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www.elsevier.com/locate/jtbi

PII: S0022-5193(13)00567-5
DOI: <http://dx.doi.org/10.1016/j.jtbi.2013.12.016>
Reference: YJTBI7541

To appear in: *Journal of Theoretical Biology*

Received date: 4 October 2013
Revised date: 12 December 2013
Accepted date: 16 December 2013

Cite this article as: Jorge Peña, Laurent Lehmann, Georg Nöldeke, Gains from switching and evolutionary stability in multi-player matrix games, *Journal of Theoretical Biology*, <http://dx.doi.org/10.1016/j.jtbi.2013.12.016>

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Gains from switching and evolutionary stability in multi-player matrix games

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Abstract

In this paper we unify, simplify, and extend previous work on the evolutionary dynamics of symmetric N -player matrix games with two pure strategies. In such games, gains from switching strategies depend, in general, on how many other individuals in the group play a given strategy. As a consequence, the gain function determining the gradient of selection can be a polynomial of degree $N - 1$. In order to deal with the intricacy of the resulting evolutionary dynamics, we make use of the theory of polynomials in Bernstein form. This theory implies a tight link between the sign pattern of the gains from switching on the one hand and the number and stability of the rest points of the replicator dynamics on the other hand. While this relationship is a general one, it is most informative if gains from switching have at most two sign changes, as is the case for most multi-player matrix games considered in the literature. We demonstrate that previous results for

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public goods games are easily recovered and extended using this observation. Further examples illustrate how focusing on the sign pattern of the gains from switching obviates the need for a more involved analysis.

Keywords:

evolutionary game theory, replicator dynamics, polynomials in Bernstein form, public goods games

1. Introduction

Game theory has been widely applied to evolutionary biology (Maynard Smith and Price, 1973; Maynard Smith, 1982; Eshel, 1996; Hofbauer and Sigmund, 1998; Rousset, 2004; Vincent and Brown, 2005; Dercole and Rinaldi, 2008; Broom and Rychtář, 2013). More specifically, the application of game-theoretic concepts has been instrumental in explaining the evolution of traits as diverse as the sex ratio (Hamilton, 1967; Frank, 1987), dispersal (Hamilton and May, 1977; Comins et al., 1980), reciprocity (Axelrod and Hamilton, 1981), group foraging (Clark and Mangel, 1986), policing (Frank, 1995), and anisogamy (Bulmer and Parker, 2002). Evolutionary models of these traits often assume “playing the field” type of interactions (Maynard Smith, 1982, p. 23), where the payoff to an individual depends on an average property of the population or the group with which it interacts.

There are many situations, however, where the payoff to an individual depends critically on the strategy profile in the population (or its group) and where the actions of different individuals cannot be averaged; that is, mass action does not apply. Typical examples involve collective action problems in moderately sized groups, where the change in behavior by a single individual

19 can result in a large, discontinuous change in payoffs to others (e.g., Boyd
 20 and Richerson, 1988). Such collective action problems have been modeled as
 21 multi-player (or multi-person) matrix games (Broom et al., 1997; Kurokawa
 22 and Ihara, 2009; Gokhale and Traulsen, 2010). Except for the very special
 23 cases in which group size is taken to be equal to two (so that the well-
 24 developed theory of two-player matrix games can be applied, cf. Weibull,
 25 1995; Hofbauer and Sigmund, 1998; Cressman, 2003) or the payoff structure
 26 is linear (as in the standard model of the N -person prisoner's dilemma), such
 27 games have proven difficult to analyze.

The intrinsic complexity of multi-player matrix games is already evident
 for the case of symmetric games with two pure strategies A and B on which
 we focus in this paper. For these games, the average payoff difference in a
 large and well-mixed population is given by the so-called *gain function* (Bach
 et al., 2006)

$$g(x) = \sum_{k=0}^n \binom{n}{k} x^k (1-x)^{n-k} d_k.$$

28 Here, n is the number of co-players of a focal player (so that $N = n + 1$ is
 29 the group size), x is the population fraction of A-strategists, and d_k is the
 30 gain a focal player would obtain if switching from strategy B to strategy A
 31 when k other group-members play A. The evolutionary solution of the game
 32 (such as the set of evolutionarily stable strategies, ESSs, or the set of stable
 33 rest points of the replicator dynamics) involves not only finding the roots of
 34 the gain function $g(x)$ (a polynomial of degree n) but also, as discussed in
 35 Broom et al. (1997), determining the behavior of $g(x)$ in the vicinity of such
 36 roots. While this is straightforward for two-player games (for which $g(x)$ is
 37 linear in x) and a full classification for three-player games (for which $g(x)$ is

38 quadratic in x) is available (Bukowski and Miękisz, 2004), payoff structures
39 in groups of size larger than five lead to polynomials of degree greater than
40 four that cannot, in general, be solved analytically (Clark, 1984).

41 In order to deal with such complexity, the vast majority of previous works
42 on multi-player matrix games has considered particular functional forms for
43 the specification of the payoffs and has resorted to lengthy algebra or numer-
44 ical methods to study the models (Joshi, 1987; Boyd and Richerson, 1988;
45 Dugatkin, 1990; Weesie and Franzen, 1998; Hauert et al., 2006; Zheng et al.,
46 2007; Cuesta et al., 2008; Pacheco et al., 2009; Archetti, 2009; Souza et al.,
47 2009; Archetti and Scheuring, 2011; van Segbroeck et al., 2012). In this
48 way, some non-linear public goods games, including multi-player extensions
49 of well-known two-person matrix games such as the stag hunt (Skyrms, 2004)
50 and the snowdrift game (Sugden, 1986), have been characterized on a case-
51 by-case basis.

52 In contrast to these efforts, Motro (1991) and Bach et al. (2006) have
53 taken a more systematic approach to the study of non-linear public goods
54 games. Both of these papers consider situations in which each contributor to
55 a public good pays a constant cost, whereas the benefit from the public good,
56 which is obtained by all players, is a function of the number of contributors.
57 Motro (1991) proves that in this case the replicator dynamics has at most
58 one interior rest point if the benefit is concave or convex in the number of
59 contributors. He also provides necessary and sufficient conditions for the
60 existence of such a rest point and characterizes the stability property of all
61 rest points. In a similar spirit, Bach et al. (2006) find sufficient conditions on
62 the shape of the benefits such that there exists a critical cost level with the

63 property that for costs below such a level the replicator dynamics has two
64 interior rest points, whereas for higher costs there is no interior rest point.

65 Gokhale and Traulsen (2010) have discussed the relationship between the
66 sign pattern of the gains from switching and the number of interior rest points
67 of the replicator dynamics. Specifically, these authors observe that the repli-
68 cator dynamics has a single interior rest point if the sequence (d_0, d_1, \dots, d_n) ,
69 which we refer to as the *gain sequence*, has exactly one sign change. Gokhale
70 and Traulsen (2010) also note that the direction of selection (as given by
71 the sign of the gain function $g(x)$) cannot have more sign changes than the
72 gain sequence. This implies that the number of sign changes of the gain
73 sequence provides an upper bound on the number of interior rest points of
74 the replicator dynamics. The latter observation is also made in Hauert et al.
75 (2006) and Cuesta et al. (2007). When $g(x)$ has no multiple roots, any upper
76 bound on the number of interior rest points translates directly into an upper
77 bound on the number of stable rest points because, as noted in Broom et al.
78 (1997, p. 939), in this case the rest points alternate between being stable
79 and unstable.

80 In this paper, we show how sign-change conditions like the ones discussed
81 by Gokhale and Traulsen (2010) can be refined by using the fact that the gain
82 function $g(x)$ is a particular kind of polynomial, known as a polynomial in
83 Bernstein form (or Bernstein polynomial), with coefficients given by the gain
84 sequence (d_0, d_1, \dots, d_n) . Our analysis rests on the variation-diminishing
85 property of Bernstein polynomials and a property that we refer to as the
86 preservation of initial and final signs. These properties provide a tight link
87 between the sign pattern of the gain sequence and the sign pattern of the gain

88 function.¹ In particular, if the gain sequence has at most two sign changes,
 89 a full characterization of the possible dynamic regimes is easily obtained.

90 For most of the collective action problems that have been modeled as
 91 multi-player matrix games it is straightforward to determine the sign pat-
 92 tern of the gain sequence. Moreover, because the gain sequences of these
 93 games have at most two sign changes, our characterization results provide all
 94 the information necessary to recover the results on the number and stability
 95 of rest points obtained in previous studies. We demonstrate these claims for
 96 two classes of public goods games, namely threshold games (e.g., Dugatkin,
 97 1990; Weesie and Franzen, 1998; Zheng et al., 2007; Souza et al., 2009) and
 98 constant cost games (e.g., Motro, 1991; Bach et al., 2006; Hauert et al., 2006;
 99 Pacheco et al., 2009; Archetti and Scheuring, 2011), and two additional ex-
 100 amples taken from Hauert et al. (2006) and van Segbroeck et al. (2012), thus
 101 supporting the claim that the approach developed here unifies, simplifies,
 102 and extends much of the previous work on multi-player matrix games.

103 2. Model

104 Interactions occur in groups of size $N = n + 1$, in which a focal individual
 105 plays a game against n co-players or opponents. Each individual can choose
 106 between one of two different pure strategies, A and B. The game is symmetric
 107 so that, from the focal's point of view, any two co-players are exchangeable.

¹The fact that the gain function $g(x)$ is a Bernstein polynomial has previously been noted by Cuesta et al. (2007). These authors also suggest that the variation diminishing property of these polynomials may make the analysis of many multi-player games straightforward, but do not pursue this idea.

