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To cite this version:
Olivier Benichou, Vincent Calvez, Nicolas Meunier, Raphael Voituriez. Front acceleration by dynamic selection in Fisher population waves. 2012. hal-00716348

HAL Id: hal-00716348
https://hal.archives-ouvertes.fr/hal-00716348
Submitted on 10 Jul 2012

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Front acceleration by dynamic selection in Fisher population waves

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(Dated: June 12, 2012)

We introduce a minimal model of population range expansion in which the phenotypes of individuals present no selective advantage and differ only in their diffusion rate. We show that such neutral phenotypic variability can yield alone phenotype segregation at the front edge even in absence of genetic noise, and significantly impact the dynamical properties of the expansion wave. We present an exact asymptotic traveling wave solution and show analytically that phenotype segregation accelerates the front propagation. The results are compatible with field observations such as invasions of cane toads in Australia or bush crickets in Britain.

PACS numbers:

I. INTRODUCTION

The combination of random mutations and natural selection plays a key role in evolution. In the case of large well-mixed populations, it is found typically that mutations that eventually become dominant must present a selective advantage (i.e. a larger effective growth rate) over the wild type [1]. More recently, it appeared that the case of expanding populations, which are a common phenomenon in biology, can lead to strikingly different behaviors. It was observed that in population range expansions even neutral mutations, i.e. mutations that present no selective advantage, can prevail at least transiently due to large fluctuations at the edge of the population traveling front, and lead to spectacular gene segregation phenomena [2–4]. Such population waves are exemplified by the hypothesized migration of humans from Africa [5], or invasions of species such as cane toads in Australia [6] or bush crickets in Britain [7], and can now be observed at a smaller scale on the example of migrations of microorganisms in a Petri dish [3]. Following these observations, numerical models have been developed in the ecology community to analyze quantitatively range expansions [8–13]; these studies give further support to the mechanism of phenotype selection, which was dubbed spatial sorting in [14].

The standard mathematical models for population waves are generally based on the Fisher-Kolmogorov (FKPP) equation [15, 16], which is the simplest non linear equation that combines diffusion and growth. Beyond population genetics, this equation has been widely used for example in ecology, epidemiology [17] or chemistry [18]. Fisher waves are still actively investigated from the theoretical point of view and despite this effort, exact results, which mostly concern the propagation speed [19, 20], remain elusive especially in the presence of noise or in space dimensions larger than one.

In this letter, we introduce a minimal model of population dynamics based on a modified FKPP equation in which mutations are neutral and affect only the mobility properties of individuals. Our approach is inspired by the observations of range expansions of cane toads in Australia [6] or bush crickets in Britain [7], as well as several numerical studies[8–14] which highlighted the following features: (i) The leading part of the expanding front is mostly populated by the fastest phenotypes, that is longer-legged individuals in the case of cane toads [6] and longer-winged individuals in the case of bush crickets. (ii) The proportion of fast phenotypes is the highest in locations of new arrivals and then declines over time. (iii) The speed of the population wave increases with time. Our analysis is based on a modified FKPP equation with an additional mutation term that allows for variations of the diffusion rate of individuals, for which we present an exact asymptotic traveling wave solution and calculate analytically the front speed. We show that such phenotypic variability, when taken into account, can yield alone the segregation of the fastest phenotypes at the front edge in absence of genetic noise, and significantly accelerate the propagation speed. Importantly, the results reproduce qualitatively the observations (i)-(iii) above made on the basis of field experiments and numerical simulations. The novelty of our approach lies on the modified FKPP equation that we propose in Eq.(1), which enables an analytical derivation of the main properties of phenotypic segregation in range expansions. We analyze in this paper this equation in its minimal form, keeping in mind that it could be modified to account more faithfully for real systems. Given the wide spectrum of situations described by the FKPP equation, potential applications for example in the context of chemical kinetics are also expected.
The function $P(D,t)$ defined by Eq. (4) can be linearized, which suggests the following ansatz

$$n(D,u) = P_*(D) \exp(-\gamma u),$$

with $\gamma > 0$. Eq. (4) then rewrites

$$0 = (\gamma v + D\gamma^2 + s)P_*(D) + \alpha \partial^2_D P_*(D),$$

(a typical width $\Delta D$). Without loss of generality, a simple choice for the density current is then

$$j = -\alpha \partial_D n(x,D,t)$$

where $\alpha$ is the mutation rate, together with reflecting boundary conditions ($j = 0$) at $D = D_\pm \equiv D_0 - \Delta D/2$ and $D = D_+ \equiv D_0 + \Delta D/2$. With this choice of $j$, the distribution $P_0$ is then uniform over the range $[D_- \cdot D_+]$ and the different phenotypes have no selective advantage. Another example of interest is given by

