Mathematical Models of Epidemics
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Introduction

The aim of these Notes is to present the essentials of probabilistic model of the propagation of epidemics. These are very serious issues concerning public health. During the 14th century, the black plague killed between 30% and 50% of Europe’s population. In 1720 a plague epidemic decimated almost half of the population of Marseille and one fourth of the population of Provence. The Spanish flu in 1918–1919 killed between 30 et 100 million humans. It resulted from a particularly virulent $H1N1$ strain. It has been probably the most severe pandemic in human history so far. Even with the progress of medicine and vaccination, several illnesses have not been eradicated (e.g. malaria), and new ones have appeared (HIV, SARS, Ebola), some of them propagating faster than in the past, due to more rapid and massive transportation. In addition, we should not forget the hospital–acquired infections (even in the cleanest hospitals of the rich part of the world), and the serious problem of antibiotic resistance of bacteria. As a matter of fact, the fight against epidemics is not a problem of the past for humanity. It is a problem of the present and the future, in particular in Africa.

A little more than one hundred years ago, Sir Ronald Ross, a british medical doctor who contributed to the understanding of malaria (together with among others the Italian Giovanni Battista Grassi and the French Laveran, Ross and Laveran won a Nobel prize) wrote: “As a matter of fact all epidemiology, concerned as it is with variation of disease from time to time and from place to place, must be considered mathematically (...) and the mathematical method of treatment is really nothing but the application of careful reasoning to the problems at hand”. As a matter of fact, Ross deduced from mathematical arguments conclusions concerning malaria, which his colleagues physicians had difficulties to accept.

The main aim of these Notes is to describe some aspects of mathematical epidemiology, with an emphasis on probabilistic models. Learning this topics is also a good way to learn mathematics and mathematical modeling. Historically, deterministic models have received most attention. But as we shall see, probabilistic modeling is essential. We shall also discuss some of the associated statistical procedures.

These notes are very much inspired by the recent monograph by Diekmann, Heesterbeek and Britton [2]. Section 14.5 is taken from [7]. We urge the reader to consult [5] for a complete treatment of the “interludes” below (except for the second one).
1 Epidemics in a closed population

We are going to assume here, and in a great part of these Notes, that the studied epidemic concerns a population of fixed size \( N \). This is justified whenever, and it is often the case, the epidemic under study has a life time which rather short, compared with the time scale of the fluctuations of the population size. Some results will be established for large \( N \) (we shall study the limit of the model as \( N \to \infty \)), others for arbitrary \( N \).

We consider the situation where 1 or a small number of infected individuals are introduced in a population of susceptibles. We discuss the following questions:

- Under which circumstances might a major epidemic start, and what is the probability that such an event occur?
- In case a major epidemic develops, at which speed does it progress?
- Which fraction of the total population will be eventually hit by the epidemic?
- How long will the epidemic last?

In order to answer those questions, we first need, in order to formulate a mathematical model of the epidemic,

- to describe the process of contacts which propagates the illness;
- to describe how the population mixes (who meets whom?), and which fraction of the contacts of an infected individual will be with “susceptible” individuals;
- to precise the probability that such a contact yields the transmission of the illness.

Let us first explain that our models will compartmental models, which means that the population under consideration will be divided into compartments, each individual belonging to one and only one of those compartments. The number of compartments depends upon the choice of a particular model and of course upon the type of illness under study. The main compartments are:
$ S $ like susceptible, it is the subset of those individuals which might contract the illness at the occasion of an encounter with an infectious individual;

$ E $ like exposed, it is the subset of those individuals who are suffering from the illness, but they are in the incubation phase, they are not contagious;

$ I $ like infectious, it is the subset of those individuals who are suffering from the illness, and are contagious;

$ R $ like removed, it is the subset of those individuals who have suffered from the illness, and have recovered; they have acquired immunity concerning the illness, they are not susceptible to contract it again. One might include dead individuals in the compartment $ R $.

The propagation of the illness is the result of an encounter between an individual from compartment $ I $ and an individual from compartment $ S $. What is the meaning of encounter? It depends upon the illness. In case of HIV, it means a sexual intercourse (or a contact of bloods, for example through an exchange of syringe). In the case of malaria, it means a bite of a human by a mosquito, where one of the two is susceptible (of type $ S $), the other one being infectious (of type $ I $). In case of the flu, SARS, contact can just mean shaking hands, or an infectious individual sneezing in the face of a susceptible (same for the whooping cough).

Let us suppose that each individual of type $ I $ meets other individuals at rate $ c $ (meaning that this individual meets in average $ c $ individuals per unit time). In other words, encounters of that individual with other individuals of the population happen according to a rate $ c $ Poisson process (see section 2 below). The next question is: whom does that infectious individual meet? We shall assume in almost all of these Notes that the population is fully mixed, which means that the individual who is met is chosen uniformly among all individuals of the population except himself. Other more realistic situations will be discussed in other courses of this school. Our assumption allows to simplify the model, and obtain first interesting results, as we shall see.
2 Interlude 1. The Poisson Process

This rather simple process will be central in what follows. Let \( \lambda > 0 \) be given. A rate \( \lambda \) Poisson (counting) process is defined as

\[
P_t = \sup\{k \geq 1, T_k \leq t\},
\]

where \( 0 = T_0 < T_1 < T_2 < \cdots < T_k < \cdots < \infty \), the r.v.’s \( \{T_k - T_{k-1}, k \geq 1\} \) being independent and identically distributed, each following the law \( \text{Exp}(\lambda) \).

We have

**Proposition 1.** For all \( n \geq 1, 0 < t_1 < t_2 < \cdots < t_n, \) the r.v.’s \( P_{t_1}, P_{t_2} - P_{t_1}, \ldots, P_{t_n} - P_{t_{n-1}} \) are independent, and for all \( 1 \leq k \leq n, P_{t_k} - P_{t_{k-1}} \sim \text{Poi}[\lambda(t_k - t_{k-1})] \).

**Proof** Let us first prove that for all \( t,s > 0, \)

\[
\mathbb{P}(P_{t+s} - P_t = 0|P_t = k,T_1,T_2,\ldots,T_k) = \exp(-\lambda s).
\]

Indeed

\[
\mathbb{P}(P_{t+s} - P_t = 0|P_t = k,T_1,T_2,\ldots,T_k) = \mathbb{P}(T_{k+1} > t + s|P_t = k,T_k)
\]
\[
= \mathbb{P}(T_{k+1} - T_k > t + s - T_k|T_{k+1} - T_k > t - T_k > 0)
\]
\[
= \mathbb{P}(T_{k+1} - T_k > s)
\]
\[
= \exp(-\lambda s).
\]

Let now \( n \geq 1. \) For \( 1 \leq i \leq n, \) we define \( X_{n,i} = 1\{P_{t+i/s} - P_{t+(i-1)/s} \geq 1\}, \) and finally \( S_n = X_{n,1} + X_{n,2} + \cdots + X_{n,n}. \) It follows from the first part of the proof that conditionally upon \( \sigma\{P_r, 0 \leq r \leq t\}, \) the r.v.’s \( X_{n,1}, X_{n,2}, \ldots, X_{n,n} \) are i.i.d., each Bernoulli with parameter \( 1 - e^{-\lambda s/n}. \) Then conditionally upon \( \sigma\{P_r, 0 \leq r \leq t\}, S_n \) is binomial with parameters \( (n,1-e^{-\lambda s/n}). \) But \( S_n \rightarrow P_{t+s} - P_t \) a.s. as \( n \rightarrow \infty, \) while its conditional law given \( \sigma\{P_r, 0 \leq r \leq t\} \) converges towards the Poisson distribution with parameter \( \lambda s, \) according to the following Lemma. The Proposition follows. \( \square \)

We have used the following well-known result. Recall the notation \( \text{B}(n,p) \) for the binomial law with parameters \( n \) and \( p, \) where \( n \geq 1 \) and \( 0 < p < 1. \)

**Lemma 2.** For all \( n \geq 1, \) let \( U_n \) be a \( \text{B}(n,p_n) \) random variable. If \( np_n \rightarrow \lambda \) as \( n \rightarrow \infty, \) with \( \lambda > 0, \) then \( U_n \) converges in law towards \( \text{Poi}(\lambda). \)
A Poisson process will be called standard if its rate is 1. If \( P \) is a standard Poisson process, then \( \{ P(\lambda t), \ t \geq 0 \} \) is a rate \( \lambda \) Poisson process.

We will also use the following

**Exercise 1.** Let \( \{ P_t, \ t \geq 0 \} \) be a rate \( \lambda \) Poisson process, and \( \{ T_k, \ k \geq 1 \} \) the random points of this Poisson process, such that for all \( t > 0 \), \( P_t = \sup \{ k \geq 1, \ T_k \leq t \} \). Let \( 0 < p < 1 \). Suppose that each \( T_k \) is selected with probability \( p \), not selected with probability \( 1 - p \), independently of the others. Let \( P'_t \) denote the number of selected points on the interval \([0, t]\). Then \( \{ P'_t, \ t \geq 0 \} \) is a rate \( \lambda p \) Poisson process.

### 3 Start of an epidemic

During the initial phase of an epidemic, there are very few infectious individuals, so that we can pretend that any encountered individual is susceptible.

We now need to precise the probability that an infectious individual infects an encountered susceptible. Consider an individual who has been infected at the initial time \( t = 0 \). He is in state \( E \) when \( 0 \leq t \leq T_1 \), in state \( I \) when \( T_1 \leq t < T_2 \), and in state \( R \) when \( t \geq T_2 \). He will infect a susceptible encountered at time \( t \) with probability

\[
\begin{cases} 
0, & \text{if } t < T_1; \\
p, & \text{if } T_1 \leq t < T_2; \\
0, & \text{if } t \geq T_2.
\end{cases}
\]

We state the

**Definition 3.** The “basic reproduction number” is the quantity \( R_0 \) defined as the mean number of susceptibles whom an infectious individual infects, during the initial phase of the epidemic.

Note that “during the initial phase of the epidemic” is an essential precision. This means “while all encountered individuals are susceptibles”. With the above notations, if we let \( \Delta T = T_2 - T_1 \), then

\[
R_0 = cp\mathbb{E}[\Delta T].
\]
4 Interlude 2. Branching processes

We describe here the basic results concerning discrete time branching processes, also called Bienaymé–Galton–Watson processes.

Consider an ancestor (at generation 0) who has $X_0$ children, such that

$$\mathbb{P}(X_0 = k) = q_k, \ k \geq 0 \ \text{et} \ \sum_{k \geq 0} q_k = 1.$$ 

Define $m = \mathbb{E}[X_0] = \sum_{k \geq 1} k q_k$.

Each child of the ancestor belongs to generation 1. The $i$-th of those children has himself $X_{1,i}$ children, where the r.v.'s $\{X_{k,i}, \ k \geq 0, i \geq 1\}$ are i.i.d., all having the same law as $X_0$. If we define $Z_n$ as the number of individuals in generation $n$, we have

$$Z_{n+1} = \sum_{i=1}^{Z_n} X_{n,i}.$$ 

Let $g$ denote the generating function of the r.v. $X_0$, i.e.

$$g(s) = \sum_{k=0}^{\infty} q_k s^k = \mathbb{E}[s^{X_0}], \ 0 \leq s \leq 1.$$ 

We have $g(0) = q_0$, $g(1) = 1$, $g'(1) = m$, $g''(s) > 0$, for all $0 \leq s \leq 1$ (we assume that $q_0 > 0$ and $q_0 + q_1 < 1$). Let us compute the generating function of $Z_n : g_n(s) = \mathbb{E}[s^{Z_n}]$.

$$g_n(s) = \mathbb{E} \left[ s^{\sum_{i=1}^{Z_{n-1}} X_{n-1,i}} \right]$$

$$= \mathbb{E} \left[ \mathbb{E} \left[ s^{\sum_{i=1}^{Z_{n-1}} X_{n-1,i}} \bigg| Z_{n-1} \right] \right]$$

$$= \mathbb{E} \left[ g(s)^{Z_{n-1}} \right]$$

$$= g_{n-1} \circ g(s).$$

If we iterate this argument, we obtain

$$g_n(s) = g \circ \cdots \circ g(s),$$

and also

$$\mathbb{P}(Z_n = 0) = g^{\circ n}(0)$$

$$= g \left[ g^{\circ (n-1)}(0) \right].$$
Figure 1: Graphs of $g$ in case $m > 1$ (left) and in case $m \leq 1$ (right).

Hence if $z_n = \mathbb{P}(Z_n = 0)$, $z_n = g(z_{n-1})$, and $z_1 = q_0$. We have $z_n \uparrow z_\infty$, where $z_\infty = \mathbb{P}(Z_n = 0$ from some $n$ on). The proof of the following Proposition is essentially clear from Figure 1.

**Proposition 4.** If $m \leq 1$, then $\mathbb{P}(Z_n = 0) \to 1$ as $n \to \infty$, and $z_\infty = 1$.

If $m > 1$, $\mathbb{P}(Z_n = 0) \to z_\infty$ as $n \to \infty$, where $z_\infty$ is the smallest solution of the equation $z = g(z)$.

In the second case, with probability $1 - z_\infty$, the branching process does not go extinct. Let us show that $W_n = m^{-n}Z_n$ is a martingale.

\[
\mathbb{E}(W_{n+1}|Z_n) = m^{-n}\mathbb{E}\left(m^{-1}\sum_{i=1}^{Z_n}X_{n,i}|Z_n\right) = m^{-n}Z_n = W_n.
\]

One can show that $W_n \to W$ a.s. as $n \to \infty$, and moreover

\[
\{W > 0\} = \{\text{the branching process does not go extinct}\}.
\]

## 5 The start of the epidemic

The start of the epidemic behaves as a BGW process, with $m = R_0$, since as long as we can pretend that any encountered individual is susceptible,
the various processes of transmission of the illness are independent. What is the relation between \( z_\infty \) and \( R_0 \)? Conditionally upon \( \Delta T \), the number of contacts of an infectious individual at the start of the epidemic is \( k \) with probability

\[
e^{-c\Delta T \frac{(c\Delta T)^k}{k!}},
\]
each of those contacts resulting in an infection with probability \( p \). We then deduce from Proposition 1 and Exercise 1 that, conditionally upon \( \Delta T \), the number of susceptibles infected by one infectious individual at the start of the epidemic is \( k \) with probability

\[
\exp (-cp\Delta T) \frac{(cp\Delta T)^k}{k!}.
\]
We now compute \( z_\infty \) in two cases.

### 5.1 \( \Delta T \) constant

Assume that this constant value is independent of the considered individual. Then \( g \) is the generating function of the Poisson distribution with parameter \( R_0 = cp\Delta T \).

\[
g(s) = \sum_{k=0}^{\infty} e^{-R_0} \frac{(R_0 s)^k}{k!} = e^{R_0(s-1)}.
\]

Hence \( z_\infty \) is the smallest solution of equation \( z = e^{R_0(z-1)} \).

### 5.2 \( \Delta T \sim \text{Exp}(\alpha) \).

In this case, the generating function \( g \) is given as

\[
g(s) = \alpha \int_0^\infty e^{cp(s-1)}e^{\alpha h} dh = \frac{\alpha}{\alpha - cp(s - 1)}.
\]
It is the generating function of a geometric distribution, in other words

\[
\mathbb{P}(X_0 = k) = \left( \frac{cp}{cp + \alpha} \right)^k \frac{\alpha}{cp + \alpha}.
\]
Here $R_0 = cp/\alpha$, $z_\infty$ is the smallest solution of the equation

$$z = \frac{1}{1 - R_0(z - 1)}.$$ 

Consequently

$$z_\infty = \frac{1}{R_0}.$$ 

We see in particular that the relation between $z_\infty$ and $R_0$ depends very much upon the details of the model.

5.3 Remark on the speed of propagation of the epidemic

That speed does not depend only upon $\Delta T$, but upon the pair $(T_1, T_2)$. If we compare the demography of a country where each women has three children between the age of 20 and the age of 25, with that of another country where each women has three children between the age of 35 and the age of 40, it is rather clear that the speed at which those two populations evolve are different.

The two quantities $cp$ and $\Delta T$ being kept constant, we can choose two pairs $(T_1^*, T_2^*)$ and $(T_1^{**}, T_2^{**})$ such that $R_0^* > R_0^{**}$ and $V^* < V^{**}$. We shall discuss this issue again later.

6 The final size of the epidemic in case of no major outbreak

Let $X_1, X_2, \ldots$ be i.i.d. $\mathbb{N}$-valued r.v.’s, all having the same law as $X_0$. Let $Z$ denote the final size of the epidemic (i.e. the total number of individuals which are infected at some stage of the epidemic, including the initially infected individual).

**Proposition 5.** For all $k \geq 1$,

$$\mathbb{P}(Z = k) = \frac{1}{k} \mathbb{P}(X_1 + X_2 + \cdots + X_k = k - 1).$$
Consider the process of depth–first search of the genealogical tree of the infected individuals. The tree is explored from the root. Suppose we have visited \( k \) vertices. The next visit will be to the leftmost still unexplored son of this individual, if any; otherwise to the leftmost unexplored son of the nearest ancestor of the last visited individual. \( X_1 \) is the number of sons of the root. \( X_k \) is the number of sons of the \( k \)-th visited individual. This exploration of the tree ends at step \( k \) if and only if \( X_1 \geq 1, \ X_1 + X_2 \geq 2, \ X_1 + X_2 + X_3 \geq 3, \ldots \ X_1 + X_2 + \cdots X_{k-1} \geq k - 1, \) and \( X_1 + X_2 + \cdots + X_k = k - 1. \) Let us rewrite those conditions. Define
\[
Y_i = X_i - 1, \ i \geq 1, \ \\
S_k = Y_1 + Y_2 + \cdots + Y_k.
\]
A trajectory \( \{Y_i, 1 \leq i \leq k\} \) explores a tree of size \( k \) iff the following conditions are satisfied
\[
(C_k) \ S_0 = 0, S_1 \geq 0, S_2 \geq 0, \ldots, S_{k-1} \geq 0, S_k = -1.
\]
The statement of the Proposition is equivalent to
\[
\mathbb{P}(Z = k) = \frac{1}{k} \mathbb{P}(Y_1 + Y_2 + \cdots + Y_k = -1).
\]
Denote by \( V_k \) the set of sequences of \( k \) integers \( \geq -1 \) which satisfy conditions \( (C_k) \), and \( U_k \) the set of sequences of \( k \) integers \( \geq -1 \) which satisfy the unique condition \( S_k = -1. \) We will use circular permutations operating on the \( Y_i \)'s. For \( 1 \leq i, \ell \leq k, \) let
\[
(i + \ell)_k = \begin{cases} 
  i + \ell, & \text{if } i + \ell \leq k; \\
  i + \ell - k, & \text{if } i + \ell > k.
\end{cases}
\]
For each \( 1 \leq \ell \leq k, \) let \( Z_i^\ell = Y_{(i+\ell)k}, \ S_j^\ell = \sum_{i=1}^j Z_i^\ell \) for \( 1 \leq i \leq k. \) Clearly \( S_k^\ell = -1 \) for all \( \ell \) as soon as \( (C_k) \) is satisfied. On the other hand \( S_k^* = S \) is the only trajectory which satisfies conditions \( (C_k). \) The other \( S^\ell \) hit the value \(-1\) before rank \( k. \) The \( Z^\ell \)'s are sequences of integers \( \geq -1 \) of length \( k, \) whose sum equals \(-1. \) Finally to each element of \( V_k \) we have associated \( k \) distinct elements of \( U_k, \) all having the same probability.

