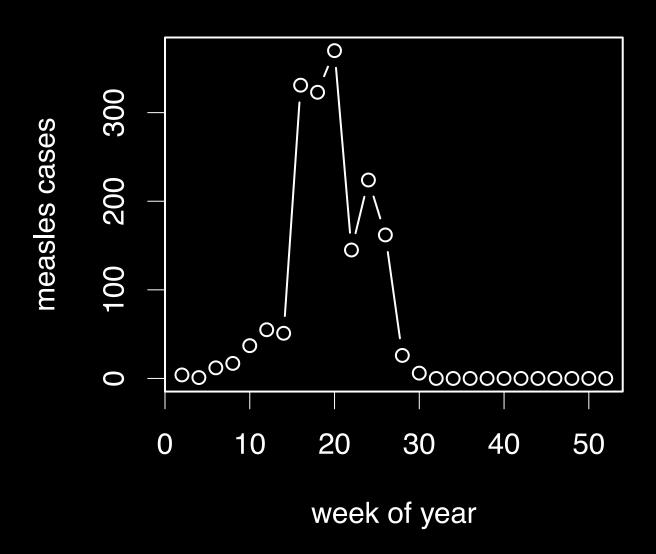
# Model Fitting: The Basic Concept



Cara Brook  $E^2M^2, \, Centre \, \, ValBio$ Ranomafana National Park, Madagascar

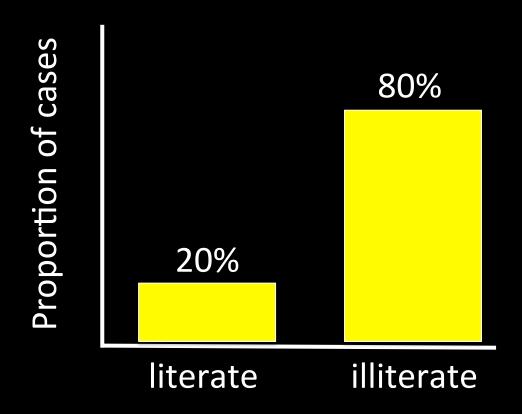
# What happened in Niamey?



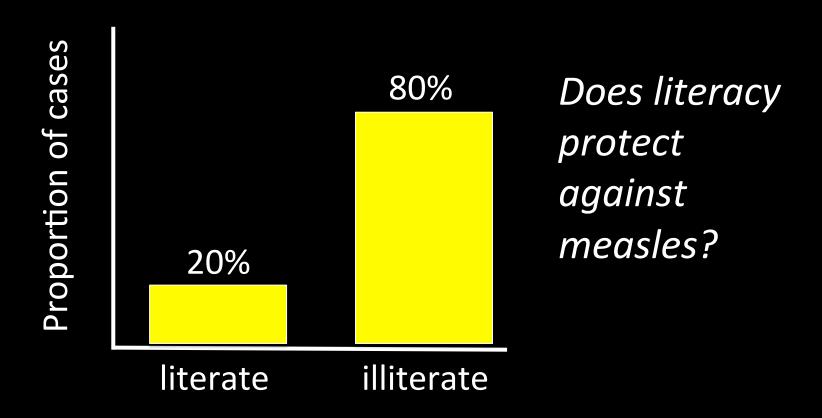
- We might ask questions about these data:
  - What proportion of cases occurred in males vs. females?
  - In winter vs. spring?
  - In Maradi vs. Dosso arrondisement?

Goal: find correlations that imply causality

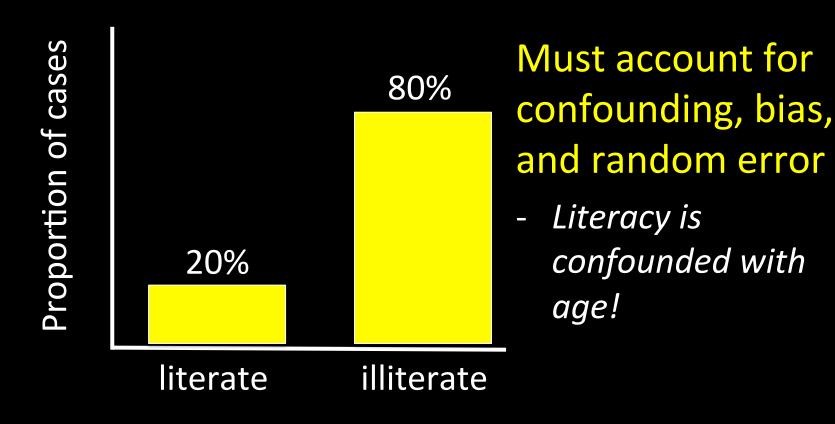
- Goal: find correlations that imply causality
- Imagine you discover:



