# **Modelling Epidemics**

#### **Introduction to Network Science**

Instructor: Michele Starnini — https://github.com/chatox/networks-scienc

# Content

- Compartmental epidemiological models
- Results on homogenepus mixing scenarios

# E[xamples: human epidem](https://en.wikipedia.org/wiki/Cocoliztli_Epidemic_of_1545-1548)i

●Influenza, measles, STIs, ...

●Smallpox and other diseases brought by Europeans to America since early 1500s

.The "Black Death" (next slide)



https://en.wikipedia.org/wiki/Cocoliztli\_Epidemic\_of\_1545-1548

The "Black Death" (Bubonic plague) 1300s

Probably originated in Central Asia, it spread throughout all of Europe between 1346 and 1353. The Black Death is estimated to have killed 30-60% of Europe's population



1347 1348  $1349$ 1346

Approximate border between the Principality of Kiev and the Golden Horde - passage prohibited for Christians.

- 2008

Land trade routes

 $1350 > 1351 > 1352$ 



Maritime trade routes

1353

# S[ARS Outbreak \(2003\)](https://en.wikipedia.org/wiki/Timeline_of_the_SARS_outbreak)

 $\cdot$ February 21st: Chinese doctor who have bee several treating "atypical pneumonia" cases check-ins into hotel in Hong Kong

–Hospitalized on Feb 22nd

–Died on March 4th

•March 1st: "Ms. E. M." returns to Singapore after visiting Hong Kong

–Graph depicts 144 out of the first 206 SARS patients in Singapore

–Ms. E. M. lived, various of her family members died





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## COVID vs Black Death

- The Black Death spread across Europe in **years**
- Covid spread across the world in **weeks**
- Covid reached even Antardide & remote islands
- What changed?

# Modeling epidemics

- **Modeling an epidemic process (dynamics):**
- Contagiousness
- Length of infectious period,
- Severity
- ...
- **Modeling underlying network substrate** (static)
- Structure of contacts in a population

# Modeling epidemics

- **Modeling Epidemic process (dynamics):**
- Branching process
- SI model
- SIR model
- SIS model
- **Modeling underlying network substrate (static)**
- Mean-field mixing (fully connected network)
- Homogeneous networks (ER networks)
- –Heterogeneous networks (SF networks)

## Diffusion of ideas vs diseases

Adopting a new idea, behavior, fashion, product, taste, may also spread from person to person: **"social contagion"**

. There is a certain agency of the receiver

●In diffusion of diseases, we assume **there is no agency: each contagion is random**

## Epidemic model: branching process

## Branching process

. Each person interacts with other *k* people

•Each interaction ends in infection with probability *beta*



Example: k=3

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# Transmission rate or "Basic reproductive number"  $R_0$

- Each person interacts with other  $k$  people
- Each interaction ends in infection with probability  $\beta$

. What is the expected number of cases caused by a single individual,  $R_0$ ?

- What do you think happens if  $R_0 < 1$ ?
- What do you think happens if  $R_0 > 1$ ?

## Epidemic threshold

- •The **basic reproductive number** is the average number of individuals that are infected by an infectious one during his infectious period, in a fully susceptible population.
- •An epidemic outbreak **can occur only if** the basic reproductive number is **larger than 1**.



# Changing *R0 = beta k*

#### ●**Sanitary practices** (to reduce what?)

#### ●**Quarantine** (to reduce what?)

#### SARS-CoV-2





## The SI model

#### ●**Susceptible node:**

#### **INFECTION**  $\beta$ **INFECTED ISICKI**

#### – each contact with an infected node can result in infection with probability **beta**

#### ●**Infected node:**

- has the disease and can spread it
- will stay sick forever

## Notation

- ●Number of susceptible S(t)
- –Fraction of susceptible  $s(t) = S(t) / N$
- ●Number of infected I(t)
- –Fraction of infected  $i(t) = I(t) / N$

 $\textsf{.s}(t) + i(t) = 1$ 

## Network substrate

- Let's assume a homogeneous mixing
- All nodes have the same number of neighbors <k>
- example: <k>=4, network is a degree regular d=4
- example: fully connected graph  $\langle k \rangle$  = N-1

#### How many susceptible neighbors a node has?

$$
\braket{k}{\frac{S(t)}{N}} = \braket{k}{s(t)}
$$

#### How many new infections are produced?

(for every infected, iterate through its susceptible neighbors, infect with probability *β*)

 $i(t) \langle k \rangle s(t) \beta$ 

infected susceptible neighbors infection probability

**Prove that** 
$$
i(t) = \frac{i_0 e^{\beta \langle k \rangle t}}{1 - i_0 + i_0 e^{\beta \langle k \rangle t}}
$$
  
\n**Begin from:**  $\frac{di(t)}{dt} = i(t) \langle k \rangle (1 - i(t)) \beta$ 

First, place all terms with  $i(t)$  on the left side

Second, use  $\frac{1}{x \cdot (1-x)} = \frac{1}{x} + \frac{1}{1-x}$ 

Third, integrate from  $t = 0$  to t and denote by  $i_0 = i(t = 0)$ 

$$
\int \frac{1}{x} dx = \log x + C \qquad \int \frac{1}{1-x} dx = -\log(1-x) + C
$$

#### Behavior in the limit  $t \rightarrow \infty$

What is the limit of 
$$
i(t) = \frac{i_0 e^{\beta \langle k \rangle t}}{1 - i_0 + i_0 e^{\beta \langle k \rangle t}}
$$
  
when  $t \rightarrow \infty$ ?

