



CHICAGO JOURNALS



The University of Chicago

Adaptive Dynamics with Interaction Structure.

Author(s): Benjamin Allen, Martin A. Nowak, and Ulf Dieckmann

Source: *The American Naturalist*, Vol. 181, No. 6 (June 2013), pp. E139-E163

Published by: [The University of Chicago Press](#) for [The American Society of Naturalists](#)

Stable URL: <http://www.jstor.org/stable/10.1086/670192>

Accessed: 13/06/2014 15:03

Your use of the JSTOR archive indicates your acceptance of the Terms & Conditions of Use, available at <http://www.jstor.org/page/info/about/policies/terms.jsp>

JSTOR is a not-for-profit service that helps scholars, researchers, and students discover, use, and build upon a wide range of content in a trusted digital archive. We use information technology and tools to increase productivity and facilitate new forms of scholarship. For more information about JSTOR, please contact support@jstor.org.



The University of Chicago Press, The American Society of Naturalists, The University of Chicago are collaborating with JSTOR to digitize, preserve and extend access to *The American Naturalist*.

<http://www.jstor.org>

Adaptive Dynamics with Interaction Structure

Benjamin Allen,^{1,2,*} Martin A. Nowak,^{2,3} and Ulf Dieckmann⁴

1. Department of Mathematics, Emmanuel College, Boston, Massachusetts 02115; 2. Program for Evolutionary Dynamics, Harvard University, Cambridge, Massachusetts 02138; 3. Department of Mathematics, Department of Organismic and Evolutionary Biology, Harvard University, Cambridge, Massachusetts 02138; 4. Evolution and Ecology Program, International Institute for Applied Systems Analysis, Schlossplatz 1, A-2361 Laxenburg, Austria

Submitted July 21, 2011; Accepted August 31, 2012; Electronically published April 26, 2013

ABSTRACT: Evolutionary dynamics depend critically on a population's interaction structure—the pattern of which individuals interact with which others, depending on the state of the population and the environment. Previous research has shown, for example, that cooperative behaviors disfavored in well-mixed populations can be favored when interactions occur only between spatial neighbors or group members. Combining the adaptive dynamics approach with recent advances in evolutionary game theory, we here introduce a general mathematical framework for analyzing the long-term evolution of continuous game strategies for a broad class of evolutionary models, encompassing many varieties of interaction structure. Our main result, the canonical equation of adaptive dynamics with interaction structure, characterizes expected evolutionary trajectories resulting from any such model, thereby generalizing a central tool of adaptive dynamics theory. Interestingly, the effects of different interaction structures and update rules on evolutionary trajectories are fully captured by just two real numbers associated with each model, which are independent of the considered game. The first, a structure coefficient, quantifies the effects on selection pressures and thus on the shapes of expected evolutionary trajectories. The second, an effective population size, quantifies the effects on selection responses and thus on the expected rates of adaptation. Applying our results to two social dilemmas, we show how the range of evolutionarily stable cooperative behaviors systematically varies with a model's structure coefficient.

Keywords: adaptive dynamics, interaction structure, evolutionary game theory, quantitative traits, cooperation.

Introduction

A key challenge for evolutionary biology is to understand how the mechanistic features of an evolutionary process affect the ultimate outcome of evolution. This is already quite challenging under the classical assumption of a well-mixed population (e.g., Dieckmann and Law 1996; Champagnat et al. 2006). Recent decades, however, have seen a

surge of interest in evolutionary processes in which ecological events are localized in some sense. Well-studied examples include evolution in spatially structured populations (Wright 1943; Kimura and Weiss 1964; Barton and Slatkin 1986; Durrett and Levin 1994; Tilman and Kareiva 1997; Dieckmann et al. 2000; Rousset 2004; Lieberman et al. 2005; Lion and van Baalen 2008), in group- or deme-structured populations (Wright 1931; Wilson 1977; Taylor 1992; Hanski and Gilpin 1997; Traulsen and Nowak 2006; Ohtsuki 2010), and in populations with active assortment by kin (Hamilton 1971; Eshel and Cavalli-Sforza 1982) or phenotype (Antal et al. 2009). These forms of localized interaction have significant consequences for the evolution of cooperation (Nowak and May 1992; Killingback and Doebeli 1996; Nakamaru et al. 1997; Mitteldorf and Wilson 2000; Santos and Pacheco 2005; Ohtsuki et al. 2006; Traulsen and Nowak 2006; Taylor et al. 2007a; Fletcher and Doebeli 2009; Helbing and Yu 2009; van Veelen et al. 2012), host-parasite interactions (Hassell et al. 1991; Herre 1993; Boots and Sasaki 1999; Haraguchi and Sasaki 2000; van Baalen 2002; Read and Keeling 2003; Boots et al. 2004), signaling (Krakauer and Pagel 1995; Werfel and Bar-Yam 2004), and life-history traits (Hanski and Gaggiotti 2004).

In the context of evolutionary game theory, notions of locality are reflected in a model's interaction structure, which designates which individuals participate in game interaction with which others, depending on the current state of the population and the environment. Notions of locality may also be reflected in a model's update rule (Ohtsuki et al. 2006), which specifies how the outcomes of game interaction affect births, deaths, and other changes to the population and the environment.

It is thus of great interest (e.g., Nowak et al. 2010a) to understand how differences in the interaction structure and update rule affect the evolution of game behavior. Historically, this question has been investigated one model at a time, revealing both common themes and intriguing differences. In the case of discrete games, Tarnita et al. (2009b, 2011) have proposed a more general approach that

* Corresponding author; e-mail: benjcallen@gmail.com.

exploits the Markov chain structure of evolutionary game models. However, no corresponding theory exists for continuous games (Brown and Vincent 1987; McGill and Brown 2007), in which payoffs depend continuously on quantitative trait values.

Here we provide a general mathematical framework for determining the evolutionary trajectories of continuous game strategies under different interaction structures and update rules. Our work merges adaptive dynamics theory (Nowak and Sigmund 1990; Hofbauer and Sigmund 1990; Dieckmann and Law 1996; Metz et al. 1996; Geritz et al. 1997)—a framework for studying quantitative trait evolution when mutations are incremental and rare—with the approach of Tarnita et al. (2009*b*, 2011).

Our main result, which we term the “canonical equation of adaptive dynamics with interaction structure,” characterizes expected evolutionary trajectories of strategies for any continuous game, interaction structure, and update rule satisfying basic assumptions. This result generalizes the canonical equation of adaptive dynamics (which describes expected trajectories of adaptive evolution in large well-mixed populations; Dieckmann and Law 1996) to a broad class of models, including established models of evolution in groups (Wright 1931; Taylor 1992; Traulsen and Nowak 2006; Ohtsuki 2010), in space (Lieberman et al. 2005; Ohtsuki et al. 2006; Taylor et al. 2007*a*; Ohtsuki et al. 2007), with tag-based assortment (Antal et al. 2009), and in finite well-mixed populations with different forms of generational structure (Nowak et al. 2004; Imhof and Nowak 2006; Lessard and Ladret 2007; Traulsen et al. 2007).

Interestingly, we find that the effects of different interaction structures and update rules on the dynamics of adaptive evolution are fully captured by just two real numbers. First, a structure coefficient (Tarnita et al. 2009*b*) quantifies the effects on selection pressures and thus on the shapes of expected evolutionary trajectories and on the strategies expected as evolutionary outcomes. Second, an effective population size quantifies the effects on selection responses and thus on the expected rates of adaptation. Importantly, these two numbers are independent of the considered evolutionary game. Varying the structure coefficient and effective population size in our equation thus enables rapid, comprehensive analysis of game-strategy trajectories for a wide variety of interaction structures and update rules, representing many different ecological contexts.

Below, we begin with a review and synthesis of relevant background and previous work on adaptive dynamics theory and evolutionary game theory. We then state the mathematical assumptions defining the class of models to which our results apply, which are further detailed in appendix A. The canonical equation of adaptive dynamics with in-

teraction structure is then presented, discussed, and applied to the analysis of two social dilemmas. Its derivation is detailed in appendix B. We conclude with an exploration of connections between our work and other approaches and highlight directions for future research.

Background and Synthesis of Previous Work

Adaptive Dynamics Theory

Adaptive dynamics theory (Nowak and Sigmund 1990; Hofbauer and Sigmund 1990; Dieckmann and Law 1996; Metz et al. 1996; Geritz et al. 1997; Dercole and Rinaldi 2008) studies the long-term evolution of one or more quantitative traits, allowing the effects of complex interactions among individuals, including those implying frequency-dependent selection, to be analyzed. These interactions unfold on a short interaction timescale. Mutations are assumed to be rare, so that the evolving population is typically monomorphic, and its evolution can be studied on two separated timescales: the intermediate demographic timescale, on which competition between phenotypes results in fixation of one of them, and the longer evolutionary timescale, on which adaptation occurs via a sequence of trait-substitution events. Mutations are also assumed to be incremental, so that mutant phenotypes can be viewed as perturbations of resident phenotypes. Trait-substitution events start with a mutation and generally conclude with the fixation of either the mutant type or the resident type (Geritz et al. 2002; Geritz 2005).

The canonical equation of adaptive dynamics (Dieckmann 1994; Dieckmann et al. 1995; Dieckmann and Law 1996) describes the expected dynamics, on the evolutionary timescale, of trait values in a large well-mixed population, as these values are updated through the successive fixation of new mutant types. In the case of a single real-valued trait x evolving under birth-death dynamics with mutation, the canonical equation can be written as

$$\frac{dx}{dt} = \frac{1}{2} N \mu \nu \left. \frac{\partial f(x'; x)}{\partial x'} \right|_{x'=x}, \quad (1)$$

where t is evolutionary time, N is the equilibrium population size, μ is the mutation probability per birth, ν is the variance in mutational step size, and $f(x'; x)$ is the invasion fitness of the mutant type x' , given that the resident type is x . Invasion fitness is defined as the exponential growth rate of the mutants when they are rare (Metz et al. 1992). The right-hand side of equation (1) equals the expected change in x per unit time, averaged over all possible trait-substitution events. The canonical equation, in setting the expected change equal to the actual change dx/dt , characterizes the expected dynamics of the trait

value through evolutionary time (Dieckmann 1994), as well as the deterministic dynamics under appropriate scaling limits (Champagnat et al. 2002).

The occurrence of the invasion fitness function $f(x'; x)$ in the canonical equation originates from the fact that, in a large well-mixed population, the probability of mutant fixation is proportional to $f(x'; x)$ when $f(x'; x) > 0$, and it is zero otherwise (Dieckmann 1994; Dieckmann and Law 1996; Dercole and Rinaldi 2008; Durinx et al. 2008). In populations that are not large or not well mixed, this proportionality may not hold (see “Discussion”). In such situations, we must fall back on a more general formulation of adaptive dynamics by using fixation probabilities directly (Proulx and Day 2002; Rousset 2004; Imhof and Nowak 2010), with expected evolutionary trajectories then being described by

$$\frac{dx}{dt} = Nu(x)v \left. \frac{\partial \rho(x'; x)}{\partial x'} \right|_{x'=x}. \quad (2)$$

Here $u(x)$ denotes the per capita rate of mutant appearance in a monomorphic population of type x , and $\rho(x'; x)$ denotes the fixation probability of a mutant of type x' in such a population—that is, the probability that a new mutant type x' , starting from a single individual, will eventually displace a resident population of type x . For large well-mixed populations, and with $b(x)$ denoting the per capita birth rate of type x , equations (1) and (2) are connected by the relations $u(x) = \mu b(x)$ and $\rho(x'; x) = \max\{0, f(x'; x)/b(x)\}$ (Dieckmann and Law 1996).

Equation (2) illustrates the separation of timescales inherent in the adaptive dynamics approach. The whole of equation (2) describes trait dynamics on the evolutionary timescale. These dynamics depend, in turn, on the outcome of resident-mutant competition on the demographic timescale, as summarized by the fixation probability $\rho(x'; x)$. To apply equation (2), one must obtain $\rho(x'; x)$ (or, at minimum, its first-order behavior near $x' = x$) from analysis of resident-mutant competition under the evolutionary model in question. This may be a difficult task, depending on the complexity of the model. One contribution of the present work is to obtain expressions for this first-order behavior of $\rho(x'; x)$ for a general class of models in evolutionary game theory.

Individual-Based Evolutionary Game (IBEG) Models

Evolutionary game theory (Maynard Smith and Price 1973; Taylor and Jonker 1978; Weibull 1997; Cressman 1992; Hofbauer and Sigmund 1998, 2003; Nowak and Sigmund 2004) is a powerful approach to studying the evolution of social behavior. A game represents a particular mode of interaction, the outcome of which is summarized

in payoffs assigned to each participant. Evolving game strategies may be either discrete or continuous. For continuous games (Brown and Vincent 1987; McGill and Brown 2007), the strategies are real numbers or vectors representing quantitative traits, and the payoffs to participants are continuous functions of these strategies.

While early formulations of evolutionary game theory focused on notions of evolutionary stability (Maynard Smith and Price 1973) and on frequency dynamics in infinite populations (Taylor and Jonker 1978), much recent attention has been devoted to individual-based evolutionary game models (IBEG models) in which births, deaths, game interactions, and other events are represented explicitly (Nowak and May 1992; Durrett and Levin 1994; Killingback and Doebeli 1996; Nakamaru et al. 1997; van Baalen and Rand 1998; Mitteldorf and Wilson 2000; Nowak et al. 2004; Santos and Pacheco 2005; Ohtsuki et al. 2006; Traulsen and Nowak 2006; Szabó and FÁth 2007; Taylor et al. 2007a; Antal et al. 2009; Tarnita et al. 2009a; Nowak et al. 2010a; Perc and Szolnoki 2010; van Veelen et al. 2012). In contrast to traditional evolutionary game theory, IBEG models allow investigation of how localized interaction, population size, and other factors affect evolutionary game competition.

