

Network SIR Model

- Network Structure: Suppose individuals are connected randomly — Erdős-Rényi graph.
 - can treat as a fixed Erdős-Rényi graph, or as a network where the edges are re-selected at each time step.
 - let probability of each edge be p_e
- Time step: synchronous
- Disease states: susceptible, infectious, recovered (immune)
- Rules: at each time step
 - Transmission: if susceptible, can be infected by each infectious neighbor at the next time step w/ probability p_i
 - Recovery: if infectious, can recover at

next time step w/ probability Pr

We can write the network model as a difference equation system

$$x_{i,t+1} = f(x_{j,t} \in N(x_i))$$

↑
state of node i

function of the states of the neighboring nodes

However, this requires one variable for each node — very high-dimensional!

We can build a meanfield model to reduce dimension.

↳ take advantage that nodes use same set of general rules, and are connected randomly.

Building the mean field model

let N be total number of nodes.

$s = \frac{S}{N}$ = fraction of nodes
who are susceptible

$i = \frac{I}{N}$ = fraction of nodes
who are infectious

$r = \frac{R}{N}$ = fraction of nodes
who are recovered

note $r = 1 - s - i$, since
the total population is
fixed.

Probability that a svsc. node j is infected by node k is:

$$p_{e \times i} \times p_i$$
 probability that $j-k$ edge exists. probability node k is infectious probability of transmission

Then $(1 - p_{e \times i} \times p_i)^{N-1}$ = probability that node j remains susceptible at the next time step.

Probability of having a $S \rightarrow S$ transition in the next timestep, i.e. the fraction of nodes we expect to see go $S \rightarrow S$

is: $S (1 - p_{e \times i} \times p_i)^{N-1}$ (not infected by other nodes)

We can work out the probabilities of each transition as:

Current state	Next state	Probability of transition
Susc.	Susc.	$s(1 - p_e i p_i)^{N-1}$
Susc.	Inf.	$s(1 - (1 - p_e i p_i)^{N-1})$
Inf.	Inf.	$i(1 - p_r)$
Inf.	Rec.	$i p_r$

Then we can write difference equations for the fraction of nodes in each state:

S_{t+1} = only people who were
S and stay S.

$$= S_t (1 - p_e i_t p_i)^{N-1} \quad (\text{from table})$$

or can write:

$$= S_t \text{ - } S \rightarrow I \text{ transitions}$$

current - loss

$$= S_t - S_t (1 - (1 - p_e i_t p_i)^{N-1})$$

$$= S_t (1 - (1 - (1 - p_e i_t p_i)^{N-1}))$$

$$= S_t (1 - p_e i_t p_i)^{N-1}$$

i_{t+1} = incoming from S + stay sick
(do not recover)

$$= S_t (1 - (1 - p_e i_t p_i)^{N-1})$$

$$+ i_t (1 - p_r)$$

(note this is also same
as $i_t + \text{incoming} - \text{recovery}$)

$$r_{t+1} = 1 - s_{t+1} - i_{t+1}$$

(or we can write out similarly)

Note the eqns above are
good, but we can simplify
further — the binomial approx
tells us that when $p_e i p_i$ is
small (i.e. probability of
connection and subsequent
transmission w/a node is small),
then $(1 - p_e i p_i)^{N-1} \approx 1 - (N-1)p_e i p_i$.

Then we can rewrite as:

$$\begin{aligned}S_{t+1} &= S_t (1 - p_e \dot{i}_t p_i)^{N-1} \\ &\approx S_t (1 - (N-1) p_e \dot{i}_t p_i) \\ &= S_t - \underbrace{(N-1) p_e p_i}_b S_t \dot{i}_t\end{aligned}$$

$$S_{t+1} = S_t - b S_t \dot{i}_t$$

and similarly

$$\begin{aligned}\dot{i}_{t+1} &= S_t (1 - (1 - p_e \dot{i}_t p_i)^{N-1}) \\ &\quad + \dot{i}_t (1 - p_r) \\ &\approx S_t (1 - (1 - (N-1) p_e \dot{i}_t p_i))\end{aligned}$$

$$\begin{aligned}
 & + i_t(1 - pr) \\
 = & S_t (N - I) p_e p_r i_t \\
 & + i_t(1 - pr) \\
 = & b s_t \bar{i}_t \quad \bar{i}_t(1 - pr)
 \end{aligned}$$

$$\dot{i}_{t+1} = \bar{i}_t + b s_t \bar{i}_t - pr \bar{i}_t$$

And

$$r_{t+1} = 1 - s_{t+1} - \dot{i}_{t+1}$$

$$= \underbrace{(1 - s_t - \bar{i}_t)}_{r_t} + \underbrace{pr \bar{i}_t}_{\text{new recoveries}}$$

number we had

new recoveries

This looks a lot like the classic SIR model! (it is)