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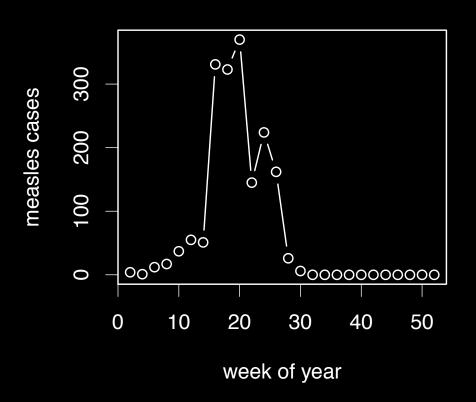
 We want to understand what happened, when it happened, and why it happened

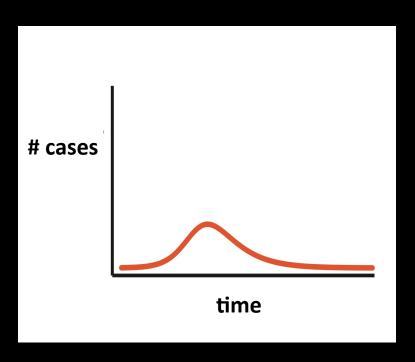
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- Allows us to scale up from individual-level processes to population-level patterns

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- We start by building a model that uses explicit processes to recover the same outcomes ("states") as our data

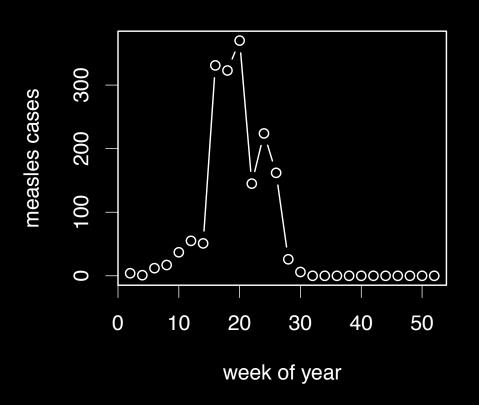
- We want to understand what happened, when it happened, and why it happened
- Allows us to scale up from individual-level processes to population-level patterns
- We start by building a model that uses explicit processes to recover the same outcomes ("states") as our data
- What state variables are captured in our data?

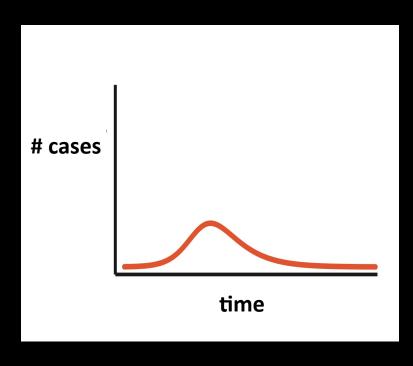
# These data give us infecteds over time...



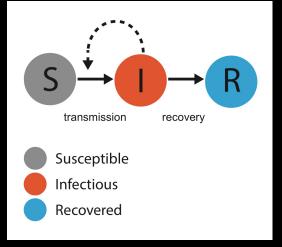


## These data give us infecteds over time...





What *processes* contribute to the "infected" state in our system?



- Build a model that uses explicit processes to recover the same states as the data.
- 2. Use any statistical tool (i.e. maximum likelihood, least squares) to ask, assuming our model is true, how likely are we to recover the observed data?
- 3. Optimize the parameters behind the processes to make the model most likely to recover the data.
- 4. If need be, restructure your model to better match your data.

1. Build a model that uses explicit processes to recover the same states as the data.

Discrete time models are simple:

$$S_{t+1} = S_t - beta*I_t*S_t/N$$

$$I_{t+1} = I_t + beta*I_t*S_t/N - gamma*I_t$$

where beta = transmission coefficient and gamma = recovery rate

 Build a model that uses explicit processes to recover the same states as the data.

If we set the timestep = 1/gamma, we can reduce the system to:

$$S_{t+1} = S_t - beta*I_t*S_t/N_t$$

$$I_{t+1} = beta*I_t*S_t/N_t$$

This means we make the assumption that there are no overlapping infectious generations.

1. Build a model that uses explicit processes to recover the same states as the data.

 $I_2 = I_1 + beta*I_1*S_1/N - gamma*I_1$ 

$$I_{2} = 1 + (.2)(1)(9/10) - (1)(1)$$

$$I_{1} = 1 \text{ person}$$

$$S_{1} = 9 \text{ persons}$$

$$N = 10$$

$$beta = .2 \text{ hour}^{-1}$$

$$gamma = 1 \text{ hour}^{-1}$$

$$\Delta t = 1 \text{ hour}$$

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$$I_{2} = 1 + (.2)(1)(9/10) - (1)(1)$$

$$I_{2} = 1 + (.2)(1)(9/10) - 1$$

$$I_{2} = .18 \text{ at 2 hours}$$

$$I_{1} = 1 \text{ person}$$

$$S_{1} = 9 \text{ persons}$$

$$N = 10$$

$$beta = .2 \text{ hour}^{-1}$$

$$gamma = 1 \text{ hour}^{-1}$$

$$\Delta t = 1 \text{ hour}$$

$$I_2 = 1 + (.2)(2)(1)(9/10) - (1)(1)(2)$$

 $I_1 = 1$  person  $S_1 = 9$  persons N = 10beta = .2 hour<sup>-1</sup> gamma = 1 hour<sup>-1</sup>  $\Delta t = 2$  hours