Let a_k denote the payoff to an individual choosing A when k opponents choose A (and hence $n - k$ co-players choose B); likewise, let b_k denote the payoff to an individual choosing B when k opponents choose A. Also let

$$d_k \equiv a_k - b_k$$

108 denote the gain the focal player makes from choosing A over B, taking the
 109 choices of other players (k playing A and $n - k$ playing B) as given. The
 110 parameters d_k , which describe the gains from switching, are collected in the
 111 gain sequence $\mathbf{d} = (d_0, d_1, \dots, d_n)$. We assume $\mathbf{d} \neq \mathbf{0}$, thus excluding the
 112 uninteresting case in which payoffs are independent of the actions chosen.

113 Evolution occurs in an infinitely large and well-mixed population with
 114 groups randomly formed by binomial sampling. Hence, if the frequency of
 115 A-strategists in the whole population is x , the average payoffs obtained by
 116 an A-strategist and a B-strategist are respectively given by

$$\pi_A(x) = \sum_{k=0}^n \binom{n}{k} x^k (1-x)^{n-k} a_k$$

117 and

$$\pi_B(x) = \sum_{k=0}^n \binom{n}{k} x^k (1-x)^{n-k} b_k.$$

118 We assume that the rules of transmission of the strategies (whether genet-
 119 ically encoded or individually or socially learned) are such that the frequency
 120 x of A-strategists in the population can be described by the replicator dy-
 121 namics (Taylor and Jonker, 1978; Hofbauer and Sigmund, 1998)

$$\frac{dx}{dt} = x(1-x)g(x), \quad (1)$$

122 where $g(x) = \pi_A(x) - \pi_B(x)$ is the gain function (Bach et al., 2006) given by

$$g(x) = \mathcal{B}_n(x; \mathbf{d}) \equiv \sum_{k=0}^n \binom{n}{k} x^k (1-x)^{n-k} d_k. \quad (2)$$

123 As we have already mentioned in the Introduction, the gain function is a *poly-*
 124 *nomial in Bernstein form* (also known as a *Bernstein polynomial*, cf. Farouki
 125 (2012)). This is made explicit by the notation we introduce in (2), where the
 126 *Bernstein operator* \mathcal{B}_n maps the vector of *Bernstein coefficients* $\mathbf{d} \in \mathbb{R}^{n+1}$
 127 into the polynomial $\sum_{k=0}^n \binom{n}{k} x^k (1-x)^{n-k} d_k$ in the variable $x \in [0, 1]$.

128 The replicator dynamics (1) has two trivial rest points at $x = 0$ (where
 129 the whole population consists of B-strategists) and $x = 1$ (where the whole
 130 population consists of A-strategists). Interior rest points $0 < x^* < 1$ are given
 131 by the solutions of the equation $g(x^*) = 0$. Because $g(x)$ is a polynomial of
 132 degree at most n (and we have assumed $\mathbf{d} \neq \mathbf{0}$) the replicator dynamics can
 133 have at most n interior rest points, corresponding to n simple roots of $g(x)$
 134 in the open interval $(0, 1)$. In the two-strategy case we analyze here, rest
 135 points of the replicator dynamics can be either (locally asymptotic) stable or
 136 unstable. Stability of a rest point x^* requires that $(x - x^*)(g(x) - g(x^*)) < 0$
 137 holds for all $x \neq x^*$ in the vicinity of x^* . Since the stable rest points of
 138 the replicator dynamics correspond to ESSs for the multi-player game (Bach
 139 et al., 2006), our following results about stable rest points of the replicator
 140 dynamics carry over to ESSs without any changes.

Remark 1. The gain function $g(x)$ given in (2) can also be interpreted as
 the selection gradient on a continuously varying mixed strategy x (denoting
 here the probability that an individual plays action A), evolving according
 to the traditional breeder's equation or the canonical equation of adaptive
 dynamics (Dieckmann and Law, 1996), so that the dynamics is of the form

$$\frac{dx}{dt} = v(x)g(x),$$

141 for some measure $v(x)$ of genetic variance (Kirkpatrick and Rousset, 2005).
142 Hence, all our subsequent results pertaining to polymorphic equilibria in
143 pure strategies can also be interpreted in terms of monomorphic equilibria
144 for mixed strategies.

145 **3. Sign patterns and (the stability of) rest points**

146 The fact that the gain function is a polynomial in Bernstein form implies
147 a tight link between the sign pattern of the gain sequence on the one hand
148 and the sign pattern and number of roots of the gain function on the other
149 hand. This is due to two properties of Bernstein polynomials, namely the
150 preservation of initial and final signs and the variation diminishing property
151 (see Properties 1 and 2 below). Because roots of the gain function correspond
152 to interior rest points of the replicator dynamics and the sign pattern of
153 the gain function informs us about changes in the direction of selection at
154 interior rest points (as well as the direction of selection at the trivial rest
155 points), general results about the number and stability of rest points follow
156 immediately (see Results 1 and 2). These results hold for any non-zero gain
157 sequence, allow for interior rest points at which the direction of selection
158 does not change, and provide more detailed information about the number
159 of rest points and stable equilibria than the observations made by Cuesta
160 et al. (2007) and Gokhale and Traulsen (2010). Results 3 to 5 summarize
161 the implications of the general results for gain sequences with at most two
162 sign changes, providing the basis for our subsequent analysis.

163 *3.1. Preliminaries*

164 To proceed, we require some terminology and notation to describe sign
 165 patterns (see Brown et al., 1981) and other relevant shape properties of gain
 166 sequences and gain functions. The same notation and terminology applies to
 167 other sequences and functions we encounter in our analysis.

168 Let $I(\mathbf{d})$ denote the sign (either $+$ or $-$) of the first non-zero entry in
 169 the sequence \mathbf{d} . Likewise, let $F(\mathbf{d})$ denote the sign of the last non-zero entry
 170 in \mathbf{d} . We refer to $I(\mathbf{d})$ and $F(\mathbf{d})$ as the initial and final signs of the gain
 171 sequence \mathbf{d} . We also denote by $S(\mathbf{d})$ the number of sign changes between
 172 consecutive entries in \mathbf{d} after zero entries have been eliminated. Obviously,
 173 $0 \leq S(\mathbf{d}) \leq n$.

174 As we have assumed $\mathbf{d} \neq \mathbf{0}$, there exists a neighborhood of $x^* = 0$ such
 175 that the sign of $g(x)$ is either $+$ or $-$ for all $x \neq 0$ in this neighborhood. We
 176 define the initial sign $I(g)$ of $g(x)$ as the sign of $g(x)$ in such neighborhood,
 177 and define the final sign $F(g)$ in an analogous way. Note that $I(g)$ coincides
 178 with the sign of $g(0)$ if $g(0) \neq 0$ holds. Similarly, if $g(1) \neq 0$ holds, then
 179 $F(g)$ coincides with the sign of $g(1)$. The number of sign changes $S(g)$ of
 180 the function $g(x)$ in the interval $(0, 1)$ is the number of times it crosses the
 181 x -axis in $(0, 1)$.

182 The notation $\Delta \mathbf{d} = (\Delta d_0, \dots, \Delta d_{n-1})$, where $\Delta d_k \equiv d_{k+1} - d_k$, denotes
 183 the (first) forward difference of the sequence \mathbf{d} . The second forward difference
 184 of the sequence \mathbf{d} is $\Delta^2 \mathbf{d} = (\Delta^2 d_0, \dots, \Delta^2 d_{n-2})$, where $\Delta^2 d_k \equiv \Delta d_{k+1} - \Delta d_k$.
 185 These forward differences can be viewed as the counterparts to the first and
 186 second derivatives of a real function and are a useful tool for describing the
 187 shape of a sequence. In particular, the sequence \mathbf{d} is increasing (resp. de-

188 creasing) if $\Delta \mathbf{d} \geq \mathbf{0}$ ($\Delta \mathbf{d} \leq \mathbf{0}$) holds, convex (resp. concave) if $\Delta^2 \mathbf{d} \geq \mathbf{0}$
 189 (resp. $\Delta^2 \mathbf{d} \leq \mathbf{0}$) holds, and unimodal (resp. anti-unimodal) if the sequence
 190 $\Delta \mathbf{d}$ has a single sign change from positive to negative (resp. from negative
 191 to positive). Corresponding definitions apply to the gain function $g(x)$. For
 192 instance, a gain function is unimodal if its first derivative $g'(x)$ has one sign
 193 change from positive to negative and is concave if its second derivative sat-
 194 isfies $g''(x) \leq 0$ for all $0 \leq x \leq 1$.

195 3.2. Stability of trivial rest points

196 One important property of the Bernstein operator \mathcal{B}_n is that it preserves
 197 end-points, i.e. $g(0) = \mathcal{B}_n(0; \mathbf{d}) = d_0$ and $g(1) = \mathcal{B}_n(1; \mathbf{d}) = d_n$ (Farouki,
 198 2012). From this, it is immediate that the initial and final signs of $g(x)$ and
 199 \mathbf{d} coincide in the case when $d_0 \neq 0$ and $d_n \neq 0$. We show in Appendix A
 200 that the same conclusion obtains in general, so that we have the following
 201 property.

Property 1 (Preservation of initial and final signs). The initial and fi-
 nal signs of $g(x)$ and \mathbf{d} coincide. That is,

$$I(g) = I(\mathbf{d}) \text{ and } F(g) = F(\mathbf{d}).$$

202 The initial sign of $g(x)$ describes the direction of selection in a vicinity of
 203 the trivial rest point $x = 0$, so that the rest point $x = 0$ is stable if and only
 204 if the initial sign of $g(x)$ is negative. Similarly, the rest point $x = 1$ is stable
 205 if and only if the final sign of $g(x)$ is positive. Hence, Property 1 implies that
 206 the initial and final signs of the gain sequence are all the information required
 207 to determine the stability of the trivial rest points. This is explicitly stated in

208 the following result, which has previously been noted by Broom et al. (1997,
209 Section 4.1).

