notes_NovoANDNetwork

Connection between Novozhilov with network models

- Novozhilov(2008)
- J.C. Miller, A.C. Slim & E.M. Volz(2011)
- I.Z. Kiss, J.C. Miller & P.L. Simon(2017)

Novozhilov's Framework for PhenHet

The Novozhilov framework provide a connection between

- heterogeneous in susceptibility and/or infectivity with some known distributions parameterized by a single variable (trait) ω_s or ω_i and time t.
 - Density of susceptible/infected individuals with trait ω_s or ω_i is $s(t,\omega_s), i(t,\omega_i)$
 - pdf $p_s(t,\omega_s)=rac{s(t,\omega_s)}{S(t)}, p_i(t,\omega_i)=rac{i(t,\omega_s)}{I(t)}$ respectively
 - The transmission rate β is determined by the traits ω_s and ω_i independently, such that $\beta = \beta(\omega_s, \omega_i) = \beta_s(\omega_s)\beta_i(\omega_i)$
 - For each traits of S, the density $s(t, \omega_s)$ in trait ω_s is governed by ODE:

$$egin{aligned} rac{\partial s(t,\omega_s)}{t} &= s(t,\omega_s)eta_s(\omega_s)\int_{\Omega_i}eta_i(\omega_i)i(t,\omega_i)d\omega_i \ &= s(t,\omega_s)eta_s(\omega_s)I(t)\int_{\Omega_i}eta_i(\omega_i)p_i(t,\omega_i)d\omega_i \ &= s(t,\omega_s)eta_s(\omega_s)I(t)ar{eta}_i(t) \end{aligned}$$

- $\psi(\omega_i,\omega_i')$ is the probability that a newly infected individual gets trait value ω_i if infected by an individual with trait value ω_i'
- Assume perfect inheriting and fidelity of infectiousness: the infectivity traits ω_i of new infected individual must be inherited from its infector, s.t.

$$\psi(\omega_i,\omega_i')=\delta(\omega_i'-\omega_i)$$

with Dirac delta function $\delta(x)$

• For each traits of I, the density $i(t,\omega_s)$ in trait ω_s is governed by ODE:

$$egin{aligned} rac{\partial i(t,\omega_i)}{t} &= \int_{\Omega_s} \int_{\Omega_i} s(t,\omega_s) eta_s(\omega_s) \psi(\omega_i,\omega_i') i(t,\omega_i') eta_i(\omega_i') d\omega_i' d\omega_s - \gamma i(t,\omega_i) \ &= \int_{\Omega_s} s(t,\omega_s) eta_s(\omega_s) d\omega_s imes \int_{\Omega_i} \delta(\omega_i - \omega_i') i(t,\omega_i') eta_i(\omega_i') d\omega_i' - \gamma i(t,\omega_i) \ &= S(t) \int_{\Omega_s} p_s(t,\omega_s) eta_s(\omega_s) d\omega_s imes eta_i(\omega_i) i(t,\omega_i) \ &= S(t) ar{eta}_s(t) eta_i(\omega_i) i(t,\omega_i) - \gamma i(t,\omega_i) \end{aligned}$$

• Homogeneous models with general nonlinear but separable transmission functions:

$$egin{aligned} rac{d}{dt}S &= -h_s(S) imes h_i(I) \ rac{d}{dt}I &= +h_s(S) imes h_i(I) + \cdots \end{aligned}$$

where h_s and h_i are determined purely by the moment generating function (mgf) of p_s and p_i at initial time t=0.

A subcase of the general framework, with ω_s and ω_i being gamma-distributed and $\beta_s(\omega_s)=\omega_s, \beta_i(\omega_s)=\omega_i \Leftrightarrow \beta=\omega_s\omega_i$ and leads to the known result with power incidence functions:

$$\beta S^p I^q$$

at least for SI model.

- For SIR model, $\beta S^p I$ form has been generated by homogeneous infectivity.
- SI S^pI^q cases are derived in the <u>original paper</u>. A further derivation of SIR case are presented in <u>I_q_and_Exponential_Incident.md</u>.

This frame provide a mechanism derivation to a more general class of non-linear incidence expression and provide a clear relationship from the "parametric" heterogeneity to the non-linear incidence term.

