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Cooperation in two-dimensional mixedgames

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Abstract

Evolutionary game theory is a common framework to study the evolution of cooperation, where it is usually assumed that the same game is played in all interactions. Here, we investigate a model where the game that is played by two individuals is uniformly drawn from a sample of two different games. Using the master equation approach we show that the random mixture of two games is equivalent to play the average game when (i) the strategies are statistically independent of the game distribution and (ii) the transition rates are linear functions of the payoffs. We also use Monte-Carlo simulations in a two-dimensional lattice and mean-field techniques to investigate the scenario when the two above conditions do not hold. We find that even outside of such conditions, several quantities characterizing the mixed-games are still the same as the ones obtained in the average game when the two games are not very different.

Keywords: game theory, mixed games, cooperation, prisoners dilemma

(Some figures may appear in colour only in the online journal)

1. Introduction

How is it possible for cooperation to exist in a system dominated by competitive and self-interested individuals? This is one of the big open questions in science [1]. It becomes more interesting when we realize that cooperation is not exclusively a human phenomena, it exists between different animal species [2], within very organized insect societies and amongst kin [3]. If we generalize cooperation, using the framework of game theory, even between cells

and RNA the concept can be used to model the emergence of organization in more complex and efficient forms [4, 5].

In competitive systems it would be reasonable to think that selfish behaviour would be the best choice of strategy, so how can cooperation spontaneously emerge? Evolutionary game theory is often used to analyze such questions, specifically using the so called dilemma games [6, 7]. In such games players can cooperate to obtain a payoff, but there's a temptation to betray the cooperator and get a better payoff.

One of the simplest cases in game theory is the two-player game with only two possible strategies. Each player can choose either to cooperate (C) or to defect (D) in each round. Although one player can interact with many other players, each round is played only by two individuals. The payoff matrix, G = (T, R, P, S), represents how much the players gain from a single game and is normally stated as: if both cooperated they receive R (Reward), if both defected they receive P (Punishment) and if they chose different strategies the defector receives T (Temptation) and the cooperator receives S (Sucker). These four parameters can describe different games [6, 8–10], being the Prisoners Dilemma (PD) the most canonical case, where T > R > P > S. This means that the best payoff is to defect when someone tries to cooperate with you. But if the whole population starts to defect the total payoff is lower than of a cooperating population. The Snow-Drift game (SD) happens when T > R > S > P. Observe that in this case it is worst when both players defect. This is a common game in animal contests where the damage of escalating conflicts is usually higher than being exploited [11]. The Stag-Hunt (SH) happens when R > T > P > S. Now it is better to imitate your opponent. This game can describe the behaviour of cooperative associations, where one can exploit the other, but the best results are achieved by cooperation [12]. The last case is the Harmony-Game (HG), characterized by R > T and S > P. In this case it is always better to cooperate.

The evolution of cooperation in well-mixed populations can be described by mean-field equations, like the replicator equation [6]. It predicts that cooperation cannot survive, a conclusion that does not seems to happen in nature [6, 8, 13]. Many mechanisms have been proposed to explain the survival of cooperation: direct and indirect reciprocity [13-16], volunteering [17], coevolution [9], spatial selection [18], multi-level and kin selection [3]. The essential feature of these mechanisms is that cooperators interact more often between themselves than with defectors. For example, spatial reciprocity happens when individuals are located in the vertices of a graph and can interact only with their neighbours. Cooperators spontaneously form clusters of cooperation that survive in a sea of defectors for a wide range of parameters [6]. Most models assume that only one game is repeatedly played in all interactions. However, there is nothing a priori precluding the coexistence of different games in a population. For example, players can have different perceptions of the rewards, giving rise to a multi-game scenario. In this case, individuals play asymmetric games, where the payoff matrix of one player is different from the payoff matrix of his co-player [19, 20]. Here, we study a model where different payoff matrices are randomly assigned to each interaction the mixed-game model [10]. Different from the multi-game model, here the game is symmetric, that is, in each round both players use the same payoff matrix.

The mixed-game model is presented in the next section. In section 3 we develop the master equation approach for mixed-games. We show that these equations are identical to the ones where the evolution is driven by a single game with the payoff given by the average of the two games (average game), when particular conditions hold. In section 4 we use Monte-Carlo simulations to investigate the evolution of the cooperators for mixed-games in a two-dimensional lattice. We also analyze how robust is the result of one cooperative game if there is a chance of a more selfish game be played and compare the mixed-game model to the

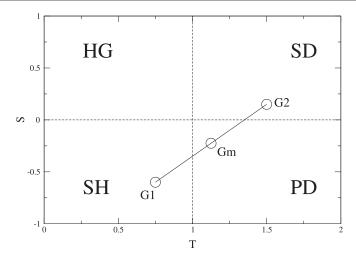


Figure 1. Parameter space in T and S for the possible payoff matrices. Using R = 1 and P = 0 we have the four games (HG, SD, PD, SH) well defined in each quadrant. For every two single games (SH = G_1 , and SD = G_2 , for example) there will be a mean game that is in the medium point between the two games. This mean game can be a totally different game (PD), as is shown in the example above.

average game. The relation of the master equation with the replicator equation is discussed in appendix A. Appendix B contains the development of the pair-approximation model used to study the mixed-games. Finally, we present our conclusions in the last section.