Here we provide a synthesis and general vocabulary for IBEG models, highlighting several critical features. We describe these models in biological terms, noting, however, that many IBEG models are applicable to cultural evolution through the spreading of behaviors or ideas. We also note that, while the term “individual-based” often carries the connotation of models analyzed through numerical analysis and computer simulation, here we use it to characterize the level at which the considered evolutionary game dynamics are defined, irrespective of the method subsequently applied for their analysis.

IBEG models appearing in the literature generally have the mathematical structure of a Markov chain. That is, they can be represented as a collection of states with transition probabilities between them. (Deterministic and/or continuous-state models can be understood as special and/or limiting cases of Markov chain models.) In this terminology, the “state” by definition provides a full snapshot of the evolutionary process, containing all information relevant to the determination of future states. At minimum, this includes the number and strategies of all living individuals. The inclusion of other information depends on the model in question. For example, in models of group-structured populations (Traulsen and Nowak 2006; Tarnita et al. 2009a; Ohtsuki 2010; Fu et al. 2012), the state includes the group affiliations of all individuals. In models with dynamic social networks (Pacheco et al. 2006a, 2006b; Perc and Szolnoki 2010; Wu et al. 2010; Fehl et al. 2011; Rand et al. 2011), the state includes the current

network topology. Models have also been studied in which the state includes individuals' ages, developmental stages, memories of past events, and/or states of health, as well as the environmental conditions each of them experiences.

An IBEG model must incorporate a scheme for designating when game interactions occur and between which pairs or sets of individuals. We call this scheme the model's interaction structure. Models of well-mixed populations (e.g., Nowak et al. 2004; Imhof and Nowak 2006; Lessard and Ladret 2007; Traulsen et al. 2007) typically employ a trivial interaction structure: at all time steps, each individual interacts equally with each of the other individuals. Other models use interaction structures that incorporate some notion of locality. For example, individuals may interact only if they are members of the same group (Taylor 1992; Traulsen and Nowak 2006; Ohtsuki 2010), neighbors on a lattice (Nowak and May 1992; Wilson et al. 1992; Durrett and Levin 1994; Killingback and Doebeli 1996; Nakamaru et al. 1997; Hauert and Doebeli 2004; Helbing and Yu 2009; Roca et al. 2009), or are connected through the (current) social network (van Baalen and Rand 1998; Santos and Pacheco 2005; Ohtsuki et al. 2006; Pacheco et al. 2006*a*, 2006*b*; Taylor et al. 2007*a*; Perc and Szolnoki 2010; Allen et al. 2012). In the most abstract sense, a model's interaction structure can be understood as a mapping from its current state to the current collection of game-interaction partners.

Once game interactions have taken place, births, deaths, movements, and other relevant events must be determined according to another scheme, depending (often stochastically) on payoffs. Following Ohtsuki et al. (2006), we call this scheme the model's "update rule." For well-mixed population models, update rules usually have the property that each individual's expected offspring number is an increasing function of its own payoff (Nowak et al. 2004; Imhof and Nowak 2006; Lessard and Ladret 2007). For models with nontrivial interaction structure, there is no canonical choice of update rule, and different choices can lead to strikingly different outcomes. For example, Ohtsuki et al. (2006) explored three different update rules for evolution on graphs. For one (birth-death), the benefits of cooperator clustering are opposed by a spatial competition effect separate from game interaction, rendering cooperation ineffective. (A similar effect appears in the group-selection model of Taylor 1992 and Ohtsuki 2010.) For two other update rules (death-birth and imitation), the spatial competition effect is weaker and cooperation can be favored. Thus, the choice of update rule is a modeling decision that must be treated with appropriate care.

Finally, the probabilities of transition between states—and hence the dynamics and outcomes of evolutionary competition—depend on the game, the interaction structure, and the update rule of the considered IBEG model.

We therefore view the game, interaction structure, and update rule as three key ingredients of any IBEG model, leaving the abstract formalization of this idea to future work. An important and attractive feature of IBEG models is that the considered game can be varied independently from the considered interaction structure and the considered update rule. Varying the latter two ingredients can elucidate, for example, how the success of a particular trait or behavior varies with the geometry of the considered environment.

Structure Coefficient

Tarnita et al. (2009*b*, 2011; see also Nowak et al. 2010*a*) have recently pioneered a general approach to studying IBEG models on the demographic timescale. This approach focuses on the Markov chain structure of these models, abstracting from the details of births, deaths, and game interactions.

Tarnita et al. (2009*b*) considered IBEG models involving pairwise games with two strategies, described by 2×2 payoff matrices,

$$\begin{array}{cc} & X & Y \\ \begin{array}{c} X \\ Y \end{array} & \begin{pmatrix} a & b \\ c & d \end{pmatrix} \end{array} \quad (3)$$

Above, X and Y are the two considered strategies, a and b are the payoffs received by X when playing against X and Y , respectively, and c and d are the payoffs received by Y when playing against X and Y , respectively. In the Markov chain representation of such an IBEG model, each transition probability between any pair of states can be considered a function of the payoff values a , b , c , and d , with this function depending in turn on the interaction structure and update rule of the considered model. Tarnita et al. (2009*b*) imposed basic differentiability and symmetry assumptions on the transition probabilities as functions of the payoff values. On this basis, they proved that any IBEG model satisfying their assumptions can be characterized by a real-valued "structure coefficient" σ . The value of σ depends on the interaction structure and update rule but is independent of the game. Given a particular game of the form (3), strategy X is favored over strategy Y —in the sense that X has greater fixation probability than Y under weak selection—if and only if

$$\sigma a + b > c + \sigma d. \quad (4)$$

The utility of this result is that, once the value of σ has been determined for a particular combination of interaction structure and update rule (see examples in table 1), condition (4) can be used to determine which strategy is favored in any 2×2 game. In short, σ quantifies how a particular combination of interaction structure and update

Table 1: Effective population size and structure coefficient for individual-based evolutionary game models

Model	Effective population size N_e	Structure coefficient σ	References
Well-mixed Wright-Fisher process	N	$\frac{N-2}{N}$	Fisher 1930; Wright 1931
Well-mixed continuous-time birth-death process	$\frac{N}{2}$	$\frac{N-2}{N}$	Dieckmann 1994
Well-mixed Moran process	$\frac{N}{2}$	$\frac{N-2}{N}$	Nowak et al. 2004
Transitive graph, birth-death updating	$\frac{N}{2}$	$\frac{N-2}{N}$	Ohtsuki et al. 2006; Taylor et al. 2007a
Transitive graph, death-birth updating	k^2	$\frac{(k+1)N-4k}{(k-1)N}$	Ohtsuki et al. 2006; Taylor et al. 2007a
Group selection without migration	$\frac{Nm+n_g-2}{2(N-1)}$	$1 + \frac{2n_g-4}{m}$	Traulsen and Nowak 2006
Group selection with migration	$\frac{N}{2(N-1)} \left[m - 1 + \frac{m(q-\lambda)(n_g-1)}{\lambda+mq} \right]$	$1 - \frac{2}{m} + \frac{2q(n_g-1)}{\lambda+mq}$	Traulsen and Nowak 2006
Infinite island model	Unknown	1	Wright 1931; Taylor 1992; Ohtsuki 2010
Phenotypic assortment	Unknown	$\frac{1+4N\nu}{2+4N\nu} \left(1 + \sqrt{\frac{3+12N\nu}{3+4N\nu}} \right)$	Antal et al. 2009

Note: Term N = population size, k = degree of graph, n_g = number of groups, m = number of individuals per group, q = group splitting probability, λ = migration rate, ν = phenotypic mutation rate. Transitive graphs are spatially homogeneous; see Taylor et al. (2007a) and Grafen and Archetti (2008). The result for the island model of Wright (1931) and Taylor (1992) applies to the limit of infinitely many islands, as analyzed by Ohtsuki (2010).

rule affects the selection pressure on game strategies. For a well-mixed population, σ approaches 1 as population size approaches infinity; thus, $\sigma = 1$ can be interpreted as a baseline value. In this case, the same-type payoffs a and d and the opposite-type payoffs b and c are of equal importance in determining evolutionary success. As σ is increased above 1, the same-type payoffs become increasingly important. Conversely, as σ is decreased below 1, the opposite-type payoffs become increasingly important. Accordingly, the former case favors cooperative strategies, whereas the latter case hinders them (Tarnita et al. 2009b).

We caution that the structure coefficient does not merely quantify assortment of strategies or genetic relatedness between interaction partners. Its value also depends, for example, on any indirect competition effects that may be subsumed in the update rule. We will revisit the interpretation of σ throughout this work. In particular, as part of our results, we derive an expression for σ in terms of the dependence of fixation probabilities on game payoffs.

Mathematical Framework

Our main result, an equation describing the evolutionary trajectories of continuous game strategies, extends the approach of Tarnita et al. (2009b) from the demographic to the evolutionary timescale, using the adaptive dynamics framework. The resultant equation applies to any IBEG model satisfying a set of basic assumptions. In this section,

we present these assumptions verbally, saving rigorous mathematical formulations for appendix A. We separate our assumptions into those applying to the interaction timescale (labeled G1, G2), the demographic timescale (D1–D6), and the evolutionary timescale (E1–E3).

Interaction Timescale: Game Strategies and Payoffs

Interactions are modeled as a continuous game with payoff function A . The possible strategies are represented as real numbers or vectors. In the two-player case, $A(x; y)$ denotes the payoff to an individual of strategy x interacting with an individual of strategy y . For n -player games, the payoff function is written $A(x; y_1, \dots, y_{n-1})$. In this case, the first argument x denotes the strategy of the focal individual receiving the payoff, and the remaining $n - 1$ arguments are the strategies of the other game participants. We place the following assumptions on A :

- G1 A is positive and twice differentiable in all of its arguments, at least in the case that all game participants use approximately the same strategy.
- G2 For games with more than two players, the arrangement of interaction partners does not matter, that is, the value of $A(x; y_1, \dots, y_{n-1})$ does not change if the arguments y_1, \dots, y_{n-1} are permuted.

Demographic Timescale: Resident-Mutant Competition

On the demographic timescale, we are interested in episodes of competition between a resident strategy x and a mutant strategy x' . Four payoff values arise in this competition, and we introduce the following shorthand notation for these:

$$\begin{aligned} a_{MM} &= A(x'; x'), & a_{MR} &= A(x'; x), \\ a_{RM} &= A(x; x'), & a_{RR} &= A(x; x). \end{aligned}$$

Following the approach of Tarnita et al. (2009b), we focus on the Markov chain representing resident-mutant competition under the considered IBEG model, abstracting from lower-level biological details. This Markov chain is initiated in a state corresponding to the appearance of a single new mutant. Eventually, the Markov chain will reach a state corresponding to either mutant fixation or extinction. We denote by ρ the probability that mutant fixation is reached.

We impose six assumptions on how the Markov chains' transition probabilities vary with respect to a_{MM} , a_{MR} , a_{RM} , and a_{RR} . These assumptions implicitly restrict the choices of interaction structures and update rules in IBEG models that can be analyzed using our approach.

- D1 The set of possible states is finite.
- D2 The Markov chain's transition probabilities are completely determined by the payoff values a_{MM} , a_{MR} , a_{RM} , and a_{RR} .
- D3 For any values of a_{MM} , a_{MR} , a_{RM} , and a_{RR} , mutant fixation is possible from any initial state corresponding to the appearance of a single new mutant.
- D4 The Markov chain's transition probabilities vary twice differentially with respect to a_{MM} , a_{MR} , a_{RM} , and a_{RR} .
- D5 The fixation probability ρ is unaffected if a_{MM} , a_{MR} , a_{RM} , and a_{RR} are each multiplied by a positive constant $K > 0$.
- D6 The fixation probability ρ is increasing in a_{MM} and a_{MR} , and decreasing in a_{RM} and a_{RR} .

Assumption D2 requires that the evolving strategies affect only the game payoffs; if the evolving strategies also affected the interaction structure or update rule, then the Markov chain's transition probabilities would not be completely determined by a_{MM} , a_{MR} , a_{RM} , and a_{RR} but would depend on x as well. Assumption D5 guarantees that the

fixation probabilities are insensitive to changes in the units in which payoffs are measured.

While these assumptions apply to a wide variety of IBEG models (a sample of which are listed in table 1), they exclude models with infinitely many states (e.g., those with infinite population size or continuous space) and models in which behaviors affecting the interaction structure (e.g., homophily) or update rule (e.g., migration) coevolve with the game strategies.

The six assumptions above are compatible with those stipulated by Tarnita et al. (2009b). Consequently, there is a structure coefficient, satisfying condition (4), associated to each combination of interaction structure and update rule satisfying these assumptions.

Evolutionary Timescale: Mutation and Trait Substitution

Following the adaptive dynamics approach, we make the following three assumptions regarding the mutation process:

- E1 Mutations are rare, so that the evolving population is typically monomorphic, and long-term evolution can be described as a sequence of trait substitutions.
- E2 Mutation is directionally unbiased.
- E3 Mutation is incremental, so that the mutant strategy x' is always close to the resident strategy x .

Main Result: Canonical Equation of Adaptive Dynamics with Interaction Structure

This section presents our central result, the canonical equation of adaptive dynamics with interaction structure. This equation describes expected evolutionary trajectories of a game strategy x for any combination of continuous game (satisfying assumptions G1 and G2), update rule and interaction structure (satisfying assumptions D1–D6), and mutation process (satisfying assumptions E1–E3).