Reciprocally, to one element of \( U_k \setminus V_k, \) choosing \( \ell = \arg\min_{1 \leq i \leq k} S_i \) and using the above transformation, we deduce that \( S^\ell \in V_k. \)
Finally, to each trajectory of $V_k$, we associate $k$ trajectories of $U_k$, who all have the same probability, and which are such that the inverse transformation gives back the same trajectory of $V_k$. The result is proved. □

Note that we have clearly
\[
\sum_{k \geq 1} \mathbb{P}(Z = k) \begin{cases} 
 1, & \text{if } R_0 \leq 1; \\
 1 - R_0, & \text{if } R_0 > 1,
\end{cases}
\]
which is not so obvious from the Proposition.

**Example 6.** Suppose that the joint law of the $X_i$’s is $\text{Poi}(\mu)$, with $0 < \mu < 1$. Then $X_1 + \cdots + X_k \sim \text{Poi}(k\mu)$, and consequently
\[
\mathbb{P}(Z = k) = \frac{1}{k} \mathbb{P}(X_1 + \cdots + X_k = k - 1)
= e^{-\mu k} \frac{(\mu k)^{k-1}}{k!}.
\]
This law of $Z$ is called the Borel distribution with parameter $\mu$. Note that
\[
\mathbb{E}Z = 1 + \mu + \mu^2 + \cdots
= \frac{1}{1 - \mu}.
\]

**Example 7.** Consider now the case where $X_i \sim \mathcal{G}(p)$, where we mean here the geometric distribution where the value 0 is taken with probability $p$. The law of $X_i + 1$ is the geometric distribution with parameter $p$ whose support is $\mathbb{N}$, in other words $\mathbb{P}(X_i + 1 > k) = (1 - p)^k$. $k + X_1 + \cdots + X_k$ follows the negative binomial distribution with parameters $(k, p)$. Hence
\[
\mathbb{P}(Z = k) = \frac{1}{k} \mathbb{P}(k + X_1 + \cdots + X_k = 2k - 1)
= \frac{1}{k} \binom{2k - 2}{k - 1} p^k (1 - p)^{k-1}
= \frac{(2k - 2)!}{k!(k - 1)!} p^k (1 - p)^{k-1}.
\]
In case $p > 1/2$, $\mathbb{E}Z = (2p - 1)^{-1}p$. 
7 The case of a major outbreak

7.1 Approximation of the initial phase of the epidemic by a branching process

In case of a major outbreak, i.e. the epidemic hits a fraction of the total population, how long can we approximate the epidemic by a branching process?

Suppose that the epidemic starts with a unique initial infected individual. The probability that the first $k$ contacts are with susceptibles is

$$1 \left(1 - \frac{1}{N}\right) \left(1 - \frac{2}{N}\right) \times \cdots \times \left(1 - \frac{k-1}{N}\right),$$

which tends to 1 as $N \to \infty$, for any fixed $k$. Suppose now that $k = k_N$ depends upon $N$. Then

$$1 \left(1 - \frac{1}{N}\right) \left(1 - \frac{2}{N}\right) \times \cdots \times \left(1 - \frac{k_N-1}{N}\right)$$

$$= 1 - \frac{k_N-1}{N} + k_N^2 O \left(\frac{1}{N^2}\right)$$

$$= 1 - \frac{k_N(k_N-1)}{N} + k_N^2 O \left(\frac{1}{N^2}\right)$$

$$\to 1,$$

if $k_N = o(\sqrt{N})$. Note the number of generations needed by a BGW process to reach the value $\sqrt{N}$ is of the order of $\log(N)$. Indeed since $m^{-n}Z_n \to W$, we expect that, in case of non extinction, for $n$ large enough, $Z_n$ is of the order of $m^n W$. But if $m^n W = \sqrt{N}$, this implies that

$$n \log(m) + \log(W) = \frac{1}{2} \log(N),$$

$$n = \frac{\log N}{2 \log m} - \frac{\log W}{\log m}.$$

This argument is of course not rigorous, since the fact that $m^{-n}Z_n \to W$ does not imply that $Z_n$ is close to $m^n W$. In fact one can show that $(Z_n)^{-1/2}[Z_n - m^n W]$ converges towards a centered Gaussian r.v. with variance $(m^2 - m)^{-1} \sigma^2$, if $\sigma^2$ is the variance of the reproduction law (provided
that variance is finite). One can also prove a law of iterated logarithm, which goes in the same direction. Those results justify the above (rather vague) statement.

### 7.2 Total size of the epidemic

Conditionally upon $\Delta T$, the probability that a given susceptible escapes infection by a given infectious individual is $\exp(-cp\Delta T/N)$. Hence the probability that a given susceptible escapes infection by a given infectious individual is

$$\mathbb{E}\left[e^{-cp\Delta T/N}\right].$$

The probability that a given susceptible escapes infection by a set of $k$ infectious individuals is

$$\mathbb{E}\left[e^{-cp(\Delta T_1 + \cdots + \Delta T_k)/N}\right].$$

If $k$ is of the order of $N$, then from the Law of Large Numbers, for $N$ large,

$$\frac{\Delta T_1 + \cdots + \Delta T_k}{N} \sim \frac{k}{N}\mathbb{E}(\Delta T).$$

Hence the probability that a given susceptible escapes infection by a fraction $\frac{k}{N}$ of infectious individuals is

$$e^{-cp\frac{k}{N}\mathbb{E}[\Delta T]} = e^{-\frac{k}{N}R_0}.$$

Let $Y$ denote the total number of individuals who are infected in the course of the epidemic. $\frac{Y}{N}$ is the fraction of the population hit by the illness. As we shall see below, the law of large numbers tells us that $Y/N$ is approximatively constant if $N$ is large. The fraction of the population which escapes the illness is $\sigma = 1 - Y/N$. Since each individual is hit by the the illness with the same probability, we have that

$$\sigma = \mathbb{P}(\text{escaping infection}) = \exp(-R_0Y/N),$$

hence

$$\sigma = e^{-R_0(1-\sigma)}.$$

Note that $\sigma = S(\infty)/N = s(\infty)$, and we have already encountered this equation.
8 The Sellke construction

We number the individuals from 0 to $N$:

$$0 \ 1 \ 2 \ 3 \ \ldots \ \ N.$$  

0 denotes the initially infected individual, and the individuals numbered from 1 to $N$ are all susceptible at time 0.

Let

$$Q_1, Q_2, \ldots, Q_N$$

be i.i.d. r.v.’s, with the law $\text{Exp}(1)$;

$$(T_{1,0}, \Delta T_0), (T_{1,1}, \Delta T_1), \ldots, (T_{1,N}, \Delta T_N)$$

i.i.d. r.v.’s, with the law $\mathbb{P}_L \otimes \mathbb{P}_I$, where $\mathbb{P}_L$ is the law of the latency period and $\mathbb{P}_I$ that of the infectious period.

Individual 0 has the latency period $T_{1,0}$ and the infectious period $\Delta T_0$. We denote below

$$L(t)$$

the number of individuals in state $E$ at time $t$;

$$I(t)$$

the number of individuals in state $I$ at time $t$.

We define the cumulated force of infection experienced by an individual, between times 0 and $t$ as

$$\Lambda_C(t) = \frac{cP}{N} \int_0^t I(s) ds.$$  

For $i = 1, \ldots, N$, individual $i$ is infected at the time when $\Lambda_C(t)$ achieves the value $Q_i$ (which might be considered as the “level of resistance to infection of individual $i$”). The $j$-th infected susceptible has the latency period $T_{1,j}$ and the infectious period $\Delta T_j$. The epidemic stops when there is no more individual in either latent of infectious state. Then $\Lambda_C(t)$ does not grow any more, $\Lambda_C(t) = \Lambda_C(\infty)$. The individuals such that $Q_i > \Lambda_C(\infty)$ escape infection.

We put the $Q_i$’s in increasing order: $Q_{(1)} < Q_{(2)} < \cdots < Q_{(N)}$. It is the order in which individuals are infected in Sellke’s model. Note that Sellke’s model respects the durations of latency and infection. In order to show that Sellke’s construction gives a process which has the same law as the process defined above, it remains to verify that the rates at which infections happen are the correct ones.
In the initial model, we assume that each infectious meets other individuals at rate $c$. Since each individual has the same probability of being the one who is met, the probability that a given individual is that one is $1/N$. Hence the rate at which a given individual is met by an infectious one is $c/N$. Each encounter between a susceptible and an infectious individual achieves an infection with probability $p$. Hence the rate at which a given individual is infected by a given infectious individual is $cp/N$. The rate at which an infectious individual infects susceptibles is then $cpS(t)/N$. Finally the epidemic propagates at rate $cpS(t)I(t)/N$.

Let us go back to Sellke’s construction. At time $t$, $S(t)$ susceptibles have not yet been infected. Each of those corresponds to $Q_i > \Lambda_C(t)$. At time $t$, the slope of the curve which represents the function $t \mapsto \Lambda_C(t)$ is $cpI(t)/N$. If $Q_i > \Lambda_C(t) = x$, then

$$\mathbb{P}(Q_i > x + y|Q_i > x) = e^{-y},$$

hence

$$\mathbb{P}(Q_i > \Lambda_C(t + s)|Q_i > \Lambda_C(t)) = \exp\left(-\frac{cp}{N} \int_t^{t+s} I(r) dr\right) = \exp\left(-\frac{cp}{N} I(t)s\right),$$

if $I$ is constant on the interval $[t, t + s]$.

Consequently, conditionally upon $Q_i > \Lambda_C(t)$,

$$Q_i - \Lambda_C(t) \sim \text{Exp}\left(\frac{cp}{N} I(t)\right).$$

The same is true for those $S(t) Q_i$ which are $> \Lambda_C(t)$. Then the first $Q_i$ to come is the minimum of those, hence the waiting time after $\Lambda_C(t)$ for the next infection follows the law $\text{Exp}\left(\frac{cp}{N} I(t)S(t)\right)$, if no removal of an infectious individual happens in the mean time, which would modify $I(t)$.

Then in Sellke’s construction, at time $t$ the next infection comes at rate

$$\frac{cp}{N} I(t)S(t),$$

as in the above described model.
9 Interlude 3. Generalization of the Poisson process

A rate \( \lambda \) Poisson process (\( \lambda > 0 \)) is a counting process \( \{ Q_t, t \geq 0 \} \) such that
\[ Q_t - \lambda t \]
is a martingale. Let \( \{ P(t), t \geq 0 \} \) be a standard Poisson process (i.e. with rate 1). Then \( P(\lambda t) - \lambda t \) is martingale, and it is not hard to show that \( \{ P(\lambda t), t \geq 0 \} \) is a rate \( \lambda \) Poisson process. Let now \( \{ \lambda(t), t \geq 0 \} \) be a measurable and locally integrable \( \mathbb{R}_+ \)-valued function. Then the process
\[ \{ Q_t := P \left( \int_0^t \lambda(s) ds \right), t \geq 0 \} \]
\( \{ Q_t := P \left( \int_0^t \lambda(s) ds \right), t \geq 0 \} \)
is called a rate \( \lambda(t) \) Poisson process. Clearly
\[ Q_t - \int_0^t \lambda(s)ds \]
is a martingale.

Let now \( \{ \lambda(t), t \geq 0 \} \) be an \( \mathbb{R}_+ \)-valued stochastic process, which at each time \( t \) depends only upon the past of \( Q \) below. Then the counting process
\[ Q_t := P \left( \int_0^t \lambda(s) ds \right), t \geq 0 \]
has again the property that
\[ Q_t - \int_0^t \lambda(s)ds \]
is a martingale.

It is sometimes called “a doubly stochastic Poisson process” or a Cox process. Of course the increments of \( Q_t \) are not Poisson distributed. In particular, the process which counts the new infections, which we have described in the preceding section, takes the form
\[ P \left( \frac{cp}{N} \int_0^t I(r)S(r)dr \right). \]

10 LLN and CLT for the final size of the epidemic

Define, for \( 0 \leq w \leq N + 1 \), with the notation \([w] = \text{integer part of } w\),
\[ J(w) = \frac{cp}{N} \sum_{i=0}^{[w]-1} \Delta T_{(i)}. \]

Note that \( i = 0 \) is the index of the initially infected individual, \( \Delta T_{(i)} \) is the latency period of individual whose resistance level is \( Q_{(i)}. \)
$\mathcal{J}(w)$ is the infection pressure produced by the first $\lfloor w \rfloor$ infected individuals (including number 0). For any integer $k$, $\mathcal{J}$ is of course constant on the interval $[k, k+1)$. Define for $v > 0$,

$$Q(v) = \sum_{i=1}^{N} \mathbf{1}_{\{Q_i \leq v\}}.$$ 

The total number of infected individuals in the epidemic is

$$Y = \min \left\{ k \geq 0 ; \; Q_{(k+1)} > \frac{pc}{N} \sum_{i=0}^{k} \Delta T_i \right\}$$

$$= \min \left\{ k \geq 0 ; \; Q_{(k+1)} > \mathcal{J}(k+1) \right\}$$

$$= \min \left\{ w \geq 0 ; \; Q(\mathcal{J}(w+1)) = w \right\}.$$ 

Suppose indeed that $Y = i$. Then according to (1),

$$\mathcal{J}(j) > Q(j), \text{ hence } Q(\mathcal{J}(j)) \geq j, \; \forall j \leq i,$n

and $\mathcal{J}(i+1) < Q_{(i+1)}$ hence $Q(\mathcal{J}(i+1)) < i + 1$.

In other words $Y = i$ iff $i$ is the smallest integer such that

$$Q(\mathcal{J}(i+1)) < i + 1, \; \text{ hence } = i.$$

### 10.1 Law of Large Numbers

Let us index $\mathcal{J}$ and $Q$ by $N$, the population size, so that they become $\mathcal{J}_N$ and $Q_N$. We now define

$$\overline{\mathcal{J}}_N(w) = \mathcal{J}_N(Nw)$$

$$\overline{Q}_N(v) = \frac{Q_N(v)}{N}.$$ 

As $N \to \infty$,

$$\overline{\mathcal{J}}_N(w) \to cpE(\Delta T)w = R_0w, \; \text{ and}$$

$$\overline{Q}_N(v) \to 1 - e^{-v} \; \text{ a.s.}$$

Hence

$$\overline{Q}_N \circ \overline{\mathcal{J}}_N(w) \to 1 - e^{-R_0w}.$$
We have

\[
\frac{Y_N}{N} = \min \left\{ \frac{w}{N} \geq 0; \ Q_N(J_N(w + 1)) = w \right\}
\]

\[
= \min \left\{ s \geq 0; \frac{1}{N} Q_N \left( J_N \left( N \left( s + \frac{1}{N} \right) \right) \right) = s \right\}
\]

\[
= \min \left\{ s \geq 0; \frac{1}{N} Q_N \left( J_N \left( s + \frac{1}{N} \right) \right) = s \right\}.
\]

Then \(Y_N/N\) converges a.s. towards the smallest positive solution of equation

\[
1 - e^{-R_0x} = x.
\]

- If \(R_0 \leq 1\), the unique solution of this equation is \(x = 0\).
- If \(R_0 > 1\), there is another solution \(x > 0\).

This solution \(0 < x < 1\) is the size (measured as the proportion of the total population) of a “significant ” epidemic, if it goes off, which happens with probability \(1 - z_\infty\).

10.2 Central Limit Theorem

From the classical CLT, as \(N \to \infty\),

\[
\sqrt{N}(J_N(w) - R_0w) = \frac{p_c\sqrt{w} \sum_{i=1}^{[Nw]} [\Delta T_i - \mathbb{E}(\Delta T_i)]}{\sqrt{Nw}} \Rightarrow A(w),
\]

where \(A(w) \sim \mathcal{N}(0, p^2 c^2 \text{Var}(\Delta T)w)\). One can in fact show that, as processes

\[
\{\sqrt{N}(J_N(w) - R_0w), \ 0 \leq w \leq 1\} \Rightarrow \{A(w), \ 0 \leq w \leq 1\},
\]

where \(\{A(w), \ 0 \leq w \leq 1\}\) is a Brownian motion (i.e. a centered Gaussian process with independent increments and continuous trajectories) such that \(\text{Var}(A(w)) = r^2 R_0^2 w\), where \(r^2 = (\mathbb{E}\Delta T)^{-2}\text{Var}(\Delta T)\). It is easy to show that for all \(k \geq 1\), all \(0 < w_1 < \cdots < w_k \leq 1\), if we define \(A_N(w) := \sqrt{N}(J_N(w) - R_0w), (A_N(w_1), \ldots, A_N(w_k)) \Rightarrow (A(w_1), \ldots, A(w_k))\).
One can show convergence in a stronger sense, but describing that would force us to introduce more complicated mathematical notions.

Consider now \( Q_N \). Again from the usual CLT,
\[
B_N(v) = \sqrt{N}(Q_N(v) - [1 - e^{-v}])
\]
\[
= \frac{1}{\sqrt{N}} \sum_{i=1}^{N} [1_{\{Q_i \leq v\}} - (1 - e^{-v})]
\]
\[
\Rightarrow B(v),
\]
where \( B(v) \sim \mathcal{N}(0, e^{-v}(1 - e^{-v})) \). We have again a functional convergence, according to the Kolmogorov–Smirnov theorem, towards a time changed Brownian bridge. In simpler words, \( \{B(v), \ v \geq 0\} \) is a centered Gaussian process with continuous trajectories whose covariance is specified by the identity \( \mathbb{E}[B(u)B(v)] = e^{-uv} - e^{-(u+v)}. \)

Recall that the above Law of Large Numbers has been obtained by taking the limit in the equation
\[
Q_N(\mathcal{J}_N(s + N^{-1})) = s.
\]
Making use of the two above CLTs, we get
\[
s = 1 - e^{-\mathcal{J}_N(s+ N^{-1})} + N^{-1/2}B_N(\mathcal{J}_N(s + N^{-1}))
\]
\[
= 1 - \exp\left(-R_0(s + N^{-1}) + N^{-1/2}A_N(s + N^{-1})\right)
\]
\[
+ N^{-1/2}B_N \left(R_0(s + N^{-1}) + N^{-1/2}A_N(s + N^{-1})\right).
\]
Let \( s = s^* + s_NN^{-1/2} + o(N^{-1/2}) \), where \( s^* \) satisfies \( e^{-R_0s^*} = 1 - s^* \). We obtain
\[
s^* + s_NN^{-1/2} + o(N^{-1/2})
\]
\[
= 1 - \exp\left(-R_0s^* - R_0s_NN^{-1/2} - A_N(s^*)N^{-1/2} + o(N^{-1/2})\right)
\]
\[
+ N^{-1/2}B_N(R_0s^*) + o(N^{-1/2})
\]
\[
= 1 - e^{-R_0s^*} + N^{-1/2}e^{-R_0s^*}(R_0s_N + A_N(s^*))
\]
\[
+ N^{-1/2}B_N(R_0s^*) + o(N^{-1/2}).
\]
We simplify this relation by making use of the equation which specifies \( s^* \). Multiplying the remaining terms by \( N^{1/2} \), we deduce
\[
[1 - (1 - s^*)R_0]s_N = B_N(R_0s^*) + (1 - s^*)A_N(s^*).
\]
Hence $s_N \Rightarrow \Xi$, where

$$\Xi \sim \mathcal{N}\left(0, \frac{s^*(1-s^*)}{(1-(1-s^*)R_0)^2} \left(1 + r^2(1-s^*)R_0^2\right)\right).$$

Finally $Y_N$ follows asymptotically the distribution

$$\mathcal{N}\left(Ns^*, N\frac{s^*(1-s^*)}{(1-(1-s^*)R_0)^2} \left(1 + r^2(1-s^*)R_0^2\right)\right).$$

11 Partially vaccinated population

Suppose that a fraction $v$ of the total population is vaccinated (with a vaccine with 100% efficiency).