1. Build a model that uses explicit processes to recover the same states as the data.

 $I_2 = I_1 + beta*I_1*S_1/N - gamma*I_1$ 

$$I_{2} = 1 + (.2)(1)(9/10) - (1)(1)$$

$$I_{2} = 1 + (.2)(1)(9/10) + 1$$

$$I_{2} = .18 \text{ at 2 hours}$$

$$I_{3} = 1 \text{ person}$$

$$I_{1} = 1 \text{ person}$$

$$I_{2} = 9 \text{ persons}$$

$$I_{3} = 9 \text{ persons}$$

$$I_{4} = 1 \text{ person}$$

$$I_{5} = 9 \text{ persons}$$

$$I_{5} = 10 \text{ beta} = .2 \text{ hour}^{-1}$$

$$I_{5} = 1 \text{ person}$$

$$I_{5} = 1 \text{ hour}^{-1}$$

$$I_{5} = 1 \text{ hour}^{-1}$$

$$I_{5} = 1 \text{ hour}^{-1}$$

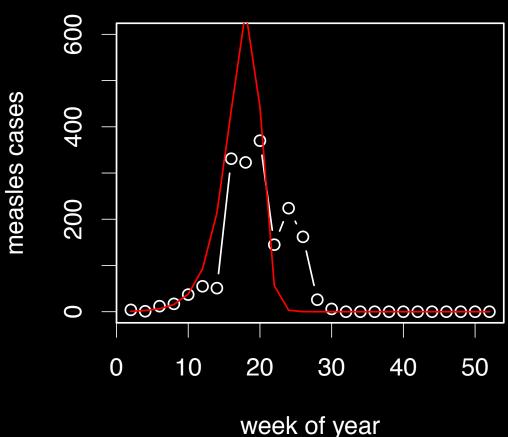
$$I_2 = 1 + (.2)(2)(1)(9/10) - (1)(1)(2)$$
 $I_2 \neq 1 + (.2)(2)(1)(9/10) + 2$ 
 $I_2 = -.64$  at 4 hours

Don't cancel!

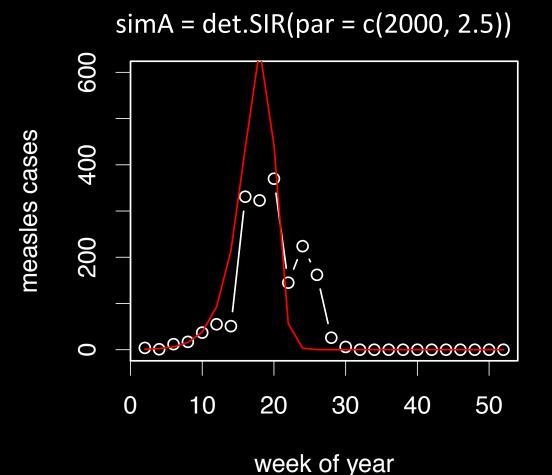
 $I_1 = 1$  person  $S_1 = 9$  persons N = 10beta = .2 hour<sup>-1</sup> gamma = 1 hour<sup>-1</sup>  $\Delta t = 2$  hours

1. Build a model that uses explicit processes to recover the same states as the data.

simA = det.SIR(par = c(2000, 2.5))



1. Build a model that uses explicit processes to recover the same states as the data.

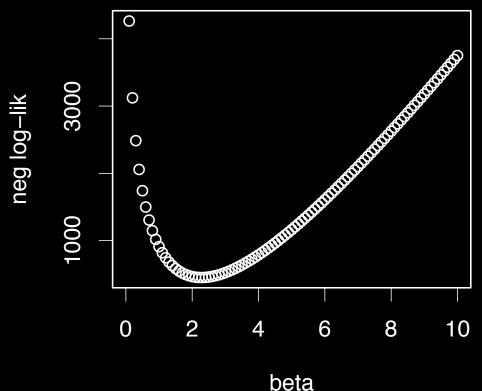


Model does a pretty good job, but but it overshoots our data by quite a bit.

What does this suggest about our guess for beta?