210 **Result 1 (Stability of trivial rest points).**

- 211 1. The rest point $x = 0$ is stable if and only if $I(\mathbf{d}) = -$.
- 212 2. The rest point $x = 1$ is stable if and only if $F(\mathbf{d}) = +$.

213 The first part of Result 1 asserts that strategy A is disadvantageous when
214 rare if and only if the first non-zero element in the gain sequence is negative.
215 The second part is the assertion that strategy A is advantageous when com-
216 mon if and only if the last non-zero element in the gain sequence is positive.

217 *3.3. Number of (stable) interior rest points*

218 Let $R(g) \geq 0$ denote the number of roots of $g(x)$ in the interval $(0, 1)$,
219 counting roots according to their multiplicity. The following is the variation
220 diminishing property of Bernstein polynomials.

221 **Property 2 (Variation diminishing property).**

- 222 1. The number of roots of $g(x)$ on $(0, 1)$ is equal to the number of sign
223 changes of \mathbf{d} or less by an even amount. That is,

$$R(g) = S(\mathbf{d}) - 2i, \text{ where } i \geq 0 \text{ is an integer.} \quad (3)$$

- 224 2. The number of sign changes of $g(x)$ is equal to the number of sign
225 changes of \mathbf{d} or less by an even amount. That is,

$$S(g) = S(\mathbf{d}) - 2j, \text{ where } j \geq i \text{ is an integer.} \quad (4)$$

226 The first part of the variation-diminishing property (see e.g. Farouki
 227 (2012)) follows from Descartes' rule of signs, which hence can be said to
 228 "carry over" to polynomials in Bernstein form. The second part follows from
 229 the first upon observing that $x \in (0, 1)$ is the location of a sign change of
 230 $g(x)$ if and only if x is a root of $g(x)$ with odd multiplicity, so that $S(g)$ is
 231 either equal to $R(g)$ or less by an even amount.

232 As the interior rest points of the replicator dynamics coincide with the
 233 roots of $g(x)$, Property 2.1 applies as stated to the interior rest points of
 234 the replicator dynamics. In particular, as noted by Cuesta et al. (2007)
 235 and Gokhale and Traulsen (2010), the number of sign changes of the gain
 236 sequence \mathbf{d} provides an upper bound on the number of interior rest points.
 237 If the number of sign changes of \mathbf{d} is odd, (3) implies that $R(g)$ is odd.
 238 Consequently, the replicator dynamics possesses at least one interior rest
 239 point in this case.

240 Stability of an interior rest point is equivalent to the requirement that
 241 the sign of $g(x)$ changes from $+$ to $-$ at the rest point. As sign changes must
 242 alternate and initial signs are preserved (Property 1), the second part of the
 243 variation diminishing property yields the following result.

244 **Result 2 (Number of stable interior rest points).** Let ℓ denote the num-
 245 ber of stable interior rest points of the replicator dynamics and let $j \geq 0$ be
 246 the integer appearing in the statement of Property 2.2.

- 247 1. If $S(\mathbf{d})$ is even, then $\ell = S(g)/2 = S(\mathbf{d})/2 - j$.
- 248 2. If $S(\mathbf{d})$ is odd and $I(\mathbf{d}) = -$, then $\ell = (S(g) - 1)/2 = (S(\mathbf{d}) - 1)/2 - j$.
- 249 3. If $S(\mathbf{d})$ is odd and $I(\mathbf{d}) = +$, then $1 \leq \ell = (S(g) + 1)/2 = (S(\mathbf{d}) +$
 250 $1)/2 - j$.

251 In the generic case in which $g(x)$ has no multiple roots, the argument
 252 yielding Result 2 reduces to the one given by Broom et al. (1997, p. 939).

253 3.4. *Special cases*

254 It will be convenient to summarize the relationship between the sign pat-
 255 terns of the gain sequence and the rest points of the replicator dynamics
 256 for the cases in which the gain sequence has at most two sign changes. We
 257 also provide simple sufficient conditions ensuring that a gain sequence has at
 258 most one, resp. two sign changes.

259 3.4.1. *Gain sequences with one or no sign change*

260 When the gain sequence has no or one sign change, the variation diminish-
 261 ing property implies that the number of roots and the number of sign changes
 262 of the gain function both coincide with the number of sign changes of the
 263 gain sequence. In particular, Result 2 holds with $j = 0$. Combining these
 264 observations with Result 1 then shows that for games with gain sequences
 265 having at most one sign change, the sign pattern of the gain sequence con-
 266 tains all the information required to determine the number and stability of
 267 rest points. For later reference we state the ensuing case distinction in the
 268 following result.

269 **Result 3 (Gain sequences with no or one sign change).**

- 270 1. If the gain sequence has no sign changes, then the replicator dynamics
 271 has no interior rest points. Moreover
 - 272 (a) If $I(\mathbf{d}) = -$, then $x = 0$ is stable and $x = 1$ is unstable.
 - 273 (b) If $I(\mathbf{d}) = +$, then $x = 0$ is unstable and $x = 1$ is stable.

274 2. If the gain sequence has a single sign change, then the replicator dy-
 275 namics has a unique interior rest point x^* . Moreover:

276 (a) If $I(\mathbf{d}) = -$, then $x = 0$ and $x = 1$ are stable, and x^* is unstable.

277 (b) If $I(\mathbf{d}) = +$, then $x = 0$ and $x = 1$ are unstable, and x^* is stable.

278 The four possible dynamical regimes appearing in Result 3 correspond
 279 to the cases that are familiar from the evolutionary analysis of symmetric
 280 two-player games with two pure strategies (see, e.g. Cressman, 2003, Section
 281 2.2). This is, of course, not a coincidence: such two-player games are nothing
 282 but the special case of our model with $n = 1$ and thus feature gain sequences
 283 with at most one sign change.

284 A simple sufficient condition for the applicability of Result 3 is that the
 285 gain sequence is monotonic, that is, either increasing or decreasing. It is
 286 clear that an increasing gain sequence can have at most one sign change and
 287 that such a sign change occurs if and only if $d_0 < 0 < d_n$. In this case, the
 288 rest points of the replicator dynamics are characterized by Result 3.2.a. The
 289 other two possibilities for an increasing gain sequence, namely $d_n \leq 0$ and
 290 $d_0 \geq 0$, are covered by Result 3.1.a and Result 3.1.b, respectively. Similarly,
 291 for a decreasing gain sequence only three of the four scenarios described in
 292 Result 3 are possible, with a stable interior rest point occurring if and only
 293 if $d_0 > 0 > d_n$.

294 3.4.2. *Gain sequences with two sign changes*

295 If the gain sequence has two sign changes, its initial and final signs coin-
 296 cide. Suppose they are both negative. Then, by the preservation of initial
 297 and final signs (Property 1), the same is true for the initial and final signs of

298 $g(x)$. In particular, as indicated by Result 1, the rest point $x = 0$ is stable
 299 and the rest point $x = 1$ is unstable. Further, the first part of the variation
 300 diminishing property implies that the replicator dynamics has either (i) two
 301 distinct interior rest points (which correspond to simple roots in which $g(x)$
 302 crosses zero), (ii) one interior rest point (corresponding to a double root in
 303 which $g(x)$ touches, but does not cross zero), or (iii) no interior rest point.
 304 In the first of these cases $g(x)$ has two sign changes and the larger of the two
 305 interior rest points is stable. In the other two cases $g(x)$ has no sign change
 306 and, consequently, no stable interior rest point. Considering the maximal
 307 value of $g(x)$ on $[0, 1]$, which we denote by \bar{g} , provides a convenient way to
 308 describe which of these three cases arises. In particular, for $\bar{g} < 0$ there
 309 is no interior rest point, for $\bar{g} = 0$ there is exactly one interior rest point,
 310 and for $\bar{g} > 0$ there are two interior rest points. Analogous reasoning can
 311 be applied for the case in which the initial and final signs are both positive.
 312 These considerations are summarized in the following result.

313 **Result 4 (Gain sequences with two sign changes).** Let $\bar{g} = \max_{0 \leq x \leq 1} g(x)$
 314 and $\underline{g} = \min_{0 \leq x \leq 1} g(x)$. Then:

- 315 1. If $S(\mathbf{d}) = 2$ and $I(\mathbf{d}) = -$ the rest point $x = 0$ is stable and the rest
 316 point $x = 1$ is unstable. Further:
 - 317 (a) if $\bar{g} < 0$, the replicator dynamics has no interior rest points.
 - 318 (b) if $\bar{g} = 0$, then the replicator dynamics has one interior rest point
 319 \hat{x} which is unstable.
 - 320 (c) if $\bar{g} > 0$, the replicator dynamics has one unstable rest point x_L
 321 and one stable rest point x_R , satisfying $0 < x_L < x_R < 1$.

- 322 2. If $S(\mathbf{d}) = 2$ and $I(\mathbf{d}) = +$ the rest point $x = 0$ is unstable and the rest
 323 point $x = 1$ is stable. Further:
- 324 (a) If $\underline{g} > 0$, the replicator dynamics has no interior rest points.
- 325 (b) If $\underline{g} = 0$, the replicator dynamics has one interior rest point \hat{x}
 326 which is unstable.
- 327 (c) If $\underline{g} < 0$, the replicator dynamics has one stable rest point x_L and
 328 one unstable rest point x_R , satisfying $0 < x_L < x_R < 1$.

329 It is evident from the case distinctions appearing in Result 4 that for gain
 330 sequences with two sign changes, information beyond the one contained in
 331 the sign pattern of the gain sequence is required to determine the number
 332 of interior rest points. However, the additional information required takes a
 333 simple form (namely, the knowledge of the maximal, resp. minimal value of
 334 the gain function), which is amenable to further analysis.

