

# Lecture 7: SIR Models

**Radu Balan**

Department of Mathematics, NWC  
University of Maryland, College Park, MD

Version: February 20, 2023

# Epidemiological Models

In this lecture we continue discussing epidemiological models. There are two main types of epidemic models:

- deterministic (or, compartmental) model
- stochastic (e.g., agent based) model

We focus on three deterministic models:

- 1 SI (Susceptible-Infected) Model
- 2 SIR (Susceptible-Infected-Removed) Model
- 3 SEIR (Susceptible-Exposed-Infected-Removed) Model

Today we discuss the SIR model.

## The SIR Model with No vitals

Assume a system with three compartments: 'Susceptible' (S), 'Infected' (I) and 'Removed' or 'Recovered' (R). At time  $t_0 = 0$  the system has a total of  $N$  individuals (initial total population). Most of them are susceptible  $S(0)$ , but some are infected,  $I(0)$  and possibly some are in the recovered state,  $R(0)$ . Our intention is to model the time evolution of these populations. We start with SI model:

$$\begin{cases} \frac{dS}{dt} = -\beta S \frac{I}{N}, & S(0) \\ \frac{dI}{dt} = \beta S \frac{I}{N}, & I(0) \end{cases}$$

where  $\beta \geq 0$  is a parameter. We append a new term to model transition from  $I \mapsto R$ , assuming a constant rate of transition  $\alpha$ :

$$\begin{cases} \frac{dS}{dt} = -\beta S \frac{I}{N}, & S(0) \\ \frac{dI}{dt} = \beta S \frac{I}{N} - \alpha I, & I(0) \\ \frac{dR}{dt} = \alpha I, & R(0) \end{cases}$$

# The SIR Model

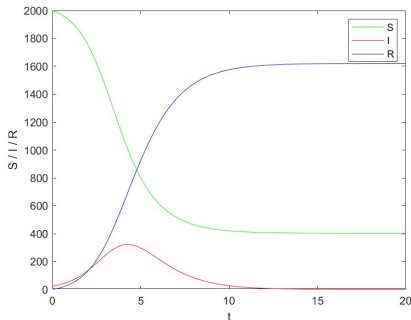
## Deterministic simulations

Simulation of the SIR model:

$$\beta = 2, \alpha = 1, S(0) = 2000, I(0) = 23, R(0) = 0$$

Results were obtained with an Euler scheme with step size  $h = 0.01$ .

Note: The infected population  $I(t)$  first increases and then decreases eventually to 0. The susceptible population decreases, but converges to some limiting value  $S(\infty) > 0$ . The removed population is monotone increasing and converges to some value  $R(\infty) < N$ . Some of the susceptible population who do not get infected are protected by the recovered population surrounding them. This is known as *herd immunity*.



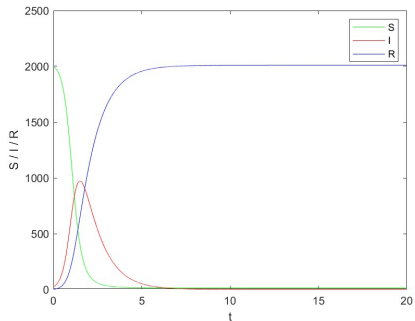
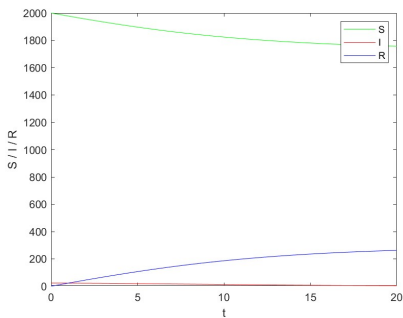
# The SIR Model

## Deterministic simulations (2)

Initial conditions:  $S(0) = 2000$ ,  $I(0) = 23$ ,  $R(0) = 0$ .