2. The model

We simulate the evolution of cooperation in populations structured on two-dimensional square lattices with first neighbour interactions and periodic boundary conditions. Initially, each individual is assigned to one strategy (C or D) with equal probability. The strategies determine the payoff obtained on pairwise interactions. We choose to parametrize R=1, P=0, $T\in[0,2]$ and $S\in[-1,1]$ to obtain an useful parameter space that contains all four games (HG, SH, PD and SD), as can be seeing in figure 1. By doing so we can analyze how the fraction of cooperators in the stationary state, $\rho(t\to\infty)$, is affected by such parameters and if there is any phase transition.

The evolutionary process of mixed-games is composed of two parts: the game phase and the imitation phase. In the *game phase* each individual plays one round of a game with each one of his four neighbours and the payoff obtained in each interaction is added to the cumulative payoff of each player. The games to be played are randomly assigned to each interaction: $G_1 = (T_1, S_1)$ with probability w or $G_2 = (T_2, S_2)$ with probability 1 - w. In the *imitation phase*, individuals may update their strategy by imitating the strategy adopted by more successful neighbours: the focal individual i randomly chooses one neighbour j to copy his strategy with probability $p(\Delta u_{ij})$, depending on the payoff difference $\Delta u_{ij} = u_j - u_i$, where u_i is the cumulative payoff of player i. For the strategy update probabilities, we use the Fermi-Dirac and the proportional imitation rule [6, 21]:

$$p(\Delta u_{ij}) = 1/(1 + e^{-(u_j - u_i)/K})$$

$$\tag{1}$$

$$p(\Delta u_{ij}) = \max[(u_i - u_i), 0]/b, \tag{2}$$

respectively. The parameter b is a normalizing factor and K is the parameter controlling the strength of selection pressure on strategies [8].

The game and the imitation phases can be implemented synchronously, partially asynchronously, or totally asynchronously [8, 22]. In the *synchronous protocol*, in each Monte Carlo step (MCS) the cumulative payoffs are set to zero, games are randomly and independently assigned to each one of the interactions, and the cumulative payoff of every player is calculated. Then all players simultaneously proceed to the imitation phase. In the *totally asynchronous protocol* the cumulative payoffs are set to zero, a randomly chosen focal player and his four neighbours play randomly and independently assigned games with their respective neighbours, the cumulative payoff of these five players is calculated, and, finally only the focal player proceeds to the imitation phase. This process is repeated until all nodes have equal chance to update their strategies at least once—this defines one MCS. In the *partially asynchronous protocol*, the games that each player will play with its neighbour are assigned only once at the beginning of each MCS—as in the synchronous protocol—and then strategies are updated asynchronously. In this work, all protocols yields the same qualitative results.

We measure the cooperation level as the average number of cooperators ρ in the stationary state. We compare the evolution of cooperation in the mixed-game case to the case where only the average game $G_m = \langle G \rangle = wG_1 + (1-w)G_2$ is played and to the cases where only G_1 or G_2 is played.

3. Master equation approach

Perhaps the first interesting question is about the equivalence between the mixed-games and the average game. In this section we investigate such equivalence for the asynchronous update. Obviously we are unable to solve the master equation exactly. However, we are able to establish conditions that allow us to show this equivalence. These conditions are: (i) the strategies are statistically independent of the game distribution and (ii) the transition rates are linear functions of the payoffs. The main result of this section can be used as a guideline for more general problems.

At each time, t, a system is characterized by the strategy configuration $\{s\} = \{s_1, s_2, ...s_N\}$, where s_i is the strategy of player i, and by the game assignment configuration $\{g\} = \{g_{12}, g_{13}, ...\}$. The variable g_{ij} is uniformly distributed over the discrete set of values $\{1, 2\}$, determining which one of the two games, G_1 or G_2 , are to be played in the interaction between player i and its next neighbour j. Note also that each link i, j is statistically independent of the other links. Let $P(\{s\}, \{g\}, t)$ be the probability to find the mixed-game system in the configuration $\{s\}$ and $\{g\}$ at time t. The time evolution of this system is given by

$$\frac{\mathrm{d}}{\mathrm{d}t}P(\{s\}, \{g\}, t) = \sum_{\{s\}', \{g\}'} P(\{s\}', \{g\}', t)W(\{s\}', \{g\}' \to \{s\}, \{g\})
- P(\{s\}, \{g\}, t)W(\{s\}, \{g\} \to \{s\}', \{g\}'),$$
(3)

where $W(\{s\}', \{g\}' \to \{s\}, \{g\})$ is the transition rate from the state $(\{s\}', \{g\}')$ to $(\{s\}, \{g\})$. If the initial game assignment probability distribution is the stationary distribution $\Theta = \prod_{ij} \theta(g_{ij}), \Theta$ is independent of time. Here we have that $\theta(g) = w\delta_{g,1} + (1 - w)\delta_{g,2}$.