The initial population of susceptibles is $N(1-v)$, instead of $N$, since $Nv$ individuals have been vaccinated and are immune.

The basic reproduction number is modified. The mean number of individuals whom an infectious individual infects during the initial phase of the epidemic is no longer $R_0 = cp\mathbb{E}[\Delta T]$, but rather

$$R_v = (1-v)R_0 = (1-v)cp\mathbb{E}[\Delta T].$$

If $R_v \leq 1$, there is no chance of a major outbreak. This inequality is equivalent to

$$v \geq 1 - \frac{1}{R_0}.$$ 

The right hand side of this inequality is the critical vaccination coverage.

Exercise 2. Suppose that the vaccine does not produce a 100% immunity, but that the probability that the encounter of an infectious and a vaccinated individual results in an infection with probability $p_v < p$, where $p$ is the probability that a susceptible be infected after an encounter with an infectious. Compute the corresponding basic reproduction number $R_v$.

12 Duration of a major epidemic

Denote by $T_N$ the duration of a major epidemic. $T_N$ takes the form

$$T_N = c_1 \log N + c_2 + c_3 \log N + X,$$
where the term $c_1 \log N$ is the duration of the initial phase, $c_2$ is the duration of the intermediate phase, which is essentially independent of the size $N$ of the population, $c_3 \log N$ is the duration of the final phase (similar to the initial phase), and $X$ is a random term, which takes into account mainly the random aspects of the initial and final phases.

### 13 Time needed by an epidemic to go extinct, and the critical population size

Let us go back to the general stochastic model, with a demographic effect.

Individuals are born at the constant rate $\mu N$, each individual lives a time $\text{Exp}(\mu)$. An individual contacts another individual of the population at rate $\gamma = cp$, and this contact results in an infection if one of the two individuals is infectious, and the other one is susceptible. We assume homogeneous mixing: any infectious infects any given susceptible at rate $\gamma/N$. The durations of the infection periods are i.i.d., with the common law $\text{Exp}(\alpha)$.

Hence $S(t)$ is a birth and death process, with births at rate $N\mu$ and deaths at rate $S(t)\mu$. The death rate is the same for susceptible and infectious individuals.

The epidemic starts with a unique infected individual which is introduced in a population of susceptibles. Its stops when there is no more infected individual. The time when this happens is called the extinction time of the epidemic. Note that we have made two simplifying assumptions in our model

1. the rate of births is not exactly proportional to the size of the population, but to its equilibrium value $N$;

2. the rate of contacts of an infectious with a given individual is $\gamma/N$, and not $\gamma$ divided by the exact population size.

Those simplifications allow us in particular to reduce our model to a 2-dimensional model.

Concerning the increase of the death rate due to the epidemic, it can be included in the rate $\alpha$ (a death caused by the infection is considered as a removed).

In our model, the pair $(S(t), I(t))$ is a continuous time Markov process.
Its rates matrix $Q$ satisfies

$$Q_{(n,m),(n',m')} = \begin{cases} 
\mu N, & \text{si } (n',m') = (n+1,m); \\
\mu n, & \text{si } (n',m') = (n-1,m); \\
\frac{\gamma}{N}nm, & \text{si } (n',m') = (n,m+1); \\
(\alpha + \mu)m, & \text{si } (n',m') = (n,m-1); \\
0, & \text{si } (n',m') \notin \{(n,m)(n+1,m)(n-1,m)(n-1,m+1)(n,m-1)\}.
\end{cases}$$

Let

$$(\mathcal{S}(t), \mathcal{I}(t)) = \left( \frac{S(t)}{N}, \frac{I(t)}{N} \right).$$

The mean infection time of an individual is $(\alpha + \mu)^{-1}$. During that period, and during the initial phase of the epidemic, an infectious infects at rate $\gamma$. Then

$$R_0 = \frac{\gamma}{\alpha + \mu}.$$

**Example 8. Realistic values of the parameters.** We may assume that $\mu^{-1} = 75$ years (mean life time of an individual). Hence $\mu = 1/75$.

The mean duration of infection depends upon the illness, say it is one week, hence $1/\alpha = 1/52$, $\alpha = 52$.

For most childhood diseases, a typical value of $R_0$ is 10. In other words

$$\gamma = R_0(\mu + \alpha) \sim 500.$$

Note that in that case, $\mu + \alpha \sim \alpha$, hence $R_0 \sim \gamma/\alpha$.

Assume that $R_0 > 1$, and we are interested in the duration of a major epidemic. As $N \to \infty$, $(\mathcal{S}(t), \mathcal{I}(t)) \to (s(t), i(t))$, solution of the ODE

$$\begin{cases} 
s'(t) = \mu(1-s(t)) - \gamma s(t)i(t), \\
i'(t) = \gamma s(t)i(t) - (\mu + \alpha)i(t).
\end{cases}$$

This ODE has two equilibria : the disease free equilibrium $(1,0)$ and the endemic equilibrium $(\hat{s}, \hat{i})$, with $\hat{s} = R_0^{-1}$, $\hat{i} = (\mu + \alpha)^{-1}\mu(1 - R_0^{-1}) = \varepsilon(1 - R_0^{-1})$, where $\varepsilon = (\mu + \alpha)^{-1}\mu \sim \mu/\alpha$.

If $R_0 > 1$, the disease free equilibrium is unstable, while the endemic equilibrium is stable.

**Example 9.** With the above data, we have $\hat{s} = 0.1$, $\hat{i} = 0.00024$. 

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More generally, if $R_0 >> 1$ and $\varepsilon << 1$, $i$ is small. The ODE gives us a clue about the time needed to reach a value close to the equilibrium. When the equilibrium is reached, the small value of $i$ makes it easy for the random fluctuations to lead the epidemic to extinction.

1st case: when $i$ is close to $\hat{i}$, the random fluctuations lead rather quickly to extinction;

2d case: there is no quick extinction, and a second epidemic starts (which hits much less individuals than the first one, since $\hat{s} << s(0)$).

etc.

The law of the extinction time is multimodal, with a first peak near the time taken to reach $\hat{i}$, and a series of much smaller peaks.

Once $\hat{i}$ has been reached, we are in the situation of a quasi–stationary distribution. The extinction time follows an exponential distribution, by the same argument as above. Denote by $\{q_{S,I}, S \geq 0, I \geq 1\}$ this quasi–stationary distribution. The parameter of this exponential distribution equals $(\alpha + \mu) \sum_S q_{S,1}$, hence

$$\mathbb{E}(T_Q) = \frac{1}{(\alpha + \mu) \sum_S q_{S,1}} = \frac{1}{\mu \sum_S q_{S,1}} \varepsilon.$$ 

How can one compute $\sum_S q_{S,1}$? For this purpose we will approximate the quasi–stationary distribution by a Gaussian law which we now specify.

Let us go back to our model. If $(\overline{S}(t), \overline{I}(t)) = \left( \frac{S(t)}{N}, \frac{I(t)}{N} \right)$,

$$\overline{S}(t) = S(0) + \frac{1}{N} P_1(\mu N t) - \frac{1}{N} P_2 \left( \mu N \int_0^t \overline{S}(r) dr \right) - \frac{1}{N} P_3 \left( \gamma N \int_0^t \overline{S}(r) \overline{I}(r) dr \right)$$

$$\overline{I}(t) = I(0) + \frac{1}{N} P_3 \left( \gamma N \int_0^t \overline{S}(r) \overline{I}(r) dr \right) - \frac{1}{N} P_4 \left( (\alpha + \mu) N \int_0^t \overline{I}(r) dr \right).$$
Hence
\[
\bar{S}(t) = \bar{S}(0) + \mu t - \mu \int_0^t \bar{S}(r)dr - \gamma \int_0^t \bar{S}(r)i(r)dr \\
+ \frac{1}{N} M_1(\mu N t) - \frac{1}{N} M_2 \left( \mu N \int_0^t \bar{S}(r)dr \right) - \frac{1}{N} M_3 \left( \gamma N \int_0^t \bar{S}(r)i(r)dr \right)
\]
\[
\bar{I}(t) = \bar{I}(0) + \gamma \int_0^t \bar{S}(r)i(r)dr - (\alpha + \mu) \int_0^t \bar{I}(r)dr \\
+ \frac{1}{N} M_3 \left( \gamma N \int_0^t \bar{S}(r)i(r)dr \right) - \frac{1}{N} M_4 \left( (\alpha + \mu) N \int_0^t \bar{I}(r)dr \right).
\]

The law of large numbers tells us that in the limit \(N \to \infty\)
\[
s'(t) = \mu (1 - s(t)) - \gamma s(t)i(t) \\
i'(t) = \gamma s(t)i(t) - (\alpha + \mu)i(t).
\]

Define
\[
\begin{pmatrix}
U_t \\
V_t
\end{pmatrix} = \lim_{N \to \infty} \sqrt{N} \begin{pmatrix}
\bar{S}(t) - s(t) \\
\bar{I}(t) - i(t)
\end{pmatrix}.
\]

The process \(\begin{pmatrix} U_t \\ V_t \end{pmatrix}\) satisfies
\[
d\begin{pmatrix} U_t \\ V_t \end{pmatrix} = \begin{pmatrix}
-\mu - \gamma i(t) & -\gamma s(t) \\
\gamma i(t) & \gamma s(t) - (\alpha + \mu)
\end{pmatrix} \begin{pmatrix} U_t \\ V_t \end{pmatrix} dt + dM_t,
\]
where \(M_t\) is a Gaussian martingale (in fact a Brownian motion) such that
\[
\mathbb{E}(M_t M'_t) = \begin{pmatrix}
\mu \int_0^t (1 + s(r))dr + \gamma \int_0^t s(r)i(r)dr \\
-\gamma \int_0^t s(r)i(r)dr
\end{pmatrix} \begin{pmatrix}
-\gamma \int_0^t s(r)i(r)dr \\
(\alpha + \mu) \int_0^t i(r)dr + \gamma \int_0^t s(r)i(r)dr
\end{pmatrix}.
\]

Since we consider our processes for large time, we can replace \(s(r)\) et \(i(r)\) by \(\hat{s}\) and \(\hat{i}\). The above system becomes
\[
d\begin{pmatrix} U_t \\ V_t \end{pmatrix} = \begin{pmatrix}
-\mu R_0 & -(\alpha + \mu) \\
\mu (R_0 - 1) & 0
\end{pmatrix} \begin{pmatrix} U_t \\ V_t \end{pmatrix} dt + dM_t,
\]
\[
\mathbb{E}(M_t M'_t) = t \begin{pmatrix}
2\mu & -\mu \left(1 - \frac{1}{R_0}\right) \\
-\mu \left(1 - \frac{1}{R_0}\right) & 2\mu \left(1 - \frac{1}{R_0}\right)
\end{pmatrix} = t\Lambda.
\]
Let
\[ A = \begin{pmatrix} -\mu R_0 & -\alpha - \mu \\ \mu(R_0 - 1) & 0 \end{pmatrix}, \quad \mathcal{V} = \frac{1}{R_0^2} \begin{pmatrix} \frac{1}{\varepsilon} + R_0 & -R_0 \\ -R_0 & R_0 - 1 + \varepsilon R_0^2 \end{pmatrix}. \]

One can show that
\[ \left( U_t \ V_t \right) \overset{\text{en loi}}{\rightarrow} \mathcal{N} \left( \begin{pmatrix} 0 \\ 0 \end{pmatrix}, \mathcal{V} \right). \]

This result follows from the identities:
\[ \frac{d}{dt} \mathbb{E} \left( U_t \ V_t \right) = A \mathbb{E} \left( U_t \ V_t \right) = 0, \]
\[ \frac{d}{dt} \text{Cov} \left( U_t \ V_t \right) = A \text{Cov} \left( U_t \ V_t \right) + \text{Cov} \left( U_t \ V_t \right) A' + \Lambda. \]

Hence \( \mathcal{V} \) is such that \( A \mathcal{V} + \mathcal{V} A' = \Lambda \). \( I \) is asymptotically Gaussian, with mean \( \mu_I = N \hat{i} \), and variance \( \sigma_I^2 = \frac{N}{R_0 - 1 + \varepsilon R_0^2}. \) We want to estimate
\[ q_{s,1} = \sum_S q_{S,1} = \lim_{t \to \infty} \mathbb{P}(I(t) = 1 | I(t) > 0), \]
which we approximate by
\[ \frac{1}{\sigma_I} \varphi \left( \frac{1 - \mu_I}{\sigma_I} \right) \left( \frac{1 - \mu_I}{\sigma_I} \right), \]
where \( \varphi \) (resp. \( \Phi \)) denotes the density (resp. the distribution function) of the law \( \mathcal{N}(0, 1) \). Note that we have approximated \( \mathbb{P}(I(t) > 0) \) by \( \mathbb{P}(I(t) > 0, 5) = 1 - \Phi(\sigma^{-1}(0.5 - \mu_I)) = \Phi(\sigma^{-1}(\mu_I - 0.5)). \) The above ratio is approximated by
\[ \frac{1}{\sigma_I} \varphi \left( \frac{\mu_I}{\sigma_I} \right) \left( \frac{\mu_I}{\sigma_I} \right), \]
Note that
\[ \frac{\mu_I}{\sigma_I} = \frac{\sqrt{N \varepsilon(R_0 - 1)}}{\sqrt{R_0 - 1 + \varepsilon R_0^2}} \simeq \sqrt{N \varepsilon^2(R_0 - 1)}. \]
Hence
\[ \mathbb{E}(T_Q) \simeq \frac{1}{\mu} \frac{\varepsilon q_{1,1}}{\Phi(\sqrt{N\varepsilon^2(R_0 - 1)})} \]
\[ \simeq \frac{1}{\mu} \frac{\varepsilon \Phi(\sqrt{N\varepsilon^2(R_0 - 1)})}{\sqrt{2\pi N(R_0 - 1)}} \exp\left(-N\varepsilon^2(R_0 - 1)/2\right) \]
\[ \simeq \frac{\sqrt{2\pi}}{\mu} \frac{\sqrt{N\varepsilon^2(R_0 - 1)}}{R_0} e^{N\varepsilon^2(R_0 - 1)/2} \Phi(\sqrt{N\varepsilon^2(R_0 - 1)}) \]
\[ \sim \exp\left(\frac{N\varepsilon^2(R_0 - 1)}{2}\right). \]

The last line gives the order of magnitude of the quantity of interest.

In case of vaccination of the proportion \( v \) of the population, we must replace \( N \) by \( N(1 - v) \) and \( R_0 \) by \( R_0(1 - v) \).

**Critical population size**  If \( N\varepsilon^2 \) is “small”, there will be no major epidemic.

If \( N\varepsilon^2 \) is “large”, a major epidemic might happen.

Note that before vaccination, measles was endemic in Great Britain, while the epidemic would go extinct after each outbreak in Iceland.

Notice that if \( Z \sim \mathcal{N}(0, 1) \), \( \operatorname{IP}(|Z| \geq 3) \simeq 0.23\% \). But
\[ \frac{\mu_I}{\sigma_I} \simeq \sqrt{N\varepsilon^2(R_0 - 1)}. \]

One may decide that the critical value \( N_c \) is such that (the factor 3 below is arbitrary)
\[ \sqrt{N_c\varepsilon^2(R_0 - 1)} = 3 \]
\[ N_c = \frac{9}{\varepsilon^2(R_0 - 1)}. \]

If we take into account vaccination, we get
\[ N_c = \frac{9}{(1 - v)\varepsilon^2(R_0(1 - v) - 1)}. \]

**Example 10.** In case of measles, \( R_0 = 15, \varepsilon = 1/3750, N_c = 9.04 \times 10^6 \).

**Example 11.** Measles with a vaccination rate \( v = 0.9 \). Then \( R_0 - 1 = 14 \) should be replaced by \( (1 - v)(R_0(1 - v) - 1) = 0.05 \). Hence we must multiply the above value of \( N_c \) by the factor 14/0.05. The result is \( 2531 \times 10^6 \).
14 Computation of $R_0$

14.1 The Perron–Frobenius theorem

The proof of the following classical result can be found e.g. in [8], see the proof of Theorem 2.7 there.

**Theorem 12.** Let $A$ be a square matrix, which is positive and irreducible. The spectral radius of $A$ is a simple eigenvalue of $A$, whose associated eigenvector has all its coordinates positive.

If moreover $A$ is aperiodic, then any other eigenvalue has a modulus which is strictly smaller than the spectral radius.

A positive: $A_{i,j} \geq 0$ for all $i, j$.

A irreducible: for all $i \neq j$, there exists $n \geq 1$ such that $(A^n)_{i,j} > 0$.

A aperiodic: there exists $n \geq 1$ such that $(A^n)_{i,j} > 0$ for all $i, j$.

The spectral radius of $A$ is defined as the $\lim_{n \to \infty} \|A^n\|^{1/n}$ (it is also the largest modulus of all eigenvalues).

14.2 Computation of $R_0$ in case of malaria

Female mosquitos strive for a fixed number of blood meals per unit time. Hence the mean number of stings which a human suffers by unit time (from female mosquitos) is proportional to the ratio of the two densities $D_{\text{mosquitos}}/D_{\text{humans}}$.

Consider an infected mosquito. Suppose that her mean infection period is $T_m$, during which she stings humans at rate $c$. Each of those stings results in the transmission of malaria with probability $p_m$. The mean number of individuals which a mosquito infects is then $cp_mT_m$.

Consider an infected human. Suppose that his mean infection time is $T_h$, during which he is stung at rate $k$, each sting resulting in the infection of the mosquito (if susceptible) with probability $p_h$. The mean number of mosquitos infected by such a human is then $kp_hT_h$.

