Population dynamics of mutualisms

Chaitanya S. Gokhale *

*New Zealand Institute for Advanced Study, Massey University, Auckland, New Zealand

Mutualistic relationships pose a conundrum for evolutionary theory. Species that exploit other species would do better than sustaining a long drawn out mutually costly relationship. However we do see mutualistic relationships amongst even the most unlikely partners . . . . Eco-evolutionary dynamics . . .

mutualism | evolutionary game theory | multiple players

Contents

    ToC: Just for us while we write to keep the big picture in mind

Introduction

As with many concepts, we can trace back the study of mutualism to Aristotle [**?**]. However formally in 1873 the Belgian zoologist Pierre van Beneden coined the term mutualism [1]. Mutualistic relationships, interspecific interactions that benefit both species, have been empirically studied for many years [2, 3, 4, 5, 6, 7, 8] and a considerable body of theory has been put forth explaining the evolution and maintenance of such relationships [9, 10, 11, 12, 13, 14, 15, 16]. Most examples of mutualisms, including the one described by Aristotle, lend themselves to the idea of direct reciprocity [17] and have thus been extensively studied using evolutionary game theory. The interactions in these models are usually dyadic: the fundamental interaction is between two individuals, one from each species, and the sum of many such interactions determines the evolutionary dynamics. However, in many cases interactions between species cannot be reduced to such dyadic encounters [18].

For example, in the interaction between ants and aphids or butterfly larvae [19, 20] many ants tend to each of the soft bodied creatures, providing them with shelter and protection from predation and parasites, in exchange for honeydew, a rich source of food for the ants [21, 18]. This is not a one-to-one interaction between a larva and an ant, but rather a one-to-many interaction from the perspective of the larva. Another well studied example of a one-to-many interaction is that of the plant-microbe mutualism wherein leguminous hosts prefer rhizobial symbionts that fix more nitrogen [7], or where plants provide more carbon resources to the fungal strains that are providing better access to nutrients [22]. Identifying and quantifying the intraspecific variation can be a daunting task [23], and the interactions within the rhizobial symbionts community itself are usually ignored in the broader picture of the between species interactions. Another possible example : bioluminescent bacteria in *Vibrio* squid, which like the legumes is a zillions-to-one mutualism.

Similarly for the interactions between the cleaner fish and their hosts [24, 25]. While the cohorts of cleaner fish together have been taken to determine the quality of a cleaning station, this can also drive variation of quality of cleaning within a cleaning station as per the interactions of individual cleaner fish amongst themselves. Furthermore, since by definition mutualistic relationships are between species, it is natural to imagine that the observed relationship may be seasonal and the interactions as not a continuous feature of the evolutionary dynamics of a single species. To assess the impact of this seasonality . . . In this manuscript we focus on this kind of – possibly – many to many interactions between two mutualistic species.

This paragraph - is it about 1-1 vs 1-many, or dyadic vs ...? Seems like it's setting up to be a "Dyadic isn't enough" para. In that case fine but the next para should be making the (main) point that intra-specific interactions are perhaps an important, and hitherto ignored, determinant of evolutionary outcomes for mutualisms.

In all, in this manuscript we look at the broader picture of mutualistic relationships and the ecology in which they are observed. We study the the cumulative effect of the within species and between species interactions and the importance of these relationships, seasonality and population dynamics taken together. To analyze how benefits are shared between the two mutualistic species, we make use of evolutionary game theory. Since we consider the interaction of two species, we resort to bimatrix games [26, 27, 28]. While initially attempting to avoid the question of how mutualisms evolve in the first place, we see that when studying the complex dynamics which are possible due to the rest of the evolutionary as well as ecological factors, we indirectly explain the evolution of interspecific mutualism as a byproduct of a complex eco-evolutionary process.

