

Foundations of dynamic modeling: The SIR Model Family

Jonathan Dushoff, McMaster University

DAIDD 2019

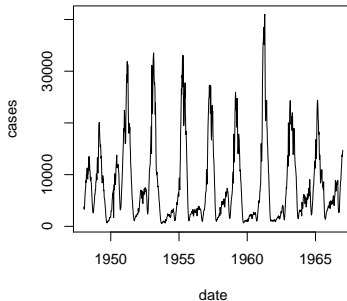
Goals

- ▶ This lecture will:
 - ▶ introduce the idea of dynamical modeling
 - ▶ explain why dynamical modeling is a key tool for understanding infectious disease
 - ▶ discuss and demonstrate simple dynamical models from the SIR model family
 - ▶ investigate some insights that can be gained from these models

Dynamical modeling connects scales



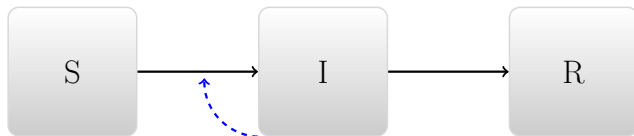
Measles reports from England and Wales



- ▶ Start with rules about how things change in short time steps
 - ▶ Usually based on *individuals*
- ▶ Calculate results over longer time periods
 - ▶ Usually about *populations*

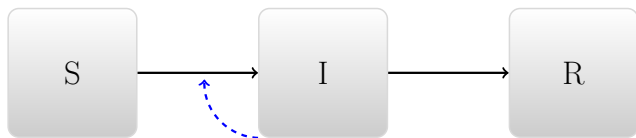
Compartmental models

Divide people into categories:



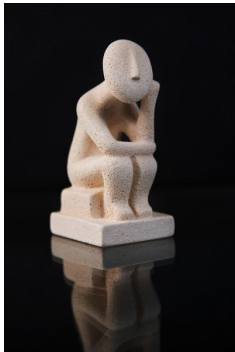
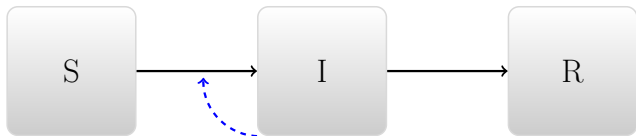
► Susceptible \rightarrow Infectious \rightarrow Recovered

What determines transition rates?

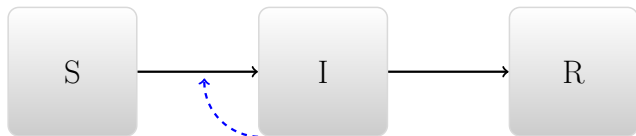


- ▶ People get better independently
- ▶ People get infected by infectious people

Conceptual modeling (present)

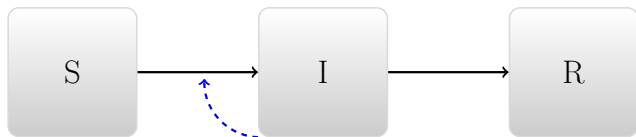


Conceptual modeling



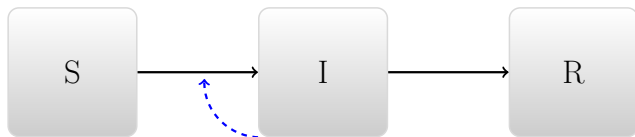
- ▶ What is the final result?
- ▶ When does disease increase, decrease?

Dynamic implementation



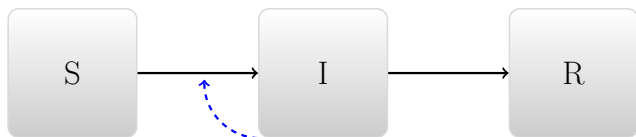
- ▶ Requires assumptions about recovery and transmission
- ▶ The *conceptually simplest* implementation uses Ordinary Differential Equations (ODEs)
 - ▶ Other options may be more realistic
 - ▶ Or simpler in practice

Recovery



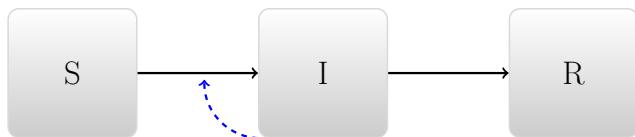
- ▶ Infectious people recover at *per capita* rate γ
 - ▶ Total recovery rate is γI
 - ▶ Mean time infectious is $D = 1/\gamma$