However, this framework is limited to following assumptions:

- 1. the infectivity traits ω_i of new infected individual must be inherited from its infector
 - $\psi(\omega_i, \omega_i')$ is the probability that a newly infected individual gets trait value ω_i if infected by an individual with trait value ω_i' and require $\psi(\omega_i, \omega_i') = \delta(\omega_i' \omega_i)$
 - This guaranteed that for each infectivity trait ω_i and time t, the population of the infection with such trait satisfy:

$$egin{aligned} rac{\partial s(t,\omega_s)}{t} &= -s(t,\omega_s) imes F(S(t),I(t),\omega_i) \ rac{\partial i(t,\omega_i)}{t} &= i(t,\omega_i) imes G(S(t),I(t),\omega_s) \end{aligned}$$

So Theorem 1 in the article could be applied.

- This is somehow justifiable if we consider the strain of pathogen or homogeneous in infectivity
- 2. The heterogeneity only applies to infectivity and susceptibility while contact rate of individuals are still assumed to be homogeneous and fully-mixed.
- 3. The transmission probability/rate $\beta = \beta(\omega_s, \omega_i)$ are determined by traits of infectivity and susceptibility independently s.t. $\beta = \beta(\omega_s, \omega_i) = \beta_s(\omega_s)\beta_i(\omega_i)$

Network Models with similar settings

Some similar approach using traits and generating functions has been applied to considering heterogeneity in contact rates with networks, by R.M. Anderson, A.L. Lloyd & R.M. May($\underline{1988}$, $\underline{1991 \text{ book}}$, $\underline{2001}$) and R. Pastor-Satorras & A. Vespignani ($\underline{2001}$, $\underline{2002}$) which release the assumption 2. In these approaches, the traits ω are replaced by the degree k of the individual/vertex.

- These network model assume that degree is invariant with time and infection events.
- Assumption 1, thus the Novozhilov's framework, cannot be applied to network model.

The system of these model can be written into following system by degree

$$egin{aligned} rac{\partial s(t,k)}{\partial t} &= -s(t,k) imes k imes eta imes \int_{j} rac{jP(j)i(t,j)}{\langle K
angle} dj \ &= -s(t,k)keta \int_{j} P_{e}(j)i(t,j)dj \ rac{\partial i(t,k)}{\partial t} &= +s(t,k) imes k imes eta imes \int_{j} rac{jP(j)i(t,j)}{\langle K
angle} dj - \gamma i(t,k) \ &= s(t,k)keta \int_{j} P_{e}(j)i(t,j)dj - \gamma i(t,k) \end{aligned}$$

- P(k) is the degree distribution of the network
- s(t,k) and i(t,k) are the density of susceptible/infected individual with degree k at time t.
- β is the transmission probability of each edge
- $\langle K
 angle = \int_k k P(k)$ is the expected degree
- $P_e(k)=rac{kP(k)}{\langle K
 angle}$ is the pdf of excess degree distribution, or the degree-biased distribution
 - ullet Also, the probability of a uniformly randomly chosen edge belongs to a vertex with degree k in the network
 - ullet Also, the probability of a randomly chosen neighbor vertex has degree k
 - Same idea with the size/length-biased sampling: the probability of such neighbor being chosen is proportional to its degree.
- Comparing with Novozhilov's framework without assumption 1,

- $\beta_s(k) = k\beta$ is determined by contact rate/degree of susceptible vertex
- $\beta_i(j) = P_e(j)$ is determined by contact rate/degree of the infected vertex
- However, ψ is now depend on trait/degree of s(t,k) instead of i(t,j), such that $\psi=\psi(k,k')=\delta(k-k')$

We can find a solution for equation of s(t, k):

$$s(k,t) = e^{-eta k \int_{-\infty}^t P_e(j) i(t',j) dt'}$$

we take $heta(t) = e^{-eta \int_{-\infty}^t P_e(j) i(t',j) dt'}$ then $s(k,t) = heta(t)^k$.

J.C. Miller, A.C. Slim & E.M. Volz(2011) has proved that with this change of notation, this model is equivalent to their percolation-based model assuming Mean-Field Social Heterogeneity (MFSH) where at each moment, every individual/vertex reform their edges uniformly randomly while keeping their assigned degree and all neighbors are interchangeable in each compartment.

Strong connection with random-network models with assumptions other than MFSH are discussed in I.Z. Kiss, J.C. Miller & P.L. Simon(2017)

Also, Newman & I.Z. Kiss, J.C. Miller & P.L. Simon(2017) has proved that in static/configuration network with large network limit (\$N\$ large enough), the final epidemic size and epidemic threshold of parametric heterogeneous transmission probability $(\tau(x))$ and recovery $(\gamma(x))$ model is the same with homogeneous model with expectation as parameter $(\tau = \langle \tau \rangle, \gamma = \langle \gamma \rangle)$.