335 **Remark 2.** If a gain sequence has more than two sign changes, Results 1
 336 and 2 still provide useful information about the possible range of dynamical
 337 scenarios, but determining which of these scenarios arises becomes much
 338 harder than in the case of at most two sign changes. To illustrate this,
 339 consider the case $S(\mathbf{d}) = 3$ and suppose $I(\mathbf{d}) = +$. We then have $F(\mathbf{d}) = -$,
 340 implying that both trivial rest points are unstable (Result 1). Furthermore,
 341 there are either one or two stable interior rest points (Result 2). In the second
 342 of these cases there must exist a single unstable interior rest point, in the first
 343 case there is either no unstable interior rest point or one unstable interior
 344 rest point which corresponds to a root of the gain function with multiplicity
 345 two (Property 2.1).

346 *3.4.3. Unimodal gain sequences*

347 Unimodality or anti-unimodality is a simple sufficient condition ensuring
 348 that a gain sequence has at most two sign changes. Furthermore, a complete
 349 classification of the possible dynamic scenarios is easily obtained. Here we
 350 demonstrate these claims for the unimodal case; the argument (and result)
 351 for the anti-unimodal case is analogous.

352 Our argument relies on the identity

$$g'(x) = n\mathcal{B}_{n-1}(x; \Delta\mathbf{d}), \quad (5)$$

353 which is a classical result in approximation theory, known as the derivative
 354 property of polynomials in Bernstein form (see e.g. Lorentz, 1986; DeVore and
 355 Lorentz, 1993; Farouki, 2012). By (5) the derivative $g'(x)$ is proportional to
 356 a Bernstein polynomial with coefficients $\Delta\mathbf{d}$. We may thus apply Properties
 357 1 and 2 to the relationship between the sign pattern of $\Delta\mathbf{d}$ and the roots and
 358 sign pattern of $g'(x)$. Recalling that for a unimodal gain sequence $\Delta\mathbf{d}$ has
 359 a single sign change from positive to negative, it follows that unimodality
 360 of the gain sequence implies unimodality of the gain function. Moreover,
 361 applying the first part of the variation diminishing property, there exists a
 362 unique $0 < \hat{x} < 1$ satisfying the first order condition $g'(\hat{x}) = 0$. Unimodality
 363 of $g(x)$ implies that \hat{x} is the unique solution to the problem $\max_{0 \leq x \leq 1} g(x)$
 364 appearing in the statement of Result 4. In particular, we have $\bar{g} = g(\hat{x})$.

365 It is clear that a unimodal gain function can have at most one sign change
 366 in its increasing part (which then must be from negative to positive) and at
 367 most one sign change in its decreasing part (which then must be from positive
 368 to negative). Moreover, a sign change in the increasing part occurs if and
 369 only if $g(0) < 0 < g(\hat{x})$ and a sign change in the decreasing part occurs if and

370 only if $g(1) < 0 < g(\hat{x})$. Combining these observations yields the following
 371 result, refining Results 3 and 4 for the unimodal case.

372 **Result 5 (Unimodal gain sequences).** If the gain sequence is unimodal,
 373 there exists a unique $0 < \hat{x} < 1$ solving the equation $g'(\hat{x}) = 0$. Moreover:

- 374 1. If $g(\hat{x}) < 0$, then the replicator dynamics has no interior rest point.
 375 The rest point $x = 0$ is stable and the rest point $x = 1$ is unstable.
- 376 2. If $g(\hat{x}) = 0$, then \hat{x} is the unique interior rest point of the replicator
 377 dynamics. The rest point $x = 0$ is stable and the rest points \hat{x} and
 378 $x = 1$ are unstable.
- 379 3. If $g(\hat{x}) > 0$ holds, then one of the following four cases applies:
 - 380 (a) If $\min\{d_0, d_n\} \geq 0$, then the replicator dynamics has no interior
 381 rest point. The rest point $x = 0$ is unstable and the rest point
 382 $x = 1$ is stable.
 - 383 (b) If $\max\{d_0, d_n\} < 0$, then the replicator dynamics has two interior
 384 rest points satisfying $x_L < \hat{x} < x_R$. The rest points $x = 0$ and x_R
 385 are stable, whereas the rest points x_L and $x = 1$ are unstable.
 - 386 (c) If $d_0 < 0$ and $d_n \geq 0$, then the replicator dynamics has a unique
 387 interior rest point $x^* < \hat{x}$. The rest points $x = 0$ and $x = 1$ are
 388 stable, whereas the rest point x^* is unstable.
 - 389 (d) If $d_0 \geq 0$ and $d_n < 0$, then the replicator dynamics has a unique
 390 interior rest point $x^* > \hat{x}$. The rest point x^* is stable, whereas the
 391 rest points $x = 0$ and $x = 1$ are unstable.

392 **Remark 3.** Using the derivative property of polynomials in Bernstein form,
 393 it can be shown that all the properties of gain sequences mentioned at the

394 end of Section 3.1 are inherited by the gain function (e.g., if the gain sequence
 395 is increasing, so is the gain function). The argument for the preservation of
 396 anti-unimodality is analogous to the one we have given for the preservation of
 397 unimodality. The other results are well known properties of Bernstein poly-
 398 nomials, namely preservation of monotonicity, and preservation of convexity
 399 (see Lorentz, 1986; Farouki, 2012). Seemingly unaware of these properties,
 400 Motro (1991) proves preservation of monotonicity and Bach et al. (2006)
 401 prove preservation of concavity (which is equivalent to preservation of con-
 402 vexity).

403 4. Public goods games

404 In this section, we apply Results 3 to 5 to two classes of public goods
 405 games, subsuming many of the models encountered in the literature of the
 406 evolution of cooperation and collective action.

407 4.1. Gain sequences for public goods games

408 In a *public goods game*, playing A means to cooperate (i.e. to contribute to
 409 the creation or maintenance of a public good) and playing B means to defect
 410 (i.e. to free ride on the contributions of others). Contributing entails a cost
 411 $c_k \geq 0$ to the focal cooperator, where k is the number of other cooperators.
 412 Defectors bear no cost. All players obtain a benefit $r_j \geq 0$ from the public
 413 good, where j is the total number of cooperators in the group. Note that
 414 for a focal cooperator $j = k + 1$, while for a focal defector $j = k$. With
 415 these assumptions, the payoff sequences for a public goods game can thus be
 416 written as

$$a_k = r_{k+1} - c_k, \quad k = 0, 1, \dots, n$$

417 and

$$b_k = r_k, \quad k = 0, 1, \dots, n$$

418 so that the gain sequence is given by

$$d_k = \Delta r_k - c_k, \quad k = 0, 1, \dots, n. \quad (6)$$

419 As it is generally considered in the literature, we assume the benefit se-
 420 quence $\mathbf{r} = (r_0, \dots, r_{n+1})$ is increasing and the cost sequence $\mathbf{c} = (c_0, \dots, c_n)$
 421 is not equal to zero.

422 If no further assumptions are imposed on the cost and benefit sequence, it
 423 is clear from (6) that any \mathbf{d} can arise as the gain sequence of a public goods
 424 game. Consequently, to obtain insights into the evolutionary dynamics of
 425 public goods games going beyond the ones summarized in Results 1 and 2,
 426 additional assumptions on the benefit or the cost sequence are required. In
 427 this light, it is not surprising that public goods games usually studied in the
 428 biological literature fall into one of the two classes that we discuss in the
 429 following subsections.

430 4.2. *Threshold games*

431 If there exists an integer m with $1 \leq m \leq n + 1$ and a constant $r > 0$
 432 such that the benefit sequence satisfies $r_j = 0$ if $j < m$ and $r_j = r$ if $j \geq m$,
 433 we say that a public goods game is a *threshold game*. This class of games
 434 describes situations in which the public good is a “step good” in the sense
 435 of Hardin (1982, p. 55): at least m cooperators are required to provide a
 436 public good for all group members, but the number of cooperators beyond the
 437 threshold m does not increase the benefit received by the players. Examples
 438 of such threshold games abound in the theoretical literature of the social

439 sciences (Hardin, 1982; Taylor and Ward, 1982; Diekmann, 1985; Sugden,
 440 1986; Weesie and Franzen, 1998; Höffler, 1999; Herold, 2012) and evolutionary
 441 biology (Dugatkin, 1990; Bach et al., 2006; Zheng et al., 2007; Archetti, 2009;
 442 Souza et al., 2009), and are sometimes referred to as volunteer's dilemmas or
 443 multi-player snowdrift games.

444 For threshold games (6) reduces to

$$d_k = \begin{cases} -c_k & \text{if } k < m - 1 \\ r - c_{m-1} & \text{if } k = m - 1 \\ -c_k & \text{if } k > m - 1 \end{cases} . \quad (7)$$

445 It is obvious that the gain sequence \mathbf{d} has no sign change when $r \leq c_{m-1}$ and
 446 that in this case defection is a dominant strategy. As illustrated in Fig. 1
 447 and discussed below, in the other cases the sign pattern of the gain sequence
 448 depends on the location of the threshold m .

449 [Figure 1 about here.]

450 4.2.1. Threshold $m = 1$

451 Threshold games with $m = 1$ represent situations in which only one co-
 452 operator is required for the provision of the public good. Such games have
 453 been considered by Dugatkin (1990), Weesie and Franzen (1998), Zheng et al.
 454 (2007), and Souza et al. (2009) for the particular case of a cost sequence satis-
 455 fying $c_k = c/(k + 1)$ for some constant $c > 0$, so that the cost to cooperators
 456 is inversely proportional to the total number of cooperators in the group.
 457 These authors have shown by algebraic manipulations or numerical simula-
 458 tions that for such games the replicator dynamics has at most one interior

459 stable rest point. Archetti (2009) shows the same result for a cost sequence
 460 satisfying $c_k = c$ for some constant $c > 0$.

