

**Introduzione alle equazioni alle derivate parziali,  
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**An application in population genetics.**

We refer to the paper by Aronson and Weinberger **Multidimensional nonlinear diffusion arising in population genetics**, Advances in Mathematics 30, pag 33-76, 1978.

Let consider a population of diploid individuals in a habitat  $\Omega$ . Assume that the gene in specific chromosome pair appears in 2 allelic forms  $a, A$ . Then the population is divided in 3 types: one with genotype  $aa$  or  $AA$  (homozygotes) and one with genotype  $aA$  (heterozygotes). Moreover we assume that the death rate in the population depends on the genotype, and we define 3 positive constants  $\tau_1, \tau_2, \tau_3 \in (0, 1)$ , which are the death rates respectively of type  $aa$ ,  $aA$  and  $AA$ . We can assume without loss of generality that one type of homozygotes has death rate bigger than the other, so we assume

$$\tau_1 \geq \tau_3.$$

Let

$\rho^1(x, t)$  = density of the population with genotype  $aa$  at place  $x$  at time  $t$ ,

$\rho^2(x, t)$  = density of the population with genotype  $aA$ ,

$\rho^3(x, t)$  = density of the population with genotype  $AA$ .

Assume that the growth rate of the population is constant, say  $r > 0$ , and that the population is spreading in the habitat  $\Omega$  with diffusion constant 1. Let  $\rho(x, t) = \rho^1(x, t) + \rho^2(x, t) + \rho^3(x, t)$  the total density. Under our assumptions, the densities satisfy the following system

$$\begin{cases} \rho_t^1 - \Delta \rho^1 = -\tau_1 \rho^1 + \frac{r}{\rho} (\rho^1 + \frac{1}{2} \rho^2)^2 \\ \rho_t^2 - \Delta \rho^2 = -\tau_2 \rho^2 + \frac{r}{\rho} (\rho^1 + \frac{1}{2} \rho^2) (\rho^3 + \frac{1}{2} \rho^2) \\ \rho_t^3 - \Delta \rho^3 = -\tau_3 \rho^3 + \frac{r}{\rho} (\rho^3 + \frac{1}{2} \rho^2)^2. \end{cases}$$

Define

$$u(x, t) = \frac{\rho^3 + \frac{1}{2} \rho^2}{\rho}$$

be the relative density of allele  $A$  at place  $x$  at time  $t$ . Under appropriate assumptions (in particular assuming that  $|\tau_1 - \tau_2| + |\tau_2 - \tau_3| + |\tau_3 - \tau_1|$  is small), it is possible to justify the fact that  $u$  solves the following semilinear problem

$$\begin{cases} u_t - \Delta u = F(u) & x \in \Omega \times (0, +\infty) \\ u(x, 0) = u_0(x) & x \in \Omega \\ u(x, t) = 0 & x \in \partial\Omega, t > 0 \end{cases}$$

where

$$F(u) = u(1 - u)[(\tau_1 - \tau_2) - (\tau_1 + \tau_3 - 2\tau_2)u].$$

The initial distribution is  $u_0(x) \in \mathcal{C}(\bar{\Omega})$ , with

$$0 \leq u_0(x) \leq 1 \quad \forall x \in \Omega$$

(since it is a density).

The euristic argument for this derivation can be found in the paper by Fischer (1937) and in previous papers by Aronson and Weinberger.

First of all observe that 0, 1 are equilibria of the system, and moreover  $0 \leq u_0 \leq 1$ . So by comparison principle, we obtain that

$$0 \leq u(x, t) \leq 1 \quad \forall (x, t) \in \bar{\Omega} \times [0, +\infty).$$

We consider 3 different cases.

**Heterozygote intermediate** Assume that  $\tau_3 \leq \tau_2 < \tau_1$ .

Under this case, note that  $F(s) > 0$  for all  $s \in (0, 1)$ ,  $F'(0) > 0$  and  $F'(1) < 0$ . In this case the state 1 is stable, whereas 0 is unstable. More precisely, we get that for every initial distribution  $u_0$  such that  $u_0 \not\equiv 0$ , then

$$\lim_{t \rightarrow +\infty} u(x, t) = 1$$

uniformly, with exponential rate depending on  $F'(1)$ .

In this case, we expect that in large time allele  $a$  will disappear.

**Heterozygote superior** Assume that  $\tau_2 < \tau_3 \leq \tau_1$ .

In this case

$$c := \frac{\tau_1 - \tau_2}{(\tau_1 - \tau_2) + (\tau_3 - \tau_2)} \in (0, 1)$$

is another equilibrium of the system in  $[0, 1]$  besides 0, 1. Moreover  $F(s) > 0$  for every  $s \in (0, c)$ ,  $F(s) < 0$  for every  $s \in (c, 1)$ ,  $F'(0) > 0$ ,  $F'(1) > 0$ ,  $F'(c) < 0$ . So  $c$  is the only stable equilibrium and that for every initial distribution  $u_0$  such that  $u_0 \not\equiv 0$ ,  $u_0 \not\equiv 1$ , then

$$\lim_{t \rightarrow +\infty} |u(x, t) - c| = 0$$

uniformly, with exponential rate depending on  $F'(c)$ .

In this case, we expect that in large time, the distribution of allele  $A$  is constant (given by  $c$ ) and then also the distribution of allele  $a$  is constant (given by  $1 - c$ ).

**Heterozygote inferior** Assume that  $\tau_3 \leq \tau_1 < \tau_2$ .

Also in this case there is another equilibrium of the system, which the same as above

$$c := \frac{\tau_2 - \tau_1}{(\tau_2 - \tau_1) + (\tau_2 - \tau_3)} \in (0, 1).$$

Differently from the previous case, we get  $F(s) < 0$  for every  $s \in (0, c)$ ,  $F(s) > 0$  for every  $s \in (c, 1)$ ,  $F'(0) < 0$ ,  $F'(1) < 0$ ,  $F'(c) > 0$ . In this case 0, 1 are two stable equilibria and  $c$  is unstable. If the initial distribution  $u_0$  is such that  $u_0 \not\equiv c$ , and  $u_0(x) \leq c$  for every  $x$ , then

$$\lim_{t \rightarrow +\infty} u(x, t) = 0$$

uniformly, with exponential rate depending on  $F'(0)$ .

If the initial distribution  $u_0$  is such that  $u_0 \not\equiv c$ , and  $u_0(x) \geq c$  for every  $x$ , then

$$\lim_{t \rightarrow +\infty} u(x, t) = 1$$

uniformly, with exponential rate depending on  $F'(1)$ .

This means that, if the initial distribution of allele  $A$  is under a given threshold ( $c$ ), then we expect that in large time allele  $A$  will disappear. On the other hand, if allele  $s$  is above the same threshold ( $c$ ), then we expect that in large time allele  $a$  will disappear.

The latest case, the case of heterozygote inferior, is the most interesting one. In this case, we have 2 stable equilibria and one unstable equilibrium. The next point will be to study which one between the two stable equilibria is more attractive.