



Strategy abundance in 2×2 games for arbitrary mutation rates

Tibor Antal^{a,*}, Martin A. Nowak^a, Arne Traulsen^b

^a Program for Evolutionary Dynamics, Harvard University, Cambridge, MA 02138, USA

^b Max-Planck-Institute for Evolutionary Biology, 24306 Plön, Germany

ARTICLE INFO

Article history:

Received 16 September 2008

Received in revised form

7 November 2008

Accepted 19 November 2008

Available online 6 December 2008

Keywords:

Evolutionary game theory

Finite populations

Stochastic effects

ABSTRACT

We study evolutionary game dynamics in a well-mixed populations of finite size, N . A well-mixed population means that any two individuals are equally likely to interact. In particular we consider the average abundances of two strategies, A and B , under mutation and selection. The game dynamical interaction between the two strategies is given by the 2×2 payoff matrix $\begin{pmatrix} a & b \\ c & d \end{pmatrix}$. It has previously been shown that A is more abundant than B , if $a(N-2) + bN > cN + d(N-2)$. This result has been derived for particular stochastic processes that operate either in the limit of asymptotically small mutation rates or in the limit of weak selection. Here we show that this result holds in fact for a wide class of stochastic birth–death processes for arbitrary mutation rate and for any intensity of selection.

© 2008 Elsevier Ltd. All rights reserved.

1. Introduction

Evolutionary dynamics describe how successful strategies spread in a population through genetic reproduction or cultural imitation. In mutation–selection processes where the fitness of each individual is constant, the competition between two types A and B is straightforward. If both fitness values are identical, then we have neutral evolution (Kimura, 1968) and the average abundances of A and B are the same. (Throughout the paper we assume symmetric mutation rates: the mutation probability from A to B is the same as from B to A .) If A is fitter than B , then A is more abundant than B in the mutation selection equilibrium. Thus, for constant selection the comparison of the abundances of the two types is trivial.

For frequency dependent selection, where fitness depends on the types and abundances of others, the situation is more complex. One way to model these systems are evolutionary games, where the reproductive success of individuals depends on their payoff derived from interaction with others (Maynard Smith and Price, 1973; Maynard Smith, 1982; Weibull, 1995; Samuelson, 1997; Hofbauer and Sigmund, 1998, 2003; Fudenberg and Tirole, 1998; Nowak and Sigmund, 2004; Szabó and Fáth, 2007; Sandholm, 2007). In the simplest case, interactions are described by a 2×2 payoff matrix,

$$\begin{matrix} & A & B \\ A & \begin{pmatrix} a & b \\ c & d \end{pmatrix} \\ B & \end{matrix} \quad (1)$$

Here, an A individual obtains a from other A individuals, but b from B individuals. Similarly, B obtains c from A , and d from other B individuals.

For $a > c$ and $b > d$, strategy A dominates strategy B : Regardless of the composition of the population, strategy A has the higher payoff. Thus, evolutionary dynamics will always lead to a population with more A individuals than B individuals. Equivalently, for $a < c$ and $b < d$, strategy B dominates. In this case, the average abundance of B will be higher than the average abundance of A at the mutation–selection equilibrium.

For $a > c$ and $b < d$, both strategies are best replies to themselves. This is a ‘coordination game’. In a population of mostly A individual, rare B mutants have a lower payoff. In a population of mostly B individuals, rare A mutants have a lower payoff. Therefore, in a mutation selection process it is not a priori clear whether strategies A or B will be more abundant at the equilibrium distribution. Two concepts are important: (i) pareto efficiency and (ii) risk-dominance. Strategy A is Pareto efficient if $a > d$. In this case, an all- A population has a higher average payoff than an all- B population. Strategy A is risk dominant if $a + b > c + d$ (Harsanyi and Selten, 1988). In this case, strategy A has a larger basin of attraction than strategy B . In a game between two players, if I do not know what the other person will do, it is less risky for me to choose the risk-dominant strategy, but it would be more rewarding for both of us to choose the Pareto efficient strategy. A coordination game is especially interesting, if the risk-dominant strategy is not Pareto efficient.