However, the dynamic will be different and <u>I.Z. Kiss, J.C. Miller & P.L. Simon(2017)</u> has discussion in Chapter 9.

New findings

MSV type network model

Network model formalized by <u>J.C. Miller, A.C. Slim & E.M. Volz(2011)</u>.

Result agreed with a recent paper by RomanescuEtAL(2023) and Novozhilov(2008)

Assumptions

- 1. Neighbors are independent
- 2. Infinite size random network: a.s. no loop network

For random distribution given a degree distribution with PDF: $\mathbb{P}(K = d) = p_d$. Following J.C. Miller, A.C. Slim & E.M. Volz(2011) at any moment t we define $\theta(t)$:

- (MSV definition) the probability that randomly chosen neighbor vertex b of a randomly chosen vertex a has not yet transmit the infectious to a.
- (JD definition) the probability of a randomly chosen edge in the network has not yet transmit infection. Let $\phi = 1 \theta$ be the probability that the infection has transmitted.

The probability generating function(PGF) of degree distribution is denoted by:

$$G_p(x) = \sum_{d=0}^{\infty} p_d x^d$$

• A useful expression would be the mean degree δ is given by:

$$\delta = \sum_{d=0}^\infty p_d d = rac{d}{dx} G_p(x)|_{x=1} = G_p'(1)$$

Also for network model, **Excess degree** is also important, as during the outbreak, any newly infected vertex with degree k could only infect at most k-1 of its susceptible neighbors, as its infection must come from one neighbor that already being infected. Based on given degree distribution with PDF p_d , we could define the distribution of excess degree with PDF denoted by q, such that:

$$\mathbb{P}(ext{excess degree} = d-1) = q_{d-1} = rac{p_d d}{\sum_{k=0}^{\infty} p_k k} = rac{p_d d}{\delta}$$

The corresponding PGF for excess degree is

$$G_q(x) = rac{G_p'(x)}{\delta}$$

Consider the effective "incidence" term using

$$ho = rac{{\cal R}_{
m eff}}{{\cal R}_0}$$

Follow JD's idea:

$$ho = rac{\mathcal{R}_{ ext{eff}}}{\mathcal{R}_0} = rac{\sigma_\phi}{\sigma_0}$$

• (??) How to connect $R_{\rm eff}$ with incidence term $\frac{dS}{dt}$?

$$R_{
m eff} = rac{rac{dS(t)}{dt}}{I(t)} imes rac{1}{\gamma}$$

• (??) How to define σ_{ϕ} : Expected number of susceptible neighbors???? Expected number of edges that can still transmit the infection for each newly infected vertex??

$$\sigma_\phi = \sum_{d=0}^\infty p_d imes d imes (1-\phi)^d = \sum_{d=0}^\infty p_d imes d imes heta^d = heta \sum_{d=0}^\infty p_d imes d imes heta^{d-1} = heta G_p'(heta)$$

• (??) How to define σ_0 :

$$\sigma_0 = \lim_{t o 0} \sigma_\phi = \lim_{ heta o 1} heta G_p'(heta) = \delta$$

Therefore,

$$ho = rac{ heta G_p'(heta)}{\delta}$$

- Follow RomanescuEtAL(2023): $\mathcal{R}_{\text{eff}}(t)$ is the expected number of secondary infections for one infected individual X_t at time t.
 - Proportion of susceptible vertices with degree k at time t in the entire population $p_k^S(t)=p_k\theta^k$
 - Total proportion of susceptible nodes at time t is $S(t) = G_p(\theta(t))$
 - Corresponding PGF is $rac{G_p(x heta)}{G_p(heta)} = rac{G_p(x heta)}{S}$
- Observe the process by which a susceptible individual becomes infected.
 - Consider a random edge that has the potential to transmit infection at time t. The uninfected individual at the end of this edge is chosen from the susceptible set, but not at random: an individual's chance of being selected is proportional to their degree, in the absence of higher-order features. Thus, the relative frequency of an individual of degree k becoming infected at the next time step is proportional to $kp_k^S(t)$.

Since neighbors are assumed to be independent, they claim

$$heta = G_p^{-1}(S)$$

Consider Poisson distribution with $p_k = k!$

For a Poisson degree distribution

$$\rho = S(1 + \frac{\log S}{\delta})$$

Agree both with Novozhilov(2008) and a recent paper by RomanescuEtAL(2023)

A generalized version to negative binomial degree distribution (which could be seen as a discrete analog of gamma)

$$ho = S(S^{\kappa} + rac{S^{\kappa} - 1}{\kappa \delta})$$

.