Transmission



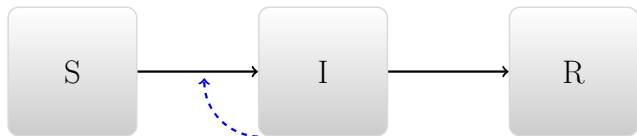
- ▶ Susceptible people get infected by:
 - ▶ Going around and contacting people (rate c)
 - ▶ Some of these people are infectious (proportion I/N)
 - ▶ Some of these contacts are effective (proportion p)
- ▶ Per capita rate of becoming infected is $cpI/N \equiv \beta I/N$
- ▶ Population-level transmission rate is $\mathcal{T} = \beta SI/N$

Another perspective on transmission



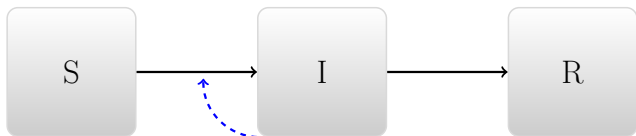
- ▶ Infectious people infect others by:
 - ▶ Going around and contacting people (rate c)
 - ▶ Some of these people are susceptible (proportion S/N)
 - ▶ Some of these contacts are effective (proportion p)
- ▶ Per capita rate of infecting others is $cpS/N \equiv \beta S/N$
- ▶ Population-level transmission rate is $\mathcal{T} = \beta SI/N$

ODE implementation

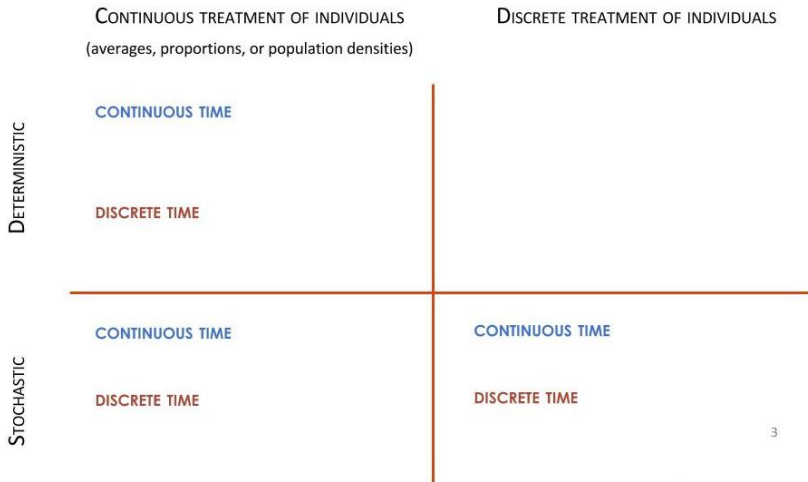


$$\begin{aligned}\frac{dS}{dt} &= -\beta \frac{SI}{N} \\ \frac{dI}{dt} &= \beta \frac{SI}{N} - \gamma I \\ \frac{dR}{dt} &= \gamma I\end{aligned}$$

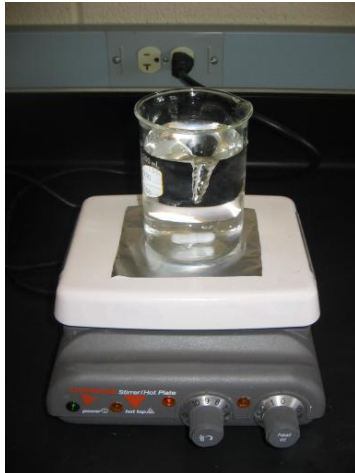
Spreadsheet implementation



Model taxonomy (present)

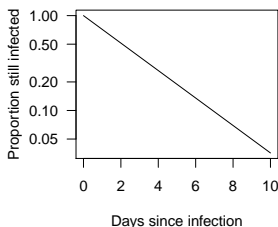
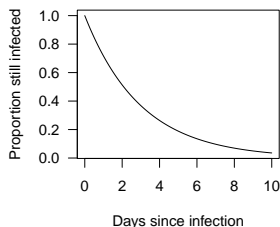


ODE assumptions



- ▶ Lots and lots of people
- ▶ Perfectly mixed

ODE assumptions



- ▶ Waiting times are exponentially distributed
- ▶ Rarely realistic

More about transmission

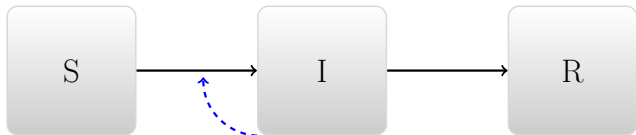


- ▶ $\beta = pc$
 - ▶ What is a contact?
 - ▶ What is the probability of transmission?
- ▶ Sometimes this decomposition is clear
- ▶ But usually it's not