$$j = -\alpha [\beta(D-D_0)n(x,D,t) + \partial_D n(x,D,t)],$$

where $\alpha$ is the mutation rate and the extra term $\beta$ accounts here for a selective advantage of phenotype $D_0$. This choice, together with the zero flux condition at $D = D_-$ and $D = D_+$, yields $P_0(D) \propto \exp[-\beta(D-D_0)^2/2]$; at stationary state in a bounded range, the trait $D_0$ is now favored, as is the case in many real situations. Alternative choices are possible to account more faithfully for field experiments, and would lead qualitatively to similar results. We will focus below on the minimal choice of Eq. (2), which captures the main features of the model and keep calculations analytically tractable.

We now turn to the case of a range expansion and assume that at $t = 0$ the population is seeded with the uniform distribution $P_0$ in a bounded interval of the $x$ axis. We show that despite the fact that mutations are neutral, a traveling wave solution characterized by a non homogeneous distribution of $D$, which we calculate in the regime $u \equiv x - vt \gg 1$, emerges. We determine the velocity $v_*$ and show that it is larger than the velocity $v_0 = 2\sqrt{sD_0}$ expected from the classical FKPP model with an homogeneous population of diffusion rate $D_0$.

Let us introduce $u \equiv x - vt$ where $v$ is to be determined, and look for a traveling front so that $N(u \to -\infty) = 1$ and $N(u \to +\infty) = 0$. Notations are kept unchanged in the co-moving frame of velocity $v$ and all functions of $x$ and $t$ are assumed to depend on $u$ only. Eq. (1) then rewrites

$$0 = v \partial_u n + D\partial^2_un + sn(1-N) + \alpha \partial^2_D n.$$
where it should be noted that $P_\ast(D)$ depends on $\gamma$. The speed $v$ of a front of prescribed profile characterized by $\gamma$ is then obtained by integration over $D$ and follows:

$$v = D_\ast \gamma + s/\gamma,$$

(7)

where $D_\ast$, which depends on $\gamma$, is determined below. Eq. (6), completed by $\partial_0 P_\ast |_{D=D_\ast} = \partial_D P_\ast |_{D=D_\ast} = 0$ for the choice (2) of the density current $j$, explicitly defines the function $P_\ast$ and shows that it is independent of $u$ in the limit $u \to \infty$, therefore proving that the factorized form of Eq. (5) holds true. We stress that this argument holds for any density current $j$, which would only modify the ordinary differential equation (6) defining $P_\ast$. In the explicit example of Eq. (2), the existence of a non zero solution for Eq. (6) implies that:

$$Ai[1, \mu(D_\ast - D_-)]Bi[1, \mu(D_\ast - D_+)] - Bi[1, \mu(D_\ast - D_-)]Ai[1, \mu(D_\ast - D_+)] = 0,$$

(8)

where $\mu = (\gamma^2/\alpha)^{1/3}$ and Ai and Bi denote Airy functions. This last equation, which can not be solved analytically to the best of our knowledge, implicitly defines $D_\ast$ as a function of $\gamma$, and completes the definition of $v$ in Eq.(7). The distribution $P_\ast$ is then given by

$$ZP_\ast = Ai[1, \mu(D_\ast - D_-)]Bi[1, \mu(D_\ast - D_+)]Bi[1, \mu(D_\ast - D_-)]Ai[1, \mu(D_\ast - D_+)],$$

(9)

where $Z$ is a normalization constant that ensures that $\int_{D_\ast}^\infty P_\ast(D)dD = 1$. This result, together with the decoupling form of Eq.(5) explicitly characterizes the asymptotics of a front of prescribed profile characterized by $\gamma$. In practice, it can be showed following classical arguments [18] that in the case of a localized initial condition in space the slowest solution is selected. The critical velocity $v_\ast$ (and the corresponding $\gamma_\ast$) is therefore obtained as the minimum of the function $v(\gamma)$ defined by Eqs.(7) and (8), and the density for $u$ large follows

$$n(D,u) \propto P_\ast(D)e^{-\gamma_\ast u},$$

(10)

While in practice $v_\ast$ can be determined by a numerical analysis of Eqs.(7) and (8), a useful and very accurate approximate of $v_\ast$, which is in fact an exact upper bound, denoted $v_\ast^\mu$, can be obtained by assuming $D_\ast$ constant in Eq.(7). Under this hypothesis the minimization of $v$ is realized for $\gamma = \gamma_\ast^\mu \equiv \sqrt{s/D_\ast}$ and yields $v_\ast^\mu = 2\sqrt{sD_\ast}$, where $D_\ast$ is determined by Eq. (8) taken at $\gamma = \gamma_\ast^\mu$.