$$\beta = 1, \quad \alpha = 1.$$

$$\beta = 5, \quad \alpha = 1.$$



## The SIR Model: The normalized form

For reasons of normalizations, we prefer to compute *fractions* of susceptible population, infected population, and of removed population:

$$s(t) = \frac{S(t)}{N} \quad , \quad i(t) = \frac{I(t)}{N} \quad , \quad r(t) = \frac{R(t)}{N}$$

In which case the model becomes:

$$\begin{cases} \frac{ds}{dt} = -\beta si \quad , \quad s(0) = \frac{S(0)}{N} \\ \frac{di}{dt} = \beta si - \alpha i \quad , \quad i(0) = \frac{I(0)}{N} \\ \frac{dr}{dt} = \alpha i \quad , \quad r(0) = \frac{R(0)}{N} \end{cases} \quad (\text{SIR Model})$$

Note  $s(t) + i(t) + r(t) = 1$  for all  $t$  (conservation of total population).  
The bad news: there is no closed form solution. The good news: some relationships can be expressed in closed form.

# The SIR Model: Numerical solution vs. Agent Based Modeling

Similar to the SI model, one way of implementing an agent based simulation is to pretend the nonlinear term is linear in  $s$   $\tilde{\beta}(t) = \beta i(t)$ .

Thus the rate matrix is given by

$$\frac{d}{dt} \begin{bmatrix} s \\ i \\ r \end{bmatrix} = A \begin{bmatrix} s \\ i \\ r \end{bmatrix}, \quad A = \begin{bmatrix} -\beta i & 0 & 0 \\ \beta i & -\alpha & 0 \\ 0 & \alpha & 0 \end{bmatrix}$$

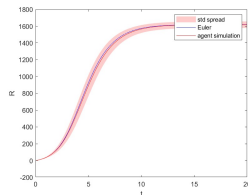
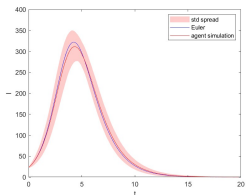
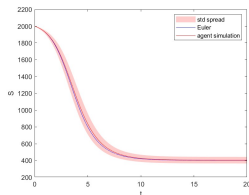
and for a discretization step  $T_0$ , at time step  $p > 0$ , the transition matrix to transition from time  $(p-1)T_0$  to  $pT_0$  with  $i_{p-1} = i((p-1)T_0)$  is given by

$$\Pi(p) = \begin{bmatrix} e^{-\beta T_0 i_{p-1}} & 0 & 0 \\ \frac{\beta i_{p-1}}{\alpha - \beta i_{p-1}} e^{-\beta T_0 i_{p-1}} - \frac{\beta i_{p-1}}{\alpha - \beta i_{p-1}} e^{-\alpha T_0} & e^{-\alpha T_0} & 0 \\ 1 - \frac{\alpha}{\alpha - \beta i_{p-1}} e^{-\beta T_0 i_{p-1}} + \frac{\beta i_{p-1}}{\alpha - \beta i_{p-1}} e^{-\alpha T_0} & 1 - e^{-\alpha T_0} & 1 \end{bmatrix}, \quad p = 1, 2, \dots$$

# The SIR Model

## Numerical Solution vs. Agent Based Modeling (2)

Here are results of a large number of simulations ( $10^3$ ) for  $T_0 = 0.01$ , with parameters  $\beta = 2$  and  $\alpha = 1$ . The numerical solution is obtained with the Euler scheme and a stepsize  $h = 0.001$ . The shaded area has semiwidth of one std of simulations

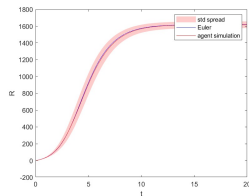
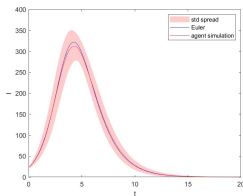
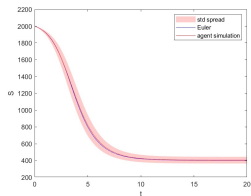




