# Basics of SIR Models

#### Some Terms

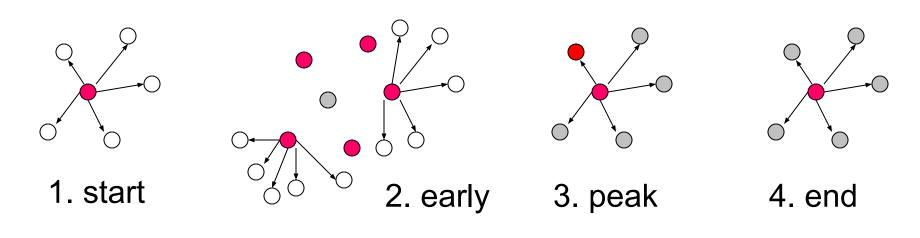
#### Closed v Open populations

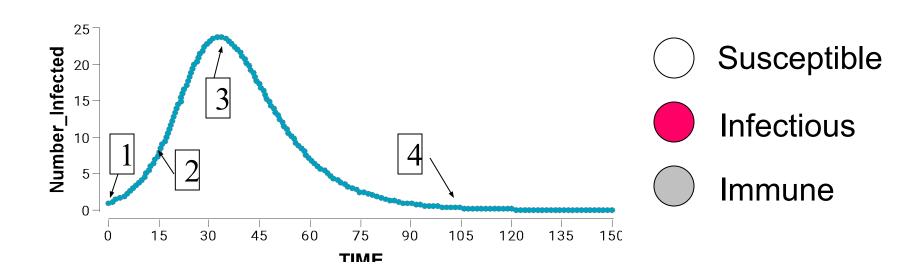
- Closed populations have no additions due to births or immigration and no losses due to death or emigration – all dynamics due *only* to infection
  - Unrealistic, but simple and a good place to start
- Open populations can have births, immigration, deaths, emigration
  - Either explicitly or implicitly

#### **Equilibrium and Transient**

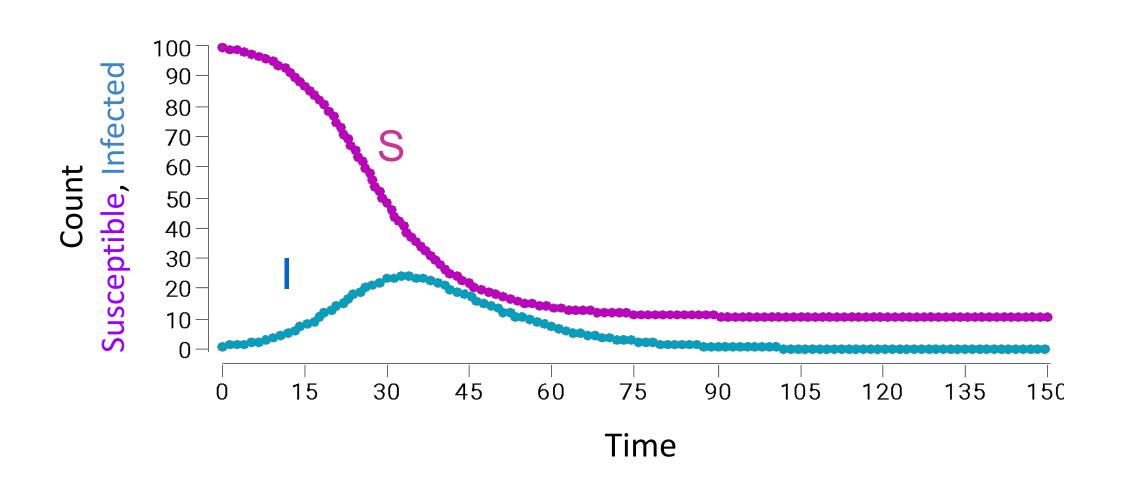
- At equilibrium, the values of all states are constant individuals may still get sick, recover, etc, but all changes balance each other out
- Dynamics are transient if the states are continuing to change ... more nuance later