Supposing the condition (i) that the strategy variables $\{s\}$ are statistically independent of the game variables $\{g\}$, we can write that $P(\{s\}, \{g\}, t) = P(\{s\}, t)P(\{g\})$. To simplify the notation, from now on we write s and g instead of $\{s\}$ and $\{g\}$. The master equation can now be written as

$$P(g)\frac{d}{dt}P(s,t) = \sum_{s',g'} P(s',t)P(g')W(s',g'\to s,g) - P(s,t)P(g)W(s,g\to s',g').$$
(4)

The mean value of a function f averaged over all game configurations is $\langle f \rangle_g = \sum_g f P(g)$. Using this, we can sum in g in both sides of equation (4) to obtain that

$$\frac{\mathrm{d}}{\mathrm{d}t}P(s,t) = \sum_{s',g} P(s',t) \left[\sum_{g'} P(g')W(s',g' \to s,g) \right] - \sum_{s',g'} P(s,t) \left[\sum_{g} P(g)W(s,g \to s',g') \right].$$

Since the quantities inside square brackets are the averaged values of W over g(g') we have that

$$\frac{\mathrm{d}}{\mathrm{d}t}P(s,t) = \sum_{s'}P(s',t) \left\langle \sum_{g}W(s',g'\to s,g) \right\rangle_{g'}$$

$$-\sum_{s'}P(s,t) \left\langle \sum_{g'}W(s,g\to s',g') \right\rangle_{g}.$$
(5)

The transition rates W will be obtained from each microscopical rule (proportional imitation, Fermi–Dirac, copy the best, etc), and will depend on the payoff differences (Δu_{ij}) of neighbouring sites, topology of the lattice and the copy mechanism. Since the dynamical process in a time step, in our model, always involves only one site, for example site i, copying the strategy of a neighbour, say j, all the transition rates in which the new g' has a link that is not connected to site i are null. Moreover all these rules are explicitly independent of the future game to be played, making the summation inside the averages lead to a finite constant κ that will only depend on the geometry of the lattice and on some aspects of the microscopic model. It follows that

$$\left\langle \sum_{g} W(s', g' \to s, g) \right\rangle_{g'} = \kappa \langle W(g', s' \to s) \rangle_{g'}. \tag{6}$$

Notice that for simplicity we hide Δu_{ij} from the transition rates $W(s', g' \to s, g; \Delta u_{ij})$ variables, although it depends on it. In regular lattices κ is the same constant for every site and can be absorbed in the time variable and we get

$$\frac{\mathrm{d}}{\mathrm{d}t}P(s,t) = \sum_{s'} P(s',t) \langle W(g,s'\to s) \rangle_g - \sum_{s'} P(s,t) \langle W(g,s\to s') \rangle_g.$$
(7)

If we use condition (ii), that the update rules are linear on the payoff, the average over the transition rules will become simply the the transition rates of the average payoff, namely

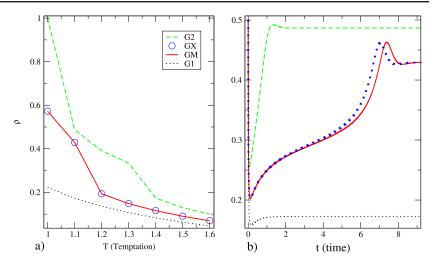


Figure 2. Integration of the mean-field, pair-approximation, ODE by a Runge–Kutta algorithm. (a) Phase transition of the stationary state value of cooperators in different single games $(G_1, G_2 \text{ and } G_m)$ and the mixed-game (G_x) . $G_m(S_m = 0)$ is the mean game of $G_1(S_1 = -0.1)$ and $G_2(S_2 = 0.1)$, GX is the model where both G_1 and G_2 are played with equal probability. (b) Time evolution of cooperators for each game, here T = 1.1. Although the mixed-game behaves differently for small times, the fraction of cooperators in the stationary state is similar.

$$\left\langle W\left(s',\,g'\to s,\,g;\,\Delta u_{ij}\right)\right\rangle_g=W\left(s',\,g'\to s,\,g;\,\left\langle\Delta u_{ij}\right\rangle_g\right).$$
 (8)

Using this relation in equation (7), we obtain that

$$\frac{\mathrm{d}}{\mathrm{d}t}P(s,t) = \sum_{s'} P(s',t)W(s'\to s;\langle G\rangle) - \sum_{s'} P(s,t)W(s\to s';\langle G\rangle), \quad (9)$$

where $\langle G \rangle$ stands for the average payoff matrix. This equation implies that the time evolution of the configuration probability in a mixed-game with asynchronous update is equal to the time evolution of a single game [6] with the transition rates evaluated by using the average payoff matrix. Note that this equation also describes the classical evolution of a single game if we use the appropriate transition rates.