Population sizes

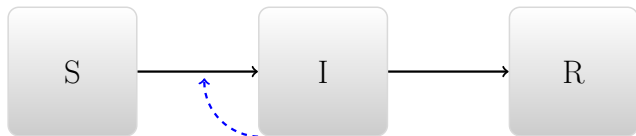
- ▶ How does β change with population size?
- ▶ Recall that β is the *per capita* rate of contacts

Population sizes (present)



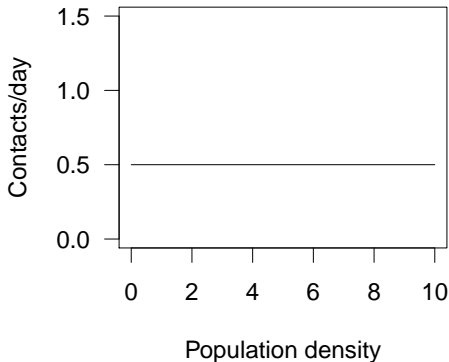
$$\begin{aligned}\frac{dS}{dt} &= -\beta \frac{SI}{N} \\ \frac{dI}{dt} &= \beta \frac{SI}{N} - \gamma I \\ \frac{dR}{dt} &= \gamma I\end{aligned}$$

Population sizes



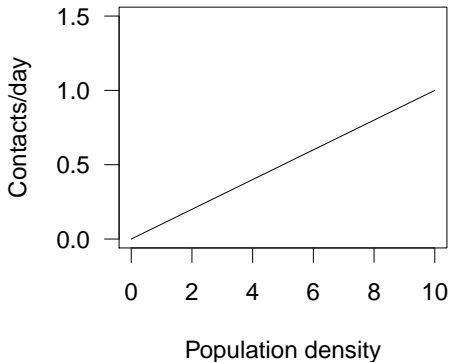
$$\begin{aligned}\frac{dS}{dt} &= -\beta(N)\frac{SI}{N} \\ \frac{dI}{dt} &= \beta(N)\frac{SI}{N} - \gamma I \\ \frac{dR}{dt} &= \gamma I\end{aligned}$$

Standard incidence



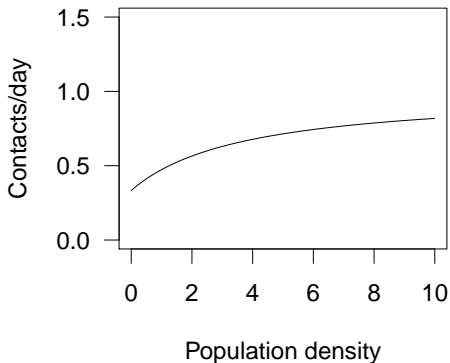
- ▶ $\beta(N) = \beta_0$
- ▶ $\mathcal{T} = \frac{\beta_0 SI}{N}$
- ▶ Also known as *frequency-dependent* transmission

Mass action



- ▶ $\beta(N) = \beta_1 N$
- ▶ $\mathcal{T} = \beta_1 SI$
- ▶ Also known as *density-dependent* transmission

General

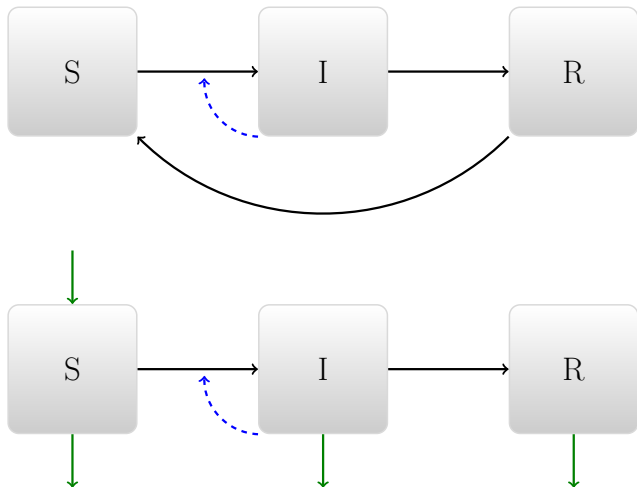


- ▶ Per-capita rate:
 - ▶ May not go to zero when N does
 - ▶ May not go to ∞ when N does