461 Considering the sign pattern of the gains from switching not only recovers
 462 this result in a simpler way, but also extends it to any strictly positive cost
 463 sequence \mathbf{c} . If $r > c_0$, the gain sequence given in (7) has exactly one sign
 464 change and $I(\mathbf{d}) = +$, so that Result 3.2.b establishes the existence of a
 465 single interior stable rest point $0 < x^* < 1$ and the instability of the trivial
 466 rest points (see Fig. 1.a). If $r \leq c_0$, Result 3.1.a applies. Hence, there is no
 467 interior rest point and $x = 0$ is the unique stable rest point.

468 4.2.2. Threshold $m = n + 1$

469 Recalling that $N = n + 1$ is group size, threshold games with $m = n + 1$
 470 represent situations in which the cooperation of all group members is required
 471 to produce the public good. For the case $m = n + 1 = 2$ and a cost sequence
 472 satisfying $0 < c_0 = c_1 < r$, Souza et al. (2009) observe that such a threshold
 473 game corresponds to a two-player stag hunt game (Skyrms, 2004) in which
 474 both trivial rest points are stable and there is a unique, unstable interior
 475 rest point. It is easy to see that this result holds more generally. Indeed,
 476 provided that the cost sequence is strictly positive and satisfies $r > c_n$, the
 477 gain sequence given in (7) is characterized by $S(\mathbf{d}) = 1$ and $I(\mathbf{d}) = -$. Then,
 478 by Result 3.2.a, it follows that the qualitative dynamics of the two-player stag
 479 hunt are recovered for every threshold game with $m = n + 1$ (see Fig. 1.b).
 480 The case $r \leq c_n$ is covered by Result 3.1.a.

481 *4.2.3. Threshold $1 < m < n + 1$*

482 Souza et al. (2009) studied a threshold game with $1 < m < n + 1$ for a
483 cost sequence of the form

$$c_k = \begin{cases} c/m & \text{if } k < m - 1 \\ c/(k + 1) & \text{if } k \geq m - 1 \end{cases} \quad (8)$$

484 for some constant $c > 0$. Their main theoretical result (Souza et al., 2009,
485 Theorem 1) uses an ingenious but rather involved argument to demonstrate
486 that in this example there exists $\bar{c} > 0$ and $0 < \bar{x} < 1$ such that (i) if $c < \bar{c}$,
487 the replicator dynamics has two interior rest points $x_L < \bar{x} < x_R$ where x_L is
488 unstable and x_R is stable (see Fig. 1.c), (ii) if $c = \bar{c}$, the replicator dynamics
489 has a unique rest point \bar{x} (which is unstable), and (iii) if $c > \bar{c}$, the replicator
490 dynamics has no interior rest point (see Fig. 1.d).²

491 In Appendix B we prove that the same result holds for any cost sequence
492 of the form $c_k = c \cdot \gamma_k$, where the strictly positive, but otherwise arbitrary,
493 sequence γ describes the shape of the cost sequence and, as in the example
494 considered by Souza et al. (2009), c shifts the level of the cost sequence.
495 Our result follows, in essence, from two observations. The first is that for
496 every threshold game with $1 < m < n + 1$ and strictly positive cost sequence
497 satisfying $0 < c_{m-1} < r$ the gain sequence has two sign changes and a negative
498 initial sign, so that the rest points of the replicator dynamics are described
499 by Result 4.1. The second observation is that the maximal value of the gain
500 function \bar{g} is strictly decreasing in the cost parameter c .

²Souza et al. (2009) express their results in terms of the cost-benefit ratio c/r . The difference is of no importance as time can always be rescaled to ensure $r = 1$.

501 Threshold games with $1 < m < n + 1$ have also been considered by Bach
 502 et al. (2006), Archetti (2009), and Archetti and Scheuring (2011). These
 503 authors assume a cost sequence satisfying $c_k = c$ for some constant $c >$
 504 0 , implying that these games fall in the class of constant cost games with
 505 sigmoid benefit functions that we discuss in Section 4.3.3.

506 4.2.4. Further threshold games

507 In economics, Höfler (1999) and Herold (2012) have studied evolutionary
 508 dynamics of threshold games which differ from the biological threshold games
 509 considered above in that cooperators pay a cost only if the threshold for the
 510 successful provision of the public good is reached. In such cases the gain
 511 sequence has the form

$$d_k = \begin{cases} 0 & \text{if } k < m - 1 \\ r - c_{m-1} & \text{if } k = m - 1 \\ -c_k & \text{if } k > m - 1 \end{cases} \quad (9)$$

512 and thus possesses at most one sign change (see Fig. 2). For $r > c_{m-1}$ and
 513 $1 \leq m < n + 1$, this gain sequence satisfies $I(\mathbf{d}) = +$ and $S(\mathbf{d}) = 1$. Applying
 514 Result 3.2.b then yields a simple direct proof of the main result obtained by
 515 Höfler (1999, Proposition 1) and Herold (2012, Proposition 1) for this class
 516 of games, namely that there exists a unique stable interior rest point.³

517 [Figure 2 about here.]

³Proposition 2 in Höfler (1999), which considers the case $m = n + 1$, is implied by our Result 3.1.b. Herold also considers the case in which cooperators only pay a cost if the threshold is *not* reached. His main result for this case (Herold, 2012, Proposition 2) is implied by our Result 3.2.a.

518 *4.3. Constant cost games*

519 If there exists a constant $c > 0$ such that $c_k = c$ holds for $k = 0, \dots, n$
 520 we say that a public goods game is a *constant cost game*. Such games have
 521 been studied, among others, by Motro (1991), Szathmáry (1993), Bach et al.
 522 (2006), Hauert et al. (2006), Pacheco et al. (2009), and Archetti and Scheur-
 523 ing (2011).

524 In the case of a constant cost game, equation (6) reduces to

$$d_k = \Delta r_k - c, \quad k = 0, 1, \dots, n. \quad (10)$$

525 It is then immediate that the gain sequence has no sign change (and hence
 526 no interior rest point) if $c \geq \max_{k=0, \dots, n} \Delta r_k$ or $\min_{k=0, \dots, n} \Delta r_k \geq c$ holds. It
 527 follows from Result 3.1 that in the former case $x = 0$ and in the latter case
 528 $x = 1$ is the unique stable rest point. In all other cases, that is whenever the
 529 inequality

$$\min_{k=0, \dots, n} \Delta r_k < c < \max_{k=0, \dots, n} \Delta r_k \quad (11)$$

530 holds, the gain sequence has at least one sign change.

531 In the following, we consider three different kinds of constant cost games,
 532 arising from three different assumptions on the shape of the benefit sequence:
 533 linear benefits (Section 4.3.1), convex or concave benefits (Section 4.3.2) and
 534 sigmoid benefits (Section 4.3.3). See Fig. 3 for a graphical illustration of
 535 these different constant cost games.

536 [Figure 3 about here.]

537 *4.3.1. Linear benefits*

538 The familiar linear public goods game is a constant cost game in which
 539 the benefit sequence is given by $r_j = jr/(n+1)$ (Sigmund, 2010). The

540 interpretation is that $r > 0$ is the amount of the public good produced by
 541 each cooperator and that this amount is split evenly among the $N = n + 1$
 542 members of the group. For such a game, we have $\Delta r_k = r/(n + 1)$, so that
 543 the gain sequence is $d_k = r/(n + 1) - c$, which is a constant independent of k .
 544 Hence \mathbf{d} has no sign change. Making the standard assumption $r < (n + 1)c$,
 545 we have $I(\mathbf{d}) = -$, so that there are no interior rest points and $x = 0$ is
 546 the unique stable rest point (see Fig. 3.a). This conclusion is, of course,
 547 well-known.

548 4.3.2. Convex or concave benefits

549 Convexity of the benefit sequence ($\Delta^2 \mathbf{r} \geq \mathbf{0}$) indicates that the incremen-
 550 tal benefit Δr_k of a further contributor is increasing in the number of other
 551 contributors k that are already present in the group. Using (10) to obtain

$$\Delta d_k = \Delta^2 r_k, \quad k = 0, 1, \dots, n - 1, \quad (12)$$

552 it is apparent that the gain sequence \mathbf{d} is increasing. As discussed in Sec-
 553 tion 3.4.1 it follows that (11) reduces to $\Delta r_0 < c < \Delta r_n$. Furthermore, if
 554 these inequalities hold, Result 3.2.a implies that there is a unique interior
 555 rest point which is unstable, whereas both trivial rest points are stable (see
 556 Fig. 3.b). Similarly, when the benefit sequence is concave ($\Delta^2 \mathbf{r} \leq \mathbf{0}$), (11)
 557 reduces to $\Delta r_n < c < \Delta r_0$ and if these inequalities hold, Result 3.2.b implies
 558 there is a unique interior rest point which is stable, whereas both trivial rest
 559 points are unstable (see Fig. 3.c).

560 The argument we have just given recovers the main results from Motro
 561 (1991). A simple illustration of a constant cost game with convex or constant
 562 benefits is provided by the model of synergy and discounting considered in

563 Hauert et al. (2006, Section 2.1). These authors consider a constant cost
 564 game with benefit function

$$r_j = \frac{r}{n+1} (1 + w + \dots w^{j-1}), \quad (13)$$

565 where $r > 0$ and $w > 0$ are parameters. For this specification we have
 566 $\Delta r_k = rw^k/(n+1)$. For $w > 1$ this benefit sequence is convex, whereas for
 567 $w < 1$ it is concave. The case $w = 1$ is the linear public goods game. We
 568 observe that the classification obtained in Section 2.2 of Hauert et al. (2006),
 569 corresponds to the one obtained from a straightforward application of our
 570 Result 3.