It is worth mentioning that this result also holds for more general game assignment distributions. Supposing that $\theta_g = \sum_{k=1}^n w_k \, \delta_{g,k}$, where w_k is the probability of play the game \mathbf{G}_k (k=1,...,n), we can easily see that the argument remains the same. The only difference is that the average payoff matrix is now given by $\langle G \rangle = \sum_{k=1}^n w_k \, G_k$.

We emphasize that equation (9) was obtained by assuming that the variables $\{s\}$ and $\{g\}$ are not correlated, condition (i). Also the equivalence between the average and the mixed-games only holds for the master equation as long as the transitions rates are linear in the payoff, condition (ii).

In appendix A we expand the replicator equation model to include mixed-games. The numerical solution of the master equation can be refined into more accurate models that consider spacial interactions. Appendix B shows the classic pair approximation [21, 23] and how we expanded it to include mixed-games in a square lattice. Our numerical solution shows that the mixed-game is equivalent to the mean game. But in the pair approximation this only

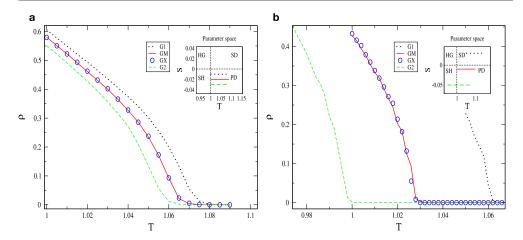


Figure 3. (a) Phase transition of cooperators in different PD games for an asynchronous model, L=100. $G_m(S=-0.02)$ is the mean game of $G_1(S=-0.01)$ and $G_2(S=-0.03)$, GX is the model where both G_1 and G_2 are played. (b) Phase transition in different games for a synchronous model, L=200. Here $G_1(T>1.05, S=0.02)$ is a SD game, $G_2(T>0.95, S=-0.05)$ a SH and the mean game $G_m(T>1, S=-0.01)$ is a PD. The mix of those two different games still behaves as a PD.

holds as long as the parameters (*T* and *S*) of the mixed-games are close enough from the mean game. The obtained results are shown in figure 2.

4. Monte Carlo simulations

Let us now study the mixed-game model with populations structured in a two-dimensional square lattice. Since the variables g_{ij} defining the game between sites i and j obeys the stationary probability distribution $\Theta = \prod_{ij} \{w\delta_{g_{ij},1} + (1-w)\delta_{g_{ij},2}\}$, they are independent of the strategy's variables. On the other hand, the probability of s_i and s_i have particular values, (C, C) for example, depends if the game between them is G_1 or G_2 . Therefore condition (i) is not obeyed. Moreover, the transitions rates are not linear when we use the payoffs defined by equations (1) and (2). This implies that condition (ii) is also not obeyed. To investigate the effect of breaking conditions (i) and (ii), we performed Monte-Carlo simulations on the square lattice. Simulations were performed on lattices with sizes (L) ranging from 100×100 to 500×500 . All quantities were averaged over 20 initial conditions and left to termalize for at least 3000 MCS for each run. It turns out that if the difference between G_1 and G_2 is small, the evolution of cooperation in a mixed-game is equivalent to the average game G_m , as shown in figure 3. Interestingly, figure 3(b) shows that cooperation may thrive in the presence of disadvantageous payoff matrix values as long as the average game is a favourable one. Notice that the mixed-game behaves as the mean game even when the two games are of different kinds, as shown in figure 3(b), where G_1 is a SD and G_2 is SH game.

Figure 4 shows the difference between the fraction of cooperators for the mixture of games $G_1 = (1.04, S)$ and $G_2 = (1.04, -S)$ and the fraction of cooperation for the corresponding average game G_m . If the difference between the games G_1 and G_2 grows, then the equivalence no longer holds. Hence, our simulations suggest that, even if the update rules are nonlinear, the equivalence between average and mixed-game may hold as long as the

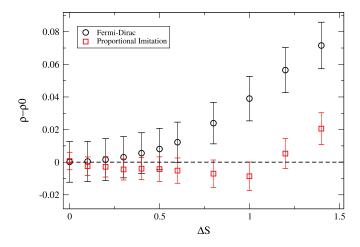


Figure 4. Difference in the cooperation from the mean game and mixed-games versus the difference from the parameter S_1 and S_2 . The dashed line corresponds to the mean game value of cooperation, squares point the data from the proportional imitation model and circles the Fermi–Dirac Until $\Delta=1$ the proportional imitation model just fluctuates around the mean game value, while the Fermi–Dirac progressive rises. The cooperator differences are of 10^{-2} order.

difference between the games G_1 and G_2 is small. Nevertheless, the fraction of cooperation differs at most 0.1. The small deviations contrast with the results obtained in [19, 20], where the authors modelled multi-games as a result of different perceptions of the interactions. In their paper the games are asymmetric, meaning that the interface between cooperators and defectors can be beneficial for cooperators, strengthening even more the expansion of cooperation clusters. In our model, the symmetric games are randomly assigned every time step and the two players always have the same perception of the game. Hence, the cooperators on the interface can thrive only if the average game is cooperative.