## Phases of an Epidemic

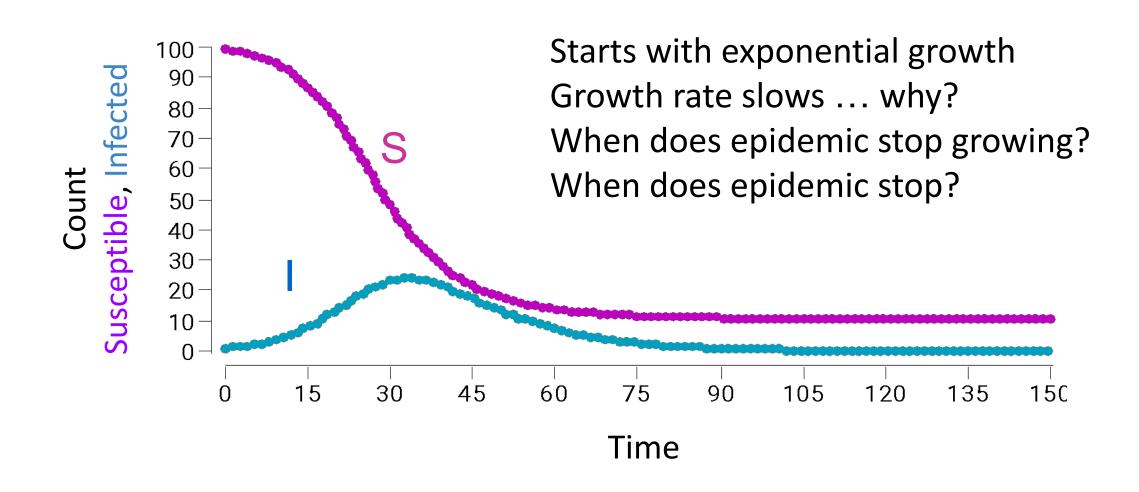




## Idealized Epidemic in a Closed Community



### Idealized Epidemic in a Closed Community



$$\frac{dS}{dt} = -\beta SI$$

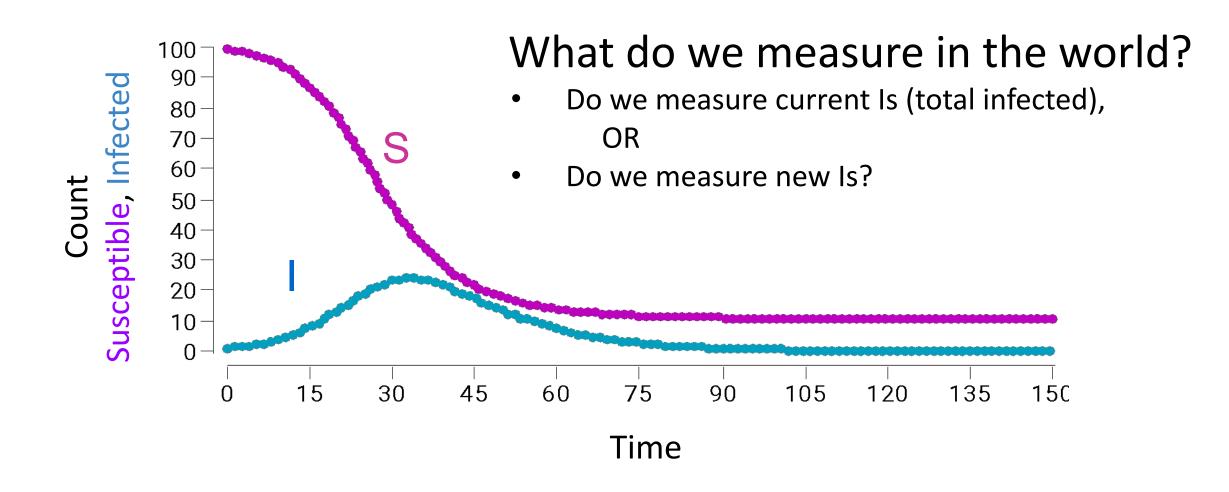
$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

