

Network Modeling for Epidemics



The workhorse of epidemic modelers And a good place to build intuition

NME Workshop

Deterministic compartmental models

- Only the aggregate count in each state ("compartment") is represented, not the individual persons
- Within each compartment, people are homogeneous
- Transitions ("flows") are represented in terms of rates
 - The fraction of the aggregate count that moves from one compartment to another at any time point



Deterministic compartmental models

- May be discrete time or continuous time
 - We will focus on discrete time in what follows because it's easier to understand
 - Most published models, and most packages (including EpiModel) solve in continuous time
- Compartmental models are usually deterministic each run gives exactly the same result
- Measures = EXPECTED counts (across an infinite number of stochastic runs)
- Compartments and flows can represent fractional persons



Here R stands for:

- Recovered with immunity
- Also sometimes called "removed" in the literature but be careful
 - *Removed* from the infection process
 - *Not removed* from the contact process

New infections per unit time (incidence) What is a reasonable expression for this quantity?



t = time

State variables

- s(t) = expected number of susceptible people at time t
- i(t) = expected number of infected people at time t
- r(t) = expected number of recovered people at time t

Parameters

- α = act rate per unit time
- τ = prob. of transmission given S-I act
- ρ = recovery rate

A new infection requires: a susceptible person to have an act with an infected person and for infection to transmit because of that act

- t = time
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Expected incidence at time t =

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Expected incidence at time $t = s(t)\alpha$

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Expected incidence at time t = s

$$s(t) \alpha \frac{i(t)}{s(t)+i(t)+r(t)}$$

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Expected incidence at time $t = s(t)\alpha \frac{i(t)}{s(t)+i(t)+r(t)}\tau$

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- n(t) = total population = s(t) + i(t) + r(t)

Expected incidence at time $t = s(t)\alpha \frac{i(t)}{s(t)+i(t)+r(t)}\tau$ = $s(t)\alpha \frac{i(t)}{n(t)}\tau$

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Expected incidence at time
$$t = s(t)\alpha \frac{i(t)}{s(t)+i(t)+r(t)}\tau$$

= $s(t)\alpha \frac{i(t)}{n(t)}\tau$
= $s(t)\alpha \frac{i(t)}{n}\tau$

Careful: only for a "closed" population can the time subscript be dropped for *n*

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t = time

s(t) = expected number of susceptible people at time t

- i(t) = expected number of infected people at time t
- r(t) = expected number of recovered people at time t
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- ρ = recovery rate

- Much simpler process: expected number of recoveries at time t equals ρ i(t)
- Reminder: Expected incidence at time $t = s(t)\alpha \frac{i(t)}{n}\tau$
- How do we turn this into a system of equations?



Add in a set of initial conditions: and a set of parameter values

s(0) = 999, i(0) = 1, r(0) = 0 α = 0.6, τ = 0.3, ρ = 0.1

And one has the full trajectory of each state over time:



Brief digression: contacts and acts

- The epi modeling literature typically uses the term "contact" so why do we use "act"?
- Because "contact" is an ambiguous word in this context
 - E.g. think of sexual activity when we say "# of contacts per year"
 - Does it mean number of sex acts?
 - Or numbers of different partners?
- To be explicit, we will make the distinction between "acts" and "partners" throughout this workshop
- This distinction matters for disease dynamics <u>when there are repeated acts with</u> <u>the same person</u>

Brief digression: contacts and acts

If multiple acts occur within partnerships, DCMs take one of two forms.



- 1. Define a contact as an act. Model each act as a separate independent event, ignoring the persistent nature of the partnerships
- 2. Define a contact as a partnership. Compress all of the acts over the partnership into a single instance in time
- We'll return to this later

- Relationship between duration of infection and recovery rate
 - Imagine a disease with a constant recovery rate of 0.2.
 - I.e., on Day 1 of infection, you have a 20% probability of recovering.
 - If you don't recover on Day 1, you have a 20% probability of recovering on Day 2. Etc.

Now, imagine 100 people who start out infected on the same day.

How many recover after being infected 1 day?	100*0.2 = 20
How many recover after being infected 2 days?	80*0.2 = 16
How many recover after being infected 3 days?	64*0.2 = 12.8
What is this distribution called?	Geometric
What is the mean (expected) duration spent infected?	1/0.2 = 5 days
	$1/\rho = D$



R₀ : A key summary metric

Definition: The expected number of secondary infections generated by the first infected case in a population that has never seen this infection before

A single number that summarizes the epidemic potential in a population

- What happens if the first infected case recovers before transmitting to someone else?
- … nothing.

R_0 and the "persistence threshold"

There is an epidemic persistence threshold at $R_0 = 1$

Value of R_0	Implication
< 1	The first infected individual will on average infect < 1 person total. In a deterministic model, the epidemic will always go extinct
> 1	The first infected individual will on average infect >1 person total. In a deterministic model, the epidemic will always grow
= 1	We are right on the threshold between an epidemic and extinction. In a deterministic model, the epidemic will just putter along

DCMs: R_0

- So, how do we calculate *R*⁰ for a DCM?
 - Intuitively, for that first case, the expected number of secondary infections generated is:

α.

duration infected x act rate per timestep x transmission rate per act

• For a simple SIR DCM: $R_0 = \frac{\alpha \tau}{\rho}$ Because $D = \frac{1}{\rho}$

τ