Consequently

$$R_0 = c^2T_mT_hp_mp_h \frac{D_{\text{mosquitos}}}{D_{\text{humans}}}.$$
14.3 Computation of $R_0$ in case of a sexually transmitted illness

Suppose that the mean number of women infected by one man is 100, and the mean number of men infected by a women is 10. Hence the transmission matrix

$$K = \begin{pmatrix} 0 & 100 \\ 10 & 0 \end{pmatrix}.$$ 

In two “generations”, the mean number of individuals of the same species infected by a given individual is 1 000. Hence $R_0 = \sqrt{1000} =$ spectral radius of $K =$ largest eigenvalue.

14.4 Computation of $R_0$ in case of a general compartmental model

Consider the ODE of the model, which we linearize in the neighborhood of $I(t) = 0$, since $R_0$ is the potential of infection of one infected individual, introduced in a fully susceptible population.

Let $x$ be the vector which describes the sizes of the infected subpopulations.

$$\dot{x} = (T + \Sigma)x,$$

where $T$ is the matrix which describes the transmission of the illness by contact; $\Sigma$ is the matrix which describes the transitions from one state to another, without contact.

**Example 13.** Suppose there are two latent states $E_1$ and $E_2$, which change to infected resp. at rates $\nu_1$ and $\nu_2$, and those infected individuals produce by contact $E_1$–type individuals at rate $p\beta$, $E_2$–type individuals at rate $(1 - p)\beta$. In addition, each individual dies at rate $\mu$, and the infected individuals heal at rate $\alpha$. In other words

$$\dot{E}_1 = p\beta I - (\nu_1 + \mu)E_1,$$

$$\dot{E}_2 = (1 - p)\beta I - (\nu_2 + \mu)E_2,$$

$$\dot{I} = \nu_1 E_1 + \nu_2 E_2 - (\alpha + \mu)I.$$

Here

$$T = \begin{pmatrix} 0 & 0 & p\beta \\ 0 & 0 & (1 - p)\beta \\ 0 & 0 & 0 \end{pmatrix}, \quad \Sigma = \begin{pmatrix} -(\nu_1 + \mu) & 0 & 0 \\ 0 & -(\nu_2 + \mu) & 0 \\ \nu_1 & \nu_2 & -(... \end{pmatrix}. $$

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Digression about the matrix $\Sigma$  Consider a jump Markov process, with the four 4 states ($E_1, E_2, T, R$), where

\[
E_1 = \text{latent state 1} \\
E_2 = \text{latent state 2} \\
I = \text{infectious} \\
R = \text{removed or dead}.
\]

The rates matrix of this process is given by

\[
Q = \begin{pmatrix}
-(\nu_1 + \mu) & 0 & \nu_1 & \mu \\
0 & -(\nu_2 + \mu) & \nu_2 & \mu \\
0 & 0 & -(\alpha + \mu) & \alpha + \mu \\
0 & 0 & 0 & 0
\end{pmatrix}
\]

The state $R$ is absorbing. Let

\[
\Sigma = Q^t = \begin{pmatrix}
-(\nu_1 + \mu) & 0 & 0 & 0 \\
0 & -(\nu_2 + \mu) & 0 & 0 \\
\nu_1 & \nu_2 & -(\alpha + \mu) & 0 \\
\mu & \mu & \alpha + \mu & 0
\end{pmatrix}
\]

Note that $(e^{t\Sigma})_{ij} = (e^{tQ})_{ji}$ is the probability to be in state $i$ at time $t$, starting from state $j$ at time 0.

In fact, if we limit ourselves to $1 \leq i, j \leq 3$, this quantity equals also

\[
(e^{t\Sigma})_{ij}, \text{ with } \Sigma = \begin{pmatrix}
-(\nu_1 + \mu) & 0 & 0 \\
0 & -(\nu_2 + \mu) & 0 \\
\nu_1 & \nu_2 & -(\alpha + \mu)
\end{pmatrix}
\]

Now for all $1 \leq i, j \leq 3$, $(e^{t\Sigma})_{ij} \to 0$ as $t \to \infty$. We have

\[
\int_0^\infty (e^{t\Sigma})_{ij} dt = \text{ mean sojourn time in state } i, \text{ starting from state } j \text{ at time 0.}
\]

But since $e^{\infty \Sigma} = 0$,

\[
\int_0^\infty \Sigma e^{t\Sigma} dt = -I,
\]

and

\[
\int_0^\infty e^{t\Sigma} dt = -\Sigma^{-1}
\]
which is a matrix with positive coefficients. Note that

\[ T_{ij} = \text{rate at which a } j \text{ produces an } i, \]

\[ (-\Sigma^{-1})_{jk} = \text{mean sojourn time of an initial } k \text{ in state } j \text{ since} \]

\[ \left( \int_0^\infty e^{t\Sigma} dt \right)_{jk} = \mathbb{E} \left( \int_0^\infty 1_{X(t)=j} dt \mid X(0) = k \right), \]

\[ (T(-\Sigma^{-1}))_{ik} = \text{mean number of } i \text{'s that an initial } k \text{ induces during his life time.} \]

Hence

\[ R_0 \text{ is the spectral radius of the matrix } K = -T\Sigma^{-1}. \]

**Back to the above example**

\[-\Sigma^{-1} = \begin{pmatrix}
\frac{1}{\nu_1+\mu} & 0 & 0 \\
0 & \frac{1}{\nu_2+\mu} & 0 \\
\frac{\nu_1}{(\nu_1+\mu)(\alpha+\mu)} & \frac{\nu_2}{(\nu_2+\mu)(\alpha+\mu)} & \frac{1}{\alpha+\mu}
\end{pmatrix}, \]

\[ K = \begin{pmatrix}
\frac{p\beta\nu_1}{(\nu_1+\mu)(\alpha+\mu)(1-p)\beta^2} & \frac{p\beta\nu_2}{(\nu_2+\mu)(\alpha+\mu)(1-p)\beta^2} & \frac{p\beta}{(\alpha+\mu)(1-p)\beta} \\
\frac{\nu_1}{(\nu_1+\mu)(\alpha+\mu)} & \frac{\nu_2}{(\nu_2+\mu)(\alpha+\mu)} & 0 \\
0 & 0 & 0
\end{pmatrix}. \]

**Exercise 3.** The two matrices \( K \) and \( K' \) below give the same \( R_0 \)

\[ K' = \begin{pmatrix}
\frac{p\beta\nu_1}{(\nu_1+\mu)(\alpha+\mu)(1-p)\beta^2} & \frac{p\beta\nu_2}{(\nu_2+\mu)(\alpha+\mu)(1-p)\beta^2} \\
\frac{\nu_1}{(\nu_1+\mu)(\alpha+\mu)} & \frac{\nu_2}{(\nu_2+\mu)(\alpha+\mu)} & 0 \\
0 & 0 & 0
\end{pmatrix}, \]

namely

\[ R_0 = \left( \frac{p\nu_1}{\nu_1+\mu} + \frac{(1-p)\nu_2}{\nu_2+\mu} \right) \frac{\beta}{\alpha+\mu} = TrK'. \]

There is a good reason to restrict oneself to the two states \( E_1 \) and \( E_2 \), which are the states of the “start of infection”.

On the other hand, if we consider the unique states \( I \), and compute the mean number of \( I \)'s which a unique \( I \) induces, we recover the same \( R_0 \), without matrix or eigenvalue!

**Exercise 4.** Recover the above \( K_{ij} \) by the interpretation “mean number of \( i \)'s produced by a \( j \)”. 

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Exercise 5. Computation of $R_0$ in a model of infection of cells by viruses

We consider the following model, where $C$ denotes the number of target cells, $C^*$ the number of infected cells, $V$ the number of viruses. We assume that this triplet satisfies the ODE

\[
\begin{align*}
\frac{dC}{dt} &= \lambda - kCV - \delta C \\
\frac{dC^*}{dt} &= kCV - (\mu + \delta)C^* \\
\frac{dV}{dt} &= pC^* - kCV - cV.
\end{align*}
\]

We consider the ODE reduced to the two last equations, which we want to put in the form

\[
\frac{d}{dt} \begin{pmatrix} C^* \\ V \end{pmatrix} = \begin{bmatrix} T + \Sigma \end{bmatrix} \begin{pmatrix} C^* \\ V \end{pmatrix},
\]

where $T$ and $-\Sigma^{-1}$ are positive matrices, such that $R_0$ is the spectral radius of the matrix $-T\Sigma^{-1}$. We consider the three following choices for the matrices $T$ and $\Sigma$:

\[
\begin{align*}
T_1 &= \begin{pmatrix} 0 & 0 \\ p & 0 \end{pmatrix}, & \Sigma_1 &= \begin{pmatrix} -\left(\mu + \delta\right) & kC \\ 0 & -(kC + c) \end{pmatrix}; \\
T_2 &= \begin{pmatrix} 0 & kC \\ p & 0 \end{pmatrix}, & \Sigma_2 &= \begin{pmatrix} -\left(\mu + \delta\right) & 0 \\ 0 & -(kC + c) \end{pmatrix}; \\
T_3 &= \begin{pmatrix} 0 & kC \\ 0 & 0 \end{pmatrix}, & \Sigma_3 &= \begin{pmatrix} -\left(\mu + \delta\right) & 0 \\ p & -(kC + c) \end{pmatrix}.
\end{align*}
\]

1. Justify the three decompositions, from the point of view of the biological interpretation (the transition from 1. to 2., a virus enters a target cell, is it a state transition of the virus, or a reproduction event ?).

2. Compute $R_0$ in the three cases.

3. Compare the three results. What do they have in common ? Can you explain the differences and the identities ?

Exercise 6. Flowers are cultivated in a field. Slips are withdrawn from these flowers at rate $\delta$. Those slips are planted in a greenhouse.
Flowers which mature in the greenhouse are replanted in the field at rate $\rho$. We admit that the system is at its equilibrium, with population sizes $N_1$ in the field, and $N_2$ in the greenhouse.

Mushrooms start to proliferate among the flowers, with the following transmission mechanism. Each infected flower in the field (resp. in the greenhouse) infects each non infected flower in the field (resp. in the greenhouse) at rate $\alpha$ (resp. $\beta$). Moreover, each time a slip is removed from the field, it is infected with probability $p$.

The infected flowers of the field (resp. of the greenhouse) die at rate $\mu$ (resp. $\gamma$). Since we are interested in the beginning of the infection, we neglect the fact that the sizes of the populations will not remain constant.

1. We denote by $C_1(t)$ (resp. $C_2(t)$) the size of the field (resp. greenhouse) population of flowers hit by the mushroom. Justify the following model for the evolution of ($C_1(t), C_2(t)$):

$$
\frac{dC_1}{dt}(t) = \alpha(N_1 - C_1(t))C_1(t) - \mu C_1(t) + \rho C_2(t)
$$
$$
\frac{dC_2}{dt}(t) = p\delta C_1(t) + \beta(N_2 - C_2(t))C_2(t) - (\gamma + \rho)C_2(t).
$$

2. Linearise the above ODE in the neighborhood of $(C_1, C_2) = (0, 0)$.

Write the linearized system in the form

$$
\frac{d}{dt} \begin{pmatrix} C_1 \\ C_2 \end{pmatrix} = \left[ T + \Sigma \right] \begin{pmatrix} C_1 \\ C_2 \end{pmatrix}.
$$

3. Compute the matrix $K = T(-\Sigma^{-1})$ and its spectral radius (do not try to simplified the formula).

4. Compute $R_0$ for this model.

Example 14. The virus of bovine diarrhea

Assume “horizontal” transmission at rate $\beta_1$, “vertical” transmission at rate $\beta_2$. Those horizontally infected go through state $E$ before going to state $I$.

A cow which was pregnant during less than 150 days before being infected, and which heals (and then is in state $Z$) before delivering, gives birth to a veal in state $P$. Those transmit infection, reproduce, at rates different from the others.
Let $\gamma$ denote the healing rate, $p_1$ the probability of ending in state $Z$, $1/\alpha$ the mean time during which an infected fetus is weared, $p_2$ the probability that an infected fetus survives, $\nu$ the transition rate from state $E$ to state $I$, $\mu$ the natural death rate, $a$ and $b$ the reduction of the reproduction rate and the increase of the death rate of infected individuals. The model reads

\[
\begin{align*}
\dot{E} &= (\beta_1 I + \beta_2 P)S - (\nu + \mu)E \\
\dot{I} &= \nu E - (\gamma + \mu)I \\
\dot{Z} &= p_1 \gamma I - (\alpha + \mu)Z \\
\dot{P} &= p_2 \alpha Z + (\mu - a)P - (\mu + b)P
\end{align*}
\]

We have

\[
\begin{align*}
k_{12} &= \frac{\beta_2}{\mu + b} \quad \text{mean number of } E \text{'s produced by a } P \\
k_{22} &= \frac{\mu - a}{\mu + b} \\
k_{11} &= \frac{\nu}{\mu + \nu} \times \frac{\beta_1}{\gamma + \mu} \\
k_{21} &= \text{mean number of } P \text{'s produced by an } E \leq 1.
\end{align*}
\]

Exercise 7. 1. Suppose in the above example that $\beta_1 = 0$. Show that in this case

\[
R_0 = \frac{1}{2} B + \frac{1}{2} \sqrt{B^2 + 4A}, \quad \text{with}
\]

\[
B = \frac{\mu - a}{\mu + b}, \quad A = \frac{\nu}{\mu + \nu} \frac{\gamma p_1}{\mu + \gamma} \frac{\alpha p_2}{\alpha + \mu} \frac{\beta_2}{\mu + b}.
\]

2. Show that $A + B \neq R_0$, but $R_0 > 1 \iff A + B > 1$.

Exercise 8. Computation of $R_0$ in Ross’ malaria model We consider the following ODE which was proposed by Ross as a model of the transmission of malaria.

\[
(*) \quad \begin{cases}
\frac{dx}{dt} = mab_1 y(1 - x) - \gamma x \\
\frac{dy}{dt} = b_2 a(1 - y)x - \mu y,
\end{cases}
\]
where $x$ (resp. $y$) is the proportion of infected individuals among humans (resp. among female anopheles – those mosquitoes which are susceptible of transmitting malaria – below we shall write mosquito instead of female anopheles). We assume that the size $H$ of the human population is constant, as well as that $V$ of mosquitoes. The parameter $m = V/H$ is the so-called “density of mosquitoes” (one says also “vectors”, since the mosquitoes are the vectors of malaria). $a$ is the mean number of stings done by a mosquito per time unit. $b_1$ is the probability that a sting of a susceptible human by an infected mosquito transmits the parasite, $b_2$ the probability that a susceptible mosquito gets infected while stinging an infected human. Mosquitos die at rate $\mu$, infected humans heal at rate $\gamma$.

1. Compute the mean number of susceptible humans which a given infected mosquito (call it $z$) infects during her life, and the mean number of infections which she generates, assuming that all humans are susceptible.

2. Assuming that almost all mosquitos are susceptible, what is the mean number of mosquitos which the humans infected by the mosquito $z$ will infect, before healing?

3. Write the linearized (in the neighborhood of $(x,y) = (0,0)$) version of equation (*)

4. Write the linearized equation in the form

$$\frac{d}{dt} \begin{pmatrix} x \\ y \end{pmatrix} = \begin{bmatrix} T + \Sigma \end{bmatrix} \begin{pmatrix} x \\ y \end{pmatrix}.$$

5. Compute the inverse matrix $\Sigma^{-1}$.

6. Compute $R_0$. Compare with the result of question 2. Discuss the notion of “generation” in the case of malaria.

**14.5 Computation of $R_0$ in case of an epidemic on a graph**

We consider an epidemic on a configuration graph. A graph $\mathcal{G}$ is a pair $(\mathcal{V}, \mathcal{E})$, where $\mathcal{V}$ is the set of vertices, and $\mathcal{E}$ the set of edges. An edge $e \in \mathcal{E}$ is a pair $(v, w)$ of vertices, i.e. $v, w \in \mathcal{V}$. Our graphs will be non oriented, which
means that \((v, w) \in E\) iff \((w, v) \in E\). Given a vertex \(v \in V\), we say that another vertex \(w\) is a neighbor of \(v\) if \((v, w) \in E\). The individuals in the population constitute the vertices of the graph, and the edges represent the social relations between individuals.

An epidemic on a graph is characterized by the fact that an infected individual might infect his neighbors on the graph, and only them.

We consider the case of an epidemic on a configuration graph. Such a graph is constructed as follows. To each vertex is associated a degree, which is an integer. The degrees of various individuals are i.i.d., with the common law \((p_k, k \geq 1)\). The degree of a vertex is the number of edges which connect this vertex to other vertices of the graph. At each vertex are attached a number of half edges equal to its degree. We construct the graph by connecting each half-edge randomly to another half-edge.

We assume that each infected individual remains infectious during a time which is exponential with parameter \(\rho\). During his infection period, each half-edge attached to the corresponding vertex transmits infection at rate \(\lambda\). This implies that, conditionally upon the fact that the infected individual is located on a vertex with degree \(k\), and that his infection time is \(y\), the number of susceptibles which this individual infects follows the \(B(k - 1, 1 - e^{-\lambda y})\) distribution. Hence the mean number of such infected, conditionally upon \(k\) and \(y\), is \((k - 1)(1 - e^{-\lambda y})\). The infection time is exponential with parameter \(\rho\). What is the law of the degree \(k\)?

The first infected individual infects an individual with degree \(k\) with probability \((\sum_{\ell \geq 1} \ell p_{\ell})^{-1}k p_{k}\). Hence if \(\xi\) is an integer–valued r.v. with the law \((p_k, k \geq 1)\),

\[
R_0 = \frac{\sum_{k \geq 1}(k - 1)kp_k}{\sum_{k \geq 1}kp_k} \rho \int_0^\infty (1 - e^{-\lambda y})e^{-\rho y}dy \\
= \frac{\mathbb{E}[\xi(\xi - 1)\xi]}{\mathbb{E}[\xi]} \left(1 - \frac{\rho}{\lambda + \rho}\right) \\
= \frac{g''(1)}{g'(1)} \frac{\lambda}{\lambda + \rho},
\]

if \(g\) denotes the generating function of the r.v. \(\xi\).

Let us now compute the probability of extinction of the branching process which approximates the start of the epidemic. This is a fixed point of the generating function of the number of individuals which a typical infectious individual infects during the start of the epidemic, in other words the solution
of the equation
\[ z = \frac{\rho}{g'(1)} \sum_{k \geq 1} \int_{0}^{\infty} (z + e^{-\lambda y}(1 - z))^{k-1} k p_k e^{-\rho y} dy \]
\[ = \frac{\rho}{g'(1)} \int_{0}^{\infty} g' \left( z + e^{-\lambda y}(1 - z) \right) e^{-\rho y} dy. \]

15 Law of Large Numbers

Suppose there is no latency period \( T_1 = 0 \) a.s. and that \( \Delta T \sim \text{Exp}(\alpha) \). Consider a \( SIR \) model with constant population size equal to \( N \). Let \( S(t) \) denote the number of susceptibles at time \( t \), \( I(t) \) the number of infectious, \( R(t) \) the number of “removed” (i.e. “healed and immune”). We could add a transition from \( R \) to \( S \), and possibly suppress the compartment \( R \). This would produce the models \( SIRS \) and \( SIS \).