571 4.3.3. Sigmoid benefits

572 A benefit sequence is sigmoid (or S-shaped) when $\Delta^2 \mathbf{r}$ has exactly one sign
 573 change from $+$ to $-$, i.e. the benefit sequence is first convex, then concave.
 574 Examples of sigmoid benefit sequences are the threshold benefit sequences
 575 with $1 < m < n+1$ considered in Section 4.2.3, the “benefit function with a
 576 hump” proposed in Szathmary (1993), and the threshold-linear and logistic
 577 benefit sequences studied respectively by Pacheco et al. (2009) and Archetti
 578 and Scheuring (2011).

579 In this case it is immediate from (12) that the gain sequence of a constant
 580 cost game with sigmoid benefits is unimodal. Consequently, the characteri-
 581 zation of the different types of dynamics that can arise in such games involves
 582 nothing more than inserting the values $d_k = \Delta r_k - c$ into our Result 5 (see
 583 Fig. 3.d for a particular example). The results of this exercise have been
 584 published by Archetti (2013).⁴

⁴Archetti (2013) ignores most of the cases in which a weak inequality occurs in Result 5

585 Sigmoid benefit sequences generalize the benefit sequences considered in
 586 Bach et al. (2006, Proposition 7), who not only assume that $\Delta^2\mathbf{r}$ has a single
 587 sign change from $+$ to $-$, but, in addition, require $\Delta^2\mathbf{r}$ to be decreasing.
 588 Using these assumptions, Bach et al. (2006) establish the existence of a
 589 $c^* > \max\{\Delta r_0, \Delta r_n\}$ such that for $c < c^*$ the replicator dynamics has two
 590 interior rest points (the larger of which is stable), whereas for $c = c^*$ there
 591 is a unique (unstable) interior rest point and for $c > c^*$ there is none. As
 592 the gain sequence (and hence the gain function and \bar{g}) for constant cost
 593 games is linearly decreasing in c , it is immediate from Result 5 that the same
 594 conclusion obtains for all sigmoid benefit sequences.

595 5. Other multi-player games

596 Up to this point our examples have considered public goods games. Here
 597 we consider two examples of other multi-player games, illustrating how fo-
 598 cusing on the shape of the gain sequence obviates the need for a more in-
 599 volved analysis. Of course, further examples could be analyzed along similar
 600 lines. For instance, it is straightforward to show that in the “shared reward
 601 dilemma” considered by Cuesta et al. (2008) the gain sequence has at most
 602 two sign changes, so that we can recover their case distinctions by applying
 603 our results.

and neglects to impose the proper sign change condition required for unimodality, but these shortcomings are easily fixed.

604 *5.1. Repeated N -person prisoner's dilemma*

605 Joshi (1987), Boyd and Richerson (1988) and van Segbroeck et al. (2012)
 606 considered a repeated N -person prisoner's dilemma with two possible strate-
 607 gies. Reciprocators (A-strategists) contribute to the public good in the first
 608 round and then contribute in each subsequent round if at least m individuals
 609 (including the focal individual) contributed in the previous move. Defectors
 610 (B-strategists) never contribute to the public good. Payoffs in each round
 611 depend on the number of contributors as in the linear public goods game
 612 considered in Section 4.3.1.

613 The gain sequence for this model is easily derived by considering the first
 614 round and the subsequent rounds separately. In the first round, the gain if
 615 switching from B to A is $r/(n+1) - c < 0$. In each subsequent round, the gain
 616 from switching is zero if $k < m - 1$ (because all players defect), $r/(n+1) - c$
 617 if $k > m - 1$ (because the other reciprocators cooperate no matter whether
 618 the focal individual contributes or not), and $mr/(n+1) - c$ if $k = m - 1$
 619 (because in this case the contribution of the focal individual in the first round
 620 is pivotal in determining the subsequent behavior of reciprocators). Setting

$$\tilde{c} = c - r/(n+1) > 0,$$

621 and

$$\tilde{r} = (m-1)r/(n+1),$$

622 the gain sequence can be written as

$$d_k = \begin{cases} -\tilde{c} & \text{if } k < m - 1 \\ T\tilde{r} - (T+1)\tilde{c} & \text{if } k = m - 1 \\ -(T+1)\tilde{c} & \text{if } k > m - 1 \end{cases}, \quad (14)$$

623 where $T > 0$ denotes the expected number of rounds after the first one.
 624 From (7) and (14) it is apparent that the model is equivalent to a threshold
 625 game with (i) the benefit $T\tilde{r}$ arising if and only if at least m reciprocators
 626 are present and (ii) costs given by $c_k = \tilde{c}$ if $k < m - 1$ and $c_k = (T + 1)\tilde{c}$
 627 otherwise. In particular, the results for the cases $m = 1$ and $m = n + 1$ are
 628 identical to the ones discussed in Sections 4.2.1 and 4.2.2. Moreover, when
 629 $T\tilde{r} - (T + 1)\tilde{c}$ is negative, it is immediate that the gain sequence is negative
 630 and Result 3.1.a applies.

631 In the remaining case, satisfying $1 < m < n + 1$ and $T\tilde{r} - (T + 1)\tilde{c} > 0$,
 632 it follows from (14) that the only non-zero elements of $\Delta\mathbf{d}$ are $\Delta d_{m-2} > 0$
 633 and $\Delta d_{m-1} < 0$. Consequently, the gain sequence is unimodal and Result 5
 634 applies with $\max\{d_0, d_n\} < 0$ to characterize the three different possible
 635 dynamical regimes. Which of these regimes arises depends on the value of
 636 $\bar{g} = g(\hat{x})$ (see Fig. 4 for an example of the case $\bar{g} > 0$). As in all applications
 637 of Results 4 and 5, a key question is whether this value can be linked to the
 638 parameters of the model.

639 [Figure 4 about here.]

640 For the parameter T this question can be answered by using the linearity
 641 of the Bernstein operator \mathcal{B}_n to write the gain function as

$$g(x) = Th(x) - \tilde{c}, \quad (15)$$

642 where $h(x) = \mathcal{B}_n(x, \mathbf{e})$ and the sequence \mathbf{e} is given by

$$e_k = \begin{cases} 0 & \text{if } k < m - 1 \\ \tilde{r} - \tilde{c} & \text{if } k = m - 1 \\ -\tilde{c} & \text{if } k > m - 1 \end{cases} .$$

643 It follows from (15) that the critical value \hat{x} satisfying the first order condition
 644 $g'(\hat{x}) = 0$ is independent of T . Further, because $I(\mathbf{e}) = +$, it follows from the
 645 preservation of initial signs that $h(\hat{x}) > 0$ holds. This in turn implies from
 646 (15) that $g(\hat{x})$ is strictly increasing in T and that the equation $\hat{T} = \tilde{c}/h(\hat{x})$
 647 identifies the critical value of T at which $g(\hat{x}) = 0$ holds. Hence, we obtain
 648 the same conclusions as van Segbroeck et al. (2012) by an application of
 649 Result 5. Namely, (i) for $T < \hat{T}$ there is no interior rest point, (ii) for $T = \hat{T}$
 650 the replicator dynamics has a single, unstable interior rest point, and (iii) for
 651 $T > \hat{T}$ two interior rest points emerge.

652 *5.2. Constant cost game with different benefit sequences for cooperators and*
 653 *defectors*

654 [Figure 5 about here.]

Hauert et al. (2006, Section 2.3.2) consider an interesting extension of constant cost games by allowing for the possibility that cooperators and defectors might obtain different benefits, say r_j^A and r_j^B , when there are j cooperators in the group (see Fig. 5). The counterpart to (12) is then $\Delta d_k = \Delta r_{k+1}^A - \Delta r_k^B$. For the particular choice of benefit sequences in Hauert et al. (2006), given by (13) for r_j^A and

$$r_j^B = \frac{r}{n+1} (1 + v^1 + \dots + v^{j-1}),$$

655 this reduces to

$$\Delta d_k = \frac{r}{n+1} (w^{k+1} - v^k), \quad (16)$$

656 where $r > 0$, $v > 0$ and $w > 0$ are parameters and $N = n + 1$ is group size.

Hauert et al. (2006) state that “only $v = w$ allows for an analytical solution [...] but in general there are [...] up to $N - 1$ equilibria [rest points] in $(0, 1)$.” Here we refine this statement and show that, as conjectured by Cuesta et al. (2007), the maximum number of interior rest points is two independently of group size. To do so, we observe that $\Delta d_k > 0$ holds if and only if

$$w > \left(\frac{v}{w}\right)^k.$$

657 Since the right side of this inequality is monotonic in k , equation (16) implies
658 the following, exhaustive case distinction:

- 659 1. if $w \geq 1$ and $w^n \geq v^{n-1}$ holds, then the gain sequence is increasing and
660 there is at most one interior rest point (see Fig. 5.a).
- 661 2. if $w \leq 1$ and $w^n \leq v^{n-1}$ holds, then the gain sequence is decreasing
662 and there is at most one interior rest point (see Fig. 5.b).
- 663 3. if $w > 1$ and $w^n < v^{n-1}$ holds, then the gain sequence is unimodal and
664 there are at most two interior rest points (see Fig. 5.c).
- 665 4. if $w < 1$ and $w^n > v^{n-1}$ holds, then the gain sequence is anti-unimodal
666 and there are at most two interior rest points (see Fig. 5.d).

667 6. Discussion

668 Bernstein polynomials were first proposed more than a century ago by Bern-
669 stein (1912) in order to provide a constructive proof of Weierstrass’s approxi-
670 mation theorem (DeVore and Lorentz, 1993). More recently, because of their
671 many shape-preserving properties, polynomials in Bernstein form have also
672 proven extremely useful in the field of computer aided geometric design (Ya-
673 maguchi and Yamaguchi, 1988; Farin and Hoschek, 2002; Prautzsch et al.,

674 2002). Here we have made the case for utilizing the shape-preserving prop-
675 erties of Bernstein polynomials in the analysis of multi-player matrix games.
676 In particular, we have used these properties to show how key insights into the
677 evolutionary dynamics of multi-player matrix games can be obtained from
678 studying the sign pattern of the gains from switching.

