tag{3}
\]

It is now straightforward to characterise the expected direction of cultural change as a function of our key parameter \( \gamma \) by determining the Nash equilibrium strategies of the matrix game \( M(\gamma) \). This can be done with standard methods [27]: Pure strategy A is a Nash equilibrium if \( (1-\gamma)a + \gamma s > (1-\gamma)c \) or equivalently...
\( \frac{\gamma}{1-\gamma} s > c - a \). Similarly, pure strategy \( B \) is a Nash equilibrium when \( \frac{\gamma}{1-\gamma} s > b - d \). Both inequalities are more easily satisfied for larger values of \( s \) or \( \gamma \) and will always hold if \( \gamma \) approaches 1. When both inequalities are reversed, \( M(\gamma) \) has a (dynamically stable) mixed equilibrium at

\[
p^* = \frac{(1-\gamma)(d-b) + \gamma s}{(1-\gamma)(a-b+c+d) + 2\gamma s}.
\]

(4)

For each of the three interaction contexts, Figure 1 illustrates how the dynamics of cultural evolution (increase or decrease in the frequency of pure strategy \( A \)) changes with the frequency of conformist learning \( \gamma \). First, consider the extreme case \( \gamma = 0 \), at the bottom of the three panels of Figure 1. Here, all learning occurs on the basis of the payoffs in matrix \( G \). In the social dilemma, \( A \) (cooperate) is disfavoured by payoff-based learning, and cultural evolution will lead to a decrease of the frequency of \( A \) (Figure 1A, bottom arrow to the left) and convergence to the sole Nash equilibrium \( p^* = 0 \). The coordination game has two pure-strategy Nash equilibria \( (p^* = 0 \text{ and } p^* = 1) \) that are separated by the dynamically unstable mixed-strategy Nash equilibrium (eq. 2). The two arrows at the bottom of Figure 1B indicate that cultural evolution will either lead to the fixation of \( A \) or to the fixation of \( B \), and that the outcome depends on initial conditions. In the evasion game, the arrows at the bottom of Figure 1C indicate that the system will converge to the mixed-strategy Nash equilibrium (eq. 2), where \( A \) and \( B \) stably coexist. Next consider the other extreme \( \gamma = 1 \), where all learning is conformism based, i.e. governed by matrix \( K \) (top of the three panels in Figure 1). Now the expected direction of change is identical for each of the three interaction contexts: since \( K \) is a coordination game, the two pure strategies are Nash equilibria and cultural evolution will either lead to the fixation of \( A \) or to the fixation of \( B \), depending on initial conditions (top arrows in all three panels). Due to conformist learning, the strategy that is initially more abundant is most likely to spread to fixation.

For intermediate frequencies of conformism \( 0 < \gamma < 1 \), we observe a gradual shift between the two extreme cases \( \gamma = 0 \) and \( \gamma = 1 \). In the social dilemma (Figure 1A), cooperation becomes a stable equilibrium as soon as conformism is sufficiently strong (i.e. if \( \frac{\gamma}{1-\gamma} s > c - a \)): when a group mainly consists of cooperators, conformist learning leads to the maintenance and fixation of this most abundant strategy, despite of its payoff disadvantage. In the coordination game (Figure 1B), the direction of change remains qualitatively unchanged, but the unstable Nash equilibrium separating the basins of attraction shifts from eq. (2) to \( p^* = 0.5 \), the mixed-strategy Nash equilibrium of the coordination game \( K \). In the evasion game (Figure 1C), \( A \) and \( B \) coexist at equilibrium when learning is mainly based on payoffs (small \( \gamma \)). High frequencies of conformism \( \gamma \) decrease the scope for this coexistence. When conformism occurs at a sufficiently high frequency (moving towards the top of the panel), the direction of expected change is reversed, and the two monomorphic equilibria become stable.