Model and Results

$$\dot{x} = r_x x \left(f_{G_1}(x, y) - \bar{f}_1(x, y) \right)$$
$$\dot{y} = r_y y \left(f_{G_2}(x, y) - \bar{f}_2(x, y) \right). \qquad [\mathbf{1}]$$

Usually when interspecies relationships such as mutualism (or antagonist relationships as in predator-prey) are considered, the within species interactions are ignored for the sake of convenience. Including intraspecies interactions can however result in qualitatively different and rich dynamics. In fact the coevolutionary dynamics between the two species is determined together by the inter as well as the intraspecific interactions.

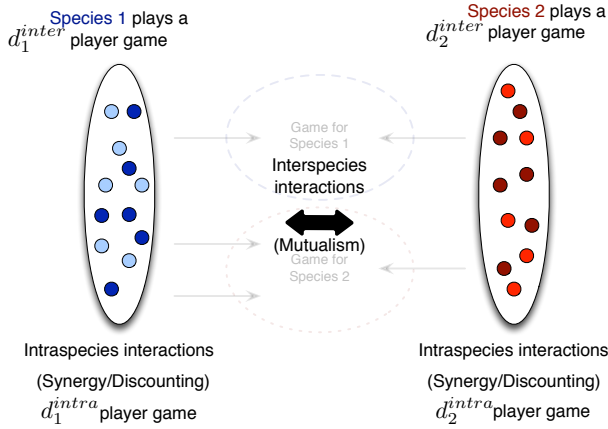
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Fig. 1. Evolutionary dynamics with combined inter-intra species dynamics. We assume the interactions between species to be mutualistic described by the snowdrift game [13, 29, 30]. Species 1 plays a d_1^{inter} player game with Species 2 while Species 2 plays a d_2^{inter} player game. Each species has two types of players "Generous" and "Selfish" who besides interacting with the members of other species, also take part in intra species dynamics. We assume a general framework of synergy and discounting for the intraspecies interactions [31, 32]

Interspecies dynamics. Since we focus on mutualism the interspecies dynamics is given by the multiplayer version of the snowdrift game [13, 29, 30]. Each species consists of two types of individuals Generous G and Selfish S. The details of the game are included in the Supplementary Material (SI), but the gist is that if everyone is "Generous" and contributing in the generation of mutual benefits then one can get away with being a bit selfish. However both species cannot be completely "selfish" by definition of mutualism. Hence the pressure is on a specie in making the partner "Generous" while getting away itself by being "Selfish". The fitness of each of the types within a species thus depends on the composition of the other species. For example if the frequency of the "Generous" types in Species 1 (G_1) is x and that in Species 2 (G_2) is y then fitness of G_1 is given by $f_{G_1}^{inter}(y)$ and that of G_2 as $f_{G_2}^{inter}(x)$

Intraspecies dynamics. We do not restrict ourselves to any particular interaction structure and thus can make use of the general multiplayer evolutionary games framework [33, 34]. Moving from the interspecies dynamics, the two types already described are "Generous" and "Selfish". Thus we already have each species containing two different types of individuals. It is possible that a different categorisation exists within a species however for the sake of simplicity we study the dynamics between "Generous" and "Selfish" types within a species. However the individuals which are "Generous" for the interspecies interaction may/may not be more giving or in a sense "Co-operators" for intraspecies dynamics. Thus we need a flexible cost-benefit framework to model the intra species dynamics which can be easily tuned to the particular situation. The cost benefit framework described in [31, 32] allows us to transition between four classic scenarios of evolutionary dynamics [35].

Describe the four outcomes

For the intra species interactions the fitness of a G_1 is then given by $f_{G_1}^{intra}(x)$ and that of G_2 is given by $f_{G_2}^{intra}(y)$ and similarly for the "Selfish" types.