In our model, two types of events happen:

1. infection of a susceptible (such an event decreases \( S(t) \) by one, and increases \( I(t) \) by one); those events happen at rate
   \[ \frac{\beta}{N} S(t) I(t), \quad \text{where } \beta = cp; \]

2. recovery of an infectious (such an event decreases \( I(t) \) by one, and increases \( R(t) \) by one); those events happen at rate
   \[ \alpha I(t). \]

Hence the following equations, with \( P_1(t) \) and \( P_2(t) \) two standard mutually independent Poisson processes:

\[ S(t) = S(0) - P_1 \left( \frac{\beta}{N} \int_{0}^{t} S(s) I(s) ds \right), \]
\[ I(t) = I(0) + P_1 \left( \frac{\beta}{N} \int_{0}^{t} S(s) I(s) ds \right) - P_2 \left( \alpha \int_{0}^{t} I(s) ds \right), \]
\[ R(t) = R(0) + P_2 \left( \alpha \int_{0}^{t} I(s) ds \right). \]
Of course $S(t) + I(t) + R(t) = S(0) + I(0) + R(0) = N$. We can clearly forget the third equation. Let us define $s_N(t) = S(t)/N$, $i_N(t) = I(t)/N$. The equations for the proportions of susceptibles and infectious are written

$$s_N(t) = s_N(0) - \frac{1}{N} P_1 \left( \beta N \int_0^t s_N(r)i_N(r)dr \right),$$

$$i_N(t) = i_N(0) + \frac{1}{N} P_1 \left( \beta N \int_0^t s_N(r)i_N(r)dr \right) - \frac{1}{N} P_2 \left( \alpha N \int_0^t i_N(r)dr \right).$$

Define the two martingales $M_1(t) = P_1(t) - t$, $M_2(t) = P_2(t) - t$. We have

$$s_N(t) = s_N(0) - \beta \int_0^t s_N(r)i_N(r)dr - \frac{1}{N} M_1 \left( \beta N \int_0^t s_N(r)i_N(r)dr \right),$$

$$i_N(t) = i_N(0) + \beta \int_0^t s_N(r)i_N(r)dr - \alpha \int_0^t i_N(r)dr + \frac{1}{N} M_1 \left( \beta N \int_0^t s_N(r)i_N(r)dr \right) - \frac{1}{N} M_2 \left( \alpha N \int_0^t i_N(r)dr \right).$$

Consider the process

$$\mathcal{M}_N(t) := \frac{1}{N} M_1 \left( \beta N \int_0^t s_N(r)i_N(r)dr \right).$$

Let $\mathcal{F}_t = \sigma \{ s_N(r), i_N(r), 0 \leq r \leq t \}$.

**Lemma 15.** $\{ \mathcal{M}_N(t), t \geq 0 \}$ is a $\mathcal{F}_t$-martingale which satisfies

$$\mathbb{E} [ \mathcal{M}_N(t) ] = 0, \quad \mathbb{E} [ |\mathcal{M}_N(t)|^2 ] = \frac{\beta}{N} \mathbb{E} \int_0^t s_N(r)i_N(r)dr.$$

**Proof** The martingale property follows from the fact that for all $0 < r < t$

$$\mathbb{E} \left[ P_1 \left( \beta N \int_0^t s_N(u)i_N(u)du \right) - P_1 \left( \beta N \int_r^t s_N(u)i_N(u)du \right) \bigg| \mathcal{F}_r \right]$$

$$= \beta N \mathbb{E} \left[ \int_r^t s_N(u)i_N(u)du \bigg| \mathcal{F}_r \right].$$

We now establish that identity.
For \( n \geq 1, 0 \leq u \leq t \), let

\[
[u]_n = \begin{cases} 
  u, & \text{if } u \leq r; \\
  r + \frac{k}{n}(t-r), & \text{if } r + \frac{k}{n}(t-r) \leq u < r + \frac{k+1}{n}(t-r).
\end{cases}
\]

Let \( \mathcal{F}_r = \sigma \{ s_N(u), i_N(u), 0 \leq u \leq t' \} \), and

\[
A_n(t') = \beta N \int_0^{t'} s_N([u]_n)i_N([u]_n)du,
\]

\[
B_n(t') = \alpha N \int_0^{t'} i_N([u]_n)du.
\]

For \( a < b \), we denote \( P_i((a, b]) = P_i(b) - P_i(a) \), and \( u_k = r + \frac{k}{n}(t-r) \). Note that

\[
\mathbb{E} \left[ P_1(A_n(t)) - P_1(A_n(r)) \mid \mathcal{F}_r \right] = \mathbb{E} \left[ \sum_{k=1}^{n} P_1((A_n(u_{k-1}), A_n(u_k))] \mid \mathcal{F}_r \right]
\]

\[
= \mathbb{E} \left[ \sum_{k=1}^{n} \mathbb{E} \left( P_1((A_n(u_{k-1}), A_n(u_{k-1}) + s_N(u_{k-1})\frac{t-r}{n})) \mid \mathcal{F}_{u_{k-1}} \right) \mid \mathcal{F}_r \right]
\]

\[
= \mathbb{E} \left[ \sum_{k=1}^{n} s_N(u_{k-1})\frac{t-r}{n} \mid \mathcal{F}_r \right]
\]

\[
= \mathbb{E} \left[ A_n(t) - A_n(r) \mid \mathcal{F}_r \right].
\]

It remains to let \( n \to \infty \) in order to deduce (1).

The martingale property implies that \( \mathbb{E}M_N(t) = 0 \). Let us compute the expectation of the square. For \( n \geq 1 \) fixed, \( 0 \leq i \leq n \), let \( t_i = it/n \). We have

\[
\mathbb{E} \left[ M_N(t)^2 \right] = \mathbb{E} \sum_{i=0}^{n-1} \left| M_N(t_{i+1}) - M_N(t_i) \right|^2.
\]

As \( n \to \infty \),

\[
\sum_{i=0}^{n-1} \left| M_N(t_{i+1}) - M_N(t_i) \right|^2 \to \sum_{0 < r \leq t} \left| \Delta M_N(r) \right|^2 \text{ a.s.,}
\]

where the above sum is taken over all jump times of \( M_N(r) \), and \( \Delta M_N(r) \) denotes the jump of the process \( M_N \) at time \( r \). It is not too hard to deduce
from a uniform integrability argument that
\[
\mathbb{E} \left[ |\mathcal{M}_N(t)|^2 \right] = \mathbb{E} \sum_{0 < r \leq t} |\Delta \mathcal{M}_N(r)|^2.
\]
Indeed, as soon as \( t_{i+1} - t_i \leq 1 \), since \( 0 \leq s_N(r), i_N(r) \leq 1 \),
\[
|\mathcal{M}_N(t_{i+1}) - \mathcal{M}_N(t_i)|^2 \leq \frac{2}{N^2} |P_N(t_{i+1}) - P_N(t_i)|^2 + \beta^2 \left( \int_{t_i}^{t_{i+1}} s_N(r)i_N(r)dr \right)^2 \leq \frac{2}{N^2} |P_N(t_{i+1}) - P_N(t_i)|^2 + \beta^2 (t_{i+1} - t_i)
\]
\[
\sum_{i=0}^{n-1} |\mathcal{M}_N(t_{i+1}) - \mathcal{M}_N(t_i)|^2 \leq \frac{2}{N^2} |P_N(t)|^2 + \beta^2 t.
\]
But
\[
\sum_{0 < r \leq t} |\Delta \mathcal{M}_N(r)|^2 = \frac{1}{N^2} P_N \left( \beta N \int_0^t s_N(r)i_N(r)dr \right).
\]
The last formula of the statement follows from the martingale property of \( \mathcal{M}_N(r) \).
\[
\square
\]
Let \( \mathcal{N}_N(t) = \frac{1}{N} M_2 \left( \alpha N \int_0^t i_N(r)dr \right) \).
We show as in the above Lemma that \( \mathcal{N}_N(t) \) is a zero mean martingale, and such that
\[
\mathbb{E} \left[ |\mathcal{N}_N(t)|^2 \right] = \frac{\alpha}{N} \mathbb{E} \int_0^t i_N(r)dr.
\]
We deduce in particular from the above results that

**Corollary 16.** As \( N \to \infty \), for all \( T > 0 \),
\[
\sup_{0 \leq t \leq T} \{|\mathcal{M}_N(t)| + |\mathcal{N}_N(t)|\} \to 0
\]
in probability.

We have in fact the a.s. convergence to 0 of the same quantities !

**Proposition 17.** As \( N \to \infty \), for all \( T > 0 \),
\[
\sup_{0 \leq t \leq T} \{|\mathcal{M}_N(t)| + |\mathcal{N}_N(t)|\} \to 0 \text{ a.s.}
\]
Proof We consider the term $M_N$. Since the proportions $s_N(t)$ et $i_N(t)$ take values in the interval $[0, 1]$,

$$\sup_{0 \leq t \leq T} |M_N(t)| \leq \frac{1}{N} \sup_{0 \leq r \leq \beta NT} |M_1(r)|.$$ 

The Law of Large Numbers for Poisson processes (see below) tells us that for all $t > 0$,

$$\frac{P_1(Nt)}{N} \to t \text{ a.s. as } N \to \infty.$$ 

Note that we have pointwise convergence of a sequence of increasing functions towards a continuous (and of course increasing) function. Consequently from the second Dini Theorem, this convergence is uniform on any compact interval, hence for all $T > 0$,

$$\frac{1}{N} \sup_{0 \leq r \leq \beta NT} |M_1(r)| \to 0 \text{ a.s.}$$

We now prove the Law of Large Numbers for Poisson processes

**Proposition 18.** Let $\{P(t), t \geq 0\}$ be a rate $\lambda$ Poisson process. Then

$$t^{-1}P(t) \to \lambda \text{ a.s. as } t \to \infty.$$ 

**Proof** Consider first for $n \geq 1$

$$n^{-1}P(n) = n^{-1} \sum_{i=1}^{n} [P(i) - P(i-1)]$$

$$\to \lambda \text{ a.s. as } n \to \infty$$

from the standard Law of Large Numbers, since the r.v.’s $P(i) - P(i-1)$, $1 \leq i \leq n$ are i.i.d., Poisson with parameter $\lambda$. Now

$$t^{-1}P(t) = \frac{[t]}{t} [t]^{-1}P([t]) + t^{-1}\{P(t) - P([t])\}$$

$$|t^{-1}P(t) - \lambda| \leq \frac{[t]}{t} [t]^{-1}P([t]) - \lambda + t^{-1}\{P([t] + 1) - P([t])\}.$$ 

But

$$t^{-1}\{P([t] + 1) - P([t])\} = t^{-1}P([t] + 1) - t^{-1}P([t])$$
is the difference of two sequences which converge towards the same limit, hence it converges to 0 a.s. □

We can now prove

Theorem 19. Law of Large Numbers If \((s_N(0), i_N(0)) \to (s_0, i_0)\) as \(N \to \infty\), then

\[
\sup_{0 \leq t \leq T} \{|s_n(t) - s(t)| + |i_N(t) - i(t)|\} \to 0
\]
a.s., where \((s(t), i(t)), t \geq 0\) is the unique solution of the ODE

\[
\begin{cases}
\frac{ds}{dt}(t) = -\beta s(t)i(t), & t > 0, \\
\frac{di}{dt}(t) = \beta s(t)i(t) - \alpha i(t), & t > 0, \\
s(0) = s_0, & i(0) = i_0.
\end{cases}
\]

Proof Define \(X(t) = \left( s(t) \atop i(t) \right)\), \(X_N(t) = \left( s_N(t) \atop i_N(t) \right)\), \(\overline{X}_N(t) = X(t) - X_N(t)\), \(Y_N(t) = \left( \mathcal{M}_N(t) \atop (\mathcal{N}_N(t) - \mathcal{M}_N(t)) \right)\), and finally \(F \left( \begin{array}{c} x \\ y \end{array} \right) = \left( \begin{array}{c} -\beta xy \\ \beta xy - \alpha y \end{array} \right)\). For \(0 \leq x, y, x', y' \leq 1\),

\[
\|F \left( \begin{array}{c} x \\ y \end{array} \right) - F \left( \begin{array}{c} x' \\ y' \end{array} \right) \| \leq C(\alpha, \beta) \| \left( \begin{array}{c} x \\ y \end{array} \right) - \left( \begin{array}{c} x' \\ y' \end{array} \right) \|.
\]

We have

\[
\overline{X}_N(t) = \overline{X}_N(0) + \int_0^t [F(X(r)) - F(X_N(r))]dr + Y_N(t).
\]

From Proposition 17, for all \(T > 0\), \(\sup_{0 \leq t \leq T} \|Y_N(t)\| \to 0\) a.s. as \(N \to \infty\). Let \(\varepsilon_N(t) = \sup_{0 \leq r \leq t} \|Y_N(r)\|\). We have

\[
\|\overline{X}_N(t)\| \leq \|\overline{X}_N(0)\| + C(\alpha, \beta) \int_0^t \|\overline{X}_N(r)\|dr + \varepsilon_N(t).
\]

It then follows from Gronwall’s Lemma (see below) that

\[
\sup_{0 \leq r \leq t} \|\overline{X}_N(r)\| \leq \left( \|\overline{X}_N(0)\| + \varepsilon_N(t) \right) \exp \left( C(\alpha, \beta) t \right).
\]
The result then follows from the assumption $\|X_N(0)\| \to 0$, plus the fact that $\varepsilon_N(t) \to 0$ a.s. as $N \to \infty$. □

Of course we can write ODEs for other epidemiological models: SEIR, SEIRS, SIRS, SIS, SIV, Malaria, etc...

Caution! This “Law of Large Numbers” approximation is only valid when $s, i > 0$, i.e. when significant fractions of the population are infectious and are susceptible. The ODE is of course of no help to compute the probability that the introduction of a unique infectious results in a major epidemic. However, as we shall see now, we can recover from the ODE the same Basic Reproduction Number $R_0$ and the dichotomy $R_0 \leq 1, R_0 > 1$.

Note that at the beginning of the epidemic $s(t) \sim 1$, and in this case the equation for $i(t)$ becomes

$$\frac{di}{dt}(t) \simeq (\beta - \alpha)i(t).$$

If $\beta \leq \alpha$, the solution of the ODE does not increase, when starting from a small value. This means that there won’t be any major epidemic, if we start from a small number of initial infected individuals. On the contrary, if $\beta > \alpha$, as soon as $i(t)$ achieves a positive value, it increases. The equilibrium $i = 0$ is unstable. Of course the ODE gives us no indication whatsoever as to what is the probability that, starting from a small number of infected individuals, the epidemic reaches a stage where a significant proportion of the population is hit.

Note that $\beta \leq \alpha$ is equivalent to $\beta / \alpha \leq 1$. In the model considered in this section, $\alpha^{-1} = \mathbb{E}\Delta T$, and $\beta = cp$. Hence $\frac{\beta}{\alpha} = R_0$! The essential parameter can be read of from the ODE.

The vast majority of the literature on mathematical models in epidemiology considers ODEs of the type of equations which we have just obtained. The probabilistic point of view is more recent.

**Lemma 20. Gronwall** Let $a, b \geq 0$ and $\varphi : [0, T] \to \mathbb{R}$ be such that for all $0 \leq t \leq T$,

$$\varphi(t) \leq a + b \int_0^t \varphi(r)dr.$$

Then $\varphi(t) \leq ae^{bt}$.

**Proof** We deduce from the assumption that

$$e^{-bt}\varphi(t) - be^{-bt} \int_0^t \varphi(r)dr \leq ae^{-bt},$$
or in other words
\[
\frac{d}{dt} \left( e^{-bt} \int_0^t \varphi(r) dr \right) \leq ae^{-bt}.
\]
Integrating this inequality, we deduce
\[
e^{-bt} \int_0^t \varphi(r) dr \leq a \frac{1 - e^{-bt}}{b}.
\]
Multiplying by \(be^{bt}\) and exploiting again the assumption yields the result. □

**Exercise 9.** Let us consider Ross’ model of malaria (see also exercise 8 below), which we rewrite in a stochastic form. Denote by \(H(t)\) the number of humans who are infected by malaria, and by \(V(t)\) the number of mosquitoes who are infected by malaria at time \(t\). Let \(N_H\) denote the total number of humans, and by \(N_V\) the total number of mosquitoes, which are supposed to be constant in time. The humans (resp. the mosquitoes) which are not infected are all supposed to be susceptibles. Let \(m = N_V / N_H\), a the mean number of stings of humans by one mosquito par time unit, \(p_1\) the probability that the sting of a susceptible human by an infected mosquito infects the human, and by \(p_2\) the probability that a susceptible mosquito gets infected while stinging an infected human. We assume that the infected humans (resp. mosquitoes) heal at rate \(\gamma\) (resp. at rate \(\mu\)).

1. What is the mean number of stings that a human suffers per time unit ?

2. Given 4 mutually independent standard Poisson processes \(P_1(t), P_2(t), P_3(t)\) et \(P_4(t)\), justify the following as a stochastic model of the propagation of malaria.

\[
H(t) = H(0) + P_1 \left( a p_1 \int_0^t V(s) \frac{N_H - H(s)}{N_H} ds \right) - P_2 \left( \gamma \int_0^t H(s) ds \right)
\]
\[
V(t) = V(0) + P_3 \left( a m p_2 \int_0^t H(s) \frac{N_V - V(s)}{N_V} ds \right) - P_4 \left( \mu \int_0^t V(s) ds \right).
\]

3. Define now (with \(N_H = N, N_V = mN\))

\[
h_N(t) = \frac{H(t)}{N_H}, \quad v_N(t) = \frac{V(t)}{N_V}.
\]
Write the equation for the pair \((h_N(t), v_N(t))\). Show that as \(N \to \infty\), with \(m\) constant, \((h_N(t), v_N(t)) \to (h(t), v(t))\), solution of Ross’ ODE :

\[
\begin{align*}
\frac{dh}{dt}(t) &= a_1mv(t)(1-h(t)) - \gamma h(t) \\
\frac{dv}{dt}(t) &= a_2h(t)(1-v(t)) - \mu v(t).
\end{align*}
\]

16 Interlude 4: Martingales

16.1 Martingales in discrete time

\((\Omega, \mathcal{F}, \mathbb{P})\) being our standing probability space, let be given an increasing sequence \(\{\mathcal{F}_n, n \geq 1\}\) of sub-\(\sigma\)-algebras of \(\mathcal{F}\).

**Definition 21.** A sequence \(\{X_n, n \geq 0\}\) of r.v.’s is called a martingale if

1. For all \(n \geq 0\), \(X_n\) is \(\mathcal{F}_n\)-measurable and integrable,
2. For all \(n \geq 0\), \(\mathbb{E}(X_{n+1}|\mathcal{F}_n) = X_n\) a.s.