3. A Stochastic Model for Cultural Evolution in Finite Populations

The approach taken above provides an intuitive understanding of the role of conformism in various types of interaction contexts. However, it is not clear whether, and to what extent, the features of conformism-based learning are captured by a coordination game. We therefore developed a dynamic model for cultural evolution where conformism is represented in a more mechanistic way. Cultural evolution takes place in finite (and often small) populations, where chance events may play an important role. Accordingly, we consider a stochastic model for cultural evolution in a finite population of fixed size \( n \). Our model is event-based, where an ‘event’ corresponds to a potential change in strategy by one population member. In each time step (i.e. when an event occurs) two individuals are chosen at random from the population, and one of them is allowed to update its behavioural strategy \((A \text{ or } B)\) by learning from the other. Updating occurs either through conformist or payoff-based learning. For each state of the population (i.e. each possible frequency of \( A \)-strategists), we calculate the probability that an \( A \)-individual switches to \( B \), and that a \( B \)-individual switches to \( A \). At the population level, each such switch corresponds to a decrease or to an increase of the number of \( A \)-individuals by one. Since we neglect the spontaneous emergence of \( A \)- or \( B \)-strategists (the cultural equivalent of genetic mutations), the stochastic process will eventually lead to an
‘absorbing state’, where all individuals have adopted either of the
two strategies. To assess the effects of conformist learning on the
outcome of cultural evolution, we evaluate how our key parameter γ affects the fixation probability and waiting time to fixation for
each of the two behavioural strategies [20].

Let i be the number of A-strategists in the population, and let
$T_i^+$ and $T_i^-$ denote the probability of gaining resp. losing one A-
individual. We model the switching dynamics by using pairwise
comparison [cf. [7]]. When an event occurs, two individuals are
chosen at random from the population. A change in strategy can
only take place when these two individuals have opposite
strategies, which occurs with probability $\frac{i(n-i)}{m(n-1)}$. In state i,
switching from A to B (probability $T_i^+$) and from B to A (probability $T_i^-$) occurs with probabilities:

$$
T_i^± = \frac{i(n-i)}{m(n-1)} \left[(1-\gamma) P_i^± + \gamma C_i^±\right]
$$

where C and P denote the probabilities of switching due to
conformist and payoff-based learning, respectively. For both forms
of social learning, we specify the switching probabilities as a
logistic function of the differences in payoffs (in case of payoff-
based learning) or frequencies (in case of conformist learning)
between strategies A and B. In state i, payoff-based switching from
B to A ($P_i^+$), and from A to B ($P_i^-$) occurs with probabilities

$$
P_i^+ = \Pr(B \rightarrow A | i) = [1 + \exp(-\beta_p (\pi_B(i) - \pi_A(i)))]^{-1} (6a)
$$

$$
P_i^- = \Pr(A \rightarrow B | i) = [1 + \exp(-\beta_p (\pi_A(i) - \pi_B(i)))]^{-1} (6b)
$$

where $\pi_A(i)$ and $\pi_B(i)$ refer to the payoffs of strategies
A and B in state i, respectively. Parameter $\beta_p$ quantifies the
strength and direction of the relation between the payoff difference
and the probability of switching. When $\beta_p = 0$, payoff-based
learning is not biased in any particular direction, and is expected
to lead to dynamics similar to genetic drift. When $\beta_p$ is large,
 payoffs are influenced by the spread of strategies with the highest payoff (see Figure S1 for an illustration).

Conformist switching is represented in a similar way. In state i, conformist switching occurs with probabilities

$$
C_i^+ = \Pr(B \rightarrow A | i) = [1 + \exp(-\beta_c f_B(i) - f_A(i))]^{-1} (7a)
$$

$$
C_i^- = \Pr(A \rightarrow B | i) = [1 + \exp(-\beta_c f_A(i) - f_B(i))]^{-1} (7b)
$$

where $f_A(i) = \frac{i}{n}$ and $f_B(i) = \frac{n-i}{n}$ refer to the relative frequencies of
strategies A and B, respectively. Parameter $\beta_c$ quantifies the
strength and direction of frequency-based social learning. When $\beta_c = 0$, such learning is not biased in any particular direction;
when $\beta_c$ is large, individuals are strongly inclined to adopt the
more frequent strategy in the population (see Figure S1).

Now we have specified $T^+$ and $T^-$ for all states i of the
population, we can use standard methods [28] to calculate fixation
probabilities $\Phi$ and waiting times to fixation, for various initial
abundances of strategies A and B, as a function of the relative
frequency of conformism $\gamma$. In File S1, section 2, we show how
explicit equations for the fixation probabilities can be derived on
the basis of a diffusion approximation of the stochastic model.