Combined dynamics. Putting together intra and interspecific dynamics provides a complete picture of the possible interactions occurring. While we are interested in mutualism at the level of the interspecies interactions there are four possible interactions within each species [35, 32]. Since the within species interactions for the two different species do not need to be the same, there are in all sixteen different possible combinations. Assuming additivity in the fitnesses of inter and intraspecies fitnesses, the combined fitness of each of the two types in the two species are given by,

$$
\begin{aligned}
f_{G_1}(x, y) &= p f_{G_1}^{inter}(y) + (1-p) f_{G_1}^{intra}(x) \\
f_{S_1}(x, y) &= p f_{S_1}^{inter}(y) + (1-p) f_{S_1}^{intra}(x) \\
f_{G_2}(x, y) &= p f_{G_2}^{inter}(x) + (1-p) f_{G_2}^{intra}(y) \\
f_{S_2}(x, y) &= p f_{S_2}^{inter}(x) + (1-p) f_{S_2}^{intra}(y)
\end{aligned}
\qquad [\mathbf{2}]
$$

The parameter p tunes the impact of each of the interactions on the actual fitness that eventually drives the evolutionary dynamics. For $p = 1$ we recover the well studied case of the Red King dynamics [30] while for $p = 0$ the dynamics of the two species are essentially decoupled and can be individually studied by the synergy/discounting framework of nonlinear social dilemmas [32]. Of interest in the continuum and the intermediate values of p. However that would mean we need to track the qualitative dynamics of sixteen possible intraspecies dynamics as p changes gradually from close to 0 to close to 1 (SI).

This approach provides us with a powerful method to incorporate a multitude of realistic concepts in the analysis. For example the number of players involved in a game, which has been shown to be a crucial factor in determining the evolutionary dynamics could be different for each interactions, inter and intra species interactions for Species 1 (d_1^{inter}, d_1^{intra}) and similarly for Species 2 (d_2^{inter}, d_2^{intra}). The interspecies interactions are proxied by the multiplayer Snowdrift game which can incorporate threshold effects. For example a certain number of "Generous" cleaner fish may be required to clean the host or a certain number of "Generous" ants required to protect larva from predators. We can have M_1 and M_2 as the thresholds in the two species. Since the interaction matrices for the inter and intra species dynamics are completely different there doesn't need to be any relationship between the costs and benefits of the four games (Two snowdrift games from the perspective of each species and the intragames within each species).

So in principle we can have a diverse and rich set of dynamics possible which brings into question the study of coevolution based on only interspecies interactions. Even if we make a large number of assumptions and even if the intraspecies dynamics accounts for only 33% $(1-p)$ of the cumulative fitness, we can see drastically different qualitative dynamics which is capable of explaining the persistence of exploiters.
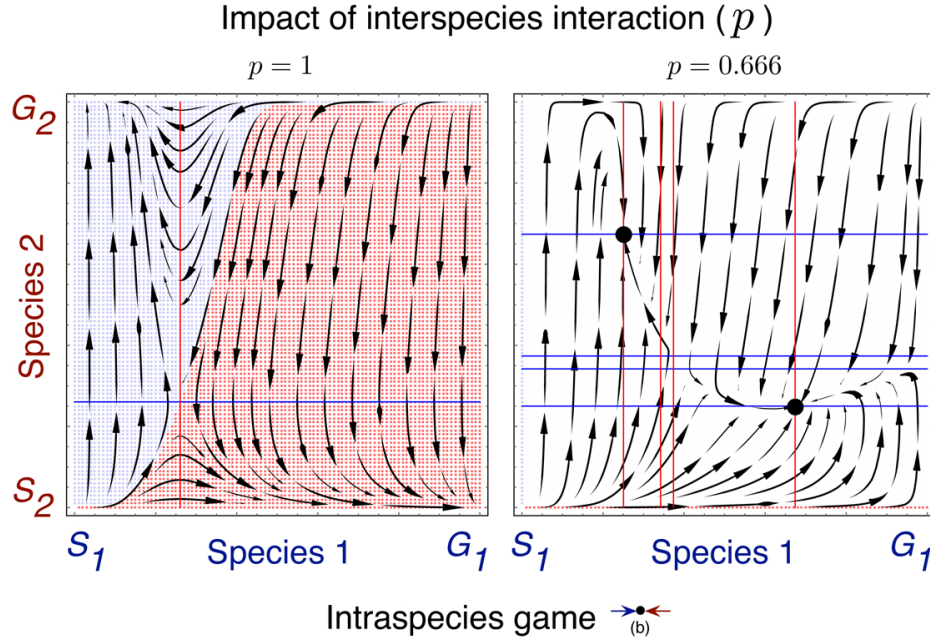
Impact of interspecies interaction (p)