# The SIR Model

## Numerical Solution vs. Agent Based Modeling (3)

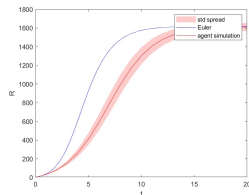
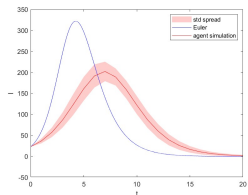
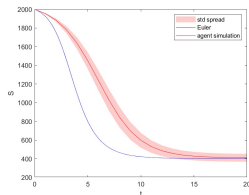
Here are results of a large number of simulations ( $10^3$ ) for  $T_0 = 0.001$ , with parameters  $\beta = 2$  and  $\alpha = 1$ . The numerical solution is obtained with the Euler scheme and a stepsize  $h = 0.001$ . The shaded area has semiwidth of one std of simulations



# The SIR Model

## Numerical Solution vs. Agent Based Modeling (4)

Here are results of a large number of simulations ( $10^3$ ) for  $T_0 = 1.0$ , with parameters  $\beta = 2$  and  $\alpha = 1$ . The numerical solution is obtained with the Euler scheme and a stepsize  $h = 0.001$ . The shaded area has semiwidth of one std of simulations



# The SIR Model

## Analytic expressions

For the normalized systems of equations, divide the equation for  $i$  by equation for  $s$  and use chain rule:

$$\frac{di}{ds} = -1 + \frac{\alpha}{\beta} \frac{1}{s}$$

The ratio  $R_0 = \frac{\beta}{\alpha}$  is known as the *reproduction ratio*, or the *contact number*. Its meaning:  $\beta$  represents the number of close contacts per day per one infected individual;  $\frac{1}{\alpha}$  is the average infectious period (or, the average number of days an infected person remains contagious). Hence  $R_0$  represents the average number of close contacts per infected individual. Use separability of this Diff Eq. and integrate both sides:

$$i(t) - i(0) = s(0) - s(t) + \frac{1}{R_0} (\log(s(t)) - \log(s(0)))$$

Thus  $i + s - \frac{1}{R_0} \log(s)$  must stay constant over time.

# The SIR Model

## Analytic calibrations

We obtained that  $i + s - \frac{1}{R_0} \log(s)$  must stay constant over time.

*Approximation:* At time  $t = 0$ ,  $s(0) \approx 1$  (assuming little infections and recovered people) and  $i(0) \approx 0$  (very few infected people compared to the total population).

$$i(t) + s(t) - \frac{1}{R_0} \log(s(t)) = 1$$

What happens for  $t \rightarrow \infty$ ?

One thing for sure:  $i(\infty) = 0$ . What happens with  $s(\infty)$ ? We obtain the following equations:

$$1 = s(\infty) - \frac{1}{R_0} \log(s(\infty))$$

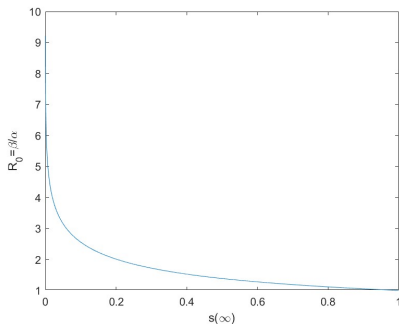
$$R_0 = \frac{\log(s(\infty))}{s(\infty) - 1}$$

# The SIR Model

## Herd Immunity

With the approximation  $i(0) = 0$ ,  $s(0) = 1$ , the plot of  $\log(s(\infty))/(s(\infty) - 1)$  as function of  $s(\infty)$  is rendered in the left figure.

For instance, if the contact number is  $R_0 = 2$ , then  $s(\infty) \approx 0.2$ . Thus 20% of population get protection from the 80% who have gotten infected and recovered.

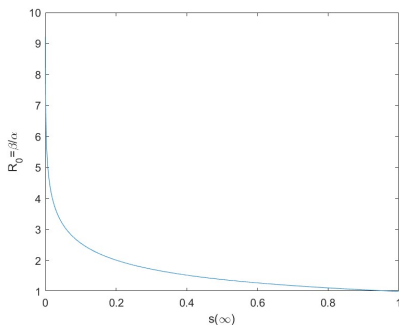


# The SIR Model

## Herd Immunity

With the approximation  $i(0) = 0$ ,  $s(0) = 1$ , the plot of  $\log(s(\infty))/(s(\infty) - 1)$  as function of  $s(\infty)$  is rendered in the left figure.