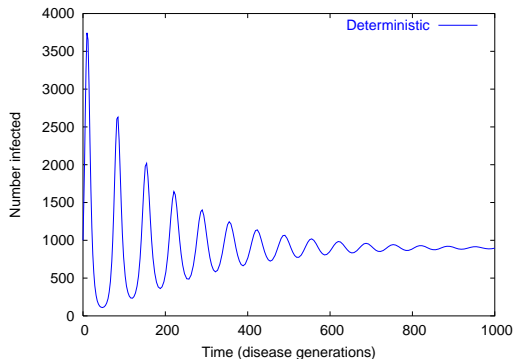
Digression – units

- ▶ $\mathcal{T} = \beta SI/N : [\text{ppl}/\text{time}]$
- ▶ $\beta : [1/\text{time}]$
 - ▶ The true β always has people in the numerator and the denominator
 - ▶ $\beta/\gamma = \beta D : [1]$
- ▶ Components of β may have different units
 - ▶ Standard incidence, $\beta_0 : [1/\text{time}]$
 - ▶ Mass-action incidence, $\beta_1 : [1/(\text{people} \cdot \text{time})]$

Closing the circle

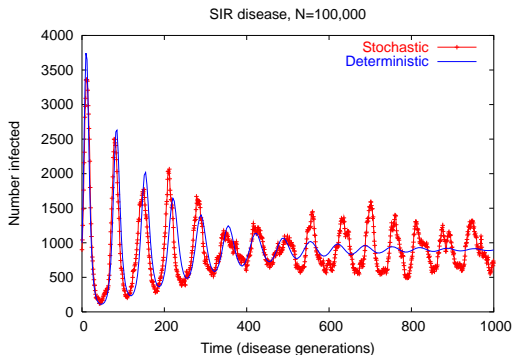


Tendency to oscillate



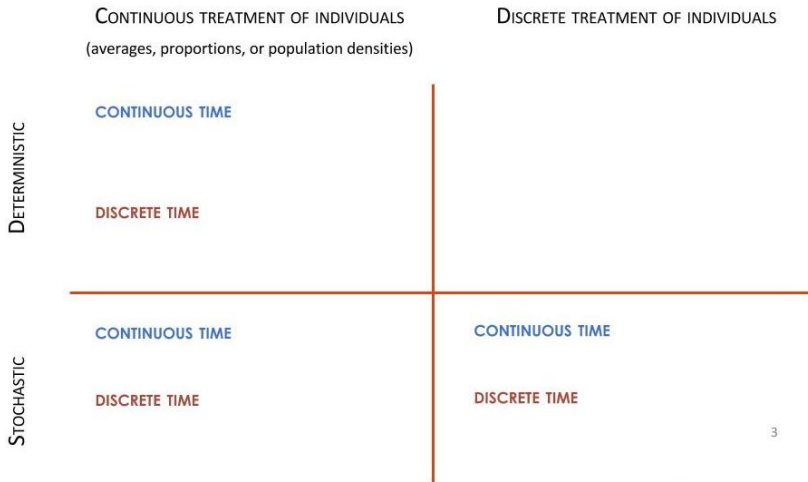
- ▶ Many susceptibles \rightarrow many infections \rightarrow few susceptibles \rightarrow few infections $\rightarrow \dots$
- ▶ Oscillations in simple models tend to be “damped”

With individuality



- ▶ Treating individuals as individuals can produce substantial oscillations even in large populations
- ▶ Interaction between random effects and the different time scales (of infection and recovery)

Model taxonomy (present)



Summary

- ▶ Dynamic models are an essential tool because they allow us to link between scales
- ▶ There are many ways to construct and implement dynamic models
- ▶ Very simple models can provide useful insights
 - ▶ Reproductive numbers and thresholds
 - ▶ Tendency for oscillation (and tendency for damping)
- ▶ More complex models can provide more detail, but also require more assumptions, and more choices
- ▶ Understanding simple models can help guide our understanding of more complicated models



This presentation is made available through a Creative Commons Attribution-Noncommercial license. Details of the license and permitted uses are available at <http://creativecommons.org/licenses/by-nc/3.0/>



© 2013–2019, International Clinics on Infectious Disease Dynamics and Data

Title: Foundations of dynamic modeling:
Attribution: Jonathan Dushoff, McMaster University, DAIDD 2019

Source URL: https://figshare.com/collections/International_Clinics_on_Infectious_Disease_Dynamics_and_Data/3788224

For further information please contact admin@ici3d.org.



AIMS

African Institute for
Mathematical Sciences
SOUTH AFRICA



SACEMA
Centre of Excellence in Systemic Modelling and Analysis



UNIVERSITY OF GEORGIA

College of Public Health