A sub-martingale is a sequence which satisfies the first condition and \(\mathbb{E}(X_{n+1}|\mathcal{F}_n) \geq X_n\). A super-martingale is a sequence which satisfies the first condition and \(\mathbb{E}(X_{n+1}|\mathcal{F}_n) \leq X_n\).

It follows readily from Jensen’s inequality for conditional expectations the

**Proposition 22.** If \(\{X_n, n \geq 0\}\) is a martingale, \(\varphi : \mathbb{R} \to \mathbb{R}\) a convex function such that \(\varphi(X_n)\) is integrable for all \(n \geq 0\), then \(\{\varphi(X_n), n \geq 0\}\) is a sub-martingale.

We shall need the notion of stopping time

**Definition 23.** A stopping time \(\tau\) is an \(\mathbb{N} \cup \{+\infty\}\)-valued r.v. which satisfies \(\{\tau = n\} \in \mathcal{F}_n\), for all \(n \geq 0\).

We have Doob’s optional sampling theorem :

**Theorem 24.** If \(\{X_n, n \geq 0\}\) is a martingale (resp. a sub-martingale), and \(\tau_1, \tau_2\) two stopping times s.t. \(\tau_1 \leq \tau_2 \leq N\) a.s., then \(X_{\tau_i} \in \mathcal{F}_{\tau_i}\) measurable and integrable, \(i = 1, 2\) and moreover

\[
\begin{align*}
\mathbb{E}(X_{\tau_2}|\mathcal{F}_{\tau_1}) &= X_{\tau_1} \\
(\text{resp. } \mathbb{E}(X_{\tau_2}|\mathcal{F}_{\tau_1}) &\geq X_{\tau_1}).
\end{align*}
\]
Proof For all $A \in \mathcal{B}$, $n \geq 0$,
\[
\{X_{\tau_i} \in A \} \cap \{\tau_i = n\} = \{X_n \in A\} \cap \{\tau_i = n\} \in \mathcal{F}_n,
\]
and moreover
\[
|X_{\tau_i}| \leq \sum_{k=1}^{N} |X_k|,
\]
which establishes the first part of the statement.

Let $A \in \mathcal{F}_{\tau_1}$. Then
\[
A \cap \{\tau_1 < k \leq \tau_2\} = A \cap \{\tau_1 \leq k - 1\} \cap \{\tau_2 \leq k - 1\}^c \in \mathcal{F}_{k-1}.
\]
Indeed, we have
\[
A \cap \{\tau_1 \leq k - 1\} = \bigcup_{j=1}^{k-1} A \cap \{\tau_1 = j\} \in \mathcal{F}_{k-1}
\]
and also $\{\tau_2 \leq k - 1\}^c \in \mathcal{F}_{k-1}$.

Let $\Delta_k = X_k - X_{k-1}$. We have
\[
\int_A (X_{\tau_2} - X_{\tau_1})d\mathbb{P} = \int_A \sum_{k=1}^{n} 1_{\{\tau_1 < k \leq \tau_2\}} \Delta_k d\mathbb{P}
\]
\[
= \sum_{k=1}^{n} \int_{A \cap \{\tau_1 < k \leq \tau_2\}} \Delta_k d\mathbb{P}
\]
\[
= 0
\]
or else $\geq 0$, depending upon whether $\{X_n, n \geq 0\}$ is a martingale or a sub–martingale. \hfill $\Box$

We have a first Doob’s inequality

**Proposition 25.** If $X_1, \ldots, X_n$ is a sub–martingale, then for all $\alpha > 0$,
\[
\mathbb{P}(\max_{1 \leq i \leq n} X_i \geq \alpha) \leq \frac{1}{\alpha} \mathbb{E}(X_n^+).
\]

**Proof** Define the stopping time $\tau = \inf\{0 \leq k \leq n, \ X_k \geq \alpha\}$ and let $M_k = \max_{1 \leq i \leq k} X_i$. We have
\[
\{M_n \geq \alpha\} \cap \{\tau \leq k\} = \{M_k \geq \alpha\} \in \mathcal{F}_k.
\]

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Hence \( \{M_n \geq \alpha\} \in \mathcal{F}_\tau \). From the optional sampling Theorem,

\[
\alpha \mathbb{P}(M_n \geq \alpha) \leq \int_{\{M_n \geq \alpha\}} X_\tau \, d\mathbb{P} \\
\leq \int_{\{M_n \geq \alpha\}} X_n \, d\mathbb{P} \\
\leq \int_{\{M_n \geq \alpha\}} X_n^+ \, d\mathbb{P} \\
\leq \mathbb{E}(X_n^+).
\]

\( \square \)

We have finally a second Doob’s inequality

**Proposition 26.** If \( M_1, \ldots, M_n \) is a martingale, then

\[
\mathbb{E}\left[ \sup_{0 \leq k \leq n} |M_k|^2 \right] \leq 4 \mathbb{E}\left[ |M_n|^2 \right].
\]

**Proof** Let \( X_k = |M_k| \). From Proposition 22, \( X_1, \ldots, X_n \) is a sub–martingale. It follows from the proof of Proposition 25 that, with the notation \( X_k^* = \sup_{0 \leq k \leq n} X_k \),

\[
\mathbb{P}(X_n^* > \lambda) \leq \frac{1}{\lambda} \mathbb{E}\left( X_n 1_{X_n^* > \lambda} \right).
\]

Consequently

\[
\int_0^\infty \lambda \mathbb{P}(X_n^* > \lambda) d\lambda \leq \int_0^\infty \mathbb{E}\left( X_n 1_{X_n^* > \lambda} \right) d\lambda \\
\mathbb{E}\left( \int_0^{X_n^*} \lambda \, d\lambda \right) \leq \mathbb{E}\left( X_n \int_0^{X_n^*} d\lambda \right) \\
\frac{1}{2} \mathbb{E}\left[ |X_n^*|^2 \right] \leq \mathbb{E}(X_n X_n^*) \\
\leq \sqrt{E(|X_n|^2)} \sqrt{E(|X_n^*|^2)},
\]

from which the result follows. \( \square \)
16.2 Martingales in continuous time

We are now given an increasing collection \( \{F_t, \ t \geq 0\} \) of sub-\( \sigma \)-algebras.

**Definition 27.** A process \( \{X_t, \ t \geq 0\} \) of r.v.'s is called a martingale if

1. for all \( t \geq 0 \), \( X_t \) is \( F_t \)-measurable and integrable;
2. for all \( 0 \leq s < t \), \( \mathbb{E}(X_t|F_s) = X_s \) a. s.

A sub-martingale is a sequence which satisfies the first condition and \( \mathbb{E}(X_t|F_s) \geq X_s \). A super-martingale is a sequence which satisfies the first condition and \( \mathbb{E}(X_t|F_s) \leq X_s \).

Suppose \( \{M_t, \ t \geq 0\} \) is a right-continuous martingale. For any \( n \geq 1 \), \( 0 = t_0 < t_1 < \cdots < t_n \), \( (M_{t_0}, M_{t_1}, \ldots, M_{t_n}) \) is a discrete time martingale, to which Proposition 26 applies. Since

\[
\sup_{0 \leq s \leq t} |M_s| = \sup_{\text{Partitions of } [0,t]} \sup_{1 \leq k \leq n} |M_{t_k}|,
\]

Consequently Proposition 26 implies readily

**Proposition 28.** If \( \{M_t, \ t \geq 0\} \) is a right-continuous martingale,

\[
\mathbb{E} \left[ \sup_{0 \leq s \leq t} |M_s|^2 \right] \leq 4 \mathbb{E} \left[ |M_t|^2 \right].
\]

17 Central Limit Theorem

We write \((s_N, i_N)\) in the form

\[
s_N(t) = s(t) + \frac{1}{\sqrt{N}} U_N(t),
\]
\[
i_N(t) = i(t) + \frac{1}{\sqrt{N}} V_N(t),
\]

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and we look for the limiting law of the process \((U_N(t), V_N(t))\) as \(N \to \infty\). It is plain that

\[
s(t) + \frac{U_N(t)}{\sqrt{N}} = s(0) - \beta \int_0^t s(r) i(r) dr - \frac{\beta}{\sqrt{N}} \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + \frac{U_N(r)V_N(r)}{\sqrt{N}} \right) dr
\]

\[
- \frac{1}{N} M_1 \left( \beta N \int_0^t s(r) i(r) + \frac{s(r) V_N(r) + i(r) U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} dr \right),
\]

\[
i(t) + \frac{V_N(t)}{\sqrt{N}} = i(0) + \beta \int_0^t s(r) i(r) dr + \frac{\beta}{\sqrt{N}} \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + \frac{U_N(r)V_N(r)}{\sqrt{N}} \right) dr
\]

\[
- \alpha \int_0^t i(r) dr - \frac{\alpha}{\sqrt{N}} \int_0^t V_N(r) dr
\]

\[
+ \frac{1}{N} M_1 \left( \beta N \int_0^t s(r) i(r) + \frac{s(r) V_N(r) + i(r) U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} \right)
\]

\[
- \frac{1}{N} M_2 \left( \alpha N \int_0^t \left( i(r) + \frac{V_N(r)}{\sqrt{N}} \right) dr \right).
\]

We use the ODE satisfied by \((s(t), i(t))\) in order to suppress the terms of order 1 in the above, and multiply the remaining terms by \(\sqrt{N}\), from which we deduce

\[
U_N(t) = -\beta \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + \frac{U_N(r)V_N(r)}{\sqrt{N}} \right) dr
\]

\[
- \frac{1}{\sqrt{N}} M_1 \left( \beta N \int_0^t s(r) i(r) + \frac{s(r) V_N(r) + i(r) U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} dr \right),
\]

\[
V_N(t) = \beta \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + \frac{U_N(r)V_N(r)}{\sqrt{N}} \right) dr - \alpha \int_0^t V_N(r) dr
\]

\[
+ \frac{1}{\sqrt{N}} M_1 \left( \beta N \int_0^t s(r) i(r) + \frac{s(r) V_N(r) + i(r) U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} \right)
\]

\[
- \frac{1}{\sqrt{N}} M_2 \left( \alpha N \int_0^t \left( i(r) + \frac{V_N(r)}{\sqrt{N}} \right) dr \right).
\]

Let

\[
M_1^N(t) = \frac{1}{\sqrt{N}} M_1 \left( \beta N \int_0^t s(r) i(r) + \frac{s(r) V_N(r) + i(r) U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} \right),
\]

\[
M_2^N(t) = \frac{1}{\sqrt{N}} M_2 \left( \alpha N \int_0^t \left( i(r) + \frac{V_N(r)}{\sqrt{N}} \right) dr \right).
\]
Let \( [\mathcal{M}_1]_t = \sum_{0 \leq s \leq t} |\Delta \mathcal{M}_1(s)|^2 \), and define analogously \( [\mathcal{M}_2]_t \). We have

\[
[\mathcal{M}^N_1]_t = \frac{1}{N} P_1 \left( \beta \int_0^t \left( s(r)i(r) + \frac{s(r)V_N(r) + i(r)U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} \right) dr \right),
\]

\[
[\mathcal{M}^N_2]_t = \frac{1}{N} P_2 \left( \alpha \int_0^t \left( i(r) + \frac{V_N(r)}{\sqrt{N}} \right) dr \right).
\]

If we define

\[
\langle \mathcal{M}^N_1 \rangle_t = \beta \int_0^t \left( s(r)i(r) + \frac{s(r)V_N(r) + i(r)U_N(r)}{\sqrt{N}} + \frac{U_N(r)V_N(r)}{N} \right) dr,
\]

\[
\langle \mathcal{M}^N_2 \rangle_t = \alpha \int_0^t \left( i(r) + \frac{V_N(r)}{\sqrt{N}} \right) dr,
\]

then with the notation

\[
\mathcal{F}^N_s = \sigma \{ s_N(r), i_N(r), \ 0 \leq r \leq t \},
\]

then we deduce by an analogous computation to that done in Lemma 15, for \( 0 \leq r < t \),

\[
\mathbb{E} \left[ |\mathcal{M}^N_1(t) - \mathcal{M}^N_1(r)|^2 \bigg| \mathcal{F}^N_r \right] = \beta \mathbb{E} \left[ \int_r^t \left( s(u)i(u) + \frac{s(u)V_N(u) + i(u)U_N(u)}{\sqrt{N}} + \frac{U_N(u)V_N(u)}{N} \right) du \bigg| \mathcal{F}^N_r \right],
\]

\[
\mathbb{E} \left[ |\mathcal{M}^N_1(t) - \mathcal{M}^N_1(r)|^2 \bigg| \mathcal{F}^N_r \right] = \alpha \mathbb{E} \left[ \int_r^t \left( i(u) + \frac{V_N(u)}{\sqrt{N}} \right) du \bigg| \mathcal{F}^N_r \right].
\]

**A priori estimate** It follows from Lemma 15 that

\[
\mathcal{M}^N_1(t) = -U_N(t) - \beta \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + \frac{U_N(r)V_N(r)}{\sqrt{N}} \right) dr,
\]

\[
\mathcal{M}^N_2(t) = \mathcal{M}^N_1(t) - V_N(t) + \beta \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + \frac{U_N(r)V_N(r)}{\sqrt{N}} \right) dr
\]

\[- \alpha \int_0^t V_N(r) dr.
\]
Moreover from the definitions of \( U_N(t) \) and \( V_N(t) \),
\[
|U_N(t)| \leq 2\sqrt{N}, \quad |V_N(t)| \leq 2\sqrt{N}.
\]
Hence we deduce from the formulas for \( \langle M_1^N \rangle_t \) and \( \langle M_2^N \rangle_t \)
\[
\mathbb{E}[(M_1^N(t))^2] \leq 9\beta t, \\
\mathbb{E}[(M_2^N(t))^2] \leq 3\alpha t, \\
\mathbb{E}(|M_1^N(t)|) \leq 3\sqrt{\beta t}, \\
\mathbb{E}(|M_2^N(t)|) \leq \sqrt{3\alpha t}
\]
But
\[
|U_N(t)| \leq \beta \int_0^t (|V_N(r)| + 3|U_N(r)|) \, dr + |M_1^N(t)|, \\
|V_N(t)| \leq \alpha \int_0^t |V_N(r)| \, dr + \beta \int_0^t (|V_N(r)| + 3|U_N(r)|) \, dr + |M_1^N(t)| + |M_2^N(t)|.
\]
Summing up those two inequalities, and taking advantage of Gronwall’s Lemma and of the above estimates of the two martingales, we deduce that for all \( T > 0 \), there exist two constants \( C_1(\alpha, \beta, T) \) and \( C_2(\alpha, \beta, T) \) (for the second estimate we take the square before taking the expectation) such that
\[
\sup_{N \geq 1, \ 0 \leq t \leq T} \mathbb{E} (|U_N(t)| + |V_N(t)|) \leq C_1(\alpha, \beta, T), \\
(1) \quad \sup_{N \geq 1, \ 0 \leq t \leq T} \mathbb{E} (|U_N(t)|^2 + |V_N(t)|^2) \leq C_2(\alpha, \beta, T).
\]
Lemma 29. For all \( T > 0 \),
\[
\sup_{N \geq 1} \mathbb{E} \left( \sup_{0 \leq t \leq T} \left[ |U_N(t)|^2 + |V_N(t)|^2 \right] \right) < \infty.
\]
Proof
\[
\sup_{0 \leq r \leq t} |U_N(r)|^2 \leq 18\beta^2 t \int_0^t (|V_N(r)|^2 + 5|U_N(r)|^2) \, dr + 2 \sup_{0 \leq r \leq t} |M_1(r)|^2.
\]
It follows from Doob’s inequality that
\[
\mathbb{E} \left( \sup_{0 \leq r \leq t} |M_1(r)|^2 \right) \leq 4\mathbb{E}(M_1)_t \\
\leq 4 \times 9\beta t.
\]
Hence the first part of the result follows from the last two inequalities combined with (1). The second part of the result follows analogously. \( \square \)
Convergence in law  The proof of convergence follows from the Lemma

**Lemma 30.** Let \( \{P(t), \ t \geq 0\} \) be a standard Poisson process. Let \( M(t) = P(t) - t \). Then for any sequence \( \{t_N, \ N \geq 1\} \) of real numbers such that \( N^{-1/2}t_N \to 0 \) as \( N \to \infty \),

\[
\left\{ \frac{M(Nt + \sqrt{N}t_N)}{\sqrt{N}}, \ t \geq 0 \right\} \Rightarrow \{B(t), \ t \geq 0\},
\]

where \( B(t) \) is a standard Brownian motion.

**Proof** We shall only prove convergence of the finite dimensional distributions. It is in fact sufficient to prove that for any \( t > 0 \) fixed,

\[
\frac{M(Nt + \sqrt{N}t_N)}{\sqrt{N}} \Rightarrow B(t),
\]

since on the left we have a process with asymptotically stationary and independent increments, and the limit has those properties.

For each \( t \geq 0 \), the convergence \( N^{-1/2}M(Nt) \Rightarrow B(t) \) follow from the usual Central Limit Theorem. Indeed

\[
\frac{M(Nt)}{\sqrt{[Nt]}} = \frac{1}{\sqrt{[Nt]}} \sum_{i=1}^{[Nt]} [M(i) - M(i - 1)] + \frac{M(Nt) - M([Nt])}{\sqrt{[Nt]}},
\]

the r.v.’s \( M(i) - M(i - 1) \) are i.i.d. centered with variance 1, and the last term above converges in probability to 0 as \( N \to \infty \), hence

\[
\frac{M(Nt)}{\sqrt{[Nt]}} \to \mathcal{N}(0,1),
\]

\[
\frac{M(Nt)}{\sqrt{N}} = \frac{\sqrt{[Nt]}}{\sqrt{N}} \times \frac{M(Nt)}{\sqrt{[Nt]}} \Rightarrow B(t),
\]

where \( B(t) \) follows the law \( \mathcal{N}(0,t) \). Now

\[
\frac{M(Nt + \sqrt{N}t_N)}{\sqrt{N}} = \frac{M(Nt)}{\sqrt{N}} + \frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}},
\]

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and it remains to prove that
\[
\frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}} \to 0
\]
in probability, as \(n \to \infty\). But
\[
\mathbb{P}\left( \left| \frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}} \right| > \varepsilon \right) \leq \frac{1}{N\varepsilon^2} \text{Var} \left( M(Nt + \sqrt{N}t_N) - M(Nt) \right)
\]
\[
= \frac{\sqrt{N}t_N}{N\varepsilon^2} \to 0,
\]
provided \(N^{-1/2}t_N \to 0\) as \(N \to \infty\).

It remains to show :

**Proposition 31.** Lemma 30 remains true with \(t_N\) random, provided \(N^{-1/2}\mathbb{E}[|t_N|] \to 0\) as \(N \to \infty\).