Univariate Traits and Pairwise Games

We begin with the simple case of a pairwise (two-player) game $A(x; y)$, in which the strategies x and y are univariate (real numbers). The canonical equation of adaptive dynamics with interaction structure in this case is

$$\frac{dx}{dt} = N_c \frac{N-1}{N} \frac{u(x)v}{A(x;x)} \times \left[\left. \frac{\partial A(x';x)}{\partial x'} \right|_{x'=x} + \frac{\sigma-1}{\sigma+1} \left. \frac{\partial A(x;x')}{\partial x'} \right|_{x'=x} \right], \quad (5)$$

where σ is the structure coefficient and N_c is the effective population size (defined below). As in equation (2), t is evolutionary time, N is the equilibrium population size, $u(x)$ is the per capita rate of mutant appearance, and v is the variance in mutational step size.

This result is obtained by first showing that expected evolutionary trajectories are related to fixation probabilities as in equation (2) and then showing that the relevant derivatives of fixation probabilities can be expressed in terms of the structure coefficient and effective population size. The full derivation is given in appendix B. We now discuss several aspects of this result in detail.

Structure Coefficient and Shapes of Evolutionary Trajectories

First we note that all effects of the interaction structure and update rule on the shape of evolutionary trajectories—in particular on the strategies corresponding to evolutionary equilibria—are described by the proportionality relation

$$\frac{dx}{dt} \propto \left. \frac{\partial A(x';x)}{\partial x'} \right|_{x'=x} + \frac{\sigma-1}{\sigma+1} \left. \frac{\partial A(x;x')}{\partial x'} \right|_{x'=x}, \quad (6)$$

where the symbol \propto means “proportional to.” The two terms of equation (6) express a mutation’s evolutionary success in terms of its effect on the payoff of its bearer (first term), as well as its effect on the payoffs of those with whom the bearer interacts (second term). In the limit of a large, well-mixed population ($\sigma \rightarrow 1$), only the bearer’s payoff matters. But with nontrivial interaction structure and/or update rule ($\sigma \neq 1$), a mutant’s effect on the payoffs of others is relevant, with the sign of $\sigma - 1$ identifying whether helpful or harmful effects are favored by the second term.

We also derive in appendix B an expression for σ in terms of the behavior of the mutant fixation probability ρ with respect to the game payoffs:

$$\sigma = \left(\frac{\partial \rho}{\partial a_{MM}} - \frac{\partial \rho}{\partial a_{RR}} \right) \left/ \left(\frac{\partial \rho}{\partial a_{MR}} - \frac{\partial \rho}{\partial a_{RM}} \right) \right., \quad (7)$$

with all derivatives taken at $a_{MM} = a_{MR} = a_{RM} =$

$a_{RR} = 1$. We note in passing that this expression also applies to the result of Tarnita et al. (2009b), with a , b , c , and d in place of a_{MM} , a_{MR} , a_{RM} , and a_{RR} , respectively. We additionally show in appendix B that σ is always positive, as a consequence of assumption D6.

The appearance of the ratio $(\sigma - 1)/(\sigma + 1)$ in equation (6) can be understood by the following argument. By condition (4), a mutant strategy x' is favored over the resident strategy x if and only if

$$\sigma A(x';x') + A(x';x) - A(x;x') - \sigma A(x;x) > 0.$$

For $x' = x$, the quantity on the left-hand side vanishes. Therefore, to first order in $x' - x$, mutant favorability is determined by the sign of

$$\begin{aligned} & \sigma \left[\left. \frac{\partial A(x';x)}{\partial x'} \right|_{x'=x} + \left. \frac{\partial A(x;x')}{\partial x'} \right|_{x'=x} \right] \\ & + \left. \frac{\partial A(x';x)}{\partial x'} \right|_{x'=x} - \left. \frac{\partial A(x;x')}{\partial x'} \right|_{x'=x} \\ & = (\sigma + 1) \left. \frac{\partial A(x';x)}{\partial x'} \right|_{x'=x} + (\sigma - 1) \left. \frac{\partial A(x;x')}{\partial x'} \right|_{x'=x}. \end{aligned}$$

Dividing by $\sigma + 1$, we obtain the coefficients in equation (6). Values of σ for a variety of IBEG models are shown in table 1.

Effective Population Size and Speed along Evolutionary Trajectories

We next observe that the rate of evolution in equation (5) is proportional to the effective population size N_c . The effective population size is defined here in terms of the selection response, that is, through the likelihood of beneficial mutations to fix. Our definition is motivated by Kimura’s (1964) result that, for a class of models amenable to one-dimensional diffusion approximation, a beneficial mutation of selective advantage s has fixation probability

$$\rho \approx \frac{1 - e^{-2sN_c/N}}{1 - e^{-2sN_c}}. \quad (8)$$

Differentiating with respect to s and solving for N_c yields

$$N_c = \frac{N^2}{N-1} \left. \frac{\partial \rho}{\partial s} \right|_{s=0}, \quad (9)$$

which we take as our definition of N_c . We note that in this work the actual and effective population sizes, N and N_c , respectively, pertain to the demographic equilibrium of the resident population rather than to its current state at a given time.

Our N_e agrees with the variance effective population size (Kimura 1964) and the eigenvalue effective population size (Ewens 1982; see also Rousset 2004) for models for which these are defined. Our definition is similar in spirit to the adaptation effective population size defined by Campos and Wahl (2009). More generally, it is proven for some models and conjectured more generally (Metz 2011) that the notion of effective population size as a quantity scaling the rate of adaptation—as in equation (6)—is equivalent to the more familiar notion of effective population size as a quantity inversely proportional to the strength of neutral drift.

A convention implicit in equation (8) is that the Wright-Fisher (nonoverlapping generations) model is taken as a baseline. Consequently, $N_e = N$ for the Wright-Fisher model, while N_e is greater than (less than) N for population structures that promote (inhibit) the fixation of beneficial mutations, relative to this model. Interestingly, many overlapping-generation IBEG models (Moran 1958; Dieckmann 1994; Nowak et al. 2004) have $N_e = N/2$. Values of N_e for a variety of models are shown in table 1.

Rates of Mutant Appearance

In the canonical equation (5), the process by which new mutations appear is deliberately left unspecified. This flexibility allows for the application of equation (5) to a wide variety of biological and cultural evolutionary processes.

For biological evolution, it is natural to set $u(x)$ equal to $\mu b(x)$, where $b(x)$ is the birth rate in a monomorphic population of strategy x , and μ is the mutation probability per birth as in equation (1). An important special case occurs when payoff is equal to birth rate in monomorphic populations, so that $b(x) = A(x; x)$. In this case the ratio $u(x)/A(x; x)$ in equation (5) simplifies to μ .

For cultural evolution, $u(x)$ can be interpreted as the per capita rate at which new ideas or behaviors are introduced. Depending on the details of the considered evolutionary process, it may then be reasonable to assume $u(x)$ to be constant (if innovation is unrelated to strategies or payoffs), proportional to $A(x; x)$ (if high payoffs induce greater levels of innovation), or inversely proportional to $A(x; x)$ (if innovation is induced by discontent with one's current payoff).

Recovery of Original Canonical Equation

Dieckmann (1994) derived the original canonical equation of adaptive dynamics, equation (1), using a continuous-time birth-death model in a large well-mixed population. For this model, $\sigma = 1$ and $N_e = N/2$. If we equate an individual's birth rate with the average payoff obtained from all other individuals and assume a constant death

rate, then $u(x) = \mu A(x; x)$, $f(x'; x) = A(x'; x) - 1$, and equation (5) reduces to the original canonical equation (1).

Multivariate Traits

For continuous games with multivariate (d -dimensional) strategies, $x = (x_1, \dots, x_d)$, evolutionary trajectories are described by the system of equations

$$\frac{dx_i}{dt} = N_e \frac{N-1}{N} \frac{u(x)v_i}{A(x; x)} \times \left[\frac{\partial A(x'; x)}{\partial x'_i} \Big|_{x'=x} + \frac{\sigma-1}{\sigma+1} \frac{\partial A(x; x')}{\partial x'_i} \Big|_{x'=x} \right], \quad (10)$$

for $i = 1, \dots, d$. Here, the mutational step size variance v_i is given the subscript i to reflect that mutations in different components i of x may be differently distributed.

Equation (10) applies when mutations in each component of x are independent. Correlations among mutations in different components can be readily accommodated, following a formalism described by Dieckmann and Law (1996, sec. 6.2).

Multilateral Interactions

Our result extends to games involving any number of players. We show in appendix C that an n -player game $B(x; y_1, \dots, y_n)$ can be replaced by the pairwise game with payoff function $A(x; y) = B(x; y, \dots, y)$ —more specifically, the payoff value $B(x; y_1, \dots, y_{n-1})$ can be replaced by the arithmetic mean of $A(x; y_j)$ for $j = 1, \dots, n-1$. In other words, one can suppose that the pairwise game A is played with each of the $n-1$ partners and the results are then averaged; this substitution leaves evolutionary trajectories unchanged.

This surprising simplification arises from the differentiability and symmetry properties, G1 and G2, imposed on the game. It does not imply that the set of interaction partners is typically monomorphic.

Range of Applications

The power of our result lies in its generality. To obtain the expected evolutionary trajectory of a particular continuous strategy under a particular IBEG model, one need only substitute the appropriate payoff function A and values of σ and N_e into the canonical equation with interaction structure, equation (10). This saves the labor of analyzing each combination of game, interaction structure, and update rule from scratch. Moreover, our result enables comparisons both within classes of models (e.g., graphs

with different topologies) and across such classes (e.g., models with predefined groups versus those with tag-based assortment). Allowing σ and N_c to vary across all positive values reveals, all at once, the range of potential trajectories a quantitative trait can follow under all conceivable ecological scenarios fulfilling the rather general assumptions above.

Examples: Social Dilemmas

We now illustrate the power of our framework by considering two social dilemmas with continuous strategies. In each case we find that, as the structure coefficient increases, new cooperative equilibria arise and existing equilibria shift toward greater levels of cooperation.

Example 1: One-Shot Continuous Prisoner’s Dilemma

We first investigate the continuous Prisoner’s Dilemma (Killingback et al. 1999; Wahl and Nowak 1999a, 1999b; Killingback and Doebeli 2002). This models a pairwise social dilemma with a continuous range of cooperativity levels, quantified by a nonnegative real number x . In each interaction, a player with cooperativity x pays a cost $C(x)$ to produce a benefit $B(x)$ for the other player. We assume that the functions $C(x)$ and $B(x)$ are twice differentiable and strictly increasing and that they satisfy $C(0) = B(0) = 0$ (so that $x = 0$ corresponds to no cooperation) and $C(x) < B(x)$ for $x > 0$. The payoff function thus is

$$A(x, y) = -C(x) + B(y).$$

Applying equation (6), we immediately obtain the cooperativity dynamics under any IBEG model satisfying our assumptions:

$$\frac{dx}{dt} \propto -C'(x) + \frac{\sigma - 1}{\sigma + 1} B'(x). \tag{11}$$

For large well-mixed populations ($\sigma = 1$), these dynamics depend only on the marginal cost $C'(x)$ of cooperation. The second term contributes when $\sigma \neq 1$ and captures how a mutant’s success depends on the marginal benefit $B'(x)$ it gives to others. We see that, depending on the interaction structure and update rule, a mutant’s success can increase ($\sigma > 1$) or decrease ($\sigma < 1$) with this marginal benefit. The ratio $(\sigma - 1)/(\sigma + 1)$ can therefore be interpreted as quantifying the “potential for altruism” (Gardner 2010) afforded by a particular combination of interaction structure and update rule.

Cooperativity increases from its current value x if and only if both $\sigma > 1$ and

$$\frac{B'(x)}{C'(x)} > \frac{\sigma + 1}{\sigma - 1}. \tag{12}$$

Conversely, if $\sigma \leq 1$ or if $B'(0)/C'(0) < (\sigma + 1)/(\sigma - 1)$, cooperativity will not increase from an initial value of zero.

The left-hand side of condition (12) is the marginal-benefit-to-marginal-cost ratio of an increase in cooperativity. Condition (12) thus is a differential version of the benefit-to-cost thresholds, such as Hamilton’s rule, that frequently arise as conditions for the evolution of cooperation (reviewed by Nowak 2006).

Cooperativity equilibria occur at values $x = x^*$ satisfying $B'(x^*)/C'(x^*) = (\sigma + 1)/(\sigma - 1)$. An equilibrium x^* is stable if

$$\left. \frac{d}{dx} \left[\frac{B'(x)}{C'(x)} \right] \right|_{x=x^*} < 0, \tag{13}$$

that is, if the marginal-benefit-to-marginal-cost ratio of cooperation is decreasing at $x = x^*$. Conversely, unstable equilibria occur when this ratio is increasing. In the latter case, x^* can be understood as a cooperativity threshold, in that cooperativity dwindles when $x < x^*$ but becomes self-reinforcing when $x > x^*$.

We can neatly summarize these results in a bifurcation diagram (fig. 1), in which the structure coefficient serves as the control parameter. This diagram shows how greater values of σ yield higher levels of evolutionarily stable cooperativity. In general, our main result enables constructing such a bifurcation diagram for any continuous game, showing how the evolutionary trajectories following equation (10) vary with a model’s structure coefficient.