For instance, if the contact number is  $R_0 = 2$ , then  $s(\infty) \approx 0.2$ . Thus 20% of population get protection from the 80% who have gotten infected and recovered.



But what happens if  $R_0 < 1$  ?

# The SIR Model

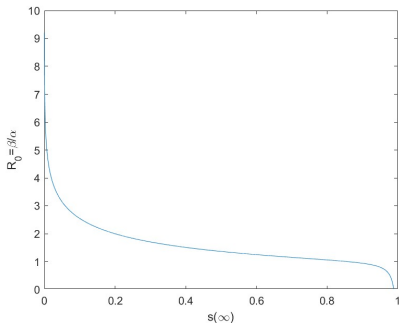
## Herd Immunity (2)

In reality,  $i(0) > 0$  and  $s(0) < 1$ . Assume that still  $r(0) = 0$ , thus  $s(0) + i(0) = 1$ , but  $\log(s(0)) < 0$ . We obtain:

$$i(t) + s(t) - \frac{1}{R_0} \log(s(t)) = 1 - \frac{\log(s(0))}{R_0} \Rightarrow R_0 = \frac{\log(s(\infty)) - \log(s(0))}{s(\infty) - 1}$$

For  $s(0) = 2000/2023$  we obtain the left plot. For  $R_0 \gg 1$ , the previous approximation is still good.

For  $R_0 < 1$ , it follows that a significant number of susceptible individual do not get infected.



# SIR Model with Two Outcomes

Since  $I(t)$ , or  $i(t)$ , is not monotone increasing sequence, the SIR model is appropriate for the time series of the *daily rates* of the number of infections. On the hand, the removed sequence  $R(t)$ , or  $r(t)$ , is monotone increasing. The “removed” compartment contains only two types of individuals: (1) individuals that recovered and gained full immunity  $X(t)$ , (so they will never get infected again), and (2) people who died,  $Y(t)$ . Thus  $R(t) = X(t) + Y(t)$ .

*Assumption:*  $Y(t) = \gamma R(t)$  for all  $t$ . In other words, a fixed fraction  $\gamma$  of people who get infected eventually die, with the same infectious period as the individuals that eventually recovered and gained immunity:

$$\left\{ \begin{array}{l} \frac{dS}{dt} = -\beta S \frac{I}{N}, \quad S(0) \\ \frac{dI}{dt} = \beta S \frac{I}{N} - \alpha I, \quad I(0) \\ \frac{dX}{dt} = (1 - \gamma)\alpha I, \quad X(0) = (1 - \gamma)R(0) \\ \frac{dY}{dt} = \gamma\alpha I, \quad Y(0) = \gamma R(0) \end{array} \right.$$

Note: For obtaining  $(S, I, X, Y)$  you need only to solve the SIR system and find  $S(t), I(t), R(t)$ , and then allocate,  $X(t) = (1 - \gamma)R(t)$  and  $Y(t) = \gamma R(t)$ .



## Two-outcome SIR Model Analysis

The two-outcome SIR model has the form:

$$\left\{ \begin{array}{l} \frac{dS}{dt} = -\beta S \frac{I}{N}, \quad S(0) \\ \frac{dI}{dt} = \beta S \frac{I}{N} - \alpha I, \quad I(0) \\ \frac{dX}{dt} = (1-\gamma)\alpha I, \quad X(0) = (1-\gamma)R(0) \\ \frac{dY}{dt} = \gamma\alpha I, \quad Y(0) = \gamma R(0) \end{array} \right.$$