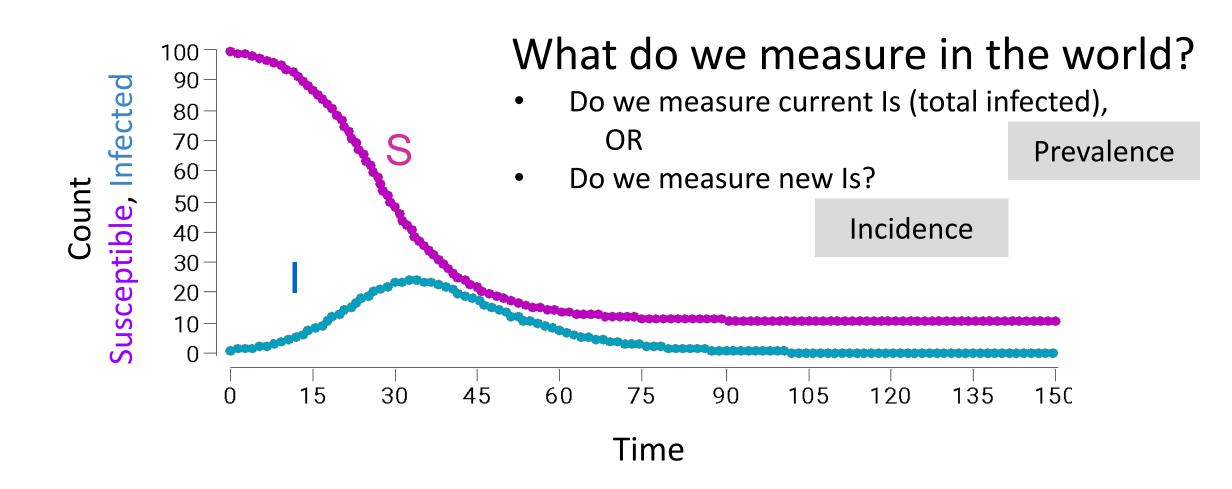
The States

S, I, and R reflect the number of individuals that are currently susceptible, infected (and infectious), and recovered, respectively

### Idealized Epidemic in a Closed Community



## Idealized Epidemic in a Closed Community



$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dI} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

#### What do we measure in the world?

- Do we measure current Is (total infected),
   OR
- Do we measure new Is?

How could we measure current Is? How could we measure new Is?

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

#### **Contact Process**

- This quantifies the rate at which susceptibles and infecteds interact
  - Increases with number (or proportion) of each
  - Creates non-linearity

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

#### **Contact Process**

• There are lots of ways to adjust this ... the most (in)famous of which is:  $S \frac{I}{N}$ 

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

#### **Contact Process**

 What other ways might contacts change with the amount of infection?

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

#### **Transmission Parameter**

- SI defines the shape of contacts.  $\beta$  turns that into infectious contacts:
  - Rate of infectious contacts (not all contacts are infectious)
  - Probability of infection given contact

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dI} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

#### Recovery Rate

- This is the rate, number per time, of recovery (or removal)
- If rate is high, many events happen per unit time and time between events is small

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dI} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

### **Recovery Rate**

- This is the rate, number per time, of recovery (or removal)
- So  $\gamma$  large means that the average duration of infection is short

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dI} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

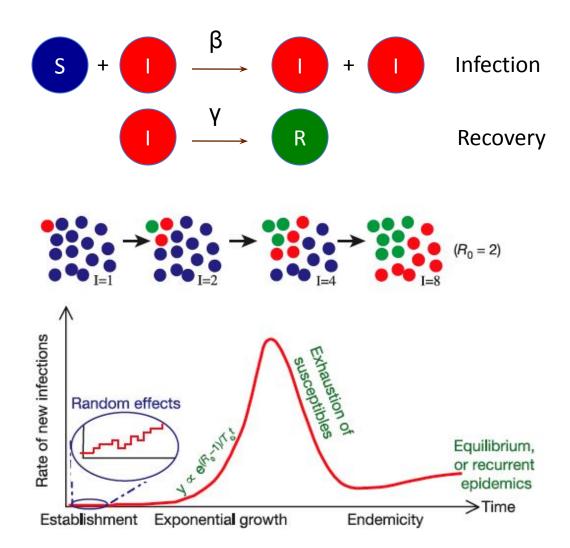
### **Recovery Rate**

- This is the rate, number per time, of recovery (or removal)
- Mean duration of infection,

L, is 
$$\frac{1}{\gamma}$$
 IF the distribution of infectious periods is exponential

How realistic is this? Why do we do this?