Kandori et al. (1993) have shown that A is chosen over B if $a(N-2) + bN > cN + d(N-2)$. This means a large population selects the risk-dominant equilibrium in the long run. They analyze a process which is stochastic in the generation of mutants, but deterministic in following the gradient of selection. Their

* Corresponding author. Tel.: +1 617 496 5551.

E-mail address: tibor_antal@harvard.edu (T. Antal).

calculation assumes asymptotically small mutation rates, but holds for a wide range of evolutionary processes, in contrast to previous approaches that make specific assumptions on the source of noise (Foster and Young, 1990; Fudenberg and Harris, 1992). However, since the dynamics follows the gradient of selection, the processes can only leave a Nash equilibrium if a sufficient number of mutations occurs simultaneously. In large populations, it is very unlikely that this happens, and the time until the population moves from one equilibrium to another becomes exponentially large in N .

If the dynamics is stochastic, a single mutation can be sufficient to leave a Nash equilibrium. Nowak et al. (2004) have studied the competition of two strategies A and B in a frequency dependent Moran process (and similar processes) which model fully stochastic evolutionary dynamics. This means that the selection steps are stochastic. They have shown that the fixation probability of A is greater than that of B if and only if $a(N - 2) + bN > cN + d(N - 2)$. This calculation assumes weak selection in a process with selection only. The result is also valid for stronger intensities of selection, if we use the pairwise comparison process (Szabó and Tóke, 1998; Traulsen et al., 2007) or a slightly modified version of the Moran process for evolutionary updating (Traulsen et al., 2008).

Here, we extend these results to both arbitrary mutation rates and arbitrary intensities of selection. We show that A is more abundant than B in the mutation–selection distribution for a wide range of stochastic processes for any mutation rate and any intensity of selection if and only if $a(N - 2) + bN > cN + d(N - 2)$. Higher mutation rates seem to be very relevant for social evolution, because humans (and higher animals) tend to try new strategies fairly often. We therefore expect that evolutionary dynamics in the cultural learning and imitation context operate under fairly high mutation (= ‘exploration’) rates. We also show that the condition holds for any payoff matrix, not only for coordination games.

2. Pairwise comparison process

The payoffs of A and B in a population of j individuals of type A and $N - j$ individuals of type B are

$$\begin{aligned} \pi_A(j) &= \frac{j-1}{N-1}a + \frac{N-j}{N-1}b, \\ \pi_B(j) &= \frac{j}{N-1}c + \frac{N-j-1}{N-1}d. \end{aligned} \quad (2)$$

Each individual interacts with $N - 1$ other individuals. The payoff difference $\Delta\pi(j)$ is a linear function of j

$$\Delta\pi(j) = \pi_A(j) - \pi_B(j) = \frac{a-b-c+d}{N-1}j + \frac{-a+bN-dN+d}{N-1}. \quad (3)$$

First, we need to specify how strategies spread in the population, depending on their payoffs. We adopt the following process: a random (focal) individual i is selected. It compares its payoff π_i to the payoff π_j of a randomly chosen role model, individual j , and takes over the strategy of that individual with probability $[1 + e^{\beta(\pi_i - \pi_j)}]^{-1}$ (Blume, 1993; Szabó and Tóke, 1998; Traulsen et al., 2007; Sandholm, 2007). This process occurs with probability $1 - \mu$. With probability μ , a mutation occurs and the focal individual produces an offspring with random strategy, A or B . The quantity β determines the intensity of selection ($0 \leq \beta < \infty$). Strong selection (large β) means that the probability to adopt a better strategy approaches one, and the probability to adopt an inferior strategy vanishes. For weak selection (small β), these two probabilities are close to $\frac{1}{2}$. The transition probabilities to increase

or decrease the number of A players by one, are given by

$$\begin{aligned} T_j^+ &= \frac{j}{N} \frac{N-j}{N} \frac{1-\mu}{1+e^{-\beta\Delta\pi(j)}} + \frac{N-j}{N} \frac{\mu}{2}, \\ T_j^- &= \frac{j}{N} \frac{N-j}{N} \frac{1-\mu}{1+e^{+\beta\Delta\pi(j)}} + \frac{j}{N} \frac{\mu}{2}. \end{aligned} \quad (4)$$