Fig. 2. Change in evolutionary dynamics due to inclusion of intraspecies dynamics. When the fitness of the "Generous" and "Selfish" types in both the species is solely determined by the interactions which occur between species (in this case mutualism, $p = 1$) then we recover the dynamics as studied previously in [30]. The colours represent the initial states which result in an outcome favourable for Species 1 (blue leading to (S_1,G_2)) and Species 2 (red, leading to (G_1,S_2)). This can result in the red King effect and other possible complexities as discussed recently in [36]. However when we start including intraspecies dynamics the picture can be very different. Even when the impact of intraspecies dynamics is only a $1/3$ on the total fitness of the "Generous" and "Selfish" types we see a very qualitatively different picture. Two fixed points are observed where both the "Generous" and "Selfish" types can co-exist in both the species. All initial states in the interior lead to either one of these fixed points (hence the lack of colours). However it is still possible to characterize the "successful" species as one of the equilibrium is favoured by one species than the other. The horizontal isoclines are for Species 1 while the vertical ones are for Species 2. The analysis was done for a 5 player game $d_1^{inter} = d_2^{inter} = d_1^{intra} = d_2^{intra} = 5$, $b = 2$, $c = 1$ and $r_x = r_y/8$ for the interspecies mutualism game while additionally $\tilde{b}_1 = \tilde{b}_2 = 10$ and $\tilde{c}_1 = \tilde{c}_2 = 1$ and $\omega_1 = \omega_2 = 3/4$ for the two intraspecies games within each species. Note that even with symmetric games within each species we can a qualitatively drastic difference when compared to the dynamics excluding intraspecies interactions. For different intraspecies interactions within each species and for varying p see SI.

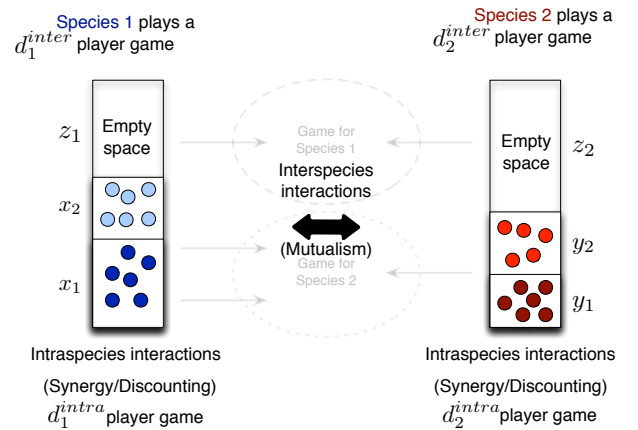## Population dynamics.

Fig. 3. Explain figure

Until now we have considered that each species consists of two types of individuals and they make up the population of that species. However populations sizes change over time. Assuming that ecological changes are fast enough that they can be averaged out, we can usually ignore their effect on the evolutionary dynamics. It is now possible to show that evolution can happen at fast timescales as well, comparable to those of the ecological dynamics add citations with examples. Hence we need to tackle not just evolutionary but eco-evolutionary dynamics together.

To include population dynamics in the previously considered scenario, we reinterpret x_1 now as the fraction of "Generous" types and x_2 as the fraction of "Selfish" types in Species 1. Also now we have $z_1 = 1 - x_1 - x_2$ as the empty spaces in the niche occupied by Species 1. Similarly we have y_1, y_2 and z_2 (Fig. 3). This approach has previously been explored in terms of social dilemmas in [37]. We adapt and modify it for the two species and hence now the dynamics of this complete system is determined by the following set of differential equations,

$$\dot{x_1} = r_x x_1 (z_1 f_{G_1} - e_1)$$
$$\dot{x_2} = r_x x_2 (z_1 f_{S_1} - e_1) \qquad [3]$$
$$\dot{z_1} = -\dot{x_1} - \dot{x_2}$$