Example 2: Iterated Prisoner’s Dilemma with Stochastic Reactive Strategies

The iterated Prisoner’s Dilemma with stochastic reactive strategies (Nowak and Sigmund 1990, 1992, 1993; Imhof and Nowak 2010) is an elegant and well-studied example of multivariate adaptive dynamics. Here we consider the Prisoner’s Dilemma game

$$\begin{matrix} & C & D \\ C & (b - c, -c) \\ D & (b, 0) \end{matrix}, \tag{14}$$

played an infinite number of times between two players. Each player’s move depends stochastically on the previous move of the other player. A strategy is characterized by a pair of probabilities, $(p, q) \in [0, 1] \times [0, 1]$. The strategy component p (respectively, q) represents the probability of cooperating in response to cooperation (respectively, defection) by the other player. Thus, the strategy $(0, 0)$ corresponds to “always defect” (ALLD), $(1, 1)$ to “always

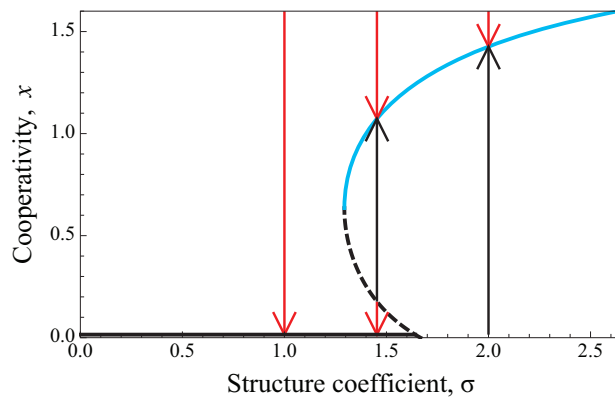


Figure 1: Bifurcation diagram showing the adaptive dynamics of cooperativity in equation (11) under different interaction structures and update rules. The structure coefficient σ serves as a control parameter. Arrows indicate the direction of cooperativity evolution. For this example, $B(x) = 4x + 4x^2$ and $C(x) = x + x^3/5$. Equilibrium points lie on the curve $\sigma = [B'(x) + C'(x)]/[B'(x) - C'(x)]$ and on the line $x = 0$. At $\sigma \approx 1.29$ a saddle-node bifurcation occurs, resulting in the creation of a stable equilibrium with positive cooperativity and an unstable cooperativity threshold. As σ increases, the cooperativity at the stable equilibrium grows and its basin of attraction expands. The leftmost arrow at $\sigma = 1$ corresponds to a large well-mixed population. The central arrows at $\sigma = 109/75 \approx 1.45$ could correspond, for example, to a 5×5 -square lattice with death-birth updating, or to 19 groups of size 75 without migration. The rightmost arrows at $\sigma = 2$ could correspond to a large cyclically structured population, or 12 groups of size 10 without migration. See table 1 for these and other interpretations of values of σ in terms of specific individual-based evolutionary game models.

cooperate” (ALLC), and (1, 0) to “tit-for-tat” (TFT). The payoff to each player is averaged over an infinite number of iterations; hence the choice of opening move is unimportant.

For this game, the payoff to a player with strategy (p_1, q_1) interacting with a player with strategy (p_2, q_2) , as calculated by Nowak and Sigmund (1990), is

$$A(p_1, q_1; p_2, q_2) = \frac{-c(q_1 + p_1q_2 - q_1q_2) + b(q_2 + p_2q_1 - q_2q_1)}{1 - (p_1 - q_1)(p_2 - q_2)}$$

We study the adaptive dynamics of the cooperation probabilities p and q using the multivariate canonical equation with interaction structure equation (10). For simplicity, we assume that mutations in p and q are independent, and of equal expected magnitude, but this assumption can readily be relaxed.

Upon computing the four partial derivatives of A as required by equation (10), it follows immediately that, for any IBEG model satisfying our assumptions, the expected evolutionary trajectories trace concentric circles around

the point $(p, q) = (1, 0)$ corresponding to TFT. It further follows that, along these trajectories, cooperation increases ($dp/dt > 0$ and $dq/dt > 0$) if

$$(\sigma + 1)[-c + b(p - q)] + (\sigma - 1)[-c(p - q) + b] > 0, \tag{15}$$

and decreases ($dp/dt < 0$ and $dq/dt < 0$) otherwise.

The observations above fully determine the evolutionary trajectories for this game under any interaction structure and update rule. Figure 2 illustrates these results, showing how the range of stable equilibria with positive p and q values increases with the structure coefficient σ .

Discussion

Summary

Although patterns of interaction and replacement can have complex and varied effects on a population’s evolution, our results here show that, for the purposes of determining the expected trajectories of continuous strategies under a broad class of evolutionary game models, these effects are fully quantified by just two real numbers, the effective population size and the structure coefficient. This remarkable simplification arises from the assumptions of rare and incremental mutation that underlie the adaptive dynamics framework.

Our results facilitate a broader approach to the study of continuous strategy evolution. Rather than studying individual combinations of a game, an interaction structure, and an update rule, one can derive the values of the structure coefficient and effective population size for a particular choice of interaction structure and update rule (as in the examples shown in table 1), and thereby immediately understand their effects on adaptive dynamics for any continuous game. Or, as in figures 1 and 2, one can focus on a single game and, by varying σ and N_e , understand how adaptive dynamics and outcomes for this game vary under different interaction structures and update rules.

Our work also unifies previous findings that spatial structure can promote cooperation in continuous-strategy social dilemmas (van Baalen and Rand 1998; Killingback et al. 1999; Le Galliard et al. 2003, 2005; Ifti et al. 2004). Our work shows that, across models and games, the strength of this spatial benefit to cooperation is quantified by the structure coefficient σ . Indeed, whenever $\sigma > 1$, the coefficient of the second term of equation (10) is positive, indicating that selection favors strategies that increase the payoffs of others. The two examples in the preceding section show how both the range and the intensity of evolutionarily stable cooperative behaviors can increase with σ .

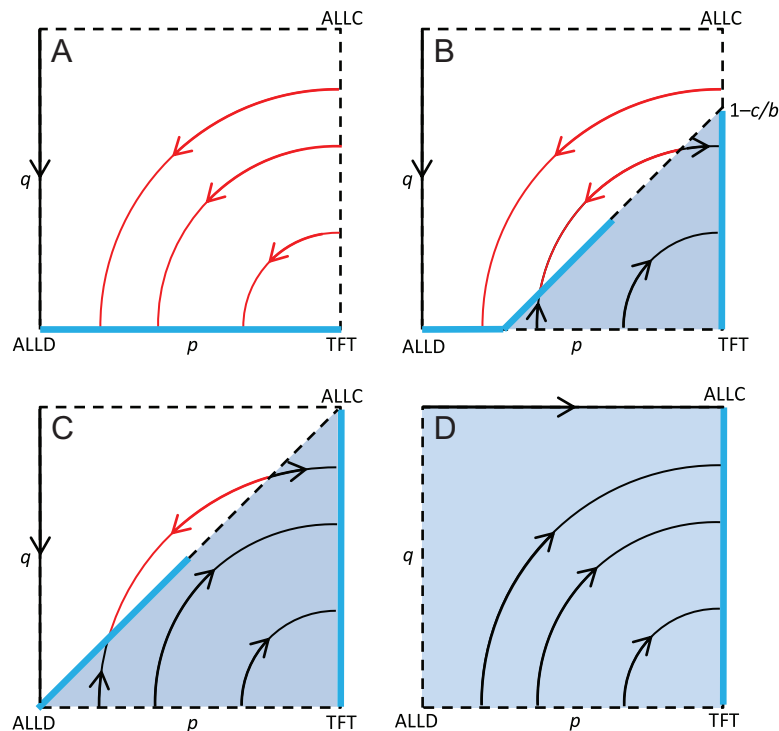


Figure 2: Adaptive dynamics of stochastic reactive strategies in the iterated Prisoner's Dilemma for different interaction structures and update rules. The value of the structure coefficient σ increases from A to D. Thin continuous lines depict evolutionary trajectories, with arrowheads and line color indicating the direction of evolutionary change. Thick continuous lines, rendered in blue, depict stable equilibria, while thin dashed lines depict unstable equilibria or boundary points. The evolutionary dynamics along the boundaries are obtained by ignoring the outward-pointing component, if any, of dx/dt . A, In the limit $\sigma \rightarrow 0$, all equilibrium strategies deterministically respond to D with D. In this case, tit-for-tat (TFT) is the only equilibrium strategy that achieves a positive payoff against itself. B–D, As σ increases, a range of “generous tit-for-tat” (GTFT; Nowak and Sigmund 1990) strategies become stable equilibria. These GTFT strategies are characterized by $p = 1$ and $q > 0$ and are therefore situated on the right-hand edge of the domain. Analysis of condition (15) shows that the GTFT strategy $(1, q)$ is a stable equilibrium if and only if $q < [2\sigma(-c + b)] / [\sigma(-c + b) + c + b]$. B, For $\sigma = 1$, corresponding to large well-mixed populations, the classical GTFT strategy $(1, 1 - c/b)$ (Nowak and Sigmund 1990) becomes stable. C, For $\sigma = (b + c)/(b - c)$, ALLC is a stable equilibrium, along with the full range of GTFT strategies. D, In the limit $\sigma \rightarrow \infty$, the set of GTFT strategies is a global evolutionary attractor.

Fixation Probability versus Invasion Fitness

We depart from most prior work on adaptive dynamics in structured populations (van Baalen and Rand 1998; Ferrière and Le Galliard 2001; Metz and Gyllenberg 2001; Durinx et al. 2008) in our use of fixation probabilities in place of invasion fitness—defined as the exponential growth rate of a rare mutant subject to the current environment (Metz et al. 1992, 1996; Rand et al. 1994; Dieckmann and Law 1996; Heino et al. 1998).

This concept is natural in well-mixed populations. Additionally, for some models with nontrivial interaction structure, it can be shown that assortment patterns are likely to equilibrate (in some sense) while a mutant is still rare (Matsuda et al. 1992; van Baalen and Rand 1998; Ferrière and Le Galliard 2001; Ohtsuki and Nowak 2008).

The invasion fitness can then be derived assuming that such equilibration has occurred. However, such an approach does not necessarily account for how the mutant's fixation probability is affected by its dynamics prior to this equilibration. It is also unclear whether the separation of timescales needed for defining an exponential phase can be expected to hold for all models of resident-mutant competition with interaction structure. Basing adaptive dynamics directly on fixation probabilities (Proulx and Day 2002; Rousset 2004; Imhof and Nowak 2010) eschews such difficulties.

The direct use of fixation probabilities is therefore attractive when these can be determined in addition to, or independently of, invasion fitness. By contrast, where fixation probabilities need to be inferred from invasion fit-

ness, they provide no marginal benefit. Moreover, fixation probabilities are less appropriate for understanding processes of evolutionary branching (Metz et al. 1992; Geritz et al. 1997; Dieckmann and Doebeli 1999), in which competing strains coexist and begin to diverge. Understanding how variations in interaction structures and update rules affect evolutionary branching will require formalizing and calculating the establishment probability (Haccou and Iwasa 1996; Metz et al. 1996; Durinx et al. 2008)—that is, the probability that an invading mutant type will reach a sustainable abundance—for models with nontrivial interaction structure.

Interpretation of Structure Coefficient

As is made precise in equation (7), the structure coefficient quantifies how the outcomes of same-type and different-type interactions affect a mutation's fixation probability. Because fixation probability is itself a summary quantity that depends in complex ways on the fundamental events (births, deaths, interactions, etc.) that drive evolution, one may wish to connect the structure coefficient more directly to these fundamental events and thereby also to other well-studied quantities such as assortment and relatedness.

How to address this challenge depends on the class of models under consideration. Here, as in the work of Tarnita et al. (2009b), evolutionary competition is abstractly represented as a Markov chain. Fixation is represented explicitly, but births, deaths, interactions, and other fundamental events are (in the general case) represented only implicitly, as aspects that may affect transitions between the Markov chain's states, without the dependence of those transitions on those events having to be made explicit or concrete. This abstraction is a strength, in that our results apply to a wide variety of models. However, it also presents a limitation, in that our results cannot be expressed in terms of these fundamental events, because they are represented only implicitly.

This limitation can be overcome by restricting our analyses to a subclass of models for which ecological events are represented more explicitly. An example is the class of models with "global updating" (Nathanson et al. 2009). In this class, the update rule has the property that an individual's reproductive fitness (the number of same-type individuals that an individual leaves after an update; Taylor et al. 2007b; Allen and Tarnita 2012) is proportional to its expected game payoff. This property can be understood ecologically as requiring that competition to reproduce be global, as local density regulation of survival or fertility otherwise break this proportionality. Notwithstanding this requirement, game interactions may still be local. This class includes most models of well-mixed populations, as well as, for example, Antal et al.'s (2009) model of evolution

with phenotypic assortment, Tarnita et al.'s (2009a), model of evolution in social sets, and Allen and Nowak's (2012) model of spatial evolution in cellular populations. For all models in this class, the structure coefficient σ can be expressed in terms of the relative frequency of same-type versus different-type game interactions (Nathanson et al. 2009), that is, as a measure of assortment.

For models without global updating, including models in evolutionary graph theory (Ohtsuki et al. 2006; Pacheco et al. 2006b; Taylor et al. 2007a; Allen et al. 2012), the structure coefficient does not merely quantify assortment in game interactions, but also reflects interactions (e.g., competition for space) that are described by the update rule. For example, the death-birth and birth-death update rules on regular graphs (Ohtsuki et al. 2006) yield the same spatial assortment pattern but result in different values of σ , due to differences in the way spatial competition is modeled (see table 1; Ohtsuki et al. 2006; Nowak et al. 2010b).

Research is ongoing to develop and investigate new classes of IBEG models, with various degrees of generality. For some of these classes, it should be possible to express σ in terms of well-studied quantities beyond fixation probabilities. The ultimate result may be a patchwork of classes, with expressions for σ that differ across classes but coincide on their intersections.

Discrete versus Continuous Strategy Evolution

The adaptive dynamics of continuous game strategies in IBEG models differs in several noteworthy ways from discrete strategy evolution. First, for discrete strategies, condition (4) is valid only under weak selection. For continuous strategies, since mutation is incremental in the adaptive dynamics framework, the difference in strategy between competing mutants and residents is assumed to be small. Consequently, fitness differences within the evolving population are small as well. Thus, no extra assumption of weak selection is needed for the canonical equation with interaction structure equation (10) to hold.