**Proof** We need to show that
\[
\frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}} \to 0
\]
in probability, as \(N \to \infty\).

\[
\mathbb{P}\left( \left| \frac{t_N}{\sqrt{N}} > \eta \right| \right) \leq \frac{1}{\eta} \mathbb{E}\left| t_N \right| \to 0,
\]
as \(N \to \infty\). Let \(\varepsilon > 0\) be fixed. We have
\[
\left\{ \left| \frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}} \right| > \varepsilon \right\} \\
\subseteq \left\{ \left| \frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}} \right| > \varepsilon \right\} \cap \left\{ 0 \leq t_N \leq \eta \sqrt{N} \right\} \\
\cup \left\{ \left| \frac{M(Nt + \sqrt{N}t_N) - M(Nt)}{\sqrt{N}} \right| > \varepsilon \right\} \cap \left\{ -\eta \sqrt{N} \leq t_N \leq 0 \right\} \cup \left\{ \frac{|t_N|}{\sqrt{N}} > \eta \right\}.
\]
The probability of the first event on the right can be estimated as follows, using Doob’s inequality
\[
\mathbb{P}\left( \sup_{0 \leq s \leq N\eta} \frac{|M(Nt + s) - M(Nt)|}{\sqrt{N}} > \varepsilon \right) \leq \frac{4}{N\varepsilon^2} \mathbb{E}\left( |M(N(t + \eta)) - M(Nt)|^2 \right) \\
\leq \frac{4\eta}{\varepsilon^2}.
\]

Estimating analogously the probability of the second event, and that of the third one by Chebychef’s inequality, we get
\[
\mathbb{P}\left( \frac{|M(Nt + \sqrt{N}t_N) - M(Nt)|}{\sqrt{N}} > \varepsilon \right) \leq \frac{8\eta}{\varepsilon^2} + \frac{16\mathbb{E}|t_N|}{\eta\sqrt{N}}.
\]

Let \( \eta = \varepsilon^3/16 \), from which we deduce
\[
\mathbb{P}\left( \frac{|M(Nt + \sqrt{N}t_N) - M(Nt)|}{\sqrt{N}} > \varepsilon \right) \leq \frac{\varepsilon}{2} + \frac{16\mathbb{E}|t_N|}{\varepsilon^3\sqrt{N}}.
\]

Then if \( N \) is large enough such that \( \mathbb{E}|t_N| \leq \varepsilon^4/32 \),
\[
\mathbb{P}\left( \frac{|M(Nt + \sqrt{N}t_N) - M(Nt)|}{\sqrt{N}} > \varepsilon \right) \leq \varepsilon.
\]

The Proposition follows since this result holds for any \( \varepsilon > 0 \). \( \square \)

We use this Proposition first with \( M = M_1 \), \( t \) being replaced by \( \beta \int_0^t s(r)i(r)dr \) and
\[
t_N = \beta \int_0^t \left( s(r)V_N(r) + i(r)U_N(r) + N^{-1/2}U_N(r)V_N(r) \right) dr
\]
which satisfies \( \mathbb{E}|t_N| \leq C \), then with \( M = M_2 \), \( t \) being replaced by \( \alpha \int_0^t i(r)dr \) and \( t_N = \alpha \int_0^t V_N(r)dr \), which again satisfies \( \mathbb{E}|t_N| \leq C \). In order to get the joint law of both limits, we exploit the fact that the product \( M_1(t)M_2(t) \) is a martingale.

Moreover one can rather easily show that the sequence \( \{(U_N(t), V_N(t)), t \geq 0\} \) is tight as a process whose trajectories belong to \( C([0, +\infty); \mathbb{R}^2) \). Hence along a subsequence
\[
\{(U_N(t), V_N(t)), t \geq 0\} \Rightarrow \{(U(t), V(t)), t \geq 0\},
\]

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where the limit satisfies

\[ U(t) = -\beta \int_0^t [s(r)V(r) + i(r)U(r)] \, dr + \sqrt{\beta} \int_0^t \sqrt{s(r)i(r)} \, dB_1(r), \]

\[ V(t) = \beta \int_0^t [s(r)V(r) + i(r)U(r)] \, dr - \alpha \int_0^t V(r) \, dr - \sqrt{\beta} \int_0^t \sqrt{s(r)i(r)} \, dB_1(r) \]

\[ + \sqrt{\alpha} \int_0^t \sqrt{i(r)} \, dB_2(r). \]

The process \( \{(U(t), V(t)), \ t \geq 0\} \) is a Gaussian process of the Ornstein–Uhlenbeck type.

The law of the limit is uniquely determined. Hence the whole sequence converges.

**Exercise 10.** Let us go back to Exercise 9. We now define \( X_N(t) \) and \( Y_N(t) \) by letting

\[ h_N(t) = h(t) + \frac{X_N(t)}{\sqrt{N}}, \quad v_N(t) = v(t) + \frac{Y_N(t)}{\sqrt{N}}. \]

Write the equation satisfied by the Ornstein–Uhlenbeck process \((X(t), Y(t)) = \lim_{N \to \infty} (X_N(t), Y_N(t))\).

### 18 Large Deviations

We consider the vector of proportions in our model as

\[ Z_N(t) = z_0 + \frac{1}{N} \sum_{j=1}^k h_j P_j \left( \int_0^t N \beta_j(Z_N(s)) \, ds \right). \]

Again, the \( P_j \)'s are mutually independent standard Poisson processes. The process \( Z_N(t) \) lives in the set

\[ A = \{ z \in \mathbb{R}_+^d; \sum_{i=1}^d z_i \leq 1 \}. \]

We shall denote by \( D_{T,A} \) the set of functions defined on \([0, T]\) with values in \( A \) which are right continuous with left limits at every \( t \), and \( AC_{T,A} \) will denote the subset of absolutely continuous functions. For \( \phi, \psi \in D_{T,A} \), we define \( \| \phi - \psi \|_T = \sup_{0 \leq t \leq T} |\phi_t - \psi_t| \). Let \( \mathbb{P}^N \) denote the law of \( Z^N \), i.e.

\[ \mathbb{P}^N(B) = \mathbb{P}(Z^N \in B), \forall B \in \mathcal{B}, \]

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where $\mathcal{B}$ denote the Borel $\sigma$–field of $D_{T,A}$.

We want to show that the collection of probability measures $\{\mathbb{P}^N, \; N \geq 1\}$ satisfies a Large Deviations Principle, in the sense that there exists a rate function $I_T$ (to be defined below) such that

$$- \inf_{\phi \in G} I_T(\phi) \leq \liminf_{N \to \infty} \frac{1}{N} \log \mathbb{P}(Z^N \in G), \; \text{if } G \subset D_{T,A} \text{ is open},$$

$$- \inf_{\phi \in F} I_T(\phi) \geq \limsup_{N \to \infty} \frac{1}{N} \log \mathbb{P}(Z^N \in F), \; \text{if } F \subset D_{T,A} \text{ is closed}.$$
Let for any \( a > 0 \)

\[ C_a = \inf_{1 \leq j \leq k} \inf_{z \in B^a} \beta_j(z). \]

It is plain that \( C_a > 0 \) for \( a > 0 \), and \( C_a \to 0 \), as \( a \to 0 \). We shall assume

**Assumptions A**

A1 The rate functions \( \beta_j \) are Lipschitz continuous with the Lipschitz constant equal to \( C \), and bounded by a constant \( \theta \).

A2 There exist two constants \( \lambda_1, \lambda_2 > 0 \) such that whenever \( z \in A \) is such that \( \beta_j(z) < \lambda_1, \beta_j(z^a) > \beta_j(z) \) for all \( 0 < a < \lambda_2 \).

A3 There exists \( \nu \in (0, 1/2) \) such that \( \lim_{a \to 0} a^\nu \log C_a = 0 \).

### 18.1 Law of Large Numbers and Girsanov theorem

We reformulate the Law of Large Numbers in the above notations

**Theorem 33.** Let \( Z^N \) be given the solution of (1). If the assumption 1 is satisfied, then for all \( T > 0 \),

\[ \|Z^N - Y\|_T \to 0 \text{ a.s. as } N \to \infty, \]

where \( Y_t \) is the unique solution of the ODE

\[ Y(t) = z_0 + \int_0^t b(Y(s))ds, \]

with \( b(z) = \sum_{j=1}^{k} \beta_j(z)h_j \).

We shall need the following Girsanov theorem. Let \( Q \) denote the number of jumps of the of \( Z^N \) in the interval \([0, T]\), \( \tau_p \) be the time of the \( p \)-th jump, and define

\[ \delta_p(j) = \begin{cases} 1, & \text{if the } p \text{-th jump is in the direction } h_j, \\ 0, & \text{otherwise.} \end{cases} \]

We shall denote \( \mathcal{F}^N_t = \sigma\{Z^N_s, 0 \leq s \leq t\} \). Consider another set of rates \( \tilde{\beta}_j(z), 1 \leq j \leq k \).
Theorem 34. Assume that \( \{x, \widehat{\beta}_j(x) = 0\} \subset \{s, \beta_j(x) = 0\} \). Let \( \overline{P}_N \) denote the law of \( Z^N \) when the rates are \( \widehat{\beta}_j \). Then on the \( \sigma \)-algebra \( \mathcal{F}_t^N \), \( \overline{P}_N|_{\mathcal{F}_t^N} \ll \overline{P}_N|_{\mathcal{F}_t^N} \), and

\[
\Delta^N_T = \frac{P^N|_{\mathcal{F}_t^N}}{\overline{P}_N|_{\mathcal{F}_t^N}} = \left( \prod_{p=1}^Q \prod_{j=1}^k \left[ \frac{\beta_j(Z^N(\tau_p^-))}{\widehat{\beta}_j(Z^N(\tau_p^-))} \right]^{\delta_p(j)} \right)^{\exp \left( N \sum_{j=1}^k \int_0^T \left[ \overline{\beta}_j(Z^N(t)) - \beta_j(Z^N(t)) \right] dt \right)}.
\]

18.2 The rate function

For any \( \phi \in \mathcal{AC}_{T,A} \), let \( \mathcal{A}_d(\phi) \) the set of vector valued Borel measurable functions \( \mu \) such that for all \( 1 \leq j \leq k \), \( \mu_j \geq 0 \) and

\[
\frac{d\phi_t}{dt} = \sum_{j=1}^k \mu_j h_j, \quad \text{t a.e.}
\]

We define the rate function

\[
I_T(\phi) = \begin{cases} 
\inf_{\mu \in \mathcal{A}_d(\phi)} I_T(\phi|\mu), & \text{if } \phi \in \mathcal{AC}_{T,A}, \\
+\infty, & \text{otherwise,}
\end{cases}
\]

where

\[
I_T(\phi|\mu) = \int_0^T \sum_{j=1}^k f(\mu_j, \beta_j(\phi_t))dt,
\]

with \( f(\nu, \omega) = \nu \log(\nu/\omega) - \nu + \omega \), where we use the convention \( \log(0/0) = +\infty \) for \( \nu > 0 \), while 0 \( \log(0/0) = 0 \log(0) = 0 \).

Another possible definition leads to

\[
\overline{I}_T(\phi) = \begin{cases} 
\inf_{\mu \in \mathcal{A}_d(\phi)} \int_0^T L(\phi_t, \phi'_t)dt, & \text{if } \phi \in \mathcal{AC}_{T,A}, \\
+\infty, & \text{otherwise,}
\end{cases}
\]

where for all \( z \in A, y \in \mathbb{R}^d \),

\[
L(x, y) = \sup_{\theta \in \mathbb{R}^d} \ell(z, y, \theta)
\]

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with
\[
\ell(z, y, \theta) = \langle \theta, y \rangle - \sum_{j=1}^{k} \beta_j(z) \left( e^{\langle \theta, h_j \rangle} - 1 \right).
\]

Recall the definition

**Definition 35.** A rate function \( I \) is a semi-continuous mapping \( I : D_{T,A} \to [0, \infty] \) (i.e. its level sets \( \Psi_I(\alpha) = \{ \phi, \ I_T(\phi) \leq \alpha \} \) are closed subsets of \( D_{T,A} \)).

A good rate function is a rate function whose level sets are compact.

We have (see Kratz, Pardoux [3], Pardoux, Samegni [6])

**Proposition 36.** \( I_T = \tilde{I}_T \) is a good rate function.

### 18.3 Preliminary Lemmas

We list here, without proofs, several Lemmas which are essential for the proofs of the results.

**Lemma 37.** Suppose that the \( \beta_j \), \( j = 1, \ldots, k \) are bounded by \( \theta \). If \( I_T(\phi|\mu) \leq s \) then for all \( 0 \leq t_1, t_2 \leq T \) such that \( t_2 - t_1 \leq 1/\theta \),
\[
\int_{t_1}^{t_2} \mu^i_t dt \leq \frac{s + 1}{-\log(\theta(t_2 - t_1))} \quad \forall j = 1, \ldots, k.
\]

For \( \phi \in D_{T,A} \) let \( \phi^a \) be defined by \( \phi^a_t = \Phi(\phi_t) \). Clearly \( \phi^a \in R^a \).

**Lemma 38.** Let \( \phi \) be such that \( I_T(\phi) < \infty \). We have \( \limsup_{a \to 0} I_T(\phi^a) \leq I_T(\phi) \).

**Lemma 39.** Let \( a > 0 \) and \( \phi \in R^a \) such that \( I_T(\phi) < \infty \). For all \( \eta > 0 \) there exists \( L > 0 \) and \( \phi^L \in R^{a/2} \) such that \( \| \phi - \phi^L \|_T < a/2 \) and \( I_T(\phi^L|\mu^L) \leq I_T(\phi) + \eta \) where \( \mu^L \in \mathcal{A}_d(\phi^L) \) such that \( \mu^L_j < L, j = 1, \ldots, k \).

Let \( \epsilon > 0 \) be such that \( T/\epsilon \in \mathbb{N} \) and let the \( \phi^\epsilon \) be the polygonal approximation of \( \phi \) defined for \( t \in [\ell \epsilon, (\ell + 1) \epsilon) \) by
\[
\phi_t^\epsilon = \phi_{\ell \epsilon} \frac{(\ell + 1)\epsilon - t}{\epsilon} + \phi_{(\ell+1)\epsilon} \frac{t - \ell \epsilon}{\epsilon}.
\]
Lemma 40. Fix $\eta > 0$. Let $a \in (0,1)$ and $\phi \in R^d$ such that $I_T(\phi) < \infty$. Suppose that $\mu \in A_d(\phi)$ such that $\mu^i_j < L, \ j = 1, \ldots, k$ for some $L > 0$ and $I_T(\phi|\mu) < \infty$ then there exists $a_\eta$ such that for all $a < a_\eta$ there exists an $\epsilon_a > 0$ such that for all $\epsilon < \epsilon_a$, $\phi^x \in R^d$ and $\|\phi - \phi^x\|_T < a/2$. Moreover, there exists $\mu^x \in A_d(\phi^x)$ such that $\mu^x_i < L, \ j = 1, \ldots, k$ and $I_T(\phi^x|\mu^x) \leq I_T(\phi|\mu) + \eta$.

The next lemma exploits a large deviation estimate for Poisson r.v.'s.

Lemma 41. Let $Y_1, Y_2, \ldots$ be independent Poisson random variables with mean $\theta \epsilon$. For all $N \in \mathbb{N}$, let

$$\bar{Y}^N = \frac{1}{N} \sum_{n=0}^{N} Y_n.$$

For any $s > 0$ there exist $K, \epsilon_0 > 0$ and $N_0 \in \mathbb{N}$ such that taking $g(\epsilon) = K \sqrt{\log^{-1}(\epsilon^{-1})}$ we have

$$\mathbb{P}^N(\bar{Y}^N > g(\epsilon)) < \exp\{-sN\}$$

for all $\epsilon < \epsilon_0$ and $N > N_0$.

Proof. We apply the Gramer’s theorem that we can find in [1] (chapter 2) to have that there exist $N_0 \in \mathbb{N}$ such that

$$\limsup_{N \to \infty} \frac{1}{N} \log(\mathbb{P}^N(\bar{Y}^N > g(\epsilon))) \leq -\inf_{x \geq g(\epsilon)} \Lambda^*_x(x)$$

where $\Lambda^*_x(x) = \sup_{\lambda \in \mathbb{R}} \{\lambda x - \Lambda_x(\lambda)\}$ with

$$\Lambda_x(\lambda) = \log(\mathbb{E}(e^{\lambda Y_1}) = \theta \epsilon (e^\lambda - 1)).$$

We deduce that

$$\Lambda^*_x(x) = x \log \frac{x}{\theta \epsilon} - x + \theta \epsilon.$$

This last function is convex then it reaches his infimuim in $x = \theta \epsilon$ and as

$$\lim_{\epsilon \to 0} \frac{g(\epsilon)}{\theta \epsilon} = +\infty$$

there exists $\epsilon_1 > 0$ such that $g(\epsilon) > \theta \epsilon$ for all $\epsilon < \epsilon_1$ and then

$$\inf_{x \geq g(\epsilon)} \Lambda^*_x(x) = g(\epsilon) \log \frac{g(\epsilon)}{\theta \epsilon} - g(\epsilon) + \theta \epsilon$$

$$= g(\epsilon) \log(g(\epsilon)) - g(\epsilon) \log(\theta \epsilon) - g(\epsilon) + \theta \epsilon$$

$$\approx K \sqrt{\log(1/\epsilon)} \to \infty \ \text{as} \ \epsilon \to 0.$$

Then there exists $\epsilon_2 > 0$ such that $\inf_{x \geq g(\epsilon)} \Lambda^*_x(x) > s$ for all $\epsilon < \epsilon_2$.

Taking $\epsilon_0 = \min\{\epsilon_1, \epsilon_2\}$, we have the lemma. □
18.4 The Lower Bound

For a path φ let \( F_\delta(\phi) = \{ \psi : \|\psi - \phi\|_T < \delta \} \). We first prove that for all fixed path φ and any \( \eta > 0, \delta > 0 \) there exists \( N_{\eta,\delta} \), such that for all \( N > N_{\eta,\delta} \)

\[
P^N(F_\delta(\phi)) = \mathbb{E}\left( \Delta^N_T \mathbf{1}_{\{Z^N \in F_\delta(\phi)\}} \right) \geq \exp\{-N(I_{T,x}(\phi) + \eta)\}.
\]

To this end, it is enough to prove (4) considering \( \phi \in \mathcal{AC}_{T,A} \) because the inequality is true when \( I_{T,x}(\phi) = \infty \). We apply some lemmas of the preceding section to show that it is enough to consider some suitable paths \( \phi \) with the \( \mu \in \mathcal{A}_d(\phi) \).