and for species 2

$$\dot{y_1} = r_y y_1 (z_2 f_{G_2} - e_2)$$
$$\dot{y_2} = r_y y_2 (z_2 f_{S_2} - e_2) \qquad [4]$$
$$\dot{z_2} = -\dot{y_1} - \dot{y_2}$$

where we have introduced e_1 and e_2 as the death rates of the two species. Setting $e_1 = \frac{z_1(x_1 f_{x_1} + x_2 f_{x_2})}{x_1 + x_2}$ and $e_2 =$

$\frac{z_2(y_1 f_{G_2} + y_2 f_{S_2})}{y_1 + y_2}$ we recover the two species replicator dynamics as in Eqs. 1. In this setup however the fitnesses need to be re-evaluated as not we need to account for the presence of empty spaces (See SI). We can reduce the dynamics by looking at only the proportion of "Generous" types in both the species thus $g_1 = x_1/(1 - z_1)$ and $g_2 = y_1/(1 - z_2)$ whose time evolution is given by,

$$\dot{g_1} = r_x z_1 g_1 (1 - g_1)(f_{G_1} - f_{S_1})$$
$$\dot{z_1} = e_1(1 - z_1) - r_x z_1(1 - z_1)(g_1 f_{G_1} - (1 - g_1)f_{S_1}) \qquad [5]$$

and

$$\dot{g_2} = r_y z_2 g_2 (1 - g_2)(f_{G_2} - f_{S_2})$$
$$\dot{z_2} = e_2(1 - z_2) - r_y z_2(1 - z_2)(g_2 f_{G_2} - (1 - g_2)f_{S_2}) \qquad [6]$$

where everywhere we have $x_1 = g_1(1 - z_1)$ (with $x_2 = (1-g_1)(1-z_1)$) and $y_1 = g_2(1-z_2)$ (with $y_2 = (1-g_2)(1-z_2)$) in the fitnesses as well.