2. Use any maximum likelihood to ask, assuming our model is true, how likely are we to recover the observed data?

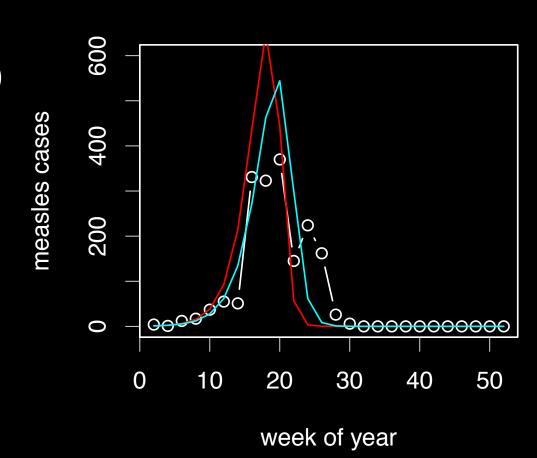
beta\_guess[which.min(II)] = 2.3



2. Use any maximum likelihood to ask, assuming our model is true, how likely are we to recover the observed data?

simB = det.SIR(par = c(2000, 2.3))

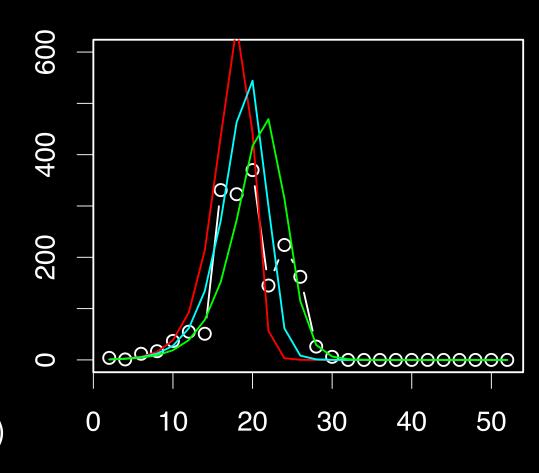
New beta fits even better!



 Optimize the parameters behind the processes to make the model most likely to recover the data.

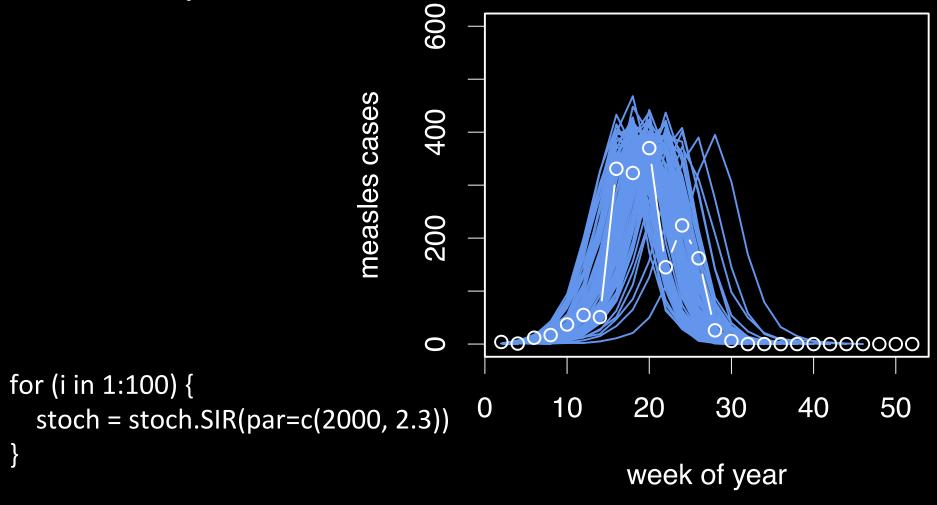
Looks great!
But why not perfect?

simC = det.SIR(par = c(2152, 2.09))

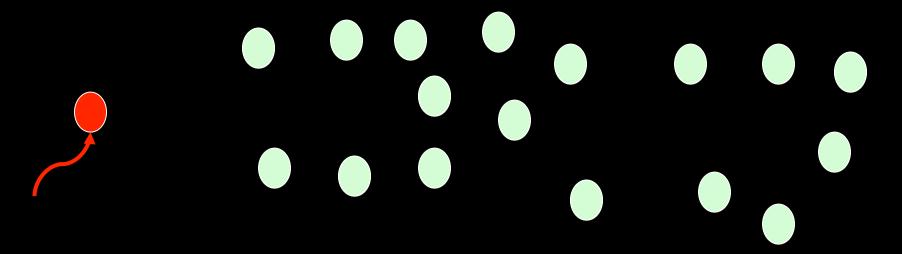


week of year

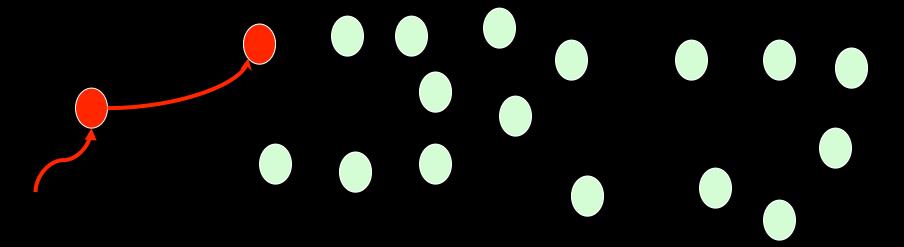
4. If need be, restructure your model to better match your data.