Second, condition (4) determines which of the two strategies X and Y is more likely to displace the other; however, for discrete strategies, it does not quantify the absolute likelihood of these fixation events. Since absolute fixation probabilities are needed to predict long-term evolutionary dynamics based on strategy substitutions, for discrete strategies, the structure coefficient and effective population size together do not fully characterize the effects of different interaction structures and update rules on long-term evolution. This situation is different for the adaptive dynamics of continuous strategies: the combined assumptions of incremental mutations and of the differentiability of the payoff function $A(x'; x)$ imply that the

first-order behavior of $\rho(x'; x)$ near $x' = x$ is fully determined by σ , N_σ , and the partial derivatives of $A(x'; x)$ at $x' = x$, as we show in appendix B.

Relation to Inclusive Fitness Theory

The form of the canonical equation with interaction structure, equation (10), is reminiscent of inclusive fitness theory (Hamilton 1964a, 1964b; Michod 1982), in that the effects of a trait are partitioned into two terms representing benefits to self and to others, with the second term weighted by a quantity related to interaction patterns.

Yet the canonical equation with interaction structure represents a different perspective than inclusive fitness theory. While no inclusive-fitness analogue of equation (10) has yet been proposed (see Rousset 2004 for some steps in this direction), inclusive fitness theory typically focuses on partitioning fitnesses rather than game payoffs (Taylor and Frank 1996; Taylor et al. 2007b).

While this distinction may at first glance seem immaterial, it does affect the utility and domains of applicability of the two approaches. To apply equation (10), an IBEG model must be specified—that is, all processes relevant to evolution must be partitioned into those represented by the game, interaction structure, and update rule. This involves an assumption that all outcomes of those processes represented by the game are adequately summarized by the payoffs assigned to the participating individuals. The advantage gained from this assumption is the clean separation of the game from the interaction structure and update rule. These ingredients can then be varied independently in equation (10), enabling simple yet comprehensive analyses of how the evolutionary trajectories of particular traits vary under different conditions (e.g., figs. 1 and 2).

In contrast, when fitness is partitioned according to inclusive fitness theory, the interaction structure and update rule affect the two fitness terms as well as the relatedness coefficient (Taylor et al. 2007b; Nowak et al. 2010b). These terms must therefore be calculated separately for every model, rendering the aforementioned type of comprehensive analysis more difficult in the inclusive fitness framework.

General Approaches to Evolutionary Theory

The results presented here apply to a class of evolutionary models, defined by a set of assumptions. Reasoning from general assumptions, rather than specific models, is a relatively new and potentially powerful approach in evolutionary theory. This approach has previously been applied to the dynamics of physiologically structured populations (Metz and Diekmann 1986; Diekmann et al. 1998, 2001,

2007; Durinx et al. 2008), evolutionary game theory (Tarnita et al. 2009b, 2011), quantitative trait evolution (Champagnat et al. 2006; Simon 2008), and models with fixed spatial structure (Allen and Tarnita 2012). The advantage of this approach is that general statements, applying to many models or systems, can be proven all at once.

There is another general approach to evolutionary theory, which is termed the “open model” approach by Gardner and West (2006), and is perhaps best typified by the work of Price (1970, 1972). The open model approach does not reason from any particular model or set of assumptions. Rather, the starting point is a collection of variables that represent quantities arising in an unspecified evolutionary process. These variables are related to each other using mathematical identities—such as the Price (1970) equation—that hold irrespective of the biological meaning associated to the variables. These identities are then rewritten into forms that can be interpreted as quantitative laws of evolution. For example, versions of Hamilton’s rule have been derived through this approach (Queller 1992; Gardner et al. 2011), as have certain formulations of multilevel selection theory (Price 1972; Oksa-sha 2006). The identities derived through this approach are valid in any system for which the variables involved have meaning but do not, on their own, determine the outcome of any particular model or the consequences of any particular assumption.

A dichotomy has been proposed (Gardner and West 2006; Gardner et al. 2007, 2011) between “closed model” (modeling in the traditional sense) and open model approaches to evolutionary theory. The approach taken here and by others mentioned above—reasoning from a set of assumptions that delineate a class of models—does not appear to fall under either category. We take the view that there is a spectrum of generality in mathematical approaches to evolutionary theory. At one extreme are detailed models representing specific systems, while the open model approach represents the opposite extreme. Our approach and others like it occupy intermediate positions in this spectrum.

Inherent in the conceptualization of this spectrum is the acknowledgement of a trade-off between generality of the domain of applicability versus specificity in results. Approaches of intermediate generality (e.g., classical mechanics as derived from Newton’s laws of motion) have been tremendously powerful in other fields of science. We believe such approaches represent an exciting growth opportunity for evolutionary theory.

Limitations

We caution that, in addition to the assumptions allowing trait evolution to be described by evolutionary game the-

ory, our results depend on the assumptions underlying the adaptive dynamics approach. In particular, our use of condition (4) and other results from evolutionary game theory requires assuming rare and incremental mutations. Relaxation of these assumptions would likely reveal a much more complex interplay between interaction structure, update rule, and game strategy evolution.

Additionally, our results assume that the evolving trait (continuous game strategy) is independent of the interaction structure and update rule. This excludes an important class of models in which behaviors affecting the interaction structure or update rule coevolve with strategy (Le Galliard et al. 2005; Pacheco et al. 2006a, 2006b; Fu et al. 2008; Helbing and Yu 2009; Perc and Szolnoki 2010; Wu et al. 2010). For such models, the value of the structure coefficient may vary with the considered game and resident strategy. Establishing the relationship between the structure coefficient and the resident strategy for different games could be an interesting research target in its own right. In such cases, however, the canonical equation with interaction structure, equation (10), loses the generality that comes from assuming the independence of the game from the interaction structure and update rule, because the aforementioned relationship may be game-specific.

Outlook

Our results have a number of implications for future research. For one, they suggest extending the focus of analysis from the detailed study of individual models to a general understanding of common features and quantifiable differences between models. Second, the canonical equation with interaction structure, equation (10), may

help illuminate how different interaction structures and update rules affect a wide range of phenomena, such as life-history evolution (Stearns 1992; Crowley and McLetchie 2002; Roff 2002), virulence evolution (Herre 1993; Boots and Sasaki 1999; Haraguchi and Sasaki 2000; Read and Keeling 2003; Boots et al. 2004), and evolutionary suicide (Ferrière 2000; Boots and Sasaki 2002; Gyllenberg et al. 2002; Lion and van Baalen 2008). Finally, the appearance of the marginal-benefit-to-marginal-cost ratio in condition (12) suggests that quantities typically considered in economic theory may be useful for understanding the adaptive dynamics of quantitative traits.

Acknowledgments

B.A. completed part of this research while participating in the Young Scientists Summer Program (YSSP) at the International Institute of Applied Systems Analysis (IIASA) in Laxenburg, Austria. He thanks his YSSP advisor A. Tarasiev, as well as Å. Brännström, B. Fath, C. Hilbe, J. A. J. Metz, and C. E. Tarnita for helpful suggestions. Participation in the IIASA Young Scientists Summer Program was made possible by a grant from the National Academy of Sciences Board on International Scientific Organizations, funded by the National Science Foundation under grant OISE-0738129. The Foundational Questions in Evolutionary Biology initiative at Harvard University is supported by a grant from the John Templeton Foundation. U.D. gratefully acknowledges support by the European Commission, the European Science Foundation, the Austrian Science Fund, the Austrian Ministry for Science and Research, and the Vienna Science and Technology Fund.

APPENDIX A

Mathematical Framework

Here we present our mathematical framework and assumptions formally. We state these in the case of a pairwise game, saving the generalization to games with more than two players for appendix C. Below, we divide our assumptions into those applying to the interaction timescale, the demographic timescale, and the evolutionary timescale. Where needed, explanations in italics provide interpretations of the preceding mathematical statements.

A1. Interaction Timescale: Game Strategies and Payoffs

We assume that the payoff function A satisfies the following condition.

G1 For each $x \in \mathbb{R}^d$, A is positive and twice differentiable in both arguments at $x' = x$.

A2. Demographic Timescale: Resident-Mutant Competition

We assume that resident-mutant competition is described by a model satisfying the following conditions.

D1 There is a finite set S with an associated probability distribution Q and a distinguished subset $F_M \subset S$ that is assigned zero probability by Q .
 S is the set of states. Q is the probability distribution of states corresponding to the appearance of a new mutant strategy. F_M is the set of states corresponding to mutant fixation.

D2 For any payoff matrix

$$\mathbf{G} = \begin{pmatrix} a_{MM} & a_{MR} \\ a_{RM} & a_{RR} \end{pmatrix},$$

there is a collection $\{p_{s'|s}\}_{s,s' \in S}$ of transition probabilities, giving S the structure of a Markov chain.

D3 The Markov chain associated to any such payoff matrix \mathbf{G} has the following properties:

- There is zero probability of transitioning from a state in F_M to a state not in F_M .
Mutant fixation is irreversible (barring further mutation).
- For any $s \in S$ that is assigned positive probability by Q , and any $s' \in F_M$, there is a positive integer n for which the probability of transitioning from s to s' in n steps is positive.
Mutant fixation is possible from any initial state.

D4 The transition probabilities $p_{s'|s}$ vary twice differentially with respect to the entries of \mathbf{G} .

D5 If the payoff matrix \mathbf{G} is multiplied by a constant $K > 0$, the probability ρ that the Markov chain hits F_M , given that its initial state is sampled from Q , is unaffected.
The evolutionary dynamics are insensitive to the choice of unit by which payoffs are quantified.

D6 The probability ρ defined in assumption D5 is increasing in a_{MM} and a_{MR} , and decreasing in a_{RM} and a_{RR} , for all values of a_{MM} , a_{MR} , a_{RM} , and a_{RR} sufficiently close to each other (that is, for all values of a_{MM} , a_{MR} , a_{RM} , and a_{RR} whose pairwise ratios are sufficiently close to 1).
Mutant fixation probability increases with mutant payoffs and decreases with resident payoffs, at least when these payoffs are close to each other.

A3. Evolutionary Timescale: Mutations and Trait Substitutions

We place the following assumptions on the adaptive dynamics of game strategies on the evolutionary timescale:

E1 Adaptive evolution follows a Markov jump process with state space \mathbb{R}^d . This process depends on the following ingredients:

- A payoff function $A(x; y)$ satisfying assumption G1,
- A positive real-valued function u on \mathbb{R}^d ,
- A probability measure U on \mathbb{R}^d , with an associated σ -algebra of measurable sets,
- A resident-mutant competition model satisfying assumptions D1–D6.

The infinitesimal transition rate to state $x' \in \mathbb{R}^d$, given that the current state is x , is equal to

$$Nu(x)dU(x' - x)\rho(x'; x).$$

Above, $\rho(x'; x)$ is the fixation probability ρ for the resident-mutant competition model, as defined in assumption D5, with the entries of \mathbf{G} given by

$$\begin{aligned} a_{MM} &= A(x'; x'), & a_{MR} &= A(x'; x), \\ a_{RM} &= A(x; x'), & a_{RR} &= A(x; x). \end{aligned}$$

A Markov jump process is a continuous-time Markov process in which the state is constant except for being punctuated by instantaneous jumps to new states. In this case, the state $x \in \mathbb{R}^d$ represents the resident strategy, and jumps represent substitution events, through which a new mutant strategy displaces the resident strategy. We regard these substitution events as instantaneous from the perspective of long-term evolution. Representing evolution in this way formalizes the assumption, often made in the adaptive dynamics approach, that mutations are rare enough that the resident population can be effectively regarded as typically monomorphic (see, e.g., Dieckmann 1994; Champagnat et al. 2002).

The function $u(x)$ gives the per capita rate at which new mutants appear in a population with resident strategy x . The probability measure U represents the distribution of possible mutational steps $x' - x$. In the case that U has an associated density function M , we can write $dU(z) = M(z)dz$. The transition rate to state x' can be understood as the rate at which type x' displaces the current resident type x , decomposed as the product of the expected number of mutants born per unit time, $Nu(x)$, the (infinitesimal) probability that a mutant is of type x' , $dU(x' - x)$, and the probability that mutant type x' will displace resident type x , should it arise, $\rho(x'; x)$.

E2 For a random variable z distributed according to U , $E[z] = 0$.
Mutation is directionally unbiased.

E3 For a random variable z distributed according to U , $E[|z|^2] \gg E[|z|^3]$.
The third moment of mutational step size is eclipsed by the second, reflecting an assumption of incremental mutation.

For notational compactness, we additionally assume that N and U do not vary with x , and that there is no covariance between the components of a mutational step (i.e., $\text{Cov}[z_i, z_j] = 0$ for all $i, j = 1, \dots, d$). These assumptions are inessential and can be relaxed in a straightforward manner, as described by Dieckmann and Law (1996).

Our derivation can also be extended to situations in which the trait space (the set of possible x values) is not the whole of \mathbb{R}^d but a domain $D \subset \mathbb{R}^d$, as in examples 1 and 2 of the main text. In this case, the canonical equation, equation (10), holds at all points x for which the probability that $x + z \notin D$ is negligible. Since the distribution U of mutational steps is presumed to be narrow, this will typically include all points x that are sufficiently far from the boundary of D . There are several schemes for extending the canonical equation to points near the boundary (Dieckmann et al. 2006). The simplest of these—which we applied in examples 1 and 2—is to assume that equation (10) holds in the whole interior of D and that the dynamics on the boundary are given by subtracting, from the vector dx/dt obtained from equation (10), the component (if any) that points orthogonally outward from D .

APPENDIX B

Derivation

We now derive our main result, equation (10), based on the mathematical framework described above. We first express the expected change in strategy in terms of the fixation probability (sec. B1), and then in terms of game payoffs (sec. B2). These two sections derive the functional form of the canonical equation. Sections B3 and B5 relate the coefficients appearing in this equation to the effective population size and structure coefficient, respectively. Sections B4 and B6 then establish formulas for, and the positivity of, the effective population size and structure coefficient, respectively.