It is well known that network topology may affect the evolution of strategies [6, 7, 24–26]. Hence, we also studied the effect of scale-free complex networks on the evolution of mixed games. We used the Redner–Krapivsky algorithm [27] to create networks of $N = 10^4$ individuals and ran the Monte-Carlo simulations for 100 different network configurations. Like in the square lattice, we observed that both the mixed and the mean game are equivalent in terms of the final fraction of cooperation.

5. Conclusion

We investigated under which circumstances the evolution of cooperation in the mixed-game model is equivalent to play a single game defined as the average between the two games in the mixed-game model. Using a master equation approach we showed that the time evolution of a mixed-game with asynchronous update is equivalent to the average game if the transition rates are linear in the payoff and if there are no correlations between the game played and the state of the player. Using both synchronous and asynchronous Monte-Carlo simulations in the square lattice, we investigated the effects of (i) nonlinear update rules and of (ii) correlations between the strategy and game assignment variables. We found that, as long as the differences between the two games are small, the mixed-game and the average game are equivalent. We

showed also that assigning different games to interactions is not the same as to say that individuals have different asymmetric perceptions of the interaction.

Acknowledgments

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Appendix A

We now derive the replicator equation for mixed-games. Suppose an infinite population of strategists where everybody interacts with everybody. Associating the mean payoff of a strategy with its capability of 'reproduction' we can obtain mean field equations for the dynamics of the strategy population. The most simple cases comes without considering spatial effects and using just a single game being played. The replicator equation for this model can be written [6, 28] in terms of games as

$$\dot{x}_i = x_i \left[f_i(\vec{x}) - \phi(\vec{x}) \right]. \tag{A1}$$

Here x_i is the fraction of players using strategy i (in our case C or D), $f_i(\vec{x})$ can be regarded as the mean payoff of strategy i, and $\phi(\vec{x})$ is the population average payoff. This mean field dynamics can be mapped into a master equation of a Markovian chain [6] that represents each player changing its state using some transition rule. It follows that

$$\dot{x}_i = \sum_{j} \left[x_j \omega_{ji} (\vec{x}) - x_i \omega_{ij} (\vec{x}) \right]. \tag{A2}$$

Here $\omega_{ij}(\vec{x})$ is the individual transition rate of a player from state i to j, and can be obtained from the microscopic rule that states how one site copies the strategy of another. Many different microscopical rules have been proposed, such as imitate the best, best response, Fermi–Dirac probability and proportional imitation rule. As can be expected, each one of this rules changes the results of the replicator dynamics, leading to different final result. The most common cases are the Maynard–Smith and Taylor–Johnsen equations [29, 30] which can be obtained from the proportional imitation and Moran process microscopical rules [6].

Now we generalize this result to mixed-games. Being able to play two different games at each interaction, the payoff matrix G, will be now a statistical quantity, with mean value G_m . This leads to a change in the transition rates of the model, where the microscopic rule now depends on a probability distribution (Θ) of games being played. Although we are changing the game being played, the replicator equation still maintain their dependence with the payoff (i.e. $f_i(\vec{x})$). If we average the replicator equation in the game assignment variables, we obtain

$$\langle \dot{x}_i \rangle_g = \left\langle \sum_j \left[x_j \omega_{ji}(\vec{x}) - x_i \omega_{ij}(\vec{x}) \right] \right\rangle_g = \sum_j \left[x_j \left\langle \omega_{ji}(\vec{x}) \right\rangle_g - x_i \left\langle \omega_{ij}(\vec{x}) \right\rangle_g \right]. \tag{A3}$$

If we use any transition rule that is linear in G, this will lead $\langle \omega_{ij}(\vec{x}) \rangle = \omega_{ij}(\vec{x}; \langle G \rangle)$ that is equivalent to playing the mean game $G_m = wG_1 + (1-w)G_2$. This description agrees with the one obtained from equation (9). Note that these results are also valid if we use the more general game distribution $\theta_g = \sum_{k=1}^n w_k \delta_{g,k}$.

Appendix B.