## **Basic Epidemic Theory**



$$\frac{dS}{dt} = -\beta SI$$

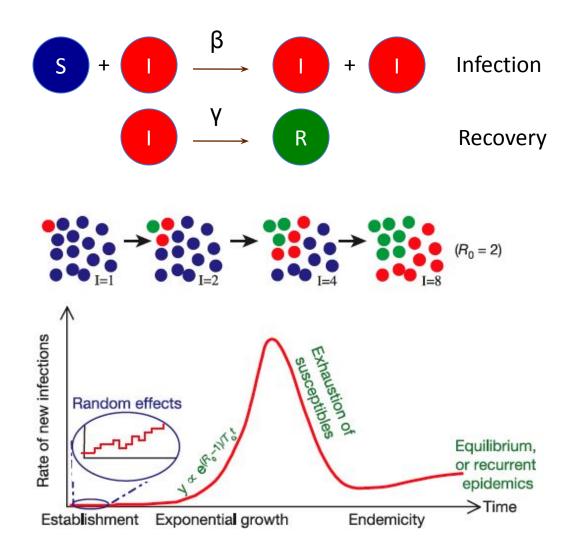
$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

At the start, when there are few infections, an epidemic grows (almost) exponentially

figures from Ferguson et al. *Nature* 2003

## **Basic Epidemic Theory**



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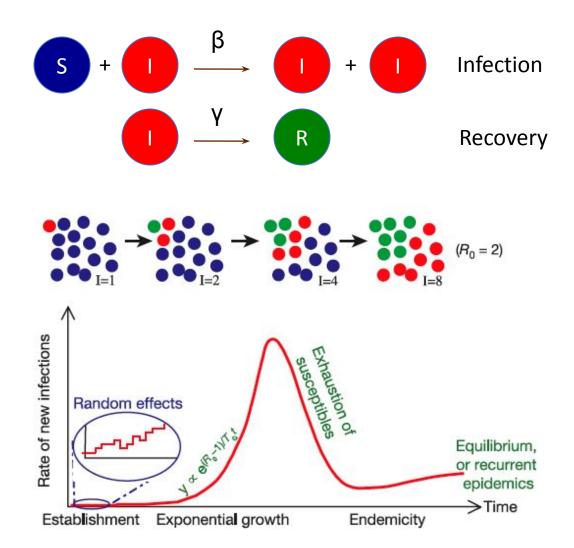
$$\frac{dR}{dt} = -\beta SI - \gamma I$$

At the start, when there are few infections, an epidemic grows (almost) exponentially

We'll use this property later to estimate the transmission rate

figures from Ferguson et al. *Nature* 2003

## **Basic Epidemic Theory**



$$\frac{dS}{dt} = -\beta SI$$

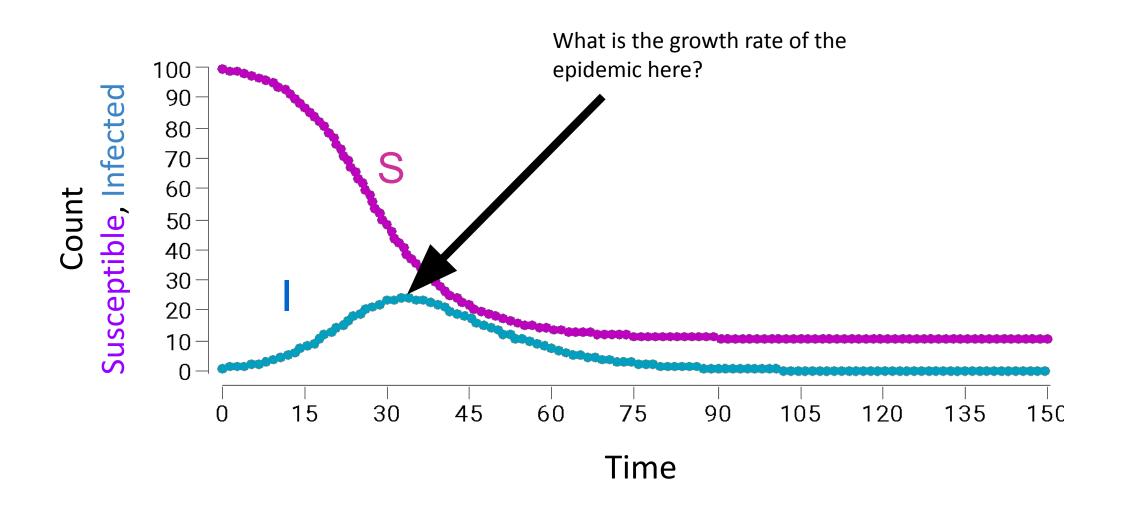
$$\frac{dI}{dt} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

As individuals recover and the number susceptible declines that growth slows because
Susceptibles are being depleted

When does epidemic stop growing?