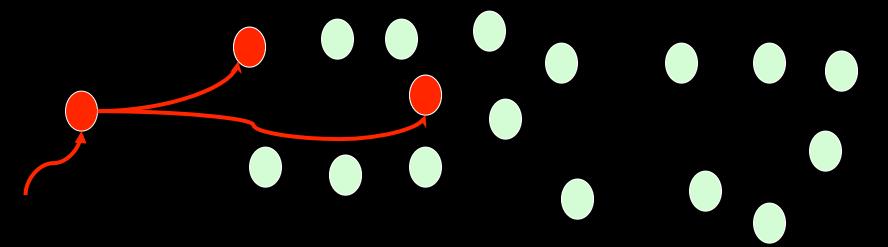
- The basic reproduction number for a pathogen
- Defined as the number of new infections generated by one existing infection in a completely susceptible host population
- In these discrete time models, R0 = beta



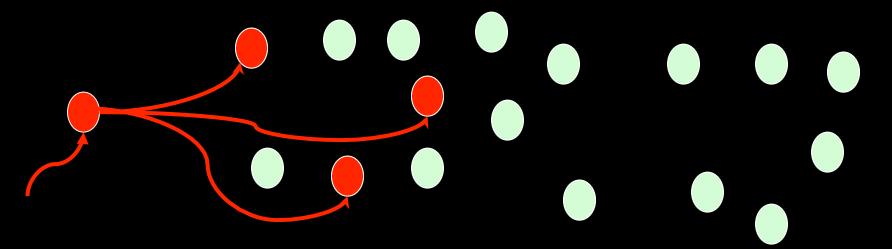
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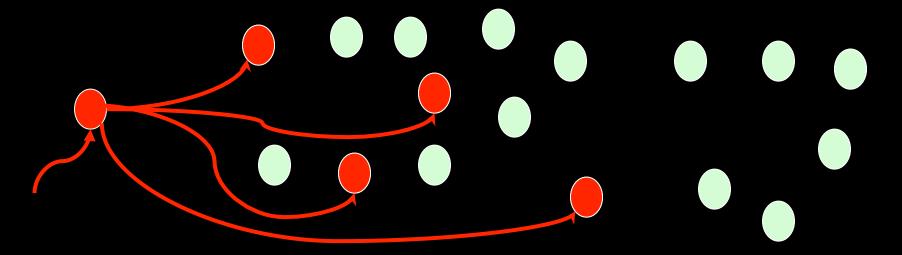
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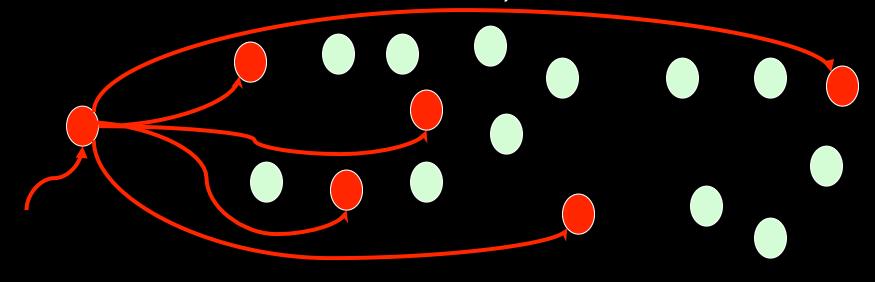
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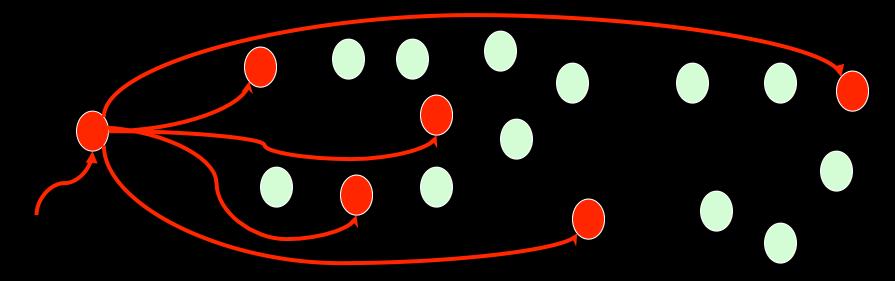