We assess the influence of the social learning rules on the
outcome of cultural evolution by comparing the fixation
probabilities $\Phi_{A} | A$ for a given initial abundance of A to that of a ‘neutral’ process, where all switching occurs randomly. It is
well known that in the latter case $\Phi_{A} | A$ equals the initial frequency $k / n$ of A [20]. Panels A to C in Figure 2 illustrate the effects
of conformism on the fixation probability of A (cooperate) in a social
dilemma. When all switching occurs on the basis of payoffs ($\gamma = 0$;
2A), cooperation is always disfavoured. Accordingly, fixation of A
is very unlikely unless the initial abundance of A is relatively large.
In fact, all fixation probabilities (red dots) are below the diagonal,
indicating that fixation of A is for all values of k less likely than
‘neutral’ updating. If social learning is partly based on conformism
($\gamma = 0.25$, 2B; $\gamma = 0.5$, 2C), the fixation curve becomes more S-
shaped. Strategy A is increasingly likely to fixate when its initial
abundance is high, which agrees with our earlier findings
(Figure 1A).

In the coordination game without conformism (Figure 2D), payoff
differences near the A equilibrium are relatively small. This means
that switching from A to B can frequently occur, despite the fact
that strategy B yields lower payoffs. When the state of the group
is close to the unstable equilibrium, such stochastic events can tip
the group into the basin of attraction of B. As conformism increases in
frequency, the pure A equilibrium tends to be more stable; the
fixation probability rises above the diagonal $k/n$ line when the
initial abundance of A is high (Figure 2E, F). This finding is again
in line with our earlier results (Figure 1B).

In the evasion game, payoff-based learning tends to favour
the spread of rare strategies. When conformist learning is absent
($\gamma = 0$, Figure 2G), fixation of B is often more likely. The
polymorphic equilibrium is located at $n/3$, and a group is
expected to spend most of the time close to this equilibrium.
Since this state is closer to the absorbing state where all individuals
play strategy B (relative to the other absorbing state, where all
individuals play strategy A), stochastic events will more likely lead
to fixation of B rather than fixation of A. When conformist
learning occurs at higher frequencies (Figure 2H, I), cultural
evolution tends to lead to fixation of the strategy that was more
abundant initially; again, this is in line with our earlier findings
(Figure 1C).

Conformism also affects the time it takes until a strategy fixates
in the evasion game (Figure 3). In an evasion game, each behavioural
strategy has higher payoff when rare. Accordingly, payoff-based
learning causes a group to spend a lot of time in polymorphic
states before it reaches one of the absorbing states. Increasing the
frequency $\gamma$ of conformist learning has two effects: first, the
frequency of payoff-based switching decreases, which ameliorates
the spread of rare strategies, thereby destabilising the coexistence
equilibrium. Second, conformist switching accelerates fixation,
because individuals preferentially adopt common strategies.

By breaking down polymorphism in the evasion game, conformism affects the average payoffs of the group members. A
simple calculation shows that the average payoff at the mixed-
strategy Nash equilibrium (eq. 2) of an evasion game is given by

$$
a d - b c + d = b c + d
$$

This can be considerably higher or lower than the
payoff $a$ in case of fixation of behavioural strategy A or the payoff $d$
in case of fixation of B.

4. Evolution of Cooperation by Cultural Group Selection

Finally, we consider a multilevel scenario in which a metapop-
ulation is subdivided into m groups of size n. Within groups,
individuals face a social dilemma. As described above, payoff-based learning within groups tends to disfavour cooperation. Once in a while, individuals from two different groups are paired for updating by comparing their payoffs. This reflects a scenario where individuals occasionally copy behaviours from groups that are performing well. It might be that individuals from other groups are considered to be healthier, or have more wealth. We assume that conformism does not play a role in between-group updating (i.e. conformism is a strictly local social learning rule, allowing to cope with local conditions). Further, in our model, updating outside the group occurs at a much lower rate than within-group updating: in between two outside-group updating events. This ‘separation of time scales’ allows us to calculate the probability of fixation of a ‘cooperative’ strategy in the metapopulation, by tracking the number of groups in the cooperative state. In each time step of this group level process, the abundance of cooperator groups can go up by one or go down by one, or can stay the same.