These asymptotic results have been checked numerically by solving Eq. (1) using a classical Euler numerical scheme, the diffusion part being considered implicitly and the reaction part explicitly, and show that the upper bound $v_\ast^\mu$ provides a very accurate estimate of the exact value $v_\ast$. Figure 2 shows an example of the obtained population wave which reaches a stationary state of constant velocity after a transient acceleration regime (Fig. 2a). Figure 2b indicates that the population is the most heterogeneous at the edge of the front where the segregation of the fastest phenotypes is observed, while variations of the trait $D$ decrease behind the front. More quantitatively, we find that after the transient accelera-
tion regime the wave advances at the predicted velocity $v_{s}$ (see Fig. 3a). Numerical results clearly validate the asymptotic result of Eq. (10) as seen in Fig. 3b,c, which show that both the dependences on $u$ and $D$ of the density current $n(D, u)$ are captured by our analysis. Remarkably, the approximate value $v_{s}$ is numerically extremely close to the exact result $v^{*}$ for all parameters analyzed. Last, Fig. 3d confirms that the segmentation of phenotypes with larger diffusion rate $D$ is more important at the leading edge of the front, where it is characterized by the predicted distribution $P_{s}(D)$, while the distribution $P_{u}(D)$ reaches its steady state value $P_{0}(D)$ (which is uniform with the choice of Eq.(2)) at positions $x$ far behind the front.

III. DISCUSSION

Several comments are in order. First, the asymptotic solution of Eq. (10) and the determination of the front velocity $v_{s}$ are exact. Similar results, and in particular the exact asymptotic decoupling of the $u$ and $D$ dependences can be obtained along the same lines for other choices of the density current $j$ defined in Eq. (2). In particular, the important case where a trait $D_{0}$ presents a selective advantage, as modeled by the current defined in Eq. (3), can be analyzed in the same way and leads to qualitatively similar results. Second, one finds that for $s/\alpha \ll 1$, $D_{s} \rightarrow D_{0}$ and $P_{s}(D) \approx P_{0}(D) = 1/\Delta D$ for all $u$. As expected, when mutations are extremely fast the population is locally well-mixed everywhere with diffusion rate $D_{0}$ and the dynamics follows the usual FKPP equation with a front speed $v_{0} = 2\sqrt{D_{0}}$. In the more realistic case of slow mutations, $s/\alpha$ is large and the dynamics selects faster individuals at the leading edge so that $P_{s}(D)$ favors larger values of $D$. In the extreme case of very slow mutations ($s/\alpha \rightarrow \infty$) one has $D_{s} \rightarrow D_{+}$ and $P_{s}(D) \approx \delta(D - D_{+})$. The front can then be significantly faster than the expected velocity $v_{0}$ for a well-mixed population of diffusion rate $D_{0}$ since $v_{s} = 2\sqrt{s(D_{0} + \Delta D/2)} > v_{0}$. Third, it should be emphasized that this segmentation of fastest phenotypes, which leads to an acceleration of the wave propagation, is purely dynamical. It is obtained in the low noise limit, and therefore strikingly differs from the mechanism of segregation of neutral mutations studied in [2–4]. Last, and importantly, our results qualitatively reproduce the observations (i)-(iii) stated in introduction. Indeed, we found that: (i) fastest phenotypes are segregated at the leading edge of the front, as shown in Figs. 2b, 3c. (ii) The proportion of fast phenotypes is higher at the edge and decreases behind the front as shown in Figs. 2b, 3d. (iii) The speed of advance of the population wave increases with time, as seen in Fig. 2a, and is larger than the expected propagation speed for a well-mixed population.

To conclude, we have proposed a minimal model of population range expansion in which mutations are neutral and affect only the mobility of individuals. We have shown that such neutral phenotypic variability can yield alone phenotype segregation at the front edge in absence of genetic noise, and significantly impact the dynamical properties of the expansion wave. An exact traveling wave solution can be obtained asymptotically, and shows that phenotype segregation accelerates the front propagation. The results are compatible with available data of invasions of cane toads in Australia or bush crickets in Britain. Applications beyond the field of population dynamics could be expected.

Acknowledgments

N.M. wishes to thank C. Mouhot for having brought his attention on this subject and for helpful discussions.