B1. Adaptive Dynamics in Terms of Fixation Probabilities

Assumption E1 implies the following equation for the expected change in strategy x per unit time,

$$\frac{dx}{dt} = Nu(x) \int_{\mathbb{R}^d} z\rho(x + z; x)dU(z).$$

Since ρ is an absorption probability of a finite Markov chain (assumptions D1–D3 and D5), and since absorption

is possible from any initial state sampled from Q (assumption D3), ρ varies smoothly with the Markov chain's transition probabilities (see, e.g., theorem 3.3 of Iosifescu 1980). Assumption D4 then implies that ρ is twice differentiable with respect to the payoff values a_{MM} , a_{MR} , a_{RM} and a_{RR} . By assumption G1, $\rho(x'; x)$ is twice differentiable in x and x' at points where $x' = x$. We can therefore replace $\rho(x + z; x)$ by its multivariate Taylor expansion around $z = 0$ up to first order,

$$\rho(x + z; x) = \rho(x; x) + \sum_{i=1}^d z_i \left. \frac{\partial \rho(x'; x)}{\partial x'_i} \right|_{x'=x} + \mathcal{O}(|z|^2),$$

yielding

$$\frac{dx}{dt} = Nu(x)\rho(x; x) \int_{\mathbb{R}^d} z dU(z) + Nu(x) \sum_{i=1}^d \left[\left. \frac{\partial \rho(x'; x)}{\partial x'_i} \right|_{x'=x} \int_{\mathbb{R}^d} z z_i dU(z) \right] + \mathcal{O} \left[\int_{\mathbb{R}^d} |z|^3 dU(z) \right]. \tag{B1}$$

By assumption E2, the first term on the right-hand side of equation (B1) is zero, and by assumption E3, the third term is negligible relative to the second. Thus equation (B1) can be replaced with

$$\frac{dx}{dt} = Nu(x) \sum_{i=1}^d \left[\left. \frac{\partial \rho(x'; x)}{\partial x'_i} \right|_{x'=x} \int_{\mathbb{R}^d} z z_i dU(z) \right].$$

Our assumption of no covariance among the components of z implies that

$$\int_{\mathbb{R}^d} z z_i dU(z) = v_i e_i,$$

where e_i is the unit vector for the i th component of \mathbb{R}^d . The expected change in x is therefore given by the system of equations

$$\frac{dx_i}{dt} = Nu(x) v_i \left. \frac{\partial \rho(x'; x)}{\partial x'_i} \right|_{x'=x}, \tag{B2}$$

for $i = 1, \dots, d$.

B2. Adaptive Dynamics in Terms of Game Payoffs

Using the multivariate chain rule, we obtain

$$\left. \frac{\partial \rho(x'; x)}{\partial x'_i} \right|_{x'=x} = \sum_{j, k \in \{M, R\}} \left. \frac{\partial \rho}{\partial a_{jk}} \right|_{G= \begin{pmatrix} A(x; x) & A(x; x) \\ A(x; x) & A(x; x) \end{pmatrix}} \left. \frac{\partial a_{jk}}{\partial x'_i} \right|_{x'=x}. \tag{B3}$$

By assumption D5, each of the payoff values a_{jk} can be divided by $a_{RR} = A(x; x)$ without changing the value of ρ . This implies that

$$\left. \frac{\partial \rho}{\partial a_{jk}} \right|_{G= \begin{pmatrix} A(x; x) & A(x; x) \\ A(x; x) & A(x; x) \end{pmatrix}} = \frac{1}{A(x; x)} \left. \frac{\partial \rho}{\partial a_{jk}} \right|_{G= \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}},$$

which, in combination with equation (B3), yields

$$\frac{\partial \rho}{\partial x'_i} \Big|_{x'=x} (x'; x) = \frac{1}{A(x; x)} \sum_{j, k \in \{M, R\}} \frac{\partial \rho}{\partial a_{jk}} \Big|_{\mathbf{G}=\begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} \frac{\partial a_{jk}}{\partial x'_i} \Big|_{x'=x}. \tag{B4}$$

By the multivariate chain rule, the partial derivatives of the a_{jk} at $x' = x$ are given by

$$\frac{\partial a_{MM}}{\partial x'_i} \Big|_{x'=x} = \frac{\partial A(x'; x)}{\partial x'_i} \Big|_{x'=x} + \frac{\partial A(x; x')}{\partial x'_i} \Big|_{x'=x},$$

$$\frac{\partial a_{MR}}{\partial x'_i} \Big|_{x'=x} = \frac{\partial A(x'; x)}{\partial x'_i} \Big|_{x'=x},$$

$$\frac{\partial a_{RM}}{\partial x'_i} \Big|_{x'=x} = \frac{\partial A(x; x')}{\partial x'_i} \Big|_{x'=x},$$

$$\frac{\partial a_{RR}}{\partial x'_i} \Big|_{x'=x} = 0.$$

We can therefore rewrite equation (B4) in the form

$$\frac{\partial \rho(x'; x)}{\partial x'_i} \Big|_{x'=x} = \frac{\kappa}{A(x; x)} \left[\frac{\partial A(x'; x)}{\partial x'_i} \Big|_{x'=x} + \kappa' \frac{\partial A(x; x')}{\partial x'_i} \Big|_{x'=x} \right]. \tag{B5}$$

Here, the quantities κ and κ' are given by

$$\kappa = \left(\frac{\partial \rho}{\partial a_{MM}} + \frac{\partial \rho}{\partial a_{MR}} \right) \Big|_{\mathbf{G}=\begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}},$$

$$\kappa' = \left[\left(\frac{\partial \rho}{\partial a_{MM}} + \frac{\partial \rho}{\partial a_{RM}} \right) \right] \Big|_{\mathbf{G}=\begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}}. \tag{B6}$$

We observe that κ and κ' are independent of A and x .

Combining equation (B5) with equation (B2) yields the dynamical equation

$$\frac{dx_i}{dt} = N \frac{u(x) v_i \kappa}{A(x; x)} \left[\frac{\partial A(x'; x)}{\partial x'_i} \Big|_{x'=x} + \kappa' \frac{\partial A(x; x')}{\partial x'_i} \Big|_{x'=x} \right]. \tag{B7}$$

It now only remains to relate κ and κ' to N_e and σ . We can establish these relationships by considering particularly simple payoff functions A and resident strategies x , and substituting these into equation (B5). Since κ , κ' , σ , and N_e are all independent of A and x , any relationship derived using a particular choice of A and x will hold generally.

B3. Appearance of Effective Population Size

As motivated in the main text, we define the effective population size for a given resident-mutant competition model as

$$N_e = \frac{N^2}{N-1} \frac{\partial \rho}{\partial s} \Big|_{s=0}, \tag{B8}$$

where ρ is the fixation probability of a mutant with selective advantage s relative to the resident population.

To relate N_e thus defined to the constant κ appearing in equation (B7), we must choose a specific payoff function

A to substitute into equation (B5). As concluded above, a particularly simple choice of A can be made without any loss of generality. We thus consider $A(x; y) = 1 + x$, describing frequency-independent selection, so that the link to selective advantage is clear. We choose $x = 0$ as the resident strategy. With these choices, mutants and residents have constant payoffs $1 + x'$ and 1 , respectively (regardless of their interaction partners), so we identify x' as the mutant's selective advantage; that is, $x' = s$.

Substituting this payoff function A and $x = 0$ in equation (B5) yields

$$\left. \frac{\partial \rho(x'; 0)}{\partial x'} \right|_{x'=0} = \kappa. \tag{B9}$$

Identifying x' with s and comparing with equation (B8), we obtain

$$\kappa = N_e \frac{N - 1}{N^2}. \tag{B10}$$

Since the values of κ and N_e do not depend on the game being played, this identity holds for all games.

B4. Alternate Expression for, and Positivity of, Effective Population Size

Equations (B10) and (B6) also imply an alternate expression for N_e :

$$N_e = \frac{N^2}{N - 1} \left(\frac{\partial \rho}{\partial a_{MM}} + \frac{\partial \rho}{\partial a_{MR}} \right) \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}}, \tag{B11}$$

We can use this expression to show the positivity of N_e . Assumption D6 implies that

$$\left. \frac{\partial \rho}{\partial a_{MM}} \right|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} > 0, \quad \left. \frac{\partial \rho}{\partial a_{MR}} \right|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} > 0.$$

In combination with equation (B11), we can thus conclude that N_e must be positive.

B5. Appearance of Structure Coefficient

To express the constant κ' appearing in equation (B7) in terms of σ , we first observe that for $x = x'$,

$$\mathbf{G} = \begin{pmatrix} A(x'; x') & A(x'; x') \\ A(x'; x') & A(x'; x') \end{pmatrix} = A(x'; x') \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}.$$

By assumption D5, the fixation probability $\rho(x'; x')$ does not depend on the value of $A(x'; x')$ and is therefore constant with respect to x' . Differentiating at $x' = 0$, we obtain

$$0 = \left. \frac{\partial \rho(x'; x')}{\partial x'} \right|_{x'=0} = \left. \frac{\partial \rho(x'; 0)}{\partial x'} \right|_{x'=0} + \left. \frac{\partial \rho(0; x')}{\partial x'} \right|_{x'=0},$$

and therefore

$$\left. \frac{\partial \rho(x'; 0)}{\partial x'} \right|_{x'=0} = - \left. \frac{\partial \rho(0; x')}{\partial x'} \right|_{x'=0}.$$

This implies that for all sufficiently small $|x'|$,

$$\rho(x'; 0) > \rho(0; x') \Leftrightarrow \left. \frac{\partial \rho(x'; 0)}{\partial x'} \right|_{x'=0} > 0. \tag{B12}$$

As above, we now choose a particular payoff function A to substitute into equation (B5). We consider the linear Prisoner's Dilemma with payoff function $A(x; y) = -cx + by + 1$, $b > c > 0$, and resident and mutant strategies $x = 0$ and $x' > 0$, respectively. The payoff matrix \mathbf{G} can then be written as

$$\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix} + x' \begin{pmatrix} b - c & -c \\ b & 0 \end{pmatrix}.$$

We observe from the form of this expression that the mutant strategy x' can be interpreted as a selection-strength parameter (Nowak et al. 2004; Tarnita et al. 2009b). By the defining condition of the structure coefficient, condition (4) in the main text, we obtain, for sufficiently small $x' > 0$,

$$\rho(x'; 0) > \rho(0; x') \Leftrightarrow \sigma(b - c) - c > b. \tag{B13}$$

Separately, substituting the chosen A and $x = 0$ into equation (B5) yields

$$\left. \frac{\partial \rho(x'; 0)}{\partial x'} \right|_{x'=0} = \kappa(-c + \kappa' b).$$

Applying equation (B12) and the positivity of κ , which follows from equation (B10) and the positivity of N_e , we obtain

$$\rho(x'; 0) > \rho(0; x') \Leftrightarrow -c + \kappa' b > 0. \tag{B14}$$

Comparing condition (B13) with condition (B14), we thus see that

$$\kappa' = \frac{\sigma - 1}{\sigma + 1}. \tag{B15}$$

Again, since the values of κ' and σ do not depend on the game being played, this identity holds for all games.

Substituting equation (B10) and equation (B15) into equation (B7) completes the derivation of the canonical equation of adaptive dynamics with interaction structure, equation (10).

B6. Explicit Expression for, and Positivity of, Structure Coefficient

We can also derive an expression for σ in terms of partial derivatives of ρ with respect to game payoffs. For this we first observe that, as a consequence of assumption D5,

$$\left(\frac{\partial \rho}{\partial a_{MM}} + \frac{\partial \rho}{\partial a_{MR}} + \frac{\partial \rho}{\partial a_{RM}} + \frac{\partial \rho}{\partial a_{RR}} \right) \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} = 0.$$

Combining this with equation (B15) and equation (B6) and solving for σ yields

$$\sigma = \left[\left(\frac{\partial \rho}{\partial a_{MM}} - \frac{\partial \rho}{\partial a_{RR}} \right) \Big/ \left(\frac{\partial \rho}{\partial a_{MR}} - \frac{\partial \rho}{\partial a_{RM}} \right) \right] \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}}. \tag{B16}$$

We can use this expression to show the positivity of σ . Assumption D6 implies that

$$\begin{aligned} \frac{\partial \rho}{\partial a_{MM}} \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} > 0, & \quad \frac{\partial \rho}{\partial a_{MR}} \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} > 0, \\ \frac{\partial \rho}{\partial a_{RM}} \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} < 0, & \quad \frac{\partial \rho}{\partial a_{RR}} \Big|_{\mathbf{G} = \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}} < 0. \end{aligned}$$

In combination with equation (B16), we can thus conclude that σ must be positive.

APPENDIX C

Generalization to Multilateral Interactions

The evolutionary dynamics of multilateral interactions, which can be described by games involving three or more players, are often more complex than dynamics arising from pairwise games (Gokhale and Traulsen 2010). One might therefore expect that the canonical equation, equation (10), would increase in complexity when applied to n -player games for $n \geq 3$. Yet, we show here that—perhaps surprisingly—this is not the case, as long as the game satisfies a reasonable symmetry property.

C1. Mathematical Framework for Multilateral Interactions

We model a multilateral interaction as an n -player continuous game with payoff function $B(x; y_1, \dots, y_{n-1})$, where $x \in \mathbb{R}^d$ is the strategy of the focal player receiving the payoff and $y_1, \dots, y_{n-1} \in \mathbb{R}^d$ are the strategies of the other participants in the game. We assume B satisfies the following two conditions:

G1 For any $x \in \mathbb{R}^d$, B is positive and twice differentiable in all arguments at $y_1 = \dots = y_{n-1} = x$.