When two individuals from different groups are chosen from the population, switching probabilities are defined analogously to updating within groups. Switching probabilities depend on the payoff difference between groups where cooperation is fixated and groups where defection is fixated. This payoff difference is given by \( a - d \). For this between-group process, we again use logistic functions to specify the relationship between payoff differences and the probability of switching. Since payoffs are higher in cooperative groups than in defector groups \((a > d)\), this process is expected to lead to the spread of cooperation between groups. Let \( j \) be the number of groups at the cooperative state, and let \( n - j \) be the number of groups at the defector state. At state \( j \), the probability of gaining \((U_j^+)\) or losing \((U_j^-)\) a cooperative group (by one defector taking over a cooperative group, or vice versa, by one cooperator taking over a defector group) can then be written as

\[
\Pr (B \rightarrow A)_G = \frac{1 + \exp(-\beta_G(a - d))}{1 + \exp(-\beta_G(d - a))}^{-1} \quad (8a)
\]

\[
\Pr (A \rightarrow B)_G = \frac{1 + \exp(-\beta_G(a - d))}{1 + \exp(-\beta_G(d - a))}^{-1} \quad (8b)
\]

Parameter \( \beta_G \) specifies the relation between the payoff difference of the members of the two different groups, and the probability of switching. Since payoffs are higher in cooperative groups than in defector groups \((a > d)\), this process is expected to lead to the spread of cooperation between groups.

Figure 2. Effects of conformism on the outcome of cultural evolution in a small group. Each panel shows the fixation probability of strategy A as a function of its initial abundance \((k)\) in a group of \( n = 30 \) individuals. Columns of panels correspond to three different frequencies \( c \) of conformism. Symbols represent fixation probabilities from the exact stochastic model, and lines represent a diffusion approximation to these fixation probabilities (see File S1, section 2). The diagonal dashed lines indicate the fixation probability of a strategy under random drift \((Q_k = k/n)\). Parameter settings: \( \beta_0 = 1 \) and \( \beta_c = 2 \); payoff matrices of the games as in Figure 1.

doi:10.1371/journal.pone.0068153.g002
Fixation probability of cooperation tends to increase with a favourable effect on the cultural evolution of cooperation, since the structures and varying frequencies of conformism per se, but rather by the associated decrease in frequency of payoff-based learning. Decreased frequencies of payoff-based learning weaken selection against cooperation within the group. 

To assess the net effects of conformism, one therefore has to compare the results of Figure 4B with a benchmark that takes this weakening of payoff-based learning into consideration. This benchmark is presented in Figure 4A where, with probability γ, individuals are imitating at random.

When the frequency of random switching within groups (as opposed to payoff-based switching) increases, the scope for cooperation increases (Fig. 4A, going from the left to the right in the panel); within groups, the selection against cooperation is weakened, whereas selection between groups is kept constant (see [29] for a general analysis of how weakening within-group selection can affect the scope for cooperation in group-structured populations). Cooperation is favoured most, when the metapopulation is structured into many small groups (Fig 4A going from the bottom to the top of the panels). When groups are small, the probability that a single cooperative strategy reaches fixation – in spite of payoff-based learning disfavouring this strategy – is relatively large. Once such fixation has happened, cooperation can spread to other groups.

Comparing Figures 4A and 4B allows us to evaluate the net effects of conformism on the cultural evolution of cooperation by group selection. It is obvious that conformism hinders rather than favours the evolution of cooperation: in the whole parameter range, cooperation spreads more easily to fixation when conformism-based updating is absent (βb=0; random updating occurs at frequency 1–γ). The solid red line reflects waiting times when payoff-based learning is absent (βb=0; βc=1; random updating occurs at frequency 1–γ). The solid blue line reflects waiting times when payoff-based learning occurs at frequency 1–γ, and conformism is absent (βb=1; βc=0; random updating occurs at frequency γ). The solid red line represents waiting times in the full model, where payoff-based learning is complemented by conformism at rate γ (βb=1; βc=1). Groups were of size n=30 and initialised at the coexistence equilibrium p=p∗ (10 A individuals). Payoff matrix as in Figure 1. doi:10.1371/journal.pone.0068153.g003