We have the

**Lemma 42.** For any \( a > 0, \epsilon > 0 \) let \( \phi \in R^a \) for \( a > 0 \). For \( \epsilon > 0 \) let \( \phi^\epsilon \) be its polygonal approximation defined by (3). Suppose that for all \( \eta > 0, \delta > 0 \) there exists \( N_{\eta,\delta} \) such that for all \( N \geq N_{\eta,\delta} \)

\[
P^N(F_\delta(\phi)) = \mathbb{P}(\|Z^N - \phi^\epsilon\|_T < \delta) \geq \exp\{-N(I_T(\phi^\epsilon|\mu^\epsilon) + \eta)\}
\]

where \( \mu^\epsilon \in \mathcal{A}_d(\phi^\epsilon) \) such that \( \mu^\epsilon_j \leq L \) for all \( j = 1, \ldots, k \) for some \( L > 0 \). Then for all fixed \( \phi \in \mathcal{AC}_{T,A} \), and any \( \eta > 0, \delta > 0 \) there exists \( N_{\eta,\delta} \) such that for all \( N > N_{\eta,\delta} \)

\[
P^N(F_\delta(\phi)) = \mathbb{P}(\|Z^N - \phi\|_T < \delta) \geq \exp\{-N(I_T(\phi) + \eta)\}.
\]

The goal of the next lemma is to show the inequality (5).

**Lemma 43.** For \( a > 0, \epsilon > 0 \), let \( \phi \in R^a \) be linear on each intervals \([\ell \epsilon, (\ell + 1) \epsilon] \), \( 0 \leq \ell \leq \frac{T}{\epsilon} \). Consider the \( \mu \in \mathcal{A}_d(\phi) \) that is constant over these time intervals and such that all the components of \( \mu \) are bounded above by some constant \( L > 0 \). Then we have that for any \( \eta > 0 \), and suitable small \( \delta > 0 \) (thus the inequality stay true for all \( \delta > 0 \)) there exists \( N_{\eta,\delta} \in \mathbb{N} \) such that for all \( N > N_{\eta,\delta} \)

\[
P(\|Z^N - \phi\|_T < \delta) \geq \exp\{-N(I_T(\phi|\mu) + \eta)\}.
\]

**Proof** Define the events \( B_j, j = 1, \ldots, k \) for controlling the likelihood ratio. For \( \xi > 0 \) let

\[
B_j = \left\{ \left| \sum_{p=1}^{Q} \delta_p(j) \log \left( \frac{\beta_j(Z^N(\tau^-_p))}{\mu_j^{\ell \epsilon}} \right) - N \sum_{\ell=1}^{T/\epsilon} \mu_{j \ell}^\epsilon \log \left( \frac{\beta_j(\phi_{\ell \epsilon})}{\mu_{j \ell}^\epsilon} \right) \right| \leq N \xi \right\}
\]
We have on \( \{Z^N \in F_\delta(\phi)\} \cap (\bigcap_{j=1}^k B_j) = \{Z^N \in F_\delta(\phi)\} \cap B \)

\[
\Delta_T^N \geq \exp \left\{ -N \int_0^T \sum_{j=1}^k \mu_j^t \log \left( \frac{\mu_j^t}{\beta_j(\phi_t)} \right) - \mu_t^t + \beta_j(\phi_t) \right\} dt - N O(\delta + \xi) \]

\[
\geq \exp \left\{ -N(I_T(\phi|\mu) + O(\delta + \xi)) \right\} \quad \text{on the event} \quad \{Z^N \in F_\delta(\phi)\} \cap B.
\]

Then for any \( \eta > 0 \), there exists \( \delta > 0 \) and \( \xi > 0 \) such that for \( N \) large enough we have

\[
\Delta_T^N \geq \exp \{ -N(I_T(\phi|\mu) + \eta/2) \}
\]

Moreover

\[
\mathbb{P}^N(F_\delta(\phi)) = \tilde{E} \left( \Delta_T^N \mathbf{1}_{\{Z^N \in F_\delta(\phi)\}} \right)
\]

\[
\geq \tilde{E} \left( \Delta_T^N \mathbf{1}_{\{Z^N \in F_\delta(\phi)\} \cap B} \right)
\]

\[
\geq \exp \{ -N(I_T(\phi|\mu) + \eta/2) \} \tilde{P}(\{Z^N \in F_\delta(\phi)\} \cap B)
\]

To finish this proof it is enough to show the following lemma:

**Lemma 44.** Let \( \phi \in R^a \) be linear over the intervals \( [\ell \epsilon, (\ell+1) \epsilon] \),

\[
\lim_{N \to \infty} \tilde{P}(\{Z^N \in F_\delta(\phi)\} \cap B) = 1
\]

We finish the proof of the lower bound by the following theorem

**Theorem 45.** For all open set \( G \in D_{T,A} \),

\[
\liminf_{N \to \infty} \frac{1}{N} \log \mathbb{P}^N(G) \geq - \inf_{\phi \in G} I_T(\phi).
\]

**Proof** It is enough to assume that (4) is true and show (45). To this end let \( I = \inf_{\phi \in G} I_T(\phi) < \infty \) then, for \( \eta > 0 \) there exists a \( \phi^\eta \in G \) such that \( I_T(\phi^\eta) \leq I + \eta \). Moreover we can choose \( \delta = \delta(\phi^\eta) \) small enough such that \( F_\delta(\phi^\eta) \subset G \). And then \( \mathbb{P}^N(F_\delta(\phi^\eta)) \leq \mathbb{P}^N(G) \). This implies from the inequality (4) that for all \( \eta > 0 \),

\[
\liminf_{N \to \infty} \frac{1}{N} \log \mathbb{P}^N(G) \geq \liminf_{N \to \infty} \frac{1}{N} \log \mathbb{P}^N(F_\delta(\phi^\eta))
\]

\[
\geq -I_T(\phi^\eta)
\]

\[
\geq -I - \eta
\]

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and then
\[ \liminf_{N \to \infty} \frac{1}{N} \log \mathbb{P}^N(G) \geq -I. \]

Specifying the starting point, we can reformulate the above result as
\[ \liminf_{N \to \infty} \frac{1}{N} \log \mathbb{P}(Z^{N,x} \in G) \geq -\inf_{\phi \in G, \phi_0 = x} I_T(\phi). \]

We need in fact the stronger statement

**Theorem 46.** For all open set \( G \in D_{T,A} \) such that all trajectories in \( G \) remain in a compact set which does not intersect the boundary \( \partial A \), for any compact set \( K \subset A \),
\[ \liminf_{N \to \infty} \frac{1}{N} \log \inf_{x \in K} \mathbb{P}(Z^{N,x} \in G) \geq -\sup_{x \in K} \inf_{\phi \in G, \phi_0 = x} I_T(\phi). \]

### 18.5 The Upper Bound

For all \( \phi \in D_{T,A} \) and \( H \subset D_{T,A} \) we define
\[ \rho_T(\phi, H) = \inf_{\psi \in H} \| \phi - \psi \|_T \]
and for all \( \delta, s > 0 \) we define the set
\[ H_\delta(s) = \{ \phi \in D_{T,A} : \rho_T(\phi, \Phi(s)) \geq \delta \} \]
where \( \Phi(s) = \{ \phi \in D_{T,A} : I_T(\phi) \leq s \} \). The main step of the proof of the upper bound consists in establishing the following Lemma.

**Lemma 47.** for any \( \delta, \eta, s > 0 \) there exists \( N_0 \in \mathbb{N} \) such that
\[ \mathbb{P}^N(H_\delta(s)) \leq \exp\{-N(s - \eta)\}\]
whenever \( N \geq N_0 \).

We then deduce

**Theorem 48.** For all closed set \( F \in D_{T,A} \),
\[ \limsup_{N \to \infty} \frac{1}{N} \log \mathbb{P}^N(F) \leq -\inf_{\phi \in F} I_T(\phi). \]
Proof All we have to show is that (7) implies the Theorem. To this end let $F \in D_{T,A}$ a closed set, choose $\eta > 0$ and put $s = \inf I_T(\phi) - \eta/2$. The closed set $F$ does not intersect the compact set $\Phi(s)$. Therefore $\delta = \inf_{\phi \in F} \inf_{\psi \in \Phi(s)} \|\phi - \psi\|_T > 0$. We use the inequality (7) to have for any $\eta, s > 0$ there exists $N_0 \in \mathbb{N}$ such that for all $N > N_0$,

$$\mathbb{P}^N(F) \leq \mathbb{P}^N(H_\delta(s)) \leq \exp\{-N(s - \eta/2)\} \leq \exp\{-N(\inf_{\phi \in F} I_T(\phi) - \eta)\}$$

then

$$\limsup_{N \to \infty} \frac{1}{N} \mathbb{P}^N(F) \leq \inf_{\phi \in F} I_T(\phi).$$

We need a slightly stronger version

Theorem 49. For all closed set $F \in D_{T,A}$ such that all trajectories in $F$ remain in a compact set which does not intersect the boundary $\partial A$, for any compact set $K \subset \hat{A}$,

$$\limsup_{N \to \infty} \frac{1}{N} \log \sup_{x \in K} \mathbb{P}(Z^N,x \in F) \leq - \inf_{x \in K} \inf_{\phi \in F, \phi_0 = x} I_T(\phi).$$

18.6 Time of exit from a domain

We let $O \subset A$ be relatively open in $A$ (with $O = \tilde{O} \cap A$ for $\tilde{O} \subset \mathbb{R}^d$ open) and $x^* \in O$ be a stable equilibrium of (2). By a slight abuse of notation, we say that

$$\tilde{\partial}O := \partial \tilde{O} \cap A$$

is the boundary of $O$. For $y, z \in A$, we define the following functionals.

$$V(x, z, T) := \inf_{\phi \in D([0,T];A), \phi(0) = x, \phi(T) = z} I_T(x)$$

$$V(x, z) := \inf_{T > 0} V(x, z)$$

$$\tilde{V} := \inf_{z \in \tilde{\partial}O} V(x^*, z).$$

In other words, $\tilde{V}$ is the minimal energy required to leave the domain $O$ when starting from $x^*$. We urge the reader to consider the two examples in section 18.6.4.
Assumptions B

B1 $x^\ast$ is the only stable equilibrium point of (2) in $O$ and the solution $Y^x$ of (2) with $x = Y^x(0) \in O$ satisfies

$$Y^x(t) \in O \text{ for all } t > 0 \text{ and } \lim_{t \to \infty} Y^x(t) = x^\ast.$$  

B2 For a solution $Y^x$ of (2) with $x = Y^x(0) \in \tilde{\partial}O$, we have

$$\lim_{t \to \infty} Y^x(t) = x^\ast.$$  

B3 $V < \infty$.

B4 For all $\rho > 0$ there exist constants $T(\rho), \epsilon(\rho) > 0$ with $T(\rho), \epsilon(\rho) \downarrow 0$ as $\rho \downarrow 0$ such that for all $z \in \tilde{\partial}O \cup \{x^\ast\}$ and all $x, y \in B(z, \rho) \cap A$ there exists an

$$\phi = \phi(\rho, x, y) : [0, T(\rho)] \to A \text{ with } \phi(0) = x, \phi(T(\rho)) = y \text{ and } I_{T(\rho)}(\phi) < \epsilon(\rho).$$

B5 For all $z \in \tilde{\partial}O$ there exists an $\eta_0 > 0$ such that for all $\eta < \eta_0$ there exists a $\tilde{z} = \tilde{z}(\eta) \in A \setminus \bar{O}$ with $|z - \tilde{z}| > \eta$.

Let us shortly comment on Assumption B By B1, $O$ is a subset of the domain of attraction of $x^\ast$. B2 is violated by the applications we have in mind: we are interested in situations where $\tilde{\partial}O$ is the characteristic boundary of $O$, i.e., the boundary separating two regions of attraction of equilibria of (2). In order to relax this assumption, we require an approximation argument later. By B3, it is possible to reach the boundary with finite energy. This assumption is always satisfied for the epidemiological models we consider. For $z = x^\ast$, B4 is also always satisfied in our models as the rates $\beta_j$ are bounded from above and away from zero in small neighborhoods of $x^\ast$; hence, the function $\phi(x, y, \rho)$ can, e.g., be chosen to be linear with speed one.

We are interested in the following quantity:

$$\tau^{N,x} := \tau^N := \inf\{t > 0 | Z^{N,x}(t) \not\in O\},$$

i.e., the first time that $Z^{N,x}$ exits $O$. 

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18.6.1 Auxiliary results

Assumptions A4 + B4 yield.

Lemma 50. Assume that Assumptions A and B hold. Then for any \( \delta > 0 \), there exists an \( \rho_0 > 0 \) such that for all \( \rho < \rho_0 \),

\[
\sup_{z \in \partial \Omega \cup x^*, x, y \in B(z, \rho)} \inf_{T \in [0, 1]} V(x, y, T) < \delta.
\]

We have moreover.

Lemma 51. Assume that Assumptions A and B hold. Then, for any \( \eta > 0 \) there exists a \( \rho_0 > 0 \) such that for all \( \rho < \rho_0 \) there exists a \( T_0 < \infty \) such that

\[
\liminf_{N \to \infty} \frac{1}{N} \log \inf_{x \in B(x^*, \rho)} \mathbb{P}[T^N \leq T_0] > -(\bar{V} + \eta).
\]

We also require the following result

Lemma 52. Assume that Assumptions A and B hold. Let \( \rho > 0 \) such that \( B(x^*, \rho) \subset O \) and

\[
\sigma^N_\rho := \inf \{ t > 0 | Z^N_\rho \in \overline{B(x^*, \rho)} \text{ or } Z^N_\rho \notin O \}.
\]

Then

\[
\lim_{t \to \infty} \limsup_{N \to \infty} \frac{1}{N} \log \sup_{x \in O} \mathbb{P}[\sigma^N_\rho > t] = -\infty.
\]

Lemma 53. Assume that Assumptions A and B hold. Let \( C \subset A \setminus O \) be closed. Then,

\[
\lim_{\rho \to 0} \limsup_{N \to \infty} \frac{1}{N} \log \sup_{x \in B(x^*, 3\rho) \setminus B(x^*, 2\rho)} \mathbb{P}[Z^N_{\sigma_\rho} \in C] \leq -\inf_{z \in C} V(x^*, z).
\]

We have moreover

Lemma 54. Assume that Assumptions A and B hold. Then, for all \( \rho > 0 \) such that \( B(x^*, \rho) \subset O \) and for all \( x \in O \),

\[
\lim_{N \to \infty} \mathbb{P}[Z^N_{\sigma_\rho} \in \overline{B(x^*, \rho)}] = 1.
\]

Lemma 55. Assume that Assumptions A and B hold. Then, for all \( \rho, c > 0 \), there exists a constant \( T = T(c, \rho) < \infty \) such that

\[
\limsup_{N \to \infty} \frac{1}{N} \log \sup_{x \in O} \mathbb{P}[ \sup_{t \in [0, T]} |Z^N_t - x| \geq \rho] < -c.
\]
18.6.2 Main results

We can now establish

Theorem 56. Assume that Assumptions A and B hold. Then, for all \( x \in O \) and \( \delta > 0 \),

\[
\lim_{N \to \infty} \mathbb{P}\left[ e^{(\tilde{V} - \delta)N} < \tau^{N,x} < e^{(\tilde{V} + \delta)N} \right] = 1.
\]

18.6.3 The case of a characteristic boundary

Since we are mainly interested in studying the time of exit from the basin of attraction of one local equilibrium to that of another, we need to consider situations which do not satisfy the above assumptions. More precisely, we want to suppress the assumptions B3 and B5, and keep assumptions B1, B2 and B4. In the examples which we have in mind, there exists a collection of open sets \( \{O_{\rho}, \rho > 0\} \) which is such that

- \( \overline{O}_{\rho} \subset O \) for any \( \rho > 0 \).
- \( d(O_{\rho}, \partial O) \to 0 \) as \( \rho \to 0 \).
- \( O_{\rho} \) satisfies assumptions B1,...,B5 for any \( \rho > 0 \).

Let then \( O \) be a domain satisfying assumptions (B1, B2 and B4, and we assume that there exist a sequence \( \{O_{\rho}, \rho > 0\} \) satisfying the three above conditions.

If we define \( \tilde{V}_{\rho} \) as \( \tilde{V} \), but with \( O \) replaced by \( O_{\rho} \), it follows from Lemma 50 that \( \tilde{V}_{\rho} \to \tilde{V} \) as \( \rho \to 0 \). By an obvious monotonicity property, the lower bound

\[
\lim_{\epsilon \to 0} \mathbb{P}[\tau^{N,x} > e^{(\tilde{V} - \delta)N}] = 1
\]

follows immediately from Theorem 56.

18.6.4 Applications

Consider the following two epidemiological models with several equilibria, both
1. the SIV model studied by Kribs–Zaleta and Velasco–Hernández:
\[
\frac{ds}{dt}(t) = \mu(1 - s(t)) + \alpha i(t) - \beta s(t)i(t) - \eta s(t) + \theta v(t), \quad t > 0,
\]
\[
\frac{di}{dt}(t) = -\mu i(t) + \beta s(t)i(t) - \alpha i(t) + \gamma \beta v(t)i(t), \quad t > 0,
\]
\[
\frac{dv}{dt}(t) = -\mu v(t) + \eta s(t) - \theta v(t) - \beta v(t)i(t), \quad t > 0;
\]

2. and the $S_0IS_1$ model of Safan, Heesterbeek and Dietz
\[
\frac{ds_0}{dt}(t) = \mu(1 - s_0(t)) - \beta s_0(t)i(t), \quad t > 0,
\]
\[
\frac{di}{dt}(t) = -\mu i(t) + \beta s_0(t)i(t) - \alpha i(t) + \gamma \beta s_1(t)i(t), \quad t > 0,
\]
\[
\frac{ds_1}{dt}(t) = -\mu s_1(t) + \alpha i(t) - \gamma \beta s_1(t)i(t), \quad t > 0.
\]

In those two above models, one can choose the parameters in such a way that both the DFE and one of the endemic equilibria are locally stable. Denote by $\mathcal{O}$ the basing of attraction of the endemic equilibrium. Let us denote by $\tau^{N,x}$ the time it takes for the stochastic system, starting from $x \in \mathcal{O}$, to exit $\mathcal{O}$ ($\approx$ the time to reach the DFE). Theorem 56 extended to the case of a characteristic boundary implies that For any $x \in \mathcal{O}$, $\delta > 0$,
\[
\lim_{N \to \infty} \Pr(e^{(\overline{V} - \delta)N} < \tau^{N,x} < e^{(\overline{V} + \delta)N}) = 1.
\]

**Numerical computation of $\overline{V}$**

1. In the SIV model with $\beta = 3.6$, $\alpha = 1$, $\theta = 0.02$, $\mu = 0.03$, $\eta = 0.3$ and $r = 0.1$, we get $\overline{V} = 0.39$.
   This gives rather astronomical values of $\tau^N$, even for $N = 100$ !

2. In the $S_0IS_1$ model with $\beta = 3$, $\alpha = 5$, $\mu = 0.015$ and $r = 2$, we get $\overline{V} = 0.0745$.
   This means that for $N = 100$, $\tau^N \approx 1720$, and for larger $N$, the value of $\tau^N$ is huge !

3. We have not yet checked how $\overline{V}$ depends upon the parameters !
It would be interesting to understand how those results would be modified if we incorporate heterogeneity (nonhomogeneous mixing, spatial dispersion, ...).

References