679 The properties of Bernstein polynomials we have used in this paper are
680 certainly not the only ones of relevance for the theoretical analysis of col-
681 lective action problems. For instance, both the effects of changes in the
682 group size (studied previously in Motro, 1991) and the group size distribu-
683 tion (studied previously in Peña, 2012) on the evolutionary dynamics can be
684 analyzed by making use of the theory of polynomials in Bernstein form. Our
685 methods can also be extended to structured populations and used to analyze
686 multi-player matrix games played between relatives.

687 **Acknowledgements**

688 We would like to thank Chaitanya S. Gokhale, Arne Traulsen and one
689 anonymous reviewer for useful comments on previous versions of the manuscript.
690 This work was supported by Swiss NSF grants PBLAP3-145860 (JP) and
691 PP00P3-123344 (LL).

692 **Appendix A. Proof of Result 1**

693 We show the result $I(g) = I(\mathbf{d})$; the argument that the final signs coincide
694 is analogous. Using the derivative property of polynomials in Bernstein form
695 (cf. equation (5)) recursively, for $0 \leq m \leq n$ the m -th derivative of the gain

696 function can be written as (Farouki, 2012)

$$g^{(m)}(x) = n(n-1)\dots(n-m+1)\mathcal{B}_{n-m}(x; \Delta^m \mathbf{d}), \quad (\text{A.1})$$

697 where (with the obvious iterative definition) $\Delta^m \mathbf{d}$ is the m -th forward differ-
698 ence of the sequence \mathbf{d} . Evaluating (A.1) at $x = 0$ we obtain

$$g^{(m)}(0) = n(n-1)\dots(n-m+1)\Delta^m d_0. \quad (\text{A.2})$$

699 Now, let ℓ be the lowest index k such that $d_\ell \neq 0$. Then $\Delta^m d_0 = 0$ holds
700 for all $m < \ell$ and $\Delta^\ell d_0 = d_\ell$. Equation (A.2) then implies that $g^{(m)}(0) = 0$
701 for all $m < \ell$ and that the sign of $g^{(\ell)}(0)$ coincides with the sign of d_ℓ which,
702 by definition, is the initial sign of \mathbf{d} . A standard Taylor-series argument as
703 given in Bach et al. (2006, Proof of Proposition 4) demonstrates that the
704 initial sign of g coincides with the sign of d_ℓ , finishing the proof.

705 **Appendix B. Proof of the generalization of Theorem 1 from Souza** 706 **et al. (2009)**

707 For any $c \geq 0$ let

$$g(x, c) = \sum_{k=0}^n \binom{n}{k} x^k (1-x)^{n-k} d_k(c), \quad (\text{B.1})$$

708 where

$$d_k(c) = \begin{cases} -c\gamma_k & \text{if } k < m-1 \\ r - c\gamma_{m-1} & \text{if } k = m-1 \\ -c\gamma_k & \text{if } k > m-1 \end{cases} \quad (\text{B.2})$$

709 and $\boldsymbol{\gamma} = (\gamma_0, \dots, \gamma_n)$ is a given, strictly positive sequence. Let $\bar{g}(c) =$
710 $\max_{0 \leq x \leq 1} g(x, c)$ denote the corresponding maximal value of the gain func-
711 tion.

712 For $0 < c < r/\gamma_{m-1}$ the gain sequence given in (B.2) satisfies $I(\mathbf{d}(c)) = -$
 713 and $S(\mathbf{d}(c)) = 2$, so that the rest points of the replicator dynamics are
 714 described by Result 4.1.

715 From (B.1) and (B.2) the function $g(x, c)$ is continuous. From the max-
 716 imum theorem (Sundaram, 1996, Theorem 9.14) this ensures continuity of
 717 $\bar{g}(c)$. Because all the Bernstein coefficients $d_k(c)$ are strictly decreasing in c ,
 718 all the summands appearing in (B.1) are strictly decreasing in c , implying
 719 that $g(x, c)$ is strictly decreasing in c . This monotonicity property obviously
 720 carries over to $\bar{g}(c)$.

721 Consider the Bernstein coefficients as given in (B.2). If $c = 0$, the only
 722 non-zero coefficient is $d_{m-1}(0) = r > 0$. It is then immediate from (B.1) that
 723 $g(x, 0) > 0$ holds for all $0 < x < 1$, ensuring $\bar{g}(0) > 0$. If $c = r/\gamma_{m-1}$, we have
 724 $d_k(c) \leq 0$ with strict inequality holding in all cases but $k = m - 1$. From
 725 (B.1) this implies $g(x, r/\gamma_{m-1}) < 0$ for all $0 \leq x \leq 1$, ensuring $\bar{g}(r/\gamma_{m-1}) < 0$.

726 Because $\bar{g}(0) > 0$ and $\bar{g}(r/\gamma_{m-1}) < 0$ hold and $\bar{g}(c)$ is continuous the in-
 727 termediate value theorem implies that there exists $0 < \bar{c} < r/\gamma_{m-1}$ satisfying
 728 $\bar{g}(\bar{c}) = 0$. By monotonicity of $\bar{g}(c)$ it follows that $\bar{g}(c) < 0$ holds for $c > \bar{c}$
 729 and $\bar{g}(c) > 0$ holds for $c < \bar{c}$. The generalized version of Theorem 1 in Souza
 730 et al. (2009) then follows from our Result 4.1 – except that it remains to
 731 establish the existence of $0 < \bar{x} < 1$ such that the interior rest points satisfy
 732 $x_L < \bar{x} < x_R$ for all $0 < c < \bar{c}$. Towards this end let \bar{x} be a solution to
 733 the problem $\max_{0 \leq x \leq 1} g(x, \bar{c})$. As $g(0, \bar{c}) < 0$ and $g(1, \bar{c}) < 0$ holds, we have
 734 $0 < \bar{x} < 1$. As $g(x, c)$ is strictly decreasing in c , we have $g(\bar{x}, c) > 0$ for all
 735 $0 < c < \bar{c}$. In conjunction with $g(0, c) < 0$ and $g(1, c) < 0$ this implies that
 736 $g(x, c)$ has at least one root in the interval $(0, \bar{x})$ and at least one root in the

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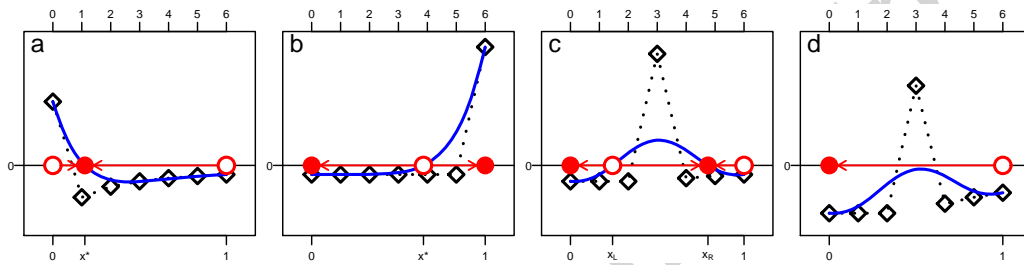


Figure 1: Gain sequence \mathbf{d} (squares, dotted line; top axis), and corresponding gain function $g(x)$ (solid line; bottom axis) and phase portrait (circles, arrows) for threshold games given by (7) and (8) with $N = 7$, $r = 2$, $c = 1$, and (a) $m = 1$ (see section 4.2.1), (b) $m = N = n + 1$ (see section 4.2.2), or (c) $m = 4$ (see section 4.2.3). Panel d illustrates the same game as in panel c, but with $c = 3$ instead of $c = 1$.