Figure 3. Effect of conformism on the persistence of behavioural polymorphism in an evolution game. Lines represent the expected number of updating events (i.e. the ‘waiting time’) until a group fixates in either A or B, as a function of the frequency of conformist updating γ. The dashed blue line indicates waiting times when all updating occurs randomly (βb=0; βc=0). The dashed red line reflects waiting times when conformism occurs at rate γ, and payoff-based learning is absent (βb=0; βc=1; random updating occurs at frequency 1–γ). The solid blue line reflects waiting times when payoff-based learning occurs at frequency 1–γ, and conformism is absent (βb=1; βc=0; random updating occurs at frequency γ). The solid red line represents waiting times in the full model, where payoff-based learning is complemented by conformism at rate γ (βb=1; βc=1).

\[
U_j^+ = \frac{j(m-j)}{m(m-1)} \cdot \Phi_1
\]

\[
U_j^- = \frac{j(m-j)}{m(m-1)} \cdot (1 - \Phi_{n-1})
\]

where \(\Phi_1\) refers to the fixation probability of a single cooperator in a group of defectors, and \(1 - \Phi_{n-1}\) reflects the fixation probability of a single defector in a group of cooperators (which is complementary to the fixation probability of cooperation starting from state n−1). Using the same techniques as before, we can calculate the probability \(\Phi_1\) that eventually all groups have reached the cooperative state, given that we start out with 1 group of cooperators. The product \(\Phi_1 \cdot \Phi_i\) then denotes the probability that cooperation reaches fixation in the metapopulation, given that we start out with one individual with the cooperative strategy.

Figure 4 gives an overview of the fixation probability of cooperation in case of a single cooperator in a population structured in m groups of fixed size n, under varying population structures and varying frequencies of conformism γ. When looking at Figure 4B only, one might conclude that conformism has a favourable effect on the cultural evolution of cooperation, since the fixation probability of cooperation tends to increase with γ. In our view, however, this conclusion would be misleading. The increasing scope for cooperation is not caused by conformism per se, but rather by the associated decrease in frequency of payoff-based learning. Decreased frequencies of payoff-based learning weaken selection against cooperation within the group.
Conformism would influence cultural evolution when individuals are characterised by a mixed strategy, which specifies a probability distribution over the pure strategies. If such probabilistic tendencies could be transmitted between individuals by social learning, the dynamics of cultural evolution could be rather different from the scenario considered here (see [30] for an example). For instance, in an evasion game, all individuals could fixate on the same mixed strategy $0 < \bar{p} < 1$, supporting a polymorphism where individuals make use of both pure strategies in a probabilistic fashion. In contrast to the findings from our analysis using two pure behaviours, conformism would be unlikely to destabilise such a behavioural polymorphism. Note, however, that the transmission of strategies by social learning depends on the degree to which individuals can evaluate the strategies of their peers. Whereas individuals might be able to evaluate the pure strategies of their peers (and possibly imitate them accordingly), it is not obvious that more complex (mixed) strategies readily transmit between individuals.

Furthermore, we assume that individuals in the population all use the same social learning strategy. Decision making experiments show that individuals tend to vary considerably in their social learning strategies (e.g. in the degree in which individuals learn based on payoffs [23]). Such individual variation in social learning strategies can affect the course of cultural evolution. To see this, consider a group in which some individuals typically learn based on payoffs and others learn based on conformism. This group may reach a stable behavioural polymorphism in an evasion game: when conformists all perform a common strategy, payoff-based learners can anticipate to that by adopting the strategy that is rare. Such an emerging differentiation, in which conformist learners perform one behaviour and payoff-based learners perform another behaviour, cannot be attained by groups that are homogeneous with respect to their social learning strategies. In our model, individuals use a mixture of conformist and payoff-based learning. As a consequence, a behavioural polymorphism is destabilised by a number of consecutive conformist learning events, potentially reducing the average payoffs of individuals in a population.

Human social learning comes in many different forms, and payoff-based and conformist learning only represent those forms that have received most attention in the social learning literature. Our analysis does not account for how other relevant forms of social learning – such as following a leader or a teacher – would affect the spread of behaviours within groups (see [31] for how leadership can affect the cultural evolution of cooperation). Also, the mechanism that spreads behaviours between groups considered our model, is only one way that this group-level mechanism might work [15]. Alternative scenarios in which groups of cooperators grow faster and split up when reaching a certain size [17] are likely to lead to different outcomes of cultural evolution. Such alternative forms of group-level selection can have different consequences for evolutionary dynamics, and can interact with within-group social learning in different ways. A more specific simulation study [32], considering other population structures and different forms and social learning and group selection, arrives at a similar conclusion: conformism can promote the cultural evolution of cooperation by group selection when groups can replace other groups, whereas cooperation cannot evolve when cooperators have to spread singly from group to group by a process of ‘infection’.