With these transition probabilities, we can determine whether A is more abundant than B or vice versa. Before discussing the problem for general mutation rates we first consider the special case of low mutation rates, because of its importance in previous papers (Kandori et al., 1993; Nowak et al., 2004; Taylor et al., 2004; Imhof and Fudenberg, 2006). We will also present a simple approximation for very high mutation rates, which illustrates the dynamics in this extreme limit.

2.1. Low mutation rates

First, we consider low mutation rates, $\mu \ll N^{-2}$. In this limit, a mutation reaches extinction or fixation before a second mutation arises. Thus, the important quantity is the probability that a single mutant takes over the population in a process without mutations (Nowak et al., 2004; Taylor et al., 2004; Imhof and Fudenberg, 2006; Antal and Scheuring, 2006). The ratio of the fixation probability ρ_A of a single A player and that of a single B player can be written as (Karlin and Taylor, 1975; Nowak, 2006)

$$\frac{\rho_B}{\rho_A} = \prod_{j=1}^{N-1} \frac{T_j^-}{T_j^+}. \quad (5)$$

Without mutations, $\mu = 0$, the ratio of the transition probabilities (4), simplifies to $T_j^-/T_j^+ = e^{-\beta\Delta\pi(j)}$ (Traulsen et al., 2006, 2007). Thus, ρ_B/ρ_A reduces to

$$\frac{\rho_B}{\rho_A} = \exp\left[-\beta \sum_{j=1}^{N-1} \Delta\pi(j)\right] = \exp\left[-\beta \left(\frac{a+b-c-d}{2}N - a + d\right)\right]. \quad (6)$$

Therefore, $\rho_A = \rho_B$ is equivalent to

$$(a+b-c-d)N = 2a - 2d. \quad (7)$$

From the payoff matrix (2) it is clear that strategy A is favored whenever a or b increases, or c or d decreases. Therefore, A is more abundant than B if

$$a(N-2) + bN > cN + d(N-2). \quad (8)$$

This condition has been derived previously in the low mutation limit for different evolutionary processes (Kandori et al., 1993; Nowak, 2006; Traulsen et al., 2008). The strategy with the higher fixation probability is the more abundant one, because more of its mutants reach fixation. Thus for low mutation rates, A has a higher abundance than B whenever condition (8) is fulfilled.

For weak selection, a wide range of processes fulfills the $\frac{1}{3}$ -rule, which states that the fixation probability of a single A mutant in a coordination game is larger than $1/N$ if $a(N-2) + b(2N-1) > c(N+1) + d(2N-4)$ (Nowak et al., 2004; Imhof and Nowak, 2006; Lessard and Ladret, 2007; Bomze and Pawlowitsch, 2008). The fixation probability for large N can thus be written as

$$\rho_A = \frac{1}{N} + \omega \left(\frac{1}{3} - \frac{d-b}{a-b-c+d}\right), \quad (9)$$

where ω is a small positive number that may depend on the payoffs and on N . Writing down the analogous equation for ρ_B and subtracting yields

$$\rho_A - \rho_B = \omega \left(\frac{a-c}{d-c-b+a} - \frac{d-b}{a-b-c+d}\right) \propto a+b-c-d. \quad (10)$$

Thus, $\rho_A > \rho_B$ is equivalent to $a + b > c + d$, which is identical to our condition (8) for large N . Lessard and Ladret (2007) have shown that any process within the domain of Kingman's (1982) coalescent fulfills the $\frac{1}{3}$ -rule (for large N and weak selection and mutation). If an evolutionary process fulfills the $\frac{1}{3}$ -rule, ρ_A and ρ_B can be written in the form of (9). Hence we find again that $\rho_A > \rho_B$ is equivalent to $a + b > c + d$. Thus, we conclude that our result is valid for any process within the domain of Kingman's coalescent for large N and weak selection and mutation.

