1

Building out a research-level model

What all does this actually entail?

Brainstorming example

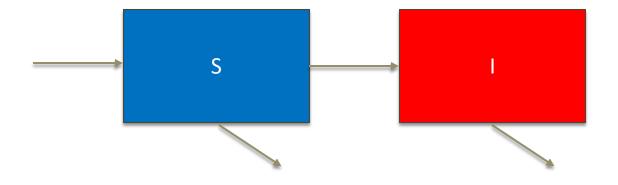
We will talk through an example involving...

HIV among men who have sex with men ©

In base EpiModel, imagine you have built out a model with:

- time step = 1 day
- a population with no attributes other than infection status
- three network models on this population main, casual, one-time
- homogeneous act rate for main, another for casual
- basic vital dynamics and epi for an SI model:
 - arrival rate
 - homogeneous transmission probability
 - departure rate for susceptibles
 - departure rate for infecteds

Brainstorming example



Adding heterogeneity

Let's first build out one attribute. What shall we chose?

Age

- What attribute(s) will we add?
- What module(s) will we add?
- What will the module(s) comprise?
- What existing model elements will be revised?
- What network statistics or model parameters do we need?
- What data do we need?

- What attribute(s) will we add?
 - age
 - but in what unit?
 - and what will we assign as starting values?
- What modules will we add?
 - aging
- What will the module(s) comprise?
 - each node ages by one day per time step

- What existing model elements will be revised?
- What network statistics or model parameters do we need?
- What data do we need?
 - Initialization need to assign everyone age (what will that look like?)
 - Arrivals need to assign an age (what will that look like?)
 - Departures can now also depend on age (what will that look like?)

- What existing model elements will be revised?
- What network statistics or model parameters do we need?
- What data do we need?
 - Networks:
 - Main effect of age how parameterize? What data?
 - Mixing on age how parametrize? What data?
 - Dissolution by age? how parametrize? What data?

- What existing model elements will be revised?
- What network statistics or model parameters do we need?
- What data do we need?
 - Act rate?
 - probably yes, but what are the details?
 - Transmission probability given SI act?

Adding testing

- Now let's add in HIV testing. We'll brainstorm:
 - A bare-minimum approach
 - A more realistic approach

Adding testing: bare minimum

- What attribute(s) will we add?
 - Diagnosis status
 - What values does it take?
 - How do we initialize it?

- What module(s) will we add?
 - Testing

Adding testing: bare minimum

- What will the module(s) comprise?
 - Each node has some probability of testing for HIV each time step
 - (Or, rather, each node who hasn't already tested HIV+)
 - Bare minimum: everyone has the exact same probability, all the time
 - We flip a virtual coin to determine who tests
 - For those that do, their current infection status becomes their diagnosis status

- What existing model elements will be revised?
 - Bare minimum: initialization and arrivals
 - (But if that's it, then what's the point?)

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Adding testing: bare minimum

- What network statistics or model parameters do we need?
 - Daily average probability that an MSM who has not previously tested positive for HIV gets an HIV test
- What data do we need?
 - Survey data about # of tests in last X months/years?
 - Clinical data about # of HIV tests in some setting?
 - (But do we know the population denominator?)

- What attribute(s) will we add?
 - Diagnosis status
 - Time since last test?

- What module(s) will we add?
 - Testing

- What will the module(s) comprise?
 - Each node who hasn't already tested for HIV+ has some probability of testing for HIV each time step
 - To be more realistic, what should this probability depend on?
 - Time since last test?
 - Age of node?
 - Recent sexual behavior?
 - We flip a virtual coin to determine who tests
 - For those that do, we determine their test result. Based on?
 - Current infection status
 - Time since infection
 - Window period of test
 - Sensitivity and specificity of test more generally

- What <u>existing</u> model elements will be revised?
 - Initialize and arrivals
 - Items to disaggregate by dx status:
 - Network:
 - Overall probability of forming a main or casual partnership (nodefactor)
 - Overall probability of breaking a main or casual partnership (nodefactor)
 - Overall probability of a one-time contact (nodefactor)
 - Partner selection for each type (serosorting; nodematch)
 - Other
 - Act rate