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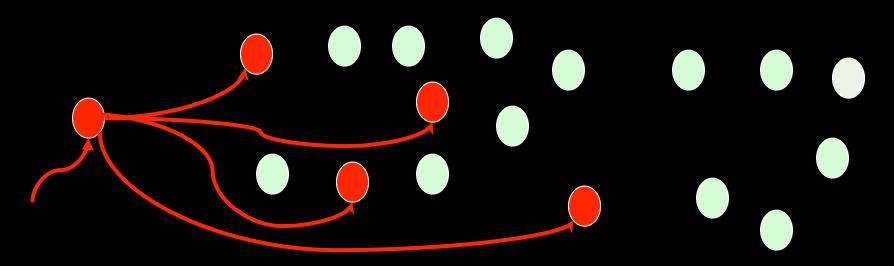
$$R_0 = 5$$

- The number of new infections generated by one existing infection in a partially susceptible host population
- Reff = R0\*(S/N)
- Changes over time!



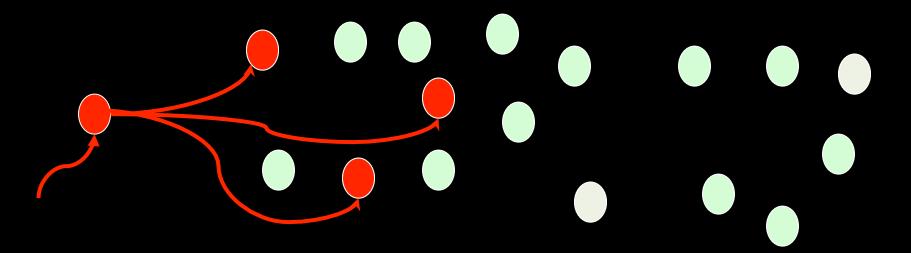
$$R_{\rm eff} = (5)*(12/17) = 3.53$$

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- Changes over time!



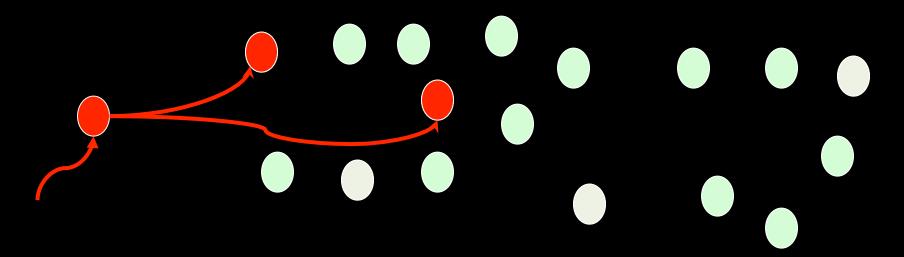
$$R_{\rm eff} = (5)*(13/17) = 3.82$$

- The number of new infections generated by one existing infection in a partially susceptible host population
- Reff = R0\*(S/N)
- Changes over time!



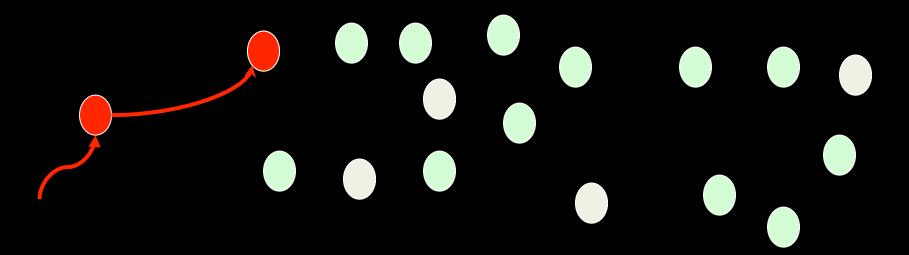
$$R_{\rm eff} = (5)*(14/17) = 4.12$$

- The number of new infections generated by one existing infection in a partially susceptible host population
- Reff = R0\*(S/N)
- Changes over time!



$$R_{\rm eff} = (5)*(15/17) = 4.41$$

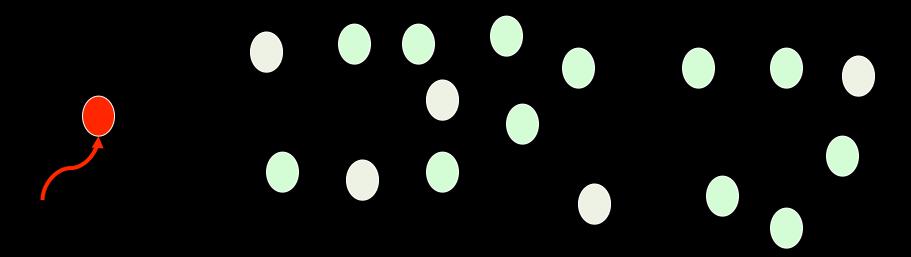
- The number of new infections generated by one existing infection in a partially susceptible host population
- Reff = R0\*(S/N)
- Changes over time!