## Idealized Epidemic in a Closed Community



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$$\frac{dI}{dt} = \beta SI - \gamma I$$

$$0 = \beta SI - \gamma I$$

$$0 = \beta S - \gamma I$$

$$\beta S = \gamma$$

$$1 = \beta S$$

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$$\beta S = \gamma$$

$$1 = \frac{\beta S}{\gamma}$$
Condition under which I doesn't change

$$\frac{dS}{dt} = -\beta SI$$

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$$\frac{dR}{dt} = -\beta SI - \gamma I$$

$$\frac{dI}{dt} = \beta SI - \gamma I$$

$$0 > \beta SI - \gamma I$$

$$0 > \beta S - \gamma$$

$$\beta S < \gamma$$

$$1 > \frac{\beta S}{\gamma}$$
Condition under which I declines ... epidemic fades

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dI} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

$$\frac{dI}{dt} = \beta SI - \gamma I$$

$$0 < \beta SI - \gamma I$$

$$0 < \beta S - \gamma$$

$$\beta S > \gamma$$

$$1 < \frac{\beta S}{\gamma}$$
Condition under which I grows ... epidemic grows

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dI}{dI} = -\beta SI - \gamma I$$

$$\frac{dR}{dt} = -\beta SI - \gamma I$$

Under what conditions can the infectious compartment grow?

$$\frac{dI}{dt} = \beta SI - \gamma I$$

$$0 < \beta SI - \gamma I$$

$$0 < \beta S - \gamma$$

$$\beta S > \gamma$$

$$1 < \beta SL$$
Recall that

Recall that  $\frac{1}{\gamma} = L$  is the mean duration of infection

# R<sub>o</sub>: The Basic Reproduction Number

$$R_0 = \frac{\beta S}{\gamma} = \beta S L$$

The expected number of new infections due to the first infection in a susceptible population

- A common currency
  - A function of the pathogen and the population (recall what  $\beta$  is)
  - Rarely observable directly
  - But closely related to many observable phenomena, as we'll see

### Estimated values of R0 for various infections

Measles	England	1947	13-14
	Nigeria	1968	16-17
	Kansas	1920	5-6
Pertussis	England	1944-78	16-18
	Canada	1912	7-8
Chickenpox	USA	1912	7-8
	USA	1944	10-11

#### What Does This Mean for Interventions?

$$R_0 = \frac{\beta S}{\gamma} = \beta S L$$

What does this suggest for interventions?

- Reduce  $\beta$
- Reduce L (increase gamma)
- Reduce S

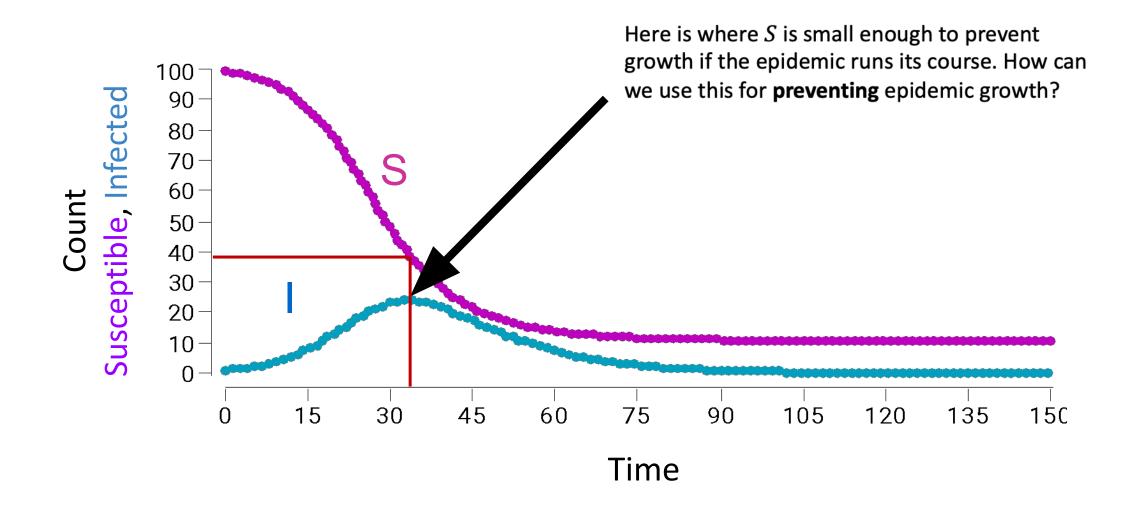
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What does this suggest for interventions?