●Hint: similar to

$$
f(t) = \frac{e^t}{1 + e^t}
$$

# Infected as a function of time (SI)

$$
i(t) = \frac{i_0 e^{\beta \langle k \rangle t}}{1 - i_0 + i_0 e^{\beta \langle k \rangle t}}
$$

Characteristic time (to infect  $1/e \approx 36\%$  of people):



### The SIS model



## The SIS model

#### ● **Susceptible node:**

– each contact with an infected node can result in infection with probability **beta**

#### ●**Infected node:**

- have the disease and can spread it
- each time step, it recovers with probability **mu** (it becomes susceptible again)



## Infection dynamics

$$
\frac{di(t)}{dt} = \beta \langle k \rangle i(t)(1 - i(t)) - \mu i(t)
$$

 $\mu$  is the recovery rate, i.e., the probability of becoming susceptible again in an unit of time

$$
i(t) = \left(1 - \frac{\mu}{\beta \langle k \rangle}\right) \frac{Ce^{(\beta \langle k \rangle - \mu)t}}{1 + Ce^{(\beta \langle k \rangle - \mu)t}}
$$

•C is a constant that depends on  $i_{0}$ 

### Behavior in the limit  $t \rightarrow \infty$

What is the limit of 
$$
i(t) = \left(1 - \frac{\mu}{\beta \langle k \rangle}\right) \frac{Ce^{(\beta \langle k \rangle - \mu)t}}{1 + Ce^{(\beta \langle k \rangle - \mu)t}}
$$

●Hint: similar to

$$
f(t) = \alpha \frac{e^t}{1 + e^t}
$$



### What happens if  $\mu > \beta \langle k \rangle$

**.Remember:**  $\frac{di(t)}{dt} = \beta \langle k \rangle i(t)(1-i(t)) - \mu i(t)$ 

# Epidemic threshold

- Assumption: early phase of the epidemic, almost all are susceptible  $I \simeq 0, S \simeq N$
- Each infected individual can transmit the disease to <*k*> people at each iteration: expected number of people infected by a single person after one iteration is *β*<*k*>
- If there are *<sup>I</sup>* infected individuals, we expect to have *Isec* <sup>=</sup> *<sup>β</sup>*<*k*>*<sup>I</sup>* new infected people after one iteration and *Irec* = *μI* recovered people
- **Threshold condition for epidemic spreading:**  *Isec* **>** *Irec*

$$
\colon \beta \langle k \rangle I > \mu I \implies R_0 = \frac{\beta}{\mu} \langle k \rangle > 1
$$

 $\bigcap$ 

- *R*<sup>0</sup> = *β*<*k*>/*μ* is the **basic reproduction number**
- If *R*<sup>0</sup> < 1, the **initial outbreak dies out in a short time** if *R*<sup>0</sup> > 1, the **epidemic keeps spreading**

#### The SIR model



# The SIR model



#### ●**Susceptible node:**

– each contact with an infected node can result in infection with probability **beta**

#### ●**Infected node:**

- –The node has the disease and can spread it
- –each time step, it recovers with probability **mu**

#### ●**Removed node:**

– has no longer the disease, and cannot catch it or propagate it again (permanent immunity -or- death)

## Infection dynamics in SIR

$$
\frac{di(t)}{dt} = \beta \langle k \rangle i(t)(1 - r(t) - i(t)) - \mu i(t)
$$
\n
$$
\frac{dr(t)}{dt} = \mu i(t)
$$
\n
$$
\frac{ds(t)}{dt} = -\frac{di(t)}{dt} - \frac{dr(t)}{dt} = -\beta \langle k \rangle i(t)(1 - r(t) - i(t))
$$

#### . No closed form solution

# Infection dynamics (SIR)

$$
\frac{di(t)}{dt} = \beta \langle k \rangle i(t)(1 - r(t) - i(t)) - \mu i(t)
$$

$$
\frac{dr(t)}{dt} = \mu i(t)
$$

$$
\frac{ds(t)}{dt} = -\beta \langle k \rangle i(t)(1 - r(t) - i(t))
$$







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#### More models…



. Can include incubation period

• non-permanent immunity (susceptible again)

• Can include a state for hospitalization

## Things to remember

●SI, SIS, SIR models

Which are the states in each process and which are the possible transitions

. Regimes under different parameters

. Practice executing by hand and write code if it helps you remember better each process

#### Sources

.A. L. Barabási (2016). Network Science – Ch

.D. Easley and J. Kleinberg (2010). Networks, and Markets — Chapter 21

. URLs cited in the footer of slides

#### Practice on your own

Under the SIS model, 
$$
i(t) = \left(1 - \frac{\mu}{\beta \langle k \rangle}\right) \frac{Ce^{(\beta \langle k \rangle - \mu)t}}{1 + Ce^{(\beta \langle k \rangle - \mu)t}}
$$

- 1. When  $\mu < \beta \langle k \rangle$  what is the limit of  $i(t)$ ?
- 2.How is this state called?
- 3. What happens when  $\mu > \beta \langle k \rangle$  ?

4. What conditions lead to large values of  $i(t)$ ?

## Practice on your own (cont.)

In the **SIRS** epidemic model, there are three possible states for a node: susceptible, infected, and recovered. Susceptible nodes can become infected, infected nodes can become recovered, and *recovered nodes can become susceptible again***.**

•During one unit of time, with probability  $\beta$  an infected node can infect one of its contacts, with probability μ, an infected node can recover, and with probability σ, a recovered node can become susceptible again.

•Let s(t) be the fraction of susceptible nodes,  $i(t)$  be the fraction of infected nodes,  $r(t)$ the fraction of recovered nodes, and <k> the average degree of the graph. Write the equations, simplifying them appropriately, for:

$$
1.\frac{di(t)}{dt}\ \ 2.\frac{dr(t)}{dt}\ \ 3.\frac{ds(t)}{dt}
$$

•4. Is  $σ$  >  $μ$  sufficient to say that the recovered will tend to zero in the long run?