Such a two species multi-type interaction system is a complicated as well as a realistic depiction of most of the mutualisms observed in nature. However even with this complexity, the reduction of variables from six to four allows us to study the eco-evolutionary dynamics of the mutualism by looking at the two species simultaneously. As shown in Figure ?? we plot the evolutionary information (fraction of "Generous" in each species) against the ecological parameter, the population density (or rather in this case the empty space which is $1-z_{1/2}$ population density).
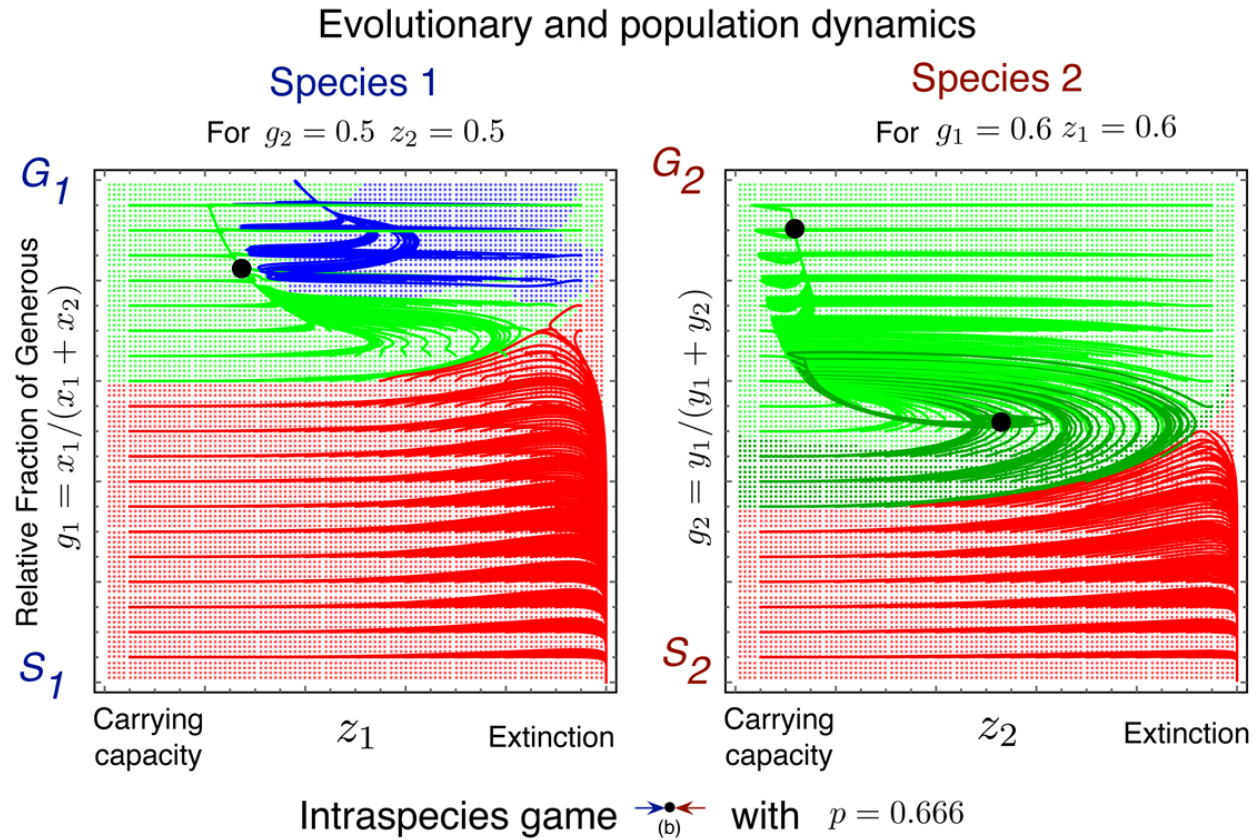
Fig. 4. **Dynamics of evolutionary strategies and population density for an intraspecies coexistence game with interspecies mutualism.** With exactly the same parameters as that of Figure 2 with symmetric death rates $e_1 = e_2 = 0.05$.

Discussion

1. Bronstein, JL (2003) in Genetic and Cultural Evolution of Cooperation, ed Hammerstein, P (MIT Press).

2. Boucher, DH (1985) in The Biology of Mutualism, ed Boucher, DH (Oxford University Press, New York), pp 1–28.

3. Hinton, HE (1951) Myrmecophilous lycaenidae and other lepidoptera - a summary. Proc. Trans. S. London Entomol. Nat. Hist. Soc. 1949-50:111–175.

4. Wilson, DS (1983) The effect of population structure on the evolution of mutualism: A field test involving burying beetles and their phoretic mites. The American Naturalist 121:851–870.

5. Bronstein, JL (1994) Our current understanding of mutualism. The Quarterly Review of Biology 69:31–51.

6. Pierce, NE et al. (2002) The Ecology and Evolution of Ant Association in the Lycaenidae (Lepidoptera). Annual Review of Entomology 47:733–770.

7. Kiers, ET, Rousseau, RA, West, SA, Denison, RF (2003) Host sanctions and the legume-rhizobium mutualism. Nature 425:78–81.

8. Bshary, RS, Bronstein, JL (2004) Game structures in mutualisms: what can the evidence tell us about the kinds of models we need? Advances in the Study of Behavior 34:59–104.

9. Poulin, R, Vickery, WL (1995) Cleaning symbiosis as an evolutionary game: to cheat or not to cheat? Journal of Theoretical Biology 175:63–70.

10. Doebeli, M, Knowlton, N (1998) The evolution of interspecific mutualisms. Proceedings of the National Academy of Sciences USA 95:8676–8680.

11. Noë, R (2001) in Economics in Nature: Social Dilemmas, Mate Choice and Biological Markets, eds Noë, R, van Hooff, JA, Hammerstein, P (Cambridge University Press).

12. Johnstone, RA, Bshary, R (2002) From parasitism to mutualism: partner control in asymmetric interactions. Ecology Letters 5:634–639.