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- What net statistics /model parameters do we need?
 And what data do we need?
 - Perhaps need to start thinking about why different people test
 - On some reasonably regular schedule?
 - As the result of a potential exposure?
 - Spontaneously when presented by an easy opportunity?
 - (In practice as part of getting on and being on PrEP but we don't have PrEP in our model yet)
 - Then each is governed by various parameters
 - Perhaps we need to consider a new attribute tester type
 - Does everybody only one type?

What net statistics /model parameters do we need?

- # of partnerships in cross-section by diagnosis status of each member
- Mean relational duration by diagnosis status of each member
- # one-time contacts by diagnosis status of each member per time unit
- Act rate within each type of partnership
- If you're starting to see some data issues arise, hang on.....

And note:

• We've already put age into the model and allowed many of these same things to vary by age – how will the two effects interact?

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- What data do we need?
 - Survey data about partnerships/contacts by dx status
 - But watch out!
 - Is my partner's/contacts diagnosis status always the same as my understanding of it?
 - Might there be systematic ways these diverge, which vary bu partner type?
 - So maybe we need to think about another module next (disclosure)

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- What data do we need?
 - Survey data about partnerships/contacts by dx status
 - But watch out!
 - Does a preponderance of HIV-concordant relationships always reflect a process of partnership selection on dx status?
 - No some of this is endogenous (i.e. p'ships involve transmission)
 - No other network processes might also impact this
 - One in our model already age mixing
 - These other phenomena are can be tricky to account for during the estimation process

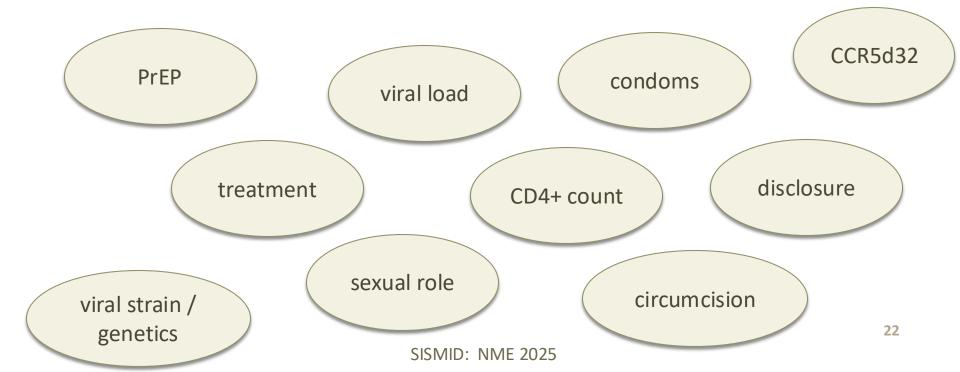
- These other phenomena are can be tricky to account for during the estimation process
 - Age mixing
 - make sure to set HIV prevalence and diagnosis status on the estimation network to reflect their observed levels by age, then have age mixing in the network model
 - Not too hard but requires more data

20

- These other phenomena are can be tricky to account for during the estimation process
 - Endogenous transmission:
 - estimate model as is
 - simulate full transmission model with estimated nodematch(dx.status)
 param nodematch statistic should end up a bit too high in practice
 - simulate full transmission model with a range of nodematch(dx.status)
 params below the estimated value and
 - check their nodematch stats
 - Pick the value that comes closest, or try new values if needed
 - This can be automated through a process known as Approxiate Bayesian
 Computation (ABC), with supporting R packages EasyABC and EpiModelABC

Adding other phenomena

- That was a lot!
- And we've only just begun.
- What other phenomena might you want to add to a reasonable researchlevel HIV model?



Adding other phenomena

- Break into groups based on one phenomenon to add
- Brainstorm all of the questions discussed
- Pay particular attention to challenges that arise that seem particularly distinctive