G2 The value of B does not change if the arguments y_1, \dots, y_{n-1} are permuted.

We suppose the strategy x evolves according to a model satisfying the analogues of assumptions D1–D6 and E1–E3 (with the payoff matrix \mathbf{G} being replaced by an n -dimensional array, as described by Gokhale and Traulsen 2010).

C2. Reduction to Pairwise Games

We now show that the adaptive dynamics of x can equivalently be described by a model based on the pairwise game A with $A(x; y) = B(x; y, \dots, y)$. This means that each instance of the game B can effectively be replaced by an interaction in which the game A is played against each interaction partner and the resulting payoffs are averaged. Thus, the adaptive dynamics of n -player games reduce to those of pairwise games, as long as the former games are symmetric under the rearrangement of interaction partners (assumption G2).

This simplification follows from combining the aforementioned symmetry with the multivariate chain rule, as we now demonstrate. By the analogues of equation (B2) and equation (B4), dx/dt depends on x only insofar as the first-order behavior, in $|x' - x|$, of the entries of the payoff array (the analog of \mathbf{G}) depends on x . By symmetry (assumption G2), each entry of the payoff array is equal to one of the quantities

$$r_j = B(x; \overbrace{x', \dots, x'}^j, \overbrace{x, \dots, x}^{n-j-1}),$$

$$m_j = B(x'; \overbrace{x', \dots, x'}^j, \overbrace{x, \dots, x}^{n-j-1}),$$

for $j = 0, \dots, n-1$. These denote, respectively, the payoff to a resident and to a mutant when interacting with j mutants and $n-j-1$ residents. Expanding r_j to first order in $|x' - x|$ yields

$$r_j = B(x; x, \dots, x) + (x' - x) \frac{\partial B}{\partial x'}(x; \overbrace{x', \dots, x'}^j, \overbrace{x, \dots, x}^{n-j-1}) \Big|_{x'=x} + \mathcal{O}(|x' - x|^2).$$

Invoking the symmetry property, the multivariate chain rule, and the definition of the pairwise game A , we obtain

$$\begin{aligned}
r_j &= B(x; x, \dots, x) + j(x' - x) \frac{\partial B}{\partial x'}(x; x', x, \dots, x) \Big|_{x'=x} + \mathcal{O}(|x' - x|^2) \\
&= B(x; x, \dots, x) + \frac{j}{n-1}(x' - x) \frac{\partial B}{\partial x'}(x; x', \dots, x') \Big|_{x'=x} + \mathcal{O}(|x' - x|^2) \\
&= A(x; x) + \frac{j}{n-1}(x' - x) \frac{\partial A(x; x')}{\partial x'} \Big|_{x'=x} + \mathcal{O}(|x' - x|^2).
\end{aligned}$$

Algebraic rearrangement yields

$$r_j = \frac{n-j-1}{n-1} A(x; x) + \frac{j}{n-1} \left[A(x; x) + (x' - x) \frac{\partial A(x; x')}{\partial x'} \Big|_{x'=x} \right] + \mathcal{O}(|x' - x|^2).$$

Using the first-order Taylor expansion of $A(x; x')$ around $x' = x$, we obtain the substitution

$$A(x; x) + (x' - x) \frac{\partial A(x; x')}{\partial x'} \Big|_{x'=x} = A(x; x') + \mathcal{O}(|x' - x|^2),$$

which finally yields

$$r_j = \frac{n-j-1}{n-1} A(x; x) + \frac{j}{n-1} A(x; x') + \mathcal{O}(|x' - x|^2).$$

By an analogous argument, we can show that

$$m_j = \frac{n-j-1}{n-1} A(x'; x) + \frac{j}{n-1} A(x'; x') + \mathcal{O}(|x' - x|^2).$$

In summary, the payoff obtained from playing the n -player game B with a set of $n-1$ interaction partners is equal, to first order in $|x' - x|$, to the arithmetic mean payoff obtained from playing the pairwise game A separately with each interaction partner. Since the canonical equation, equation (10), depends only on the first derivatives of the payoff function at $x' = x$, the adaptive dynamics of x based on the n -player game B is therefore fully described by a model based on the pairwise game A with averaged payoffs.

Literature Cited

- Allen, B., and M. A. Nowak. 2012. Evolutionary shift dynamics on a cycle. *Journal of Theoretical Biology* 311:28–39.
- Allen, B., and C. Tarnita. 2012. Measures of success in a class of evolutionary models with fixed population size and structure. *Journal of Mathematical Biology*, doi:10.1007/s00285-012-622-x.
- Allen, B., A. Traulsen, C. E. Tarnita, and M. A. Nowak. 2012. How mutation affects evolutionary games on graphs. *Journal of Theoretical Biology* 299:97–105.
- Antal, T., H. Ohtsuki, J. Wakeley, P. D. Taylor, and M. A. Nowak. 2009. Evolution of cooperation by phenotypic similarity. *Proceedings of the National Academy of Sciences of the USA* 106:8597–8600.
- Barton, N. H., and M. Slatkin. 1986. A quasi-equilibrium theory of the distribution of rare alleles in a subdivided population. *Heredity* 56:409–415.
- Boots, M., P. Hudson, and A. Sasaki. 2004. Large shifts in pathogen virulence relate to host population structure. *Science* 303:842–844.
- Boots, M., and A. Sasaki. 1999. “Small worlds” and the evolution of virulence: infection occurs locally and at a distance. *Proceedings of the Royal Society B: Biological Sciences* 266:1933–1938.
- . 2002. Parasite-driven extinction in spatially explicit host-parasite systems. *American Naturalist* 159:706–713.
- Brown, J., and T. Vincent. 1987. A theory for the evolutionary game. *Theoretical Population Biology* 31:140–166.
- Campos, P. R. A., and L. M. Wahl. 2009. The effects of population bottlenecks on clonal interference, and the adaptation effective population size. *Evolution* 63:950–958.
- Champagnat, N., R. Ferrière, and G. Ben Arous. 2002. The canonical equation of adaptive dynamics: a mathematical view. *Selection* 2: 73–83.
- Champagnat, N., R. Ferrière, and S. Méléard. 2006. Unifying evolutionary dynamics: from individual stochastic processes to macroscopic models. *Theoretical Population Biology* 69:297–321.
- Cressman, R. 1992. *The stability concept of evolutionary game theory: a dynamic approach*. Springer, Berlin.
- Crowley, P. H., and D. N. McLetchie. 2002. Trade-offs and spatial life-history strategies in classical metapopulations. *American Naturalist* 159:190–208.

- Dercole, F., and S. Rinaldi. 2008. Analysis of evolutionary processes: the adaptive dynamics approach and its applications. Princeton University Press, Princeton, NJ.
- Dieckmann, U. 1994. Coevolutionary dynamics of stochastic replicator systems. Jül-3018. Reports of the Research Center Jülich, Jülich, Germany.
- Dieckmann, U., and M. Doebeli. 1999. On the origin of species by sympatric speciation. *Nature* 400:354–357.
- Dieckmann, U., M. Heino, and K. Parvinen. 2006. The adaptive dynamics of function-valued traits. *Journal of Theoretical Biology* 241:370–389.
- Dieckmann, U., and R. Law. 1996. The dynamical theory of coevolution: a derivation from stochastic ecological processes. *Journal of Mathematical Biology* 34:579–612.
- Dieckmann, U., R. Law, and J. A. J. Metz, eds. 2000. The geometry of ecological interactions: simplifying spatial complexity. Cambridge University Press, Cambridge.
- Dieckmann, U., P. Marrow, and R. Law. 1995. Evolutionary cycling in predator-prey interactions: population dynamics and the Red Queen. *Journal of Theoretical Biology* 176:91–102.
- Diekmann, O., M. Gyllenberg, H. Huang, M. Kirkilionis, J. A. J. Metz, and H. R. Thieme. 2001. On the formulation and analysis of general deterministic structured population models II. Nonlinear theory. *Journal of Mathematical Biology* 43:157–189.
- Diekmann, O., M. Gyllenberg, and J. A. J. Metz. 2007. Physiologically structured population models: towards a general mathematical theory. Pages 5–20 in Y. Takeuchi, Y. Iwasa, and K. Sato, eds. *Mathematics for ecology and environmental sciences, biological and medical physics, biomedical engineering*. Springer, Berlin.
- Diekmann, O., M. Gyllenberg, J. A. J. Metz, and H. R. Thieme. 1998. On the formulation and analysis of general deterministic structured population models I. Linear theory. *Journal of Mathematical Biology* 36:349–388.
- Durinx, M., J. A. J. Metz, and G. Meszéna. 2008. Adaptive dynamics for physiologically structured population models. *Journal of Mathematical Biology* 56:673–742.
- Durrett, R., and S. Levin. 1994. The importance of being discrete (and spatial). *Theoretical Population Biology* 46:363–394.
- Eshel, I., and L. L. Cavalli-Sforza. 1982. Assortment of encounters and evolution of cooperativeness. *Proceedings of the National Academy of Sciences of the USA* 79:1331–1335.
- Ewens, W. J. 1982. On the concept of the effective population size. *Theoretical Population Biology* 21:373–378.
- Fehl, K., D. J. van der Post, and D. Semmann. 2011. Co-evolution of behaviour and social network structure promotes human cooperation. *Ecology Letters* 14:546–551.
- Ferrière, R. 2000. Adaptive responses to environmental threats: evolutionary suicide, insurance, and rescue. *Options Spring*:12–16.
- Ferrière, R., and J.-F. Le Galliard. 2001. Invasion fitness and adaptive dynamics in spatial population models. Pages 57–79 in J. Clobert, A. Dhondt, E. Danchin, and J. Nichols, eds. *Dispersal*. Oxford University Press, Oxford.
- Fisher, R. A. 1930. The genetical theory of natural selection. Clarendon Press, Oxford.
- Fletcher, J. A., and M. Doebeli. 2009. A simple and general explanation for the evolution of altruism. *Proceedings of the Royal Society B: Biological Sciences* 276:13–19.
- Fu, F., C. Hauert, M. A. Nowak, and L. Wang. 2008. Reputation-based partner choice promotes cooperation in social networks. *Physical Review E* 78:026117.
- Fu, F., C. E. Tarnita, N. A. Christakis, L. Wang, D. G. Rand, and M. A. Nowak. 2012. Evolution of in-group favoritism. *Scientific Reports* 2:460.
- Gardner, A. 2010. Sex-biased dispersal of adults mediates the evolution of altruism among juveniles. *Journal of Theoretical Biology* 262:339–345.
- Gardner, A., and S. A. West. 2006. Demography, altruism, and the benefits of budding. *Journal of Evolutionary Biology* 19:1707–1716.
- Gardner, A., S. A. West, and A. S. Griffin. 2007. Is bacterial persistence a social trait? *PLoS ONE* 2:e752.
- Gardner, A., S. A. West, and G. Wild. 2011. The genetical theory of kin selection. *Journal of Evolutionary Biology* 24:1020–1043.
- Geritz, S. A. H. 2005. Resident-invader dynamics and the coexistence of similar strategies. *Journal of Mathematical Biology* 50:67–82.
- Geritz, S. A. H., M. Gyllenberg, F. J. A. Jacobs, and K. Parvinen. 2002. Invasion dynamics and attractor inheritance. *Journal of Mathematical Biology* 44:548–560.
- Geritz, S. A. H., É. Kisdi, G. Meszéna, and J. A. J. Metz. 1997. Evolutionarily singular strategies and the adaptive growth and branching of the evolutionary tree. *Evolutionary Ecology* 12:35–57.
- Gokhale, C. S., and A. Traulsen. 2010. Evolutionary games in the multiverse. *Proceedings of the National Academy of Sciences of the USA* 107:5500–5504.
- Grafen, A., and M. Archetti. 2008. Natural selection of altruism in inelastic viscous homogeneous populations. *Journal of Theoretical Biology* 252:694–710.
- Gyllenberg, M., K. Parvinen, and U. Dieckmann. 2002. Evolutionary suicide and evolution of dispersal in structured metapopulations. *Journal of Mathematical Biology* 45:79–105.
- Haccou, P., and Y. Iwasa. 1996. Establishment probability in fluctuating environments: a branching process model. *Theoretical Population Biology* 50:254–280.
- Hamilton, W. D. 1964a. The genetical evolution of social behaviour. I. *Journal of Theoretical Biology* 7:1–16.
- . 1964b. The genetical evolution of social behaviour. II. *Journal of Theoretical Biology* 7:17–52.
- . 1971. Geometry for the selfish herd. *Journal of Theoretical Biology* 31:295–311.
- Hanski, I., and O. E. Gaggiotti. 2004. *Ecology, genetics, and evolution of metapopulations*. Elsevier, Amsterdam.
- Hanski, I., and M. E. Gilpin. 1997. *Metapopulation biology: ecology, genetics, and evolution*. Academic Press, San Diego, CA.
- Haraguchi, Y., and A. Sasaki. 2000. The evolution of parasite virulence and transmission rate in a spatially structured population. *Journal of Theoretical Biology* 203:85–96.
- Hassell, M. P., H. N. Comins, and R. M. May. 1991. Spatial structure and chaos in insect population dynamics. *Nature* 353:255–258.
- Hauert, C., and M. Doebeli. 2004. Spatial structure often inhibits the evolution of cooperation in the snowdrift game. *Nature* 428:643–646.
- Heino, M., J. A. J. Metz, and V. Kaitala. 1998. The enigma of frequency-dependent selection. *Trends in Ecology and Evolution* 13:367–370.
- Helbing, D., and W. Yu. 2009. The outbreak of cooperation among success-driven individuals under noisy conditions. *Proceedings of the National Academy of Sciences of the USA* 106:3680–3685.
- Herre, E. 1993. Population structure and the evolution of virulence in nematode parasites of fig wasps. *Science* 259:1442–1445.