The simplest way to describe a repeated game is by using a mean-field approximation. Here it is supposed that all individuals play with all other players in an infinite population. By doing this one can expect to obtain the mean payoff of a cooperator (u_c) or defector (u_d) using the payoff matrix and the abundance of other defectors or cooperators. In this simple model we have:

$$u_c = R\rho_c + S\rho_d, \tag{B1}$$

$$u_d = T\rho_c + P\rho_d, (B2)$$

where ρ_c and ρ_d are the fraction of cooperators and defectors of the population. Inserting these payoffs in the replicator equation (A1) as the fitness of each strategy, we obtain the time evolution of such strategies as predicted by a mean-field approximation. This framework does not account for spacial effects such as lattices topologies or number of neighbours and results in a scenario where only one specie survives, depending on the parameters. There is no coexistence. Notice that in this model the payoff is a linear function of the parameters, so it is equivalent to play a mean game G_m and averaging the payoffs of two mixed-games G_1 and G_2 .

The simplest way to take into account the lattice is by using the pair-approximation [6, 21, 23, 31]. Here we no longer suppose that all individuals are connected to everyone else. Instead we use a square lattice with interaction between the first neighbours and analyze a cluster of two focal sites (i and j) and their nearest next neighbours (x, y, z and u, v, w) as it is shown in figure B1 . Sites i and j have their payoff calculated exactly as they play games with cluster sites, while the neighbouring sites have an exact payoff calculated with i or j and the other three games (the second neighbours sites) payoffs are calculated by a mean value using a one site approximation.

To obtain the dynamics for this system we use the master equation (3). Now, instead of using $P(\{s\}, \{g\}, t)$ as the main variable, we use the probability that a connection between two sites is in the current state $\Gamma_{\alpha,\beta}$ (α and β can be C or D in this case). $W(\{s\}, \{g\} \rightarrow \{s\}', \{g\}')$ become the rates that each link changes from one state to another $(cd \rightarrow cc)$ for example). And we further simplify for the single game case ($\{g\} = \{g\}'$). Later we analyze the mixed-game models. This gives us three coupled ODE's that states how the fraction of possible connections in the cluster evolves:

$$\dot{\Gamma}_{cc} = \sum_{\{k\}} P(\{k\}; cd) W_{cd \to cc} \{k\} - P(\{k\}; cc) W_{cc \to cd} \{k\},$$
(B3)

$$\dot{\Gamma}_{cd} = \sum_{\{k\}} P(\{k\}; cc) W_{cc \to cd} \{k\} + P(\{k\}; dd) W_{dd \to cd} \{k\}$$

$$-P(\{k\}; cd)W_{cd\to cc}\{k\} - P(\{k\}; cd)W_{cd\to dd}\{k\},$$
(B4)

$$\dot{\Gamma}_{dd} = \sum_{\{k\}} P(\{k\}; cd) W_{cd \to dd} \{k\} - P(\{k\}; dd) W_{dd \to cd} \{k\}.$$
(B5)

Here $\{k\}$ is the configuration which describes the sites strategies, $P(\{k\}; \alpha\beta)$ is the probability that the system is found in configuration $\{k\}$, given that exists at least one connection of the type $\alpha\beta$ and the summation occurs over every possible configuration that allows each one of the given transitions. The summation occurs over every possible cluster state, although in certain states the rate $W_{\alpha\beta\to\gamma\beta}$ that a pair can change is zero. We use the system constrains to reduce the problem to the resolution of only two ODE's, namely $\Gamma_{cc} + \Gamma_{cd} + \Gamma_{dd} = 1$, $\rho_c = \Gamma_{cc} + \Gamma_{cd}/2$ and $\rho_c + \rho_d = 1$. Each rate W is calculated by counting every possible transition of the system from one state to the other and weighting it

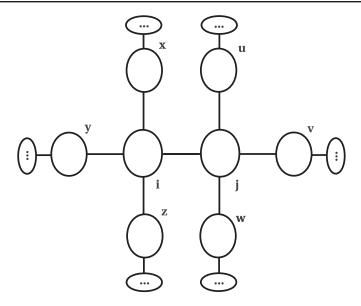


Figure B1. The cluster structure used in the mean field-calculations. Although all seven main connections can give exact payoff between the sites, the second order neighbours (shown here as '…') payoffs needs to be calculated as a mean value using one site approximation. All eight sites can be either C or D and each connection can be in the states Γ_{cc} , Γ_{cd} and Γ_{dd} .

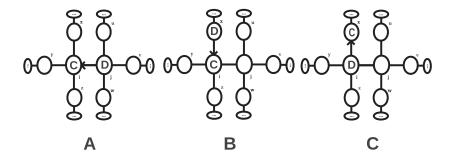


Figure B2. Example of the three process that can change any link Γ_{cd} to Γ_{dd} in the cluster. Notice that this figure is just an example using process for the rate $W_{cd \to dd}$. Each of the other rates $W_{\alpha\beta \to \gamma\beta}$ have three similar process, exchanging α , β and γ for the appropriate C or D.

by the transition probability $p(u_{ij})$. Notice that as we look for changes in the links, not only i and j can change, but any of the seven links composing the cluster. In a broad sense each term in the ODE's is composed of only three terms:

$$P(\{k\}; \alpha\beta) W_{\alpha\beta \to \gamma\beta} = A_{\alpha\beta} + B_{\alpha\beta} + C_{\alpha\beta}.$$
(B6)