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- Reduce S

### Idealized Epidemic in a Closed Community



# R<sub>F</sub>: The Effective Reproduction Number

$$R_E = \frac{\beta pS}{\gamma} = \beta pSL$$

p is the fraction susceptible1-p is the fraction immune

What does this suggest for interventions?

- Reduce  $\beta$
- Reduce L (increase gamma)
- Reduce S

The expected number of new infections due to each infection in a population with some immunity

$$R_{0} = \frac{\beta S}{\gamma} = \beta SL$$

$$R_{0} = \beta SL$$

$$1 = \frac{\beta SL}{R_{0}}$$

$$1 = \frac{1}{R_{0}}S\beta L$$

What fraction of Susceptibles need to be immune in order for

$$\frac{1}{R_0}S$$

to remain?

$$R_{0} = \frac{\beta S}{\gamma} = \beta SL$$

$$R_{0} = \beta SL$$

$$1 = \frac{\beta SL}{R_{0}}$$

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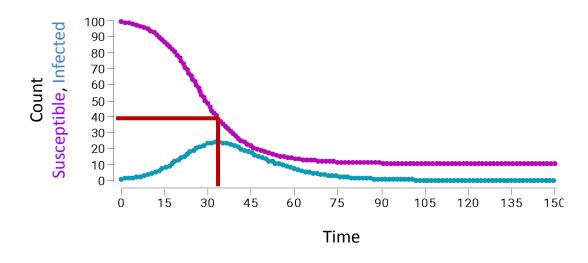
What fraction of Susceptibles need to be immune in order for

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to remain?

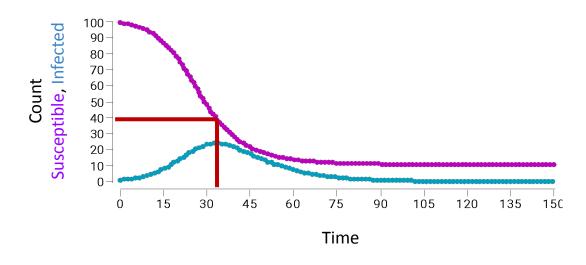
$$T_c = 1 - \frac{1}{R_0}$$

If  $T_c = 1 - \frac{1}{R_0}$  are immune **before** an outbreak then it won't be able to grow (on average)



If an outbreak takes off it **WILL NOT** stop when  $T_c = 1 - \frac{1}{R_0}$  are immune

If  $T_c = 1 - \frac{1}{R_0}$  are immune **before** an outbreak then it won't be able to grow (on average)



If an outbreak takes off it **WILL NOT** stop when  $T_c = 1 - \frac{1}{R_0}$ are immune

Why not?

#### Final Size Calculation

$$R_{\infty} = 1 - e^{-R_0 R_{\infty}}$$

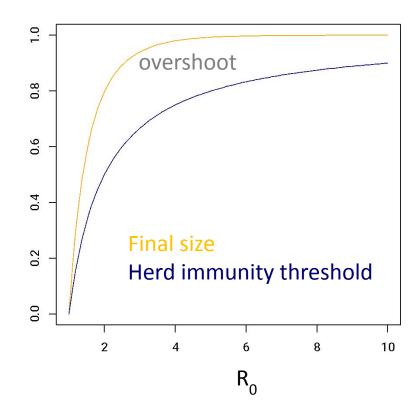
Where  $R_{\infty}$  is the proportion of the population infected at the end of the epidemic (the proportion in the R class at the end)

Citation:https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3506030/

# Comparing T<sub>c</sub> and Final Size

Many more individuals will become infected in an epidemic (on average) than need to be immunized **BEFORE** an epidemic

Herd Immunity is a relevant concept throughout an epidemic (and helps stop them), the Herd Immunity Threshold is only relevant for preventing, not stopping outbreaks.