- Hofbauer, J., and K. Sigmund. 1990. Adaptive dynamics and evolutionary stability. *Applied Mathematics Letters* 3:75–79.
- . 1998. *Evolutionary games and replicator dynamics*. Cambridge University Press, Cambridge.
- . 2003. Evolutionary game dynamics. *Bulletin of the American Mathematical Society* 40:479–520.
- Ifti, M., T. Killingback, and M. Doebeli. 2004. Effects of neighbourhood size and connectivity on the spatial continuous Prisoner's Dilemma. *Journal of Theoretical Biology* 231:97–106.
- Imhof, L. A., and M. A. Nowak. 2006. Evolutionary game dynamics in a Wright-Fisher process. *Journal of Mathematical Biology* 52: 667–681.
- . 2010. Stochastic evolutionary dynamics of direct reciprocity. *Proceedings of the Royal Society B: Biological Sciences* 277:463–468.
- Iosifescu, M. 1980. *Finite Markov processes and their applications*. Wiley Series in Probability and Mathematical Statistics. Wiley, New York.
- Killingback, T., and M. Doebeli. 1996. Spatial evolutionary game theory: hawks and doves revisited. *Proceedings of the Royal Society B: Biological Sciences* 263:1135–1144.
- . 2002. The continuous Prisoner's Dilemma and the evolution of cooperation through reciprocal altruism with variable investment. *American Naturalist* 160:421–438.
- Killingback, T., M. Doebeli, and N. Knowlton. 1999. Variable investment, the continuous Prisoner's Dilemma, and the origin of cooperation. *Proceedings of the Royal Society B: Biological Sciences* 266:1723–1728.
- Kimura, M. 1964. Diffusion models in population genetics. *Journal of Applied Probability* 1:177–232.
- Kimura, M., and G. H. Weiss. 1964. The stepping stone model of population structure and the decrease of genetic correlation with distance. *Genetics* 49:561–576.
- Krakauer, D. C., and M. Pagel. 1995. Spatial structure and the evolution of honest cost-free signalling. *Proceedings of the Royal Society B: Biological Sciences* 260:365–372.
- Le Galliard, J.-F., R. Ferrière, and U. Dieckmann. 2003. The adaptive dynamics of altruism in spatially heterogeneous populations. *Evolution* 57:1–17.
- . 2005. Adaptive evolution of social traits: origin, trajectories, and correlations of altruism and mobility. *American Naturalist* 165:206–224.
- Lessard, S., and V. Ladret. 2007. The probability of fixation of a single mutant in an exchangeable selection model. *Journal of Mathematical Biology* 54:721–744.
- Lieberman, E., C. Hauert, and M. A. Nowak. 2005. Evolutionary dynamics on graphs. *Nature* 433:312–316.
- Lion, S., and M. van Baalen. 2008. Self-structuring in spatial evolutionary ecology. *Ecology Letters* 11:277–295.
- Matsuda, H., N. Ogita, A. Sasaki, and K. Sato. 1992. Statistical mechanics of population. *Progress in Theoretical Physics* 88:1035–1049.
- Maynard Smith, J., and G. R. Price. 1973. The logic of animal conflict. *Nature* 246:15–18.
- McGill, B. J., and J. S. Brown. 2007. Evolutionary game theory and adaptive dynamics of continuous traits. *Annual Review of Ecology, Evolution, and Systematics* 38:403–435.
- Metz, J. A. J. 2011. Effective population sizes and the canonical equation of adaptive dynamics. *In* Workshop on stochastic processes in cell and population biology. Mathematical Biosciences Institute, Ohio State University, Columbus.
- Metz, J. A. J., and O. Diekmann. 1986. The dynamics of physiologically structured populations. Vol. 68. *Lecture Notes in Biomathematics*. Springer, Berlin.
- Metz, J. A. J., S. A. H. Geritz, G. Meszéna, F. A. Jacobs, and J. S. van Heerwaarden. 1996. Adaptive dynamics: a geometrical study of the consequences of nearly faithful reproduction. Pages 183–231 *in* S. J. van Strien and S. M. V. Lunel, eds. *Stochastic and spatial structures of dynamical systems*. Royal Dutch Academy of Sciences Memoirs (KNAW Verhandelingen), Amsterdam.
- Metz, J. A. J., and M. Gyllenberg. 2001. How should we define fitness in structured metapopulation models? including an application to the calculation of evolutionarily stable dispersal strategies. *Proceedings of the Royal Society B: Biological Sciences* 268:499–508.
- Metz, J. A. J., R. Nisbet, and S. Geritz. 1992. How should we define fitness for general ecological scenarios? *Trends in Ecology and Evolution* 7:198–202.
- Michod, R. E. 1982. The theory of kin selection. *Annual Review of Ecology and Systematics* 13:23–55.
- Mitteldorf, J., and D. S. Wilson. 2000. Population viscosity and the evolution of altruism. *Journal of Theoretical Biology* 204:481–496.
- Moran, P. A. P. 1958. Random processes in genetics. *Mathematical Proceedings of the Cambridge Philosophical Society* 54:60–71.
- Nakamaru, M., H. Matsuda, and Y. Iwasa. 1997. The evolution of cooperation in a lattice-structured population. *Journal of Theoretical Biology* 184:65–81.
- Nathanson, C. G., C. E. Tarnita, and M. A. Nowak. 2009. Calculating evolutionary dynamics in structured populations. *PLoS Computational Biology* 5:e1000615.
- Nowak, M., and K. Sigmund. 1990. The evolution of stochastic strategies in the Prisoner's Dilemma. *Acta Applicandae Mathematicae* 20:247–265.
- . 1993. Chaos and the evolution of cooperation. *Proceedings of the National Academy of Sciences of the USA* 90:5091–5094.
- Nowak, M. A. 2006. Five rules for the evolution of cooperation. *Science* 314:1560–1563.
- Nowak, M. A., and R. M. May. 1992. Evolutionary games and spatial chaos. *Nature* 359:826–829.
- Nowak, M. A., A. Sasaki, C. Taylor, and D. Fudenberg. 2004. Emergence of cooperation and evolutionary stability in finite populations. *Nature* 428:646–650.
- Nowak, M. A., and K. Sigmund. 1992. Tit for tat in heterogeneous populations. *Nature* 355:250–253.
- . 2004. Evolutionary dynamics of biological games. *Science* 303:793–799.
- Nowak, M. A., C. E. Tarnita, and T. Antal. 2010a. Evolutionary dynamics in structured populations. *Philosophical Transactions of the Royal Society B: Biological Sciences* 365:19–30.
- Nowak, M. A., C. E. Tarnita, and E. O. Wilson. 2010b. The evolution of eusociality. *Nature* 466:1057–1062.
- Ohtsuki, H. 2010. Evolutionary games in Wright's island model: kin selection meets evolutionary game theory. *Evolution* 64:3344–3353.
- Ohtsuki, H., C. Hauert, E. Lieberman, and M. A. Nowak. 2006. A simple rule for the evolution of cooperation on graphs and social networks. *Nature* 441:502–505.
- Ohtsuki, H., and M. A. Nowak. 2008. Evolutionary stability on graphs. *Journal of Theoretical Biology* 251:698–707.
- Ohtsuki, H., M. A. Nowak, and J. M. Pacheco. 2007. Breaking the

- symmetry between interaction and replacement in evolutionary dynamics on graphs. *Physical Review Letters* 98:108106.
- Okasha, S. 2006. *Evolution and the levels of selection*. Oxford University Press, Oxford.
- Pacheco, J. M., A. Traulsen, and M. A. Nowak. 2006a. Active linking in evolutionary games. *Journal of Theoretical Biology* 243:437–443.
- . 2006b. Coevolution of strategy and structure in complex networks with dynamical linking. *Physical Review Letters* 97:258103.
- Perc, M., and A. Szolnoki. 2010. Coevolutionary games: a mini review. *BioSystems* 99:109–125.
- Price, G. R. 1970. Selection and covariance. *Nature* 227:520–521.
- . 1972. Extension of covariance selection mathematics. *Annals of Human Genetics* 35:485–490.
- Proulx, S. R., and T. Day. 2002. What can invasion analyses tell us about evolution under stochasticity in finite populations? *Selection* 2:2–15.
- Queller, D. 1992. A general model for kin selection. *Evolution* 46:376–380.
- Rand, D. A., H. B. Wilson, and J. M. McGlade. 1994. Dynamics and evolution: evolutionarily stable attractors, invasion exponents and phenotype dynamics. *Philosophical Transactions of the Royal Society B: Biological Sciences* 343:261–283.
- Rand, D. G., S. Arbesman, and N. A. Christakis. 2011. Dynamic social networks promote cooperation in experiments with humans. *Proceedings of the National Academy of Sciences of the USA* 108:19193–19198.
- Read, J. M., and M. J. Keeling. 2003. Disease evolution on networks: the role of contact structure. *Proceedings of the Royal Society B: Biological Sciences* 270:699–708.
- Roca, C. P., J. A. Cuesta, and A. Sánchez. 2009. Effect of spatial structure on the evolution of cooperation. *Physical Review E* 80:046106.
- Roff, D. A. 2002. *Life history evolution*. Sinauer Associates, Sunderland, MA.
- Rousset, F. 2004. *Genetic structure and selection in subdivided populations*. Princeton University Press, Princeton, NJ.
- Santos, F. C., and J. M. Pacheco. 2005. Scale-free networks provide a unifying framework for the emergence of cooperation. *Physical Review Letters* 95:98104.
- Simon, B. 2008. A stochastic model of evolutionary dynamics with deterministic large-population asymptotics. *Journal of Theoretical Biology* 254:719–730.
- Stearns, S. C. 1992. *The evolution of life histories*. Oxford University Press, Oxford.
- Szabó, G., and G. Fáth. 2007. Evolutionary games on graphs. *Physics Reports* 446:97–216.
- Tarnita, C. E., T. Antal, H. Ohtsuki, and M. A. Nowak. 2009a. Evolutionary dynamics in set structured populations. *Proceedings of the National Academy of Sciences of the USA* 106:8601–8604.
- Tarnita, C. E., H. Ohtsuki, T. Antal, F. Fu, and M. A. Nowak. 2009b. Strategy selection in structured populations. *Journal of Theoretical Biology* 259:570–581.
- Tarnita, C. E., N. Wage, and M. A. Nowak. 2011. Multiple strategies in structured populations. *Proceedings of the National Academy of Sciences of the USA* 108:2334–2337.
- Taylor, P. D. 1992. Altruism in viscous populations: an inclusive fitness model. *Evolutionary Ecology* 6:352–356.
- Taylor, P. D., T. Day, and G. Wild. 2007a. Evolution of cooperation in a finite homogeneous graph. *Nature* 447:469–472.
- . 2007b. From inclusive fitness to fixation probability in homogeneous structured populations. *Journal of Theoretical Biology* 249:101–110.
- Taylor, P. D., and S. A. Frank. 1996. How to make a kin selection model. *Journal of Theoretical Biology* 180:27–37.
- Taylor, P. D., and L. B. Jonker. 1978. Evolutionary stable strategies and game dynamics. *Mathematical Biosciences* 40:145–156.
- Tilman, D., and P. M. Kareiva. 1997. *Spatial ecology: the role of space in population dynamics and interspecific interactions*. Princeton University Press, Princeton, NJ.
- Traulsen, A., and M. A. Nowak. 2006. Evolution of cooperation by multilevel selection. *Proceedings of the National Academy of Sciences of the USA* 103:10952–10955.
- Traulsen, A., J. M. Pacheco, and M. A. Nowak. 2007. Pairwise comparison and selection temperature in evolutionary game dynamics. *Journal of Theoretical Biology* 246:522–529.
- van Baalen, M. 2002. Contact networks and the evolution of virulence. Pages 85–103 in U. Dieckmann, J. A. J. Metz, M. W. Sabelis, and K. Sigmund, eds. *Adaptive dynamics of infectious diseases: in pursuit of virulence management*. Cambridge University Press, Cambridge.
- van Baalen, M., and D. A. Rand. 1998. The unit of selection in viscous populations and the evolution of altruism. *Journal of Theoretical Biology* 193:631–648.
- van Veelen, M., J. García, D. G. Rand, and M. A. Nowak. 2012. Direct reciprocity in structured populations. *Proceedings of the National Academy of Sciences of the USA* 109:9929–9934.
- Wahl, L. M., and M. A. Nowak. 1999a. The continuous Prisoner's Dilemma II. Linear reactive strategies with noise. *Journal of Theoretical Biology* 200:323–338.
- . 1999b. The continuous Prisoner's Dilemma I. Linear reactive strategies. *Journal of Theoretical Biology* 200:307–321.
- Weibull, J. W. 1997. *Evolutionary game theory*. MIT Press, Cambridge, MA.
- Werfel, J., and Y. Bar-Yam. 2004. The evolution of reproductive restraint through social communication. *Proceedings of the National Academy of Sciences of the USA* 101:11019–11024.
- Wilson, D. S. 1977. Structured demes and the evolution of group-advantageous traits. *American Naturalist* 111:157–185.
- Wilson, D. S., G. Pollock, and L. Dugatkin. 1992. Can altruism evolve in purely viscous populations? *Evolutionary Ecology* 6:331–341.
- Wright, S. 1931. Evolution in Mendelian populations. *Genetics* 16:97–159.
- . 1943. Isolation by distance. *Genetics* 28:114–138.
- Wu, B., D. Zhou, F. Fu, Q. Luo, L. Wang, A. Traulsen, and O. Sporns. 2010. Evolution of cooperation on stochastic dynamical networks. *PLoS ONE* 5:1560–1563.

Associate Editor: Peter D. Taylor
Editor: Mark A. McPeck