These terms represent the only possible processes that can change the links of the cluster in one interaction: (A) the focal site copies the strategy of the other focal site, (B) the focal site copies the strategies of a second order neighbour, (C) the second order neighbour copies the strategy of focal site. This can be seen in figure B2. These processes have a general form that depends on the two sites that will remain fixed in the summation (i.e. α , β being fixed C or D

depending on each rate $W_{\alpha\beta\to\gamma\beta}$)

$$A_{(i=\alpha,j=\beta)} = \sum_{x,y,zv,w,u} \eta_{\beta} \left[\frac{\Gamma_{ix}\Gamma_{iy}\Gamma_{iz}}{\Gamma_{i}^{3}} \Gamma_{ij} \frac{\Gamma_{ju}\Gamma_{jv}\Gamma_{jw}}{\Gamma_{j}^{3}} \right] p(\Delta u_{ij}), \tag{B7}$$

$$B_{(i=\alpha,x=\beta)} = 3 \sum_{y,z} \sum_{y,w,u} \sum_{i} \eta_{\beta} \left[\frac{\Gamma_{ix} \Gamma_{iy} \Gamma_{iz}}{\Gamma_{i}^{3}} \Gamma_{ij} \frac{\Gamma_{ju} \Gamma_{jv} \Gamma_{jw}}{\Gamma_{i}^{3}} \right] p(\Delta u_{ix}), \tag{B8}$$

$$C_{(i=\beta,x=\alpha)} = 3\sum_{y,z}\sum_{v,w,u}\sum_{j} \left[\frac{\Gamma_{ix}\Gamma_{iy}\Gamma_{iz}}{\Gamma_{i}^{3}}\Gamma_{ij}\frac{\Gamma_{ju}\Gamma_{jv}\Gamma_{jw}}{\Gamma_{j}^{3}} \right] p(\Delta u_{xi}).$$
 (B9)

Here we multiplied by three the terms B and C to account for the number of equivalent repetitions of the same configuration that can occur due to symmetries. η_{β} is the number of neighbours of the current site in the same state as the site being copied (i.e. the number of links that will change if the current site changes its configuration). η_{β} is always one for the C process, as it always changes just one link in the cluster. The quantity inside the square brackets gives the probability that the entire cluster will be in a given configuration $\{k\} = (x, y, z, v, w, u, i, j)$. Lastly $p(\Delta u_{ab})$ is given by the chosen microscopic transition rule, and gives the probability (given such configuration of the cluster) that the selected site (a) will copy its neighbour's (b) strategy, based on their payoff differences Δu_{ab} . The payoff of any is easy to be exactly obtained $u_i = G(i, x) + G(i, y) + G(i, z) + G(i, j)$ (G is the game matrix). For the payoff of any neighbouring site x, y or z we need to get the mean payoffs using a one site approximation, namely

$$\bar{u}_c = S + \rho_d^3(3S) + 3\rho_d^2\rho_c(2S + R) + 3\rho_d\rho_c^2(S + 2R) + \rho_c^3(3R),$$
 (B10)

$$\bar{u}_d = T + \rho_d^3(3\rho) + 3\rho_d^2\rho_c(2\rho + T) + 3\rho_d\rho_c^2(\rho + 2T) + \rho_c^3(3T).$$
 (B11)

Here \bar{u}_c is the mean payoff when the neighbour site is a cooperator and the focal site is a defector, and \bar{u}_d is the payoff of the opposite situation.

The counting of every possibility of a site changing its strategy is complicated and give rise to about 2^8 separated terms. Nonetheless, after determining a closed form for A, B and C for every rate W, it is a simple computational task to obtain all terms of $W_{\alpha\beta\to\gamma\beta}$. By doing so we obtain two coupled ODE's that can be numerically solved. We emphasize that this kind of mean-field results were already obtained in the literature [6, 21, 23, 31].

Returning to the mixed-game problem, we used two possible games $(G_1 \text{ and } G_2)$ being played at each interaction. To insert this in the mean-field model with cluster approximation we have now to reconsider all the possible site payoffs. Instead of having just one payoff matrix being used for each interaction, we now have several different matrix weighted by their probability. This means that we have to sum over all the possible payoffs obtained with each site's neighbour in each game. It is equivalently, in our rate terms W, to change the payoff function $p(u_{\alpha\beta})$ into a sum over all possible games $\left(\sum_{\{g\}} f_g(u_{\alpha\beta})\right)$ between the sites to be considered and its neighbours. It is important to notice however that the summation occurs over every neighbour of α and β as the payoff of each site is determined by the sum of games played with every neighbour.