Our study leads to the conclusion that conformism has only a marginal effect in the context of coordination games, that it tends to erode the polymorphic equilibrium in evasion games, and that it does not favour cooperation in a social dilemma. In other words, our evaluation of the role of conformism is considerably less favourable than the opinion of other scholars of cultural evolution [3,8,11,14,15]. To put our conclusion into context, we would like to stress that our analysis did not consider potential intrinsic benefits of conformism. Two such benefits may be relevant in a variety of situations. First, conformism could homogenise groups with respect to a diversity of norms and habits, potentially increasing social cohesion and facilitating the establishment of trust, thus making it easier to resolve internal conflicts and to get cooperation off the ground. Second, conformism could be a beneficial strategy in environments with considerable spatial variation in payoffs and/or behavioural norms. In a situation like that, newcomers in a local group or environment could profit from

Figure 4. Effect of conformism on the evolution of cooperation by cultural group selection. Panels show fixation probability of cooperation in a group-structured population where initially only a single individual cooperates. Within groups, payoff-based learning favours defection, but cooperation can spread between groups by occasional learning outside of the local group, because individuals in cooperative groups have higher payoffs. The number of groups $m$ was varied between 10 and 200, in steps of 5, holding the metapopulation size constant at $m n = 1000$. The frequency of conformism updating was varied between 0 and 1, in steps of 0.01. In (A), $\beta_C$ is equal to zero, corresponding to random updating, while the conformism-based updating has a strong effect ($\beta_C = 1$) in (B). Hence, panel (A) should mainly be viewed as a reference for panel (B). We used an interpolation procedure (using the R-package ‘akima’) to smoothen the plots. Colours facilitate comparison to the fixation probability of a selectively neutral mutant $1/(m n)$. The payoff matrix coincides with that of the social dilemma game in Figure 1.

doi:10.1371/journal.pone.0068153.g004
imitating local habits, thus quickly adopting locally superior strategies or adapting to local behavioural equilibria. More sophisticated models accounting for spatial variation and the mechanisms underlying decision making in groups might therefore arrive at a more positive judgement of the role of conformism for the cultural evolution of social behaviour.

Supporting Information

Figure S1 Illustration of the logistic function for three values of β. (TIFF)

File S1. In the first part of this supplement, we provide a graphical illustration of the logistic functions used in our stochastic model to translate information on payoff or frequency differences into the probability of switching to another strategy. In the second part, we derive a diffusion approximation of our stochastic model that yields, by good approximation, analytical expressions for the fixation probability of a strategy in a finite population as a function of the strategy’s initial frequency. (DOCX)

Acknowledgments

We would like to thank Jorge Peña, Pieter van den Berg, Andri Manser and the Theoretical Biology group Groningen for helpful discussions. Furthermore, we thank Christian Hilbe and one anonymous reviewer for their comments and suggestions.

Author Contributions

Conceived and designed the experiments: LM IP FJW. Analyzed the data: LM IP FJW. Contributed reagents/materials/analysis tools: LM IP FJW. Wrote the paper: LM IP FJW.