2.2. High mutation rates

For high mutation rates $\mu \rightarrow 1$, mutations dominate the process, and drive the system towards equal abundance of A and B , that is close to $j = N/2$. Hence the relevant transition probabilities are $T_{N/2}^+$ and $T_{N/2}^-$, and it is plausible to assume that $T_{N/2}^+ > T_{N/2}^-$ implies that A is more abundant than B in the stationary state. From (4) we obtain

$$T_{N/2}^+ - T_{N/2}^- = \frac{1 - \mu}{4} \tanh\left(\frac{\beta}{2} \Delta\pi(N/2)\right), \tag{11}$$

where the payoff difference is given by

$$\Delta\pi(N/2) = \frac{(a + b - c - d)N - 2(a - d)}{2(N - 1)}. \tag{12}$$

Since $\Delta\pi(N/2) > 0$ implies $T_{N/2}^+ - T_{N/2}^- > 0$, it also implies a more abundant A , for arbitrary intensity of selection β . Thus, we have again the same condition $a(N - 2) + bN > cN + d(N - 2)$ for the dominance of A .

2.3. Arbitrary mutation rates

So far we have shown that both for low and high mutation rates the same condition determines whether A or B is more abundant, regardless of the intensity of selection. Now, we turn to general mutation rates. Whenever Eq. (7) holds, the payoff difference (3) has the following symmetry property:

$$\Delta\pi(j) = -\Delta\pi(N - j). \tag{13}$$

This antisymmetry in the payoff differences then implies the following symmetry in our transition probabilities (4),

$$T_j^+ = T_{N-j}^-. \tag{14}$$

For symmetric transition probabilities, however, the stationary probabilities p_j of having j number of A players, are also symmetric, $p_j = p_{N-j}$. This can be shown by first writing the stationary probability distribution explicitly (Kampen, 1997; Claussen and Traulsen, 2005)

$$p_j = p_0 \prod_{i=0}^{j-1} \frac{T_i^+}{T_{i+1}^-}, \tag{15}$$

where p_0 follows from the normalization $\sum_{j=0}^N p_j = 1$. For symmetric transition probabilities (14) and for $j < N - j$ we can write

$$\frac{p_{N-j}}{p_j} = \prod_{i=j}^{N-j-1} \frac{T_i^+}{T_{i+1}^-} = \prod_{i=j}^{N-j-1} \frac{T_i^+}{T_{N-i-1}^+} = 1. \tag{16}$$

Hence the distribution is symmetric $p_j = p_{N-j}$.

For such a symmetric distribution, the number of A and B players are of course the same, since

$$\langle j \rangle = \sum_{j=0}^N j p_j = \sum_{j=0}^N j p_{N-j} = \sum_{j=0}^N (N - j) p_j = \langle N - j \rangle. \tag{17}$$

Hence we have demonstrated that when (7) is fulfilled, then the average abundance of A players and B players are the same.

On the other hand, when condition (8) is fulfilled, then the average abundance of A players is higher than the average abundance of B players. This is quite obvious by looking at the payoff matrix (1): increasing a or b favors A , while increasing c or d favors B . More formally, we could say that from (2)–(4) it is clear that T_j^+ and $-T_j^-$ are monotone increasing functions of a and b , and decreasing functions of c and d . This implies condition (8). Interestingly, other features of the game do not matter, e.g. whether a single A player has a larger disadvantage in a B population than a single B player in an A population.

Condition (8) can also be written in the form

$$a\left(1 - \frac{2}{N}\right) + b > c + d\left(1 - \frac{2}{N}\right), \tag{18}$$

which highlights the fact that there is a $1/N$ correction compared to simple risk dominance ($a + b > c + d$), which is the $N \rightarrow \infty$ limit. This implies that a strategy can be more abundant than the other strategy for large N , but less abundant for small N . For example for the payoff matrix $\begin{pmatrix} 1 & 0 \\ 0 & 1 \end{pmatrix}$, strategy A is more abundant in large populations ($N \geq 19$), but B is more abundant in small populations ($N \leq 17$). They are equally abundant for $N = 18$. In general, there is a threshold population size $N^* = 2(a - d)/(a - d + b - c)$. If $N^* > 2$, then A is more abundant than B either for $N > N^*$ or for $N < N^*$.