As each of the two sites being considered for each rate W have four neighbours, this give rise to 2^6 possibilities of different game configurations on the cluster. Notice however that the ODE's remains covariant as Γ and the symmetry constants do not explicitly depend on the

game being played. We can average over $\{g\}$, a vector that gives the configuration of each game being played, using it is probability distribution $\Theta(g)$. By doing so we obtain the new rates

$$W_{\alpha\beta\to\gamma\beta}(\{k\};f) \Rightarrow \sum_{\{g\}} \Theta(g) W'_{\alpha\beta\to\gamma\beta} \Big(\{k\};f_g\Big). \tag{B12}$$

Making this change in the ODE and using a fourth order Runge–Kutta integration, we obtained the numerical results displayed in figure 2. Notice that as the payoff of each site depends on four other neighbours we need to consider every configuration of games being played by the sites in each term of W. This is why we cannot just sum up linearly the two different game matrices to obtain a mean payoff.

References

- [1] Pennisi E 2005 What don't we know? Science 309 93
- [2] Doebeli M and Knowlton N 1998 The evolution of interspecific mutualisms *Proc. Natl Acad. Sci.* 95 8676–80
- [3] Traulsen A, Shoresh N and Nowak M A 2008 Analytical results for individual and group selection of any intensity Bull. Math. Biol. 70 1410–24
- [4] Turner P E and Chao L 1999 Prisoner's dilemma in an rna virus Nature 398 441-3
- [5] Szathmáry E and Maynard Smith J 1995 The major evolutionary transitions Nature 374 227-32
- [6] Szabó G and Fath G 2007 Evolutionary games on graphs *Phys. Rep.* **446** 97–216
- [7] Nowak M and Sigmund K 1990 The evolution of stochastic strategies in the prisoner's dilemma *Acta Applicandae Math.* **20** 247–65
- [8] Nowak M A 2006 Evolutionary Dynamics (Cambridge, MA: Harvard University Press)
- [9] Perc M and Szolnoki A 2010 Coevolutionary games a mini review Biosystems 99 109-25
- [10] Wardil L and da Silva J K L 2013 The evolution of cooperation in mixed games Chaos Solitons Fractals 56 160-5
- [11] Rapoport A 1999 Two-person Game Theory (New York: Dover)
- [12] Skyrms B 2004 The Stag Hunt and the Evolution of Social Structure (Cambridge: Cambridge University Press)
- [13] Axelrod R 1984 The Evolution of Cooperation (New York: Basic Books)
- [14] Sigmund K 2012 Moral assessment in indirect reciprocity J. Theor. Biol. 299 25-30
- [15] Wardil L and da Silva J K L 2009 Adoption of simultaneous different strategies against different opponents enhances cooperation *Europhys. Lett.* **86** 38001
- [16] Wardil L and da Silva J K L 2010 Distinguishing the opponents promotes cooperation in well-mixed populations *Phys. Rev.* E 81 036115
- [17] Hauert C, De Monte S, Hofbauer J and Sigmund K 2002 Volunteering as red queen mechanism for cooperation in public goods games *Science* 296 1129–32
- [18] Nowak M A and May R M 1992 Evolutionary games and spatial chaos Nature 359 29
- [19] Wang Z, Szolnoki A and Perc M 2014 Different perceptions of social dilemmas: evolutionary multigames in structured populations *Phys. Rev.* E **90** 032813
- [20] Szolnoki A and Perc M 2014 Coevolutionary success-driven multigames Europhys. Lett. 108 28004
- [21] Hauert C and Szabó G 2005 Game theory and physics Am. J. Phys. 73 405-14
- [22] Huberman B A and Glance N S 1993 Evolutionary games and computer simulations *Proc. Natl Acad. Sci.* 90 7716–8
- [23] Matsuda H, Ogita N, Sasaki A and Satō K 1992 Statistical mechanics of population the lattice Lotka–Volterra model Prog. Theor. Phys. 88 1035–49
- [24] Szolnoki A, Perc M and Danku Z 2008 Towards effective payoffs in the prisoners dilemma game on scale-free networks *Physica A* 387 2075–82
- [25] Albert R and Barabási A L jan 2002 Statistical mechanics of complex networks Rev. Mod. Phys. 74 47–97
- [26] Santos F C and Pacheco J M 2006 A new route to the evolution of cooperation J. Evolutionary Biol. 19 726–33

- [27] Krapivsky P L and Redner S May 2001 Organization of growing random networks Phys. Rev. E 63 066123
- [28] Schuster P and Sigmund K 1983 Replicator dynamics J. Theor. Biol. 100 533-8
- [29] Maynard Smith J 1974 The theory of games and the evolution of animal conflicts *J. Theor. Biol.* 47 209–21
- [30] Taylor P D and Jonker L B 1978 Evolutionary stable strategies and game dynamics *Math. Biosci.* 40 145–56
- [31] Dieckmann U, Law R and Metz J A J 2000 *The Geometry of Ecological Interactions: Simplifying Spatial Complexity* (Cambridge: Cambridge University Press)