References

1. Rendell L, Fogarty L, Hoppitt WJF, Morgan TJH, Webster MM, et al. (2011) Cognitive culture: Theoretical and empirical insights into social learning strategies. Trends Cogn Sci 15: 68–76.
2. Laland KN (2004) Social learning strategies. Learn behav 32: 4–14.
3. Boyd R, Richerson PJ (1985) Culture and the evolutionary process. Chicago: University of Chicago Press.
4. Henrich J, Gil-White FJ (2001) The evolution of prestige: Freely conferred deference as a mechanism for enhancing the benefits of cultural transmission. Evol Hum Behav 22: 165–196.
5. Henrich J, Boyd R (1998) The evolution of conformist transmission and the emergence of between-group differences. Evol Hum Behav 19: 215–241.
6. Vílone D, Ramasco JJ, Sánchez A, San Miguel M (2012) Social and strategic imitation: The way to consensus. Sci Rep 2: 686.
7. Traulsen A, Nowak MA, Pacheco JM (2006) Stochastic dynamics of invasion and fixation. Phys Rev E 74: 011909.
8. Andrés Guzmán R, Rodríguez-Sickert C, Rowthorn R (2007) When in Rome, do as the Romans do: the coevolution of altruistic punishment, conformist learning, and cooperation. Evol Hum Behav 28: 112–117.
9. Nowak MA, Sasaki A, Taylor C, Fudenberg D (2004) Emergence of cooperation and evolutionary stability in finite populations. Nature 428: 646–650.
10. Peña J, Volken H, Pestelacci E, Tomasini M (2009) Conformity hinders the evolution of cooperation on scale-free networks. Phys Rev E 80: 016110.
11. Henrich J, Boyd R (2001) Why people punish defectors. J Theor Biol 208: 79–89.
12. Skyrms B (2005) Dynamics of conformist bias. The Monist 88: 260–269.
13. Peña J, Volken H, Pestelacci E, Tomasini M (2009) Conformity hinders the evolution of cooperation on scale-free networks. Phys Rev E 80: 016110.
14. Henrich J, Boyd R (1998) The evolution of conformist transmission and the emergence of between-group differences. Evol Hum Behav 19: 215–241.
15. Henrich J (2004) Cultural group selection, coevolutionary processes and the emergence of between-group differences. Evol Hum Behav 19: 215–241.
16. Boyd R, Gintis H, Bowles S, Richerson PJ (2003) The evolution of altruistic helping. Am Nat 172: 12–24.
17. Henrich J (2004) Cultural group selection, coevolutionary processes and the emergence of between-group differences. Evol Hum Behav 19: 215–241.
18. Boyd R, Richerson PJ (2002) Group beneficial norms can spread rapidly in a random group formation. Theor Popul Biol 73: 506–516.
19. Lehmann L, Feldman MW (2008) The co-evolution of culturally inherited altruistic helping and cultural transmission under random group formation. Theor Popul Biol 73: 506–516.
20. Lehmann L, Feldman MW, Foster KR (2008) Cultural transmission can inhibit the evolution of altruistic helping. Proc Nat Acad Sci 105: 12370–12375.
21. Asch SE (1956) Studies of independence and conformity: I. A minority of one against a unanimous majority. Psychol Monogr: General and Applied 70: 1–70.
22. Efferson C, Lalive R, Richerson PJ, McElreath R, Lubell M (2008) Conformists and mavericks: The empirics of frequency-dependent cultural transmission. Evol Hum Behav 29: 56–64.
23. McElreath R, Lubell M, Richerson PJ, Waring TM, Baum W, et al. (2005) Applying evolutionary models to the laboratory study of social learning. Evol Hum Behav 26: 483–501.
24. Traulsen A, Semmann D, Sommerfeld RD, Kranbeck HJ, Milinski M (2010) Human strategy updating in evolutionary games. Proc Nat Acad Sci 107: 2962–2966.
25. McElreath R, Bell AV, Efferson C, Lubell M, Richerson PJ, et al. (2008) Beyond existence and aiming outside the laboratory: Estimating frequency-dependent and pay-off-biased social learning strategies. Phil Trans Roy Soc B 363: 3515–3521.
26. Mesoudi A (2011) An experimental comparison of human social learning strategies: Pay-off-biased social learning is adaptive but underused. Evol Hum Behav 32: 334–342.
27. Osborne MJ, Rubinstein A (1994) Course in game theory. Cambridge: The MIT press.
28. Karlin S, Taylor HE (1975) A first course in stochastic processes. New York: Academic press.
29. Hauert C, Imhof LA (2012) Evolutionary games in deme structured, finite populations. J Theor Biol 299: 106–112.
30. Gargiulo F, Ramasco JJ (2012) Influence of opinion dynamics on the evolution of games. PloS One 7: e48916.
31. Lehmann L, Feldman MW (2008) The co-evolution of culturally inherited altruistic helping and cultural transmission under random group formation. Theor Popul Biol 73: 506–516.
32. Molleman L, Quintones AE, Weissing FJ (2012) Cultural evolution of cooperation: the interplay between forms of social learning and group selection. Evol Hum Beh. In press.