#### What Happens When Births Are Added?

$$\frac{dS}{dt} = \delta N - \beta SI - \gamma I - \alpha I - \delta S$$

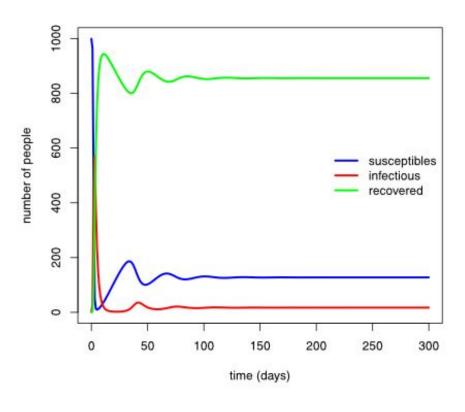
$$\frac{dI}{dI} = \delta N - \beta SI - \gamma I - \alpha I - \delta I$$

$$\frac{dR}{dt} = \delta N - \beta SI - \gamma I - \alpha R - \delta R$$

 $\delta$  is birth and death rate  $\alpha$  is disease induced death rate

# **Dynamics Over Time**

Note that after initial overshoot, susceptibles build back up until a new outbreak occurs, the second is smaller, and the third smaller than that



#### Some Terms

#### Closed v Open populations

- Closed populations have no additions due to births or immigration and no losses due to death or emigration – all dynamics due *only* to infection
  - Unrealistic, but simple and a good place to start
- Open populations can have births, immigration, deaths, emigration
  - Either explicitly or implicitly

#### **Equilibrium and Transient**

- At equilibrium, the values of all states are constant individuals may still get sick, recover, etc, but all changes balance each other out
- Dynamics are transient if the states are continuing to change ... more nuance later (really)

#### When Does / Stop Growing?

$$\frac{dI}{dt} = \beta SI - \gamma I - \alpha I - \delta I$$

$$0 < \beta SI - \gamma I - \alpha I - \delta I$$

$$\beta SI > \gamma I + \alpha I + \delta I$$

$$\beta S > \gamma + \alpha + \delta$$

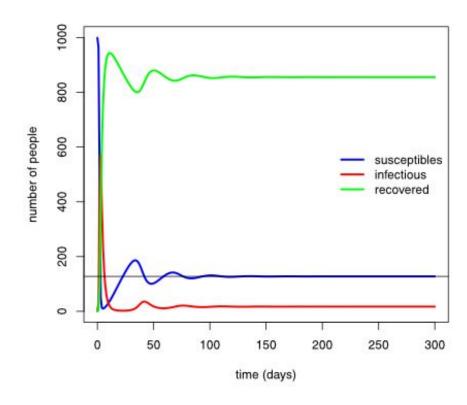
$$1 < \frac{\beta S}{\gamma + \alpha + \delta} \equiv R_0$$

 $\delta$  is birth and death rate  $\alpha$  is disease induced death rate

# **Equilibrium Dynamics**

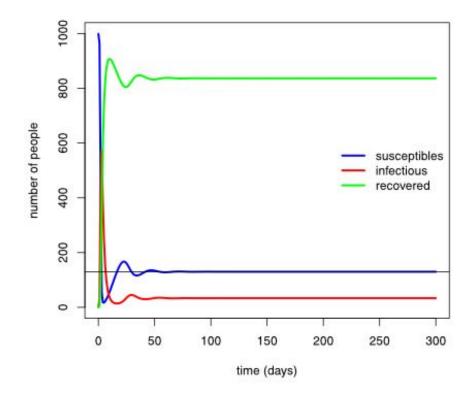
The stable equilibrium proportion susceptible is

$$\sim 1 - \frac{1}{R_0}$$



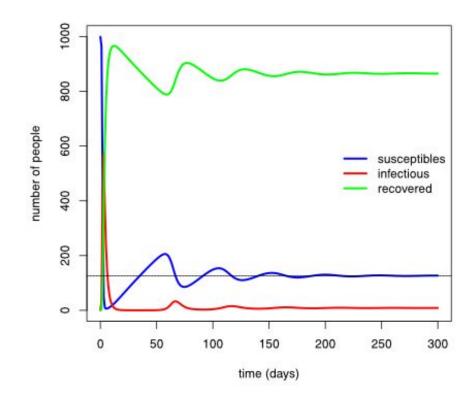
## Birth Rate Changes the Speed to Equilibrium