13. Bergstrom, CT, Lachmann, M (2003) The Red King Effect: When the slowest runner wins the coevolutionary race. Proceedings of the National Academy of Sciences USA 100:593–598.

14. Hoeksema, JD, Kummel, M (2003) Ecological persistence of the plant-mycorrhizal mutualism: A hypothesis from species coexistence theory. The American Naturalist 162:S40–S50.

15. Akçay, E, Roughgarden, J (2007) Negotiation of mutualism: rhizobia and legumes. Proceedings of the Royal Society B 274:25–32.

16. Bshary, R, Grutter, AS, Willener, AST, Leimar, O (2008) Pairs of cooperating cleaner fish provide better service quality than singletons. Nature 455:964–967.

17. Trivers, RL (1971) The evolution of reciprocal altruism. The Quarterly Review of Biology 46:35–57.

18. Stadler, B, Dixon, AFG (2008) Mutualism: Ants and their Insect partners (Cambridge University Press).

19. Pierce, NE, Kitching, RL, Buckley, RC, Taylor, MFJ, Benbow, KF (1987) The costs and benefits of cooperation between the australian lycaenid butterfly, Jalmenus evagoras, and its attendant ants. Behavioral Ecology and Sociobiology 21:237–248.

20. Hölldobler, B, Wilson, EO (1990) The Ants (Belknap Press).

21. Hill, CJ, Pierce, NE (1989) The effect of adult diet on the biology of butterflies 1. the common imperial blue, Jalmenus evagoras. Oecologia 81:249–257.

22. Kiers, ET et al. (2011) Reciprocal rewards stabilize cooperation in the mycorrhizal symbiosis. Science 333:880–882.

23. Behm, JE, Kiers, ET (2014) A phenotypic plasticity framework for assessing intraspecific variation in arbuscular mycorrhizal fungal traits. Journal of Ecology 102:315–327.

24. Bshary, R, Schäffer, D (2002) Choosy reef fish select cleaner fish that provide high-quality service. Animal Behaviour 63:557–564.

25. Bshary, RS, Noë, R (2003) in Genetic and Cultural Evolution of Cooperation, ed Hammerstein, P (MIT Press), pp 167–184.

26. Weibull, JW (1995) Evolutionary Game Theory (MIT Press, Cambridge).

27. Hofbauer, J (1996) Evolutionary dynamics for bimatrix games: A Hamiltonian system? Journal of Mathematical Biology 34:675–688.

28. Hofbauer, J, Sigmund, K (1998) Evolutionary Games and Population Dynamics (Cambridge University Press, Cambridge, UK).

29. Souza, MO, Pacheco, JM, Santos, FC (2009) Evolution of cooperation under n-person snowdrift games. Journal of Theoretical Biology 260:581–588.

30. Gokhale, CS, Traulsen, A (2012) Mutualism and evolutionary multiplayer games: revisiting the Red King. Proceedings of the Royal Society B 279:4611–4616.

31. Eshel, I, Motro, U (1988) The three brothers' problem: kin selection with more than one potential helper. 1. The case of immediate help. American Naturalist pp 550–566.

32. Hauert, C, Michor, F, Nowak, MA, Doebeli, M (2006) Synergy and discounting of cooperation in social dilemmas. Journal of Theoretical Biology 239:195–202.

33. Gokhale, CS, Traulsen, A (2010) Evolutionary games in the multiverse. Proceedings of the National Academy of Sciences USA 107:5500–5504.

34. Gokhale, CS, Traulsen, A (2014) Evolutionary multiplayer games. Dynamic Games and Applications.

35. Nowak, MA, Sigmund, K (2004) Evolutionary dynamics of biological games. Science 303:793–799.

36. Gao, L, Li, YT, Wang, RW (2015) The shift between the Red Queen and the Red King effects in mutualisms. Scientific reports 5:8237.

37. Hauert, C, Holmes, M, Doebeli, M (2006) Evolutionary games and population dynamics: maintenance of cooperation in public goods games. Proceedings of the Royal Society B 273:2565–2570.