2.4. Including self-interactions

So far, we have adopted the usual convention that individuals cannot interact with themselves. If instead we allow individuals to derive a payoff from self-interaction, we obtain

$$\begin{aligned} f_A(j) &= \frac{j}{N}a + \frac{N-j}{N}b, \\ f_B(j) &= \frac{j}{N}c + \frac{N-j}{N}d. \end{aligned} \tag{19}$$

Hence, the payoff difference is $\Delta\pi(j) = (a - b - c + d)j/N + b - d$. Now for all mutation rates and all intensities of selection the condition for the average abundance of A to exceed the average abundance of B is simple risk dominance:

$$a + b > c + d. \tag{20}$$

We conclude that the finite N correction to risk dominance in (18) results from the exclusion of self-interactions.

3. Frequency dependent Moran process

Although condition (8) is valid for a large class of evolutionary processes (see Section 4), it does not always hold if we depart from weak selection. As an example we discuss the frequency dependent Moran process (Nowak et al., 2004; Taylor et al., 2004; Antal and Scheuring, 2006). As we shall see, for this model our condition (8) is only valid in the limit of weak selection.

Let the fitness be a convex combination of a background fitness (which we set to 1) and the payoff, $f_A(j) = 1 - w + w\pi_A(j)$ and $f_B(j) = 1 - w + w\pi_B(j)$. Here, w is the intensity of selection ($0 \leq w \leq 1$ for payoff matrices with positive entries). An individual is selected for reproduction at random, but proportional to fitness. The selected individual produces an offspring, which replaces a randomly chosen individual. Mutation can occur during reproduction. This leads to the transition probabilities

$$\begin{aligned} T_j^+ &= \frac{j f_A(j)}{F(j)} \frac{N-j}{N} (1 - \mu) + \frac{(N-j) f_B(j)}{F(j)} \frac{N-j}{N} \mu, \\ T_j^- &= \frac{(N-j) f_B(j)}{F(j)} \frac{j}{N} (1 - \mu) + \frac{j f_A(j)}{F(j)} \frac{j}{N} \mu. \end{aligned} \tag{21}$$

Here $F(j) = jf_A(j) + (N - j)f_B(j)$ is the total fitness of the whole population. These transition probabilities depend on the fitness values only through the ratio $f_A(j)/f_B(j)$. Yet, there is no simple condition for the equilibrium abundance of A players. In the weak selection limit, $w \ll 1$, however, only the payoff differences enter into the transition probabilities (21),

$$\frac{f_A(j)}{f_B(j)} = 1 + w\Delta\pi(j) + \mathcal{O}(w^2). \quad (22)$$

Thus, for weak selection, condition (8) again ensures the higher abundance of A .

We can also consider a variant of the frequency dependent Moran process where fitness is an exponential function of payoff, $f_A(j) = \exp[+w\pi_A(j)]$ and $f_B(j) = \exp[+w\pi_B(j)]$ (Traulsen et al., 2008). In this case, the transition probabilities again only depend on the payoff differences, because $f_A(j)/f_B(j) = \exp(w\Delta\pi(j))$. Hence, the condition for abundance is again (8) for any intensity of selection and any mutation rate.