We can even go further and simplify to a set of ODE's,

To do this, we will need to make our timestep smaller and smaller — currently $\Delta t = 1$, let's rewrite w/ Δt explicit:

$$i_{t+\Delta t} = \hat{i}_t + b s_t i_t - p r_t$$

$$\underbrace{i_{t+\Delta t} - i_t}_{\substack{\text{change in } i \\ \Delta i}} = b s_t \dot{i}_t - p r r_t$$

$$\frac{i_{t+\Delta t} - i_t}{\Delta t} = \frac{b s_t \dot{i}_t - p r r_t}{\Delta t}$$

$$\frac{\Delta i}{\Delta t} = \frac{b s_t \dot{i}_t - p r r_t}{\Delta t}$$

take the limit as $\Delta t \rightarrow 0$
and we get a derivative!

What about RHS? Need to
think about how the

probabilities change as Δt gets smaller. A few want to do this - won't get into details now, but we can choose something "reasonable" (e.g. assuming constant rates of events etc.)

That will give us:

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

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

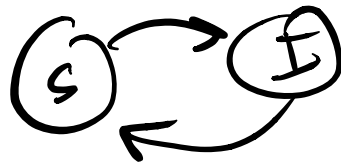
$$\frac{dR}{dt} = \gamma I$$

i.e. classic SIR.

But for now, let's stick w/
the difference eqns in
above.

The difference eqns or ODE's that
we get have a very standard
form - if 2 ^(or more) nodes interact,
usually get a product of the
state variables, and if a node
can independently do something w/
constant probability, typically
turns out linear.

So, for example, if an SIS model, i.e.



recovery returns to S (not immune), then we would

get:

$$S_{t+1} = S_t - b S_t i_t + p r i_t$$

$$i_{t+1} = i_t + b S_t i_t - p r i_t$$

note $S_t + i_t = 1$ so only
need one eqn for this model.

How can the mean-field model help us?

- simpler to code
 - easier to analyze
-

SIR equilibria

We can find equilibria by solving for where $s_{t+1} = s_t$,

$$i_{t+1} = i_t \quad , \quad r_{t+1} = r_t$$

(all of these must be true to be at eq.)

This gives us:

$$i_t = 0$$

$$s_t + r_t = 1$$

Note any of these s.t. $s_t + r_t = 1$
will work.

But this shows us that eventually
the disease must die out.

SIS Equilibria

Similarly, need $s_{t+1} = s_t$,
 $i_{t+1} = i_t$. But since $s = 1 - i$,
only need to check for one eqn.

$$i_{t+1} = i_t + b s_t i_t - p r i_t$$

"
 i_t



$$0 = b s_t i_t - p r i_t$$

$$0 = (bs_t - pr) \dot{i}_t$$

two possible equilibria:

Disease free

$$\dot{i}_t = 0$$

$$s_t = 1$$

Endemic

$$s_t = pr/b$$

$$\dot{i}_t = 1 - pr/b$$

We can look at stability
at each of these equilibria -

Note that

$$\dot{i}_{t+1} - \dot{i}_t = \text{change in } \dot{i}$$

if +, \dot{i} grows

if -, \dot{i} declines

$$i_{t+1} - i_t = \underbrace{(b s_t - p_r)}_{\text{this will control the sign}} i_t$$

So, if we start @ disease free equilibrium (DFE) then we will grow more i 's when $b s_t - p_r > 0$ and we will lose i 's when $b s_t^{\approx 1} - p_r < 0$.
If we lose i 's then it will go back to DFE.

If we grow, it will
go to endemic equl.

We can rewrite this condition

as:

$\frac{b}{p_r} < 1 \Rightarrow$ DFE stable
EE unstable
disease dies
out

$\frac{b}{p_r} > 1 \Rightarrow$ DFE unstable
EE stable
disease persists

This is R_0 !

$$b = \underbrace{(N-1)}_{\text{average degree}} \underbrace{p_i p_i}_{P(\text{infection})}$$

expected cases in next time step
generated by 1 infected person
in a fully susc. population
($s \approx 1$)

$1/p_r$ = approximate number of steps
we expect to stay infected.
(this comes from classic probability)*

$\frac{b}{p_r}$ = total cases generated
by 1 infectious individual
in a susc. pop.

Okay, so how does the
mean field actually
match up?

* take an infected node. probability of recovery @ next time = pr

memoryless, so we can write:

$$E[X] = 1 + pr \cdot 0 + (1-pr) E[X]$$

↑ time to recover ↑ I have to take one step. ↑ keep going if didn't. ↑ expected value next time is same as now since memoryless

Solve for $E[X]$:

$$E[X] (1 - (1-pr)) = 1$$

$$E[X] = \frac{1}{pr}$$