If we increase the birth rate, it takes less time to reach equilibrium under the assumption that the population isn't growing



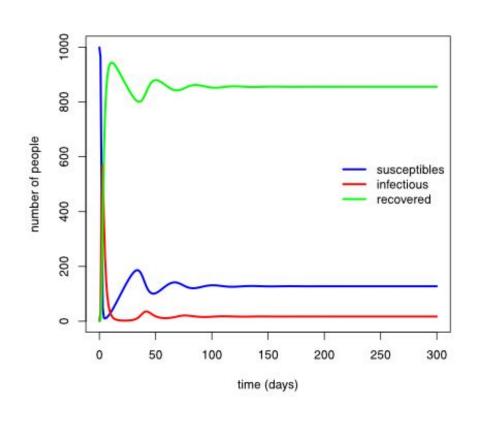
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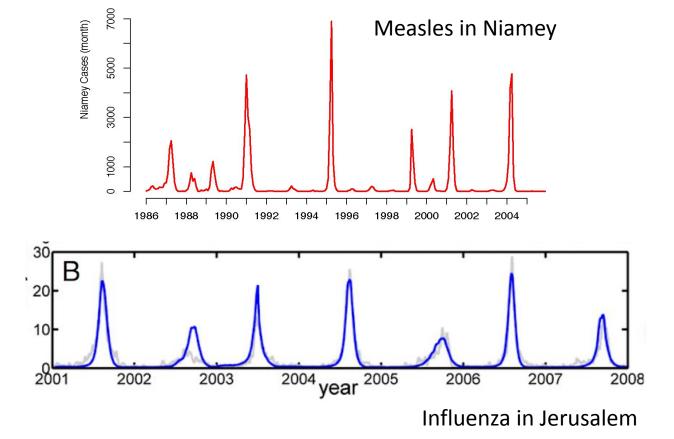
If we decrease the birth rate, it takes longer to reach equilibrium under the assumption that the population isn't growing



### What about growing populations?

- Growing populations have more susceptibles added than recovereds being taken away (by death)
  - So a greater fraction susceptible, less indirect protection, and more transmission
- More of those susceptibles are young, so if young and old have different contact rates, then transmission and dynamics will differ in young vs. old populations ...





#### Some Terms

#### Closed v Open populations

- Closed populations have no additions due to births or immigration and no losses due to death or emigration – all dynamics due *only* to infection
  - Unrealistic, but simple and a good place to start
- Open populations can have births, immigration, deaths, emigration
  - Either explicitly or implicitly

#### **Equilibrium and Transient**

- At equilibrium, the values of all states are constant individuals may still get sick, recover, etc, but all changes balance each other out
- An *attractor* is collection of states towards which a system tends it's regular and predictable, but not static.
- Dynamics are transient if the states are continuing to change

- Long term dynamics often exhibit regular fluctuations around the equilibrium levels because of seasonal changes in
  - Environmental conditions
  - Behavior
  - Population movement/aggregation
  - Vector seasonality

#### **Examples**

Influenza

Lassa fever

Legionellosis

Leptospirosis

Meningococcal meningitis

Polio

Typhoid

 Long term dynamics often exhibit regular fluctuations around the equilibrium levels because of seasonal changes in

- Environmental conditions
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#### **Examples**

Chickenpox

Measles

Pertussis

Rubella

 Long term dynamics often exhibit regular fluctuations around the equilibrium levels because of seasonal changes in

_	Environmental	l conditions
		COMMITTEE

- Behavior
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Examples	<b>Examples</b>	
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Leptospirosis	Rubella	
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Modeled as a temporal change in  $\beta$ 

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#### **Examples**

Measles

Meningococcal meningitis

Modeled as a temporal change in  $\beta$  or S

 Long term dynamics often exhibit regular fluctuations around the equilibrium levels because of seasonal changes in

- Environmental conditions
- Behavior
- Population movement/aggregation
- Vector seasonality

#### **Examples**

Chikungunya

Dengue

Malaria

Trypanosomiasis

West Nile Virus

Yellow Fever

Requires a new compartment for the vector populations