4. General birth–death processes

Here, we show that our finding holds for a wide range of birth–death processes. We need two requirements to be fulfilled. The first is that the payoffs received from other players should be additive. In this case the payoffs $\pi_A(j)$, $\pi_B(j)$, and also the payoff difference $\Delta\pi(j)$ are linear functions of j . Consequently,

$$\Delta\pi(j) + \Delta\pi(N - j) = \lambda(a, b, c, d, N) \quad (23)$$

is independent of j . Hence, when $\lambda(a, b, c, d, N) = 0$ holds, we have an antisymmetric payoff function

$$\Delta\pi(j) = -\Delta\pi(N - j). \quad (24)$$

The second requirement is that the difference between the two types of players should manifest itself in the transition probabilities T_j^+ and T_j^- , only through the payoff difference $\Delta\pi(j)$. Hence, we can write $T_j^- = T_j^+[\Delta\pi(j), \mu]$, and the probability of decreasing the number of mutants is

$$T_j^- = T_j^+[\Delta\pi(j), \mu] = T_{N-j}^+[-\Delta\pi(j), \mu]. \quad (25)$$

That is, the only difference between the two types is the change of sign of the payoff difference $\Delta\pi(j)$. Otherwise, the transition probabilities are the same, but with the number of the opposite type of individuals $N - j$. Now, simply setting the index to $N - j$ in (25), we have

$$T_{N-j}^- = T_j^+[-\Delta\pi(N - j), \mu] = T_j^+[\Delta\pi(j), \mu] = T_j^+. \quad (26)$$

For the second equality we have used (24). These symmetric transition probabilities then imply an equal abundance of the two types of players in any birth–death process, as it has been shown in Section 2.3. The condition for a higher abundance of A compared to B is then $\lambda(a, b, c, d, N) > 0$. This general condition takes the form of (8) or (20) in our examples when self-interaction are excluded or included, respectively. Note that similar arguments appear in Claussen (2007).

5. Discussion

In this paper, we have shown that $a(N - 2) + bN > cN + d(N - 2)$ is the crucial condition for A to be favored over B in wide range of evolutionary processes for any mutation rate, any intensity of selection and any finite population size. By A being favored over B we mean that A is more abundant than B in the mutation–selection equilibrium of the stochastic process.

The relative abundance in the stationary state is a natural way of comparing two strategies. In the low mutation limit, when the population is typically homogeneous, a strategy being more abundant is equivalent of having a larger fixation probability. Hence, our study of abundance can be viewed as a generalization of that concept to arbitrary mutation rates. This is of special interest for cultural dynamics, where mutation rates are not necessarily small.

In particular, we have generalized the famous result of Kandori et al. (1993) to any mutation rate. We have generalized the result of Nowak et al. (2004) to any intensity of selection. Our results are valid for any mutation rate and any intensity of selection.

It turns out that for our result to hold, a birth–death process has to fulfill two requirements: (i) additive payoffs, and (ii) that the evolutionary dynamics depend only on the payoff differences (the players are identical otherwise). These requirements hold, for example, for the pairwise comparison process described by Traulsen et al. (2006) and for a frequency dependent Moran process with exponential fitness function (Traulsen et al., 2008). For the standard frequency dependent Moran process with linear fitness function these requirements only hold in the limit of weak selection.

Finally, we note that in coordination games with strong selection both the fixation probability of A and the fixation probability of B are very small. For small mutation rates, the system will stay in one equilibrium for a very long time. Therefore, it can take a very long time for the system to obtain a representative sample of the stationary mutation–selection distribution.

Acknowledgements

We are grateful for financial support from NIH Grant R01GM078986, The John Templeton Foundation, Jeffrey Epstein (T.A. and M.A.N.) and the Emmy-Noether program of the DFG (A.T.).