$$R_{\rm eff} = (5)*(16/17) = 4.71$$

#### Insights from fitting dynamic models: R-effective

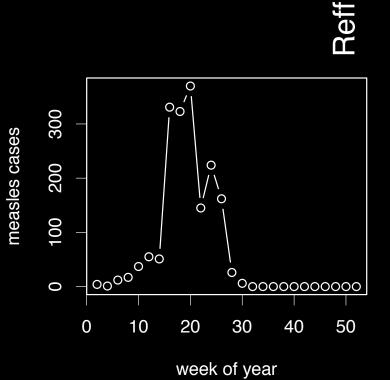
- The number of new infections generated by one existing infection in a partially susceptible host population
- Reff = R0\*(S/N)
- Changes over time!

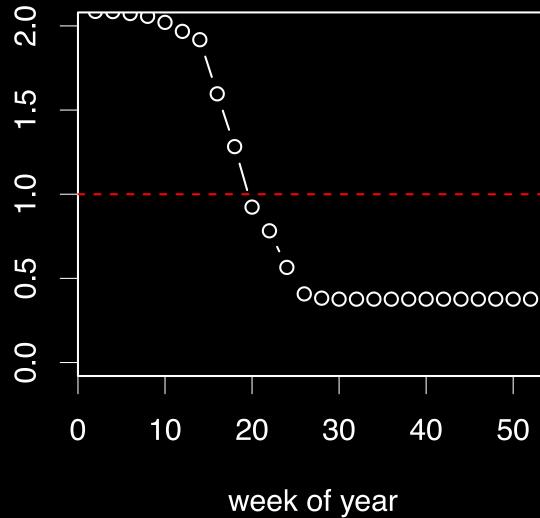


$$R_{\text{eff}} = (5)*(17/17) = 5$$

# Insights from fitting dynamic models: R-effective

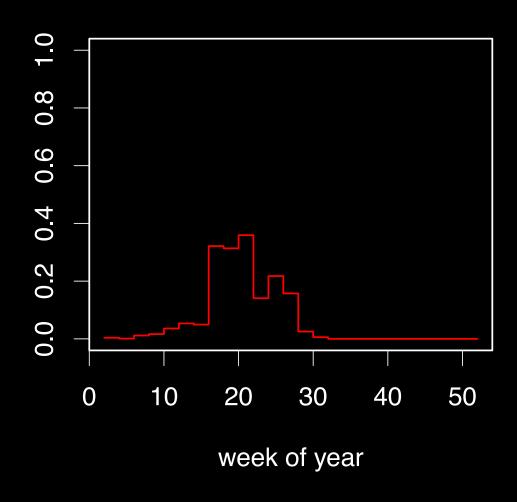
An epidemic spreads when Reff > 1 and declines when Reff < 1





# Insights from fitting dynamic models: Force of Infection

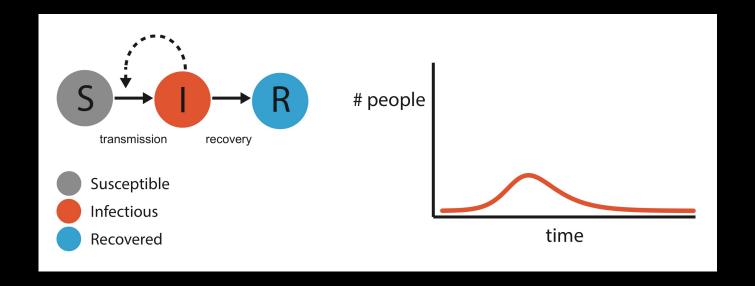
- Defined as the rate at which susceptibles become infected
- FOI = R0\*(I/N)
- Changes over time, but constant across each timestep



- We want to understand what happened, when it happened, and why it happened
- Allows us to scale up from individual-level processes to population-level patterns
- We start by building a model that uses explicit processes to recover the same outcomes ("states") as our data

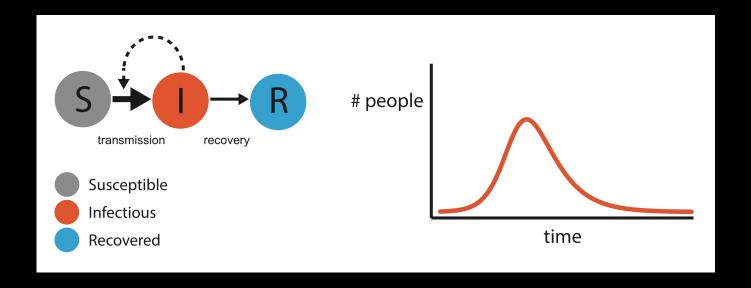
Test "what if" scenarios not amenable to experimentation