Its normalized form in variables  $s, i, r, x = \frac{X}{N}, y = \frac{Y}{N}$  is given by:

$$\left\{ \begin{array}{l} \frac{ds}{dt} = -\beta si, \quad s(0) = \frac{S(0)}{N} \\ \frac{di}{dt} = \beta si - \alpha i, \quad i(0) = \frac{I(0)}{N} \\ \frac{dx}{dt} = (1-\gamma)\alpha i, \quad x(0) = (1-\gamma)\frac{R(0)}{N} \\ \frac{dy}{dt} = \gamma\alpha i, \quad y(0) = \gamma\frac{R(0)}{N} \end{array} \right.$$

Note the conservation laws:  $N = S(t) + I(t) + X(t) + Y(t)$ ,  
 $s(t) + i(t) + x(t) + y(t) = 1$ . The deterministic system is initialized by  
 $(S(0), I(0), R(0))$  and its evolution is determined by the choice of three  
 parameters:  $(\alpha, \beta, \gamma)$ .  $X(t)$  and  $Y(t)$  are computed from  $R(t)$

# SIR Model Analysis

The three parameters have the following meaning:

- 1  $\beta$  represents the number of close contacts per day per one infected individual; differently said, it is the probability of disease transmission per contact (dimensionless) times the number of contacts per unit of time. Unit:  $day^{-1}$
- 2  $\alpha$  is the removing rate of infectious individuals; its reciprocal is the infectious period. If no death,  $\alpha$  represents the recovery rate from infections. Unit:  $day^{-1}$
- 3  $\gamma$  represents the probability of fatal infection once an individual gets infected. Unit: dimensionless.

In addition, we defined  $R_0 = \frac{\beta}{\alpha}$  as the reproduction ratio (or, the contact number) that represents the average number of infections caused by one infected individual.

Note:  $\alpha$  and  $\gamma$  are parameters that characterize the infectious disease and cannot be controlled. Instead,  $\beta$  characterizes human interactions, and therefore can be controlled by individuals (e.g., during the shut-down of 2020,  $\beta \approx 0$ ).

# SIR Model Calibration

For calibration and testing we are using two pieces of measured data: the *daily infection rates*,  $\{I(0), \dots, I(T_{max})\}$ , and the time series of *cumulative deaths*,  $\{Y(0), \dots, Y(T_{max})\}$ . Note that, if we know  $\gamma$  and  $N$  we can compute  $R(0) = \frac{Y(0)}{\gamma}$  and  $S(0) = N - I(0) - R(0)$ . At the onset of an infectious disease it is likely the case that  $Y(0) = 0$  and  $I(0)$  can be neglected in which case,  $S(0) = N$  (regardless of  $\gamma$ ).

The least-squares estimator (LSE) tries to find parameters  $\alpha, \beta, \gamma$ , and  $N$  that minimize:

$$\text{minimize}_{N \in \mathbb{N}} \quad I(\alpha, \beta, \gamma; N) := \sum_{t=0}^{T_{max}} (I(t) - I_{sim}(t))^2 + (Y(t) - \gamma R_{sim}(t))^2$$

$$\alpha, \beta, \gamma \geq 0, \gamma \leq 1$$

where  $(S_{sim}(t), I_{sim}(t), R_{sim}(t))$  are given by a numerical solver of the SRI model with parameters  $(\alpha, \beta, \gamma)$  and total population  $N$  initialized at  $(S(0), I(0), R(0))$ .

## SIR Model with Vitals

A simple modification of the SIR vanilla model is to consider vital signals, such as births and deaths at separate processes. In normalized form this becomes:

$$\begin{cases} \frac{ds}{dt} = \frac{\Lambda}{N} - \beta si - \mu s & , \quad s(0) = \frac{S_0}{N} \\ \frac{di}{dt} = \beta si - \alpha i - \mu i & , \quad i(0) = \frac{I_0}{N} \\ \frac{dr}{dt} = \alpha i - \mu r & , \quad r(0) = \frac{R_0}{N} \end{cases} \quad (\text{SIR Model})$$

where  $\Lambda \geq 0$  is the constant source of births (=number of births/day) and  $\mu \geq 0$  is the natural death rate (i.e., in the absence of this virus). Its reciprocal  $1/\mu$  represents the average life expectancy.