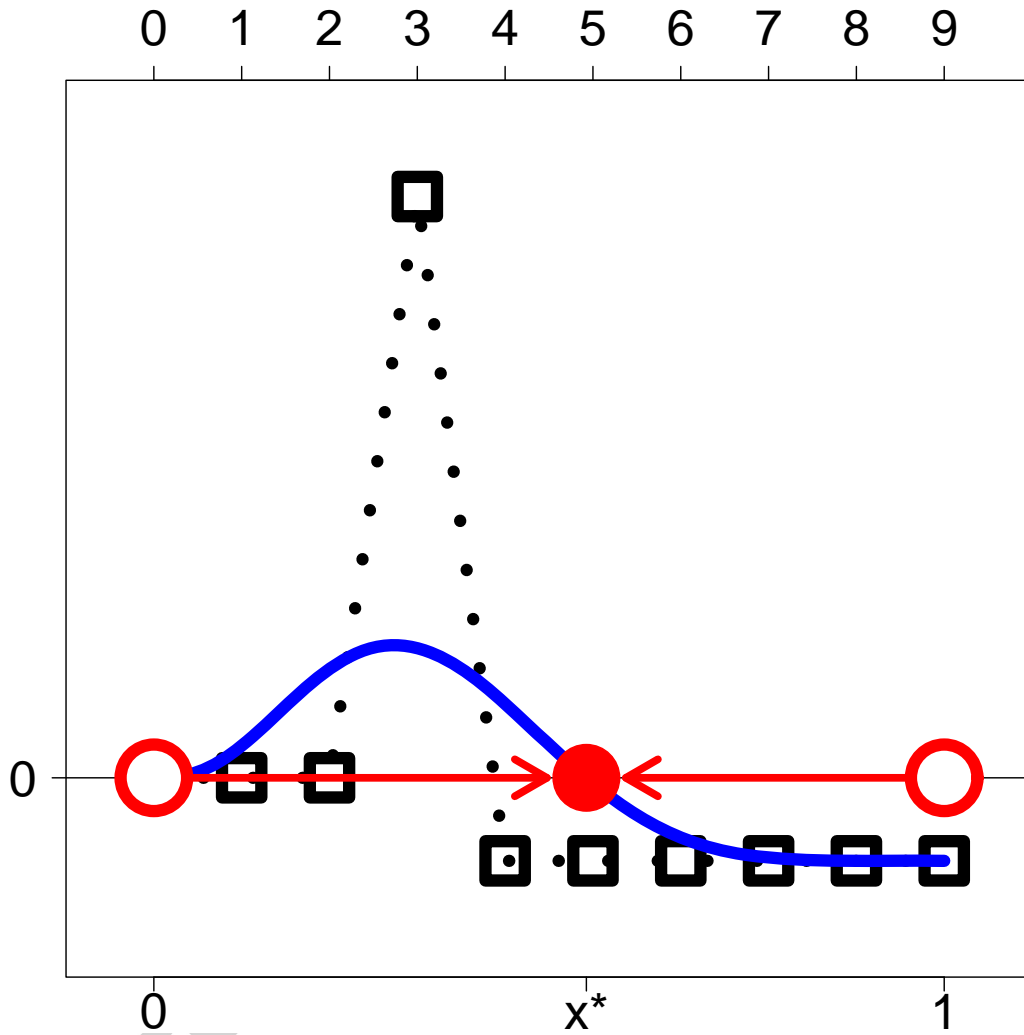


Figure 2: Gain sequence \mathbf{d} (squares, dotted line; top axis), and corresponding gain function $g(x)$ (solid line; bottom axis) and phase portrait (circles, arrows) for the threshold game given by (9) with $N = 10$, $r = 2$, $m = 4$, and $c_k = 1/4$ for all $k \geq 3$.

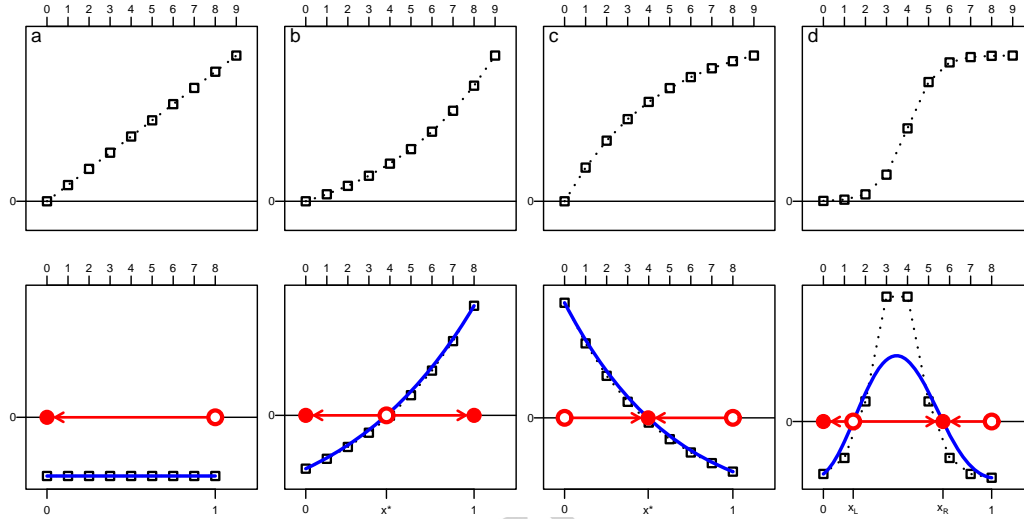


Figure 3: Examples of constant cost games with $N = n + 1 = 9$ and $c = 1/2$ for different benefit sequences. The first row shows the benefit sequence r_j ; the second row shows the gain sequence \mathbf{d} (squares, dotted line; top axis), and corresponding gain function $g(x)$ (solid line; bottom axis) and phase portrait (circles, arrows). (a) Linear benefits (see Section 4.3.1) with $r = 5$ and $c = 1$. (b) Convex benefits (see Section 4.3.2) as given by (13) with $r = 5$ and $w = 1.2$. (c) Concave benefits (see Section 4.3.2) as given by (13) with $r = 20$ and $w = 0.8$. (d) Sigmoid benefits (see Section 4.3.3) as studied by Archetti and Scheuring (2011) with $r_j = r/[1 + \exp(-s(j - m))]$, $r = 20$, $m = 4$, and $s = 1.5$.

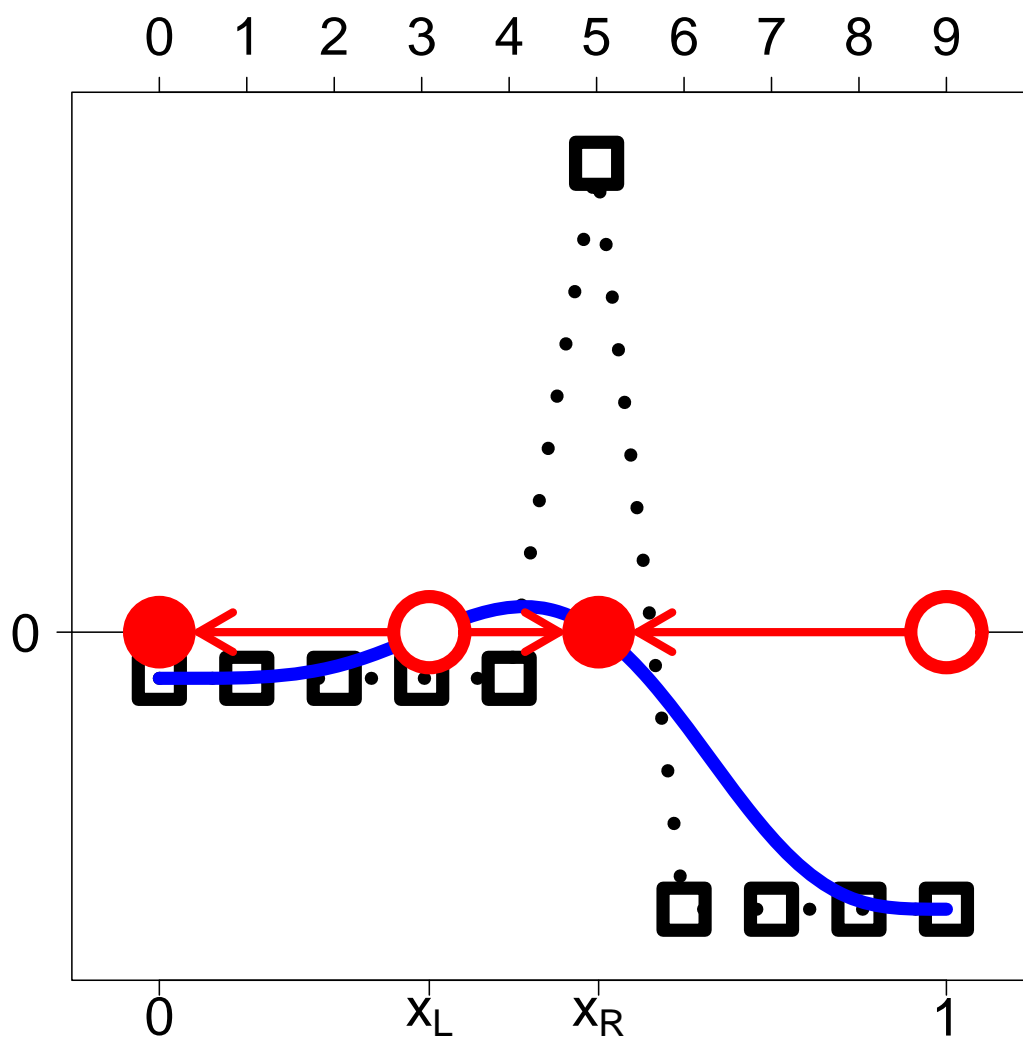


Figure 4: Gain sequence \mathbf{d} (squares, dotted line; top axis), and corresponding gain function $g(x)$ (solid line; bottom axis) and phase portrait (circles, arrows) for the repeated N -person prisoner's dilemma given by (14) with $N = 10$, $r = 7$, $c = 2$, $T = 5$, and $m = 6$.

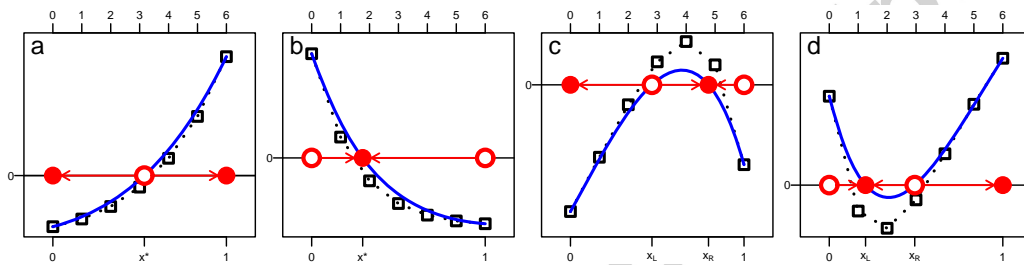


Figure 5: Gain sequence \mathbf{d} (squares, dotted line; top axis), and corresponding gain function $g(x)$ (solid line; bottom axis) and phase portrait (circles, arrows) of the game considered in Section 5.2 for $N = 7$ and different values of the parameters w, v, r and c . (a) $w = 1.3, v = 1.2, r = 1, c = 3$. (b) $w = 0.6, v = 0.57, r = 2, c = 1$. (c) $w = 1.3, v = 1.4, r = 2, c = 3.4$. (d) $w = 0.75, v = 0.6, r = 1.55, c = 1.25$.