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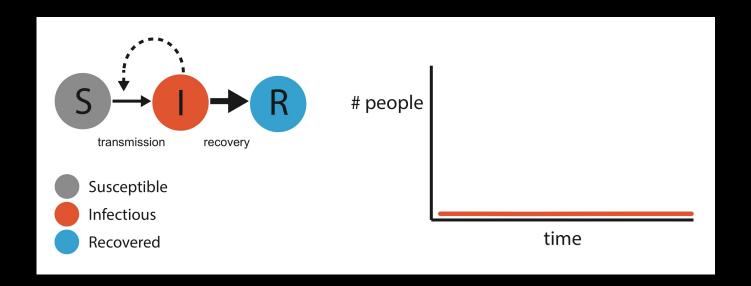
Test "what if" scenarios not amenable to experimentation

What if each person exposed 50% more people?

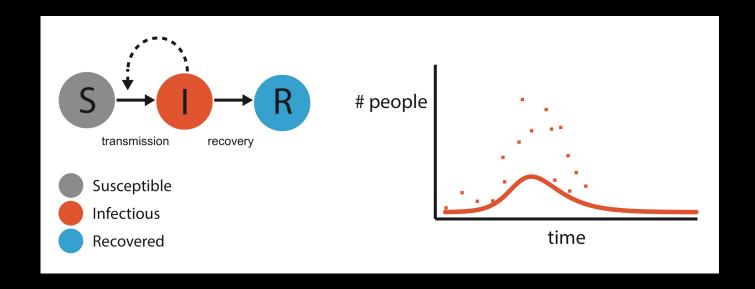


Test "what if" scenarios not amenable to experimentation

What if we treated people and doubled the rate of recovery?

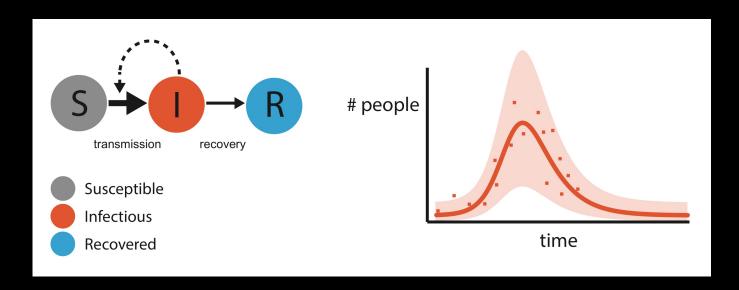


- Test "what if" scenarios not amenable to experimentation
- Estimate parameters that are difficult to measure by fitting models to available data

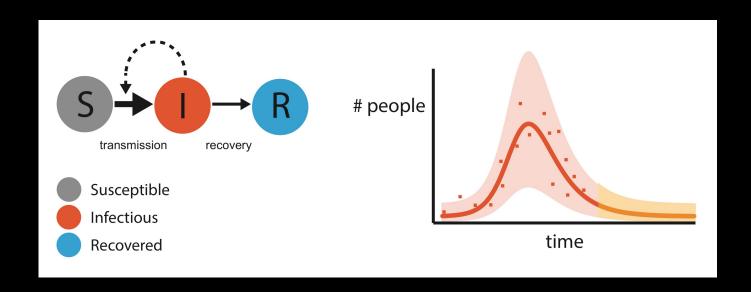


- Test "what if" scenarios not amenable to experimentation
- Estimate parameters that are difficult to measure by fitting models to available data

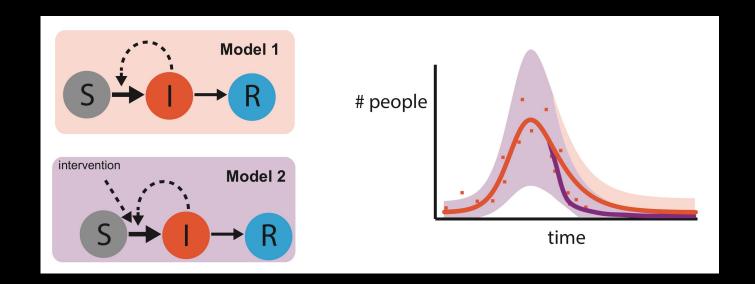
Estimate transmission rate or other model parameters (with confidence intervals)



- Test "what if" scenarios not amenable to experimentation
- Estimate parameters that are difficult to measure by fitting models to available data
- Forecast forward in time



- Test "what if" scenarios not amenable to experimentation
- Estimate parameters that are difficult to measure by fitting models to available data
- Forecast forward in time
- Select between models of differing hypotheses



- Estimate: time series of state variables/
   parameters of interest → DATA
- Inference: Build model to recapture data. Fit to optimize parameters and "infer" the process underlying the data
- Model assessment: Assess plausibility or model comparison
- End goal: explain observed patterns or predict