References

- Antal, T., Scheuring, I., 2006. Fixation of strategies for an evolutionary game in finite populations. *Bull. Math. Biol.* 68, 1923–1944.
- Blume, L.E., 1993. The statistical mechanics of strategic interaction. *Games and Economic Behavior* 4, 387–424.
- Bomze, I., Pawłowski, C., 2008. One-third rules with equality: second-order evolutionary stability conditions in finite populations. *J. Theor. Biol.* 254, 616–620.
- Claussen, J.C., 2007. Drift reversal in asymmetric coevolutionary conflicts: influence of microscopic processes and population size. *Eur. Phys. J. B* 60, 391–399.
- Claussen, J.C., Traulsen, A., 2005. Non-Gaussian fluctuations arising from finite populations: exact results for the evolutionary Moran process. *Phys. Rev. E* 71, 025101(R).
- Foster, D., Young, P., 1990. Stochastic evolutionary game dynamics. *Theor. Popul. Biol.* 38, 219–232.
- Fudenberg, D., Harris, C., 1992. Evolutionary dynamics with aggregate shocks. *J. Econ. Theory* 57, 420–441.
- Fudenberg, D., Tirole, J., 1998. *Game Theory*, sixth ed. MIT Press, Cambridge, MA.
- Harsanyi, J.C., Selten, R., 1988. *A General Theory of Equilibrium Selection in Games*. MIT Press, Cambridge, MA.
- Hofbauer, J., Sigmund, K., 1998. *Evolutionary Games and Population Dynamics*. Cambridge University Press, Cambridge.
- Hofbauer, J., Sigmund, K., 2003. Evolutionary game dynamics. *Bull. Am. Math. Soc.* 40, 479–519.
- Imhof, L.A., Fudenberg, D., 2006. Imitation process with small mutations. *J. Econ. Theory* 131, 251–262.
- Imhof, L.A., Nowak, M.A., 2006. Evolutionary game dynamics in a Wright Fisher process. *J. Math. Biol.* 52, 667–681.
- Kampen, N.G.V., 1997. *Stochastic Processes in Physics and Chemistry*, second ed. Elsevier, Amsterdam.
- Kandori, M., Mailath, G.J., Rob, R., 1993. Learning, mutation, and long run equilibria in games. *Econometrica* 61, 29–56.

- Karlin, S., Taylor, H.M.A., 1975. *A First Course in Stochastic Processes*, second ed. Academic, London.
- Kimura, M., 1968. DNA and the neutral theory. *Philos. Trans. R. Soc. London B* 312, 343–354.
- Kingman, J.F.C., 1982. The coalescent. *Stochastic Process. Appl.* 13, 235–248.
- Lessard, S., Ladret, V., 2007. The probability of fixation of a single mutant in an exchangeable selection model. *J. Math. Biol.* 54, 721–744.
- Maynard Smith, J., 1982. *Evolution and the Theory of Games*. Cambridge University Press, Cambridge.
- Maynard Smith, J., Price, G.R., 1973. The logic of animal conflict. *Nature* 246, 15–18.
- Nowak, M.A., 2006. *Evolutionary Dynamics*. Harvard University Press, Cambridge, MA.
- Nowak, M.A., Sigmund, K., 2004. Evolutionary dynamics of biological games. *Science* 303, 793–799.
- Nowak, M.A., Sasaki, A., Taylor, C., Fudenberg, D., 2004. Emergence of cooperation and evolutionary stability in finite populations. *Nature* 428, 646–650.
- Samuelson, L., 1997. *Evolutionary Games and Equilibrium Selection*. MIT Press, Cambridge, MA.
- Sandholm, W.H., 2007. *Population Games and Evolutionary Dynamics*. MIT Press, Cambridge, MA.
- Szabó, G., Fáth, G., 2007. Evolutionary games on graphs. *Phys. Rep.* 446, 97–216.
- Szabó, G., Tóke, C., 1998. Evolutionary prisoner's dilemma game on a square lattice. *Phys. Rev. E* 58, 69.
- Taylor, C., Fudenberg, D., Sasaki, A., Nowak, M.A., 2004. Evolutionary game dynamics in finite populations. *Bull. Math. Biol.* 66, 1621–1644.
- Traulsen, A., Nowak, M.A., Pacheco, J.M., 2006. Stochastic dynamics of invasion and fixation. *Phys. Rev. E* 74, 11909.
- Traulsen, A., Pacheco, J.M., Nowak, M.A., 2007. Pairwise comparison and selection temperature in evolutionary game dynamics. *J. Theor. Biol.* 246, 522–529.
- Traulsen, A., Shresh, N., Nowak, M.A., 2008. Analytical results for individual and group selection of any intensity. *Bull. Math. Biol.* 63, 363.
- Weibull, J., 1995. *Evolutionary Game Theory*. MIT Press, Cambridge, MA.