EPIDEMIC PROPAGATION ON NETWORKS: A DIFFERENTIAL EQUATION APPROACH

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- $S \rightarrow I$, rate: $k\tau$, k is the number of I neighbours.
- $I \rightarrow S$, rate: γ

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Known models:

- Mean-field equation
- Pairwise model
- Compact pairwise model
- ...

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(R has no effect on the propagation)

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- $S \rightarrow I$, rate: $k\tau$, k is the number of I neighbours.
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Example: Influenza in Hungary in 2016. Weekly number of new reported cases for 100,000 persons.



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Further models: SEIR (E stands for exposed), SIRS, SEIRS, ...

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For simplicity, we present the theory for the SIS.

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State space for a triangle graph



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- Infection: $SIS \rightarrow IIS$
- Recovery: $SIS \rightarrow SSS$

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Master equations are formulated for the probabilities of states.

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 $X_{SIS}(t)$ is the probability of state SIS at time t.

MASTER EQUATIONS

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$$\begin{split} \dot{X}_{SSS} &= \gamma (X_{SSI} + X_{SIS} + X_{ISS}), \\ \dot{X}_{SSI} &= \gamma (X_{SII} + X_{ISI}) - (2\tau + \gamma) X_{SSI}, \\ \dot{X}_{SIS} &= \gamma (X_{SII} + X_{IIS}) - (2\tau + \gamma) X_{SIS}, \\ \dot{X}_{ISS} &= \gamma (X_{ISI} + X_{IIS}) - (2\tau + \gamma) X_{ISS}, \\ \dot{X}_{ISI} &= \gamma X_{III} + \tau (X_{SSI} + X_{SIS}) - 2(\tau + \gamma) X_{SII}, \\ \dot{X}_{ISI} &= \gamma X_{III} + \tau (X_{SSI} + X_{ISS}) - 2(\tau + \gamma) X_{ISI}, \\ \dot{X}_{IIS} &= \gamma X_{III} + \tau (X_{SIS} + X_{ISS}) - 2(\tau + \gamma) X_{IIS}, \\ \dot{X}_{III} &= -3\gamma X_{III} + 2\tau (X_{SII} + X_{ISI}) + X_{IIS}), \end{split}$$

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 2^N equations for a graph with N nodes

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The size of the system can be reduced by using the automorphisms of the graph:

Simon, P.L., Taylor, M., Kiss., I.Z., Exact epidemic models on graphs using graph-automorphism driven lumping, J. Math. Biol., 62 (2011).

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Approximating differential equation for [/]

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This is the well-known compartmental model, which does not give accurate result for networks.

Reason: the approximation assumes random distribution of infected nodes.

Better idea: derive a differential equation for [*SI*], this leaded to the pairwise model.

Keeling, M.J., The effects of local spatial structure on epidemiological invasions, *Proc. R. Soc.* Lond. B 266 (1999), 859-867.

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Approximation:

$$[ABC] \approx \frac{n-1}{n} \frac{[AB][BC]}{[B]}, \quad n \text{ average degree}$$

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M. Taylor, P. L. Simon, D. M. Green, T. House, I. Z. Kiss, From Markovian to pairwise epidemic models and the performance of moment closure approximations, *J. Math. Biol.* 64 (2012), 1021-1042.

Regular random graph with N = 1000 nodes, average degree n = 20, $\gamma = 1$, critical value of τ from compartmental model: $\tau_{cr} = \gamma/n$

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Mean-field: dashed, Pairwise: continuous Simulation (average of 200 runs): grey thick curve

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 $\tau = \tau_{cr} \Leftrightarrow$ basic reproduction number $R_0 = 1$.

Bimodal random graph with N = 1000 nodes, average degree n = 20, $\gamma = 1$, $\tau = 2\tau_{cr} = 2\gamma/n$

N/2 nodes have degree d_1 , N/2 nodes have degree d_2 .

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Reason of inaccuracy: in the closure $[ABC] \approx \frac{n-1}{n} \frac{[AB][BC]}{[B]}$ it is assumed that each node has the same degree *n*.

Mean-field model at the level of singles:

$$\dot{[I]} = \tau \frac{n}{N} [I] (N - [I]) - \gamma [I].$$

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If $\tau(n-1) < \gamma$, then the disease-free steady state is globally stable.

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Closure: $[S_k I]$ can be expressed in terms of singles, $[S_k]$,

or in terms of pairs, [SI], and singles.

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Compact pairwise model: K + 3 equations

Bimodal random graph with N = 1000 nodes, average degree $n_1 = 20$, $\gamma = 1$, $\tau = 3\gamma n_1/n_2$, $n_i = \sum d_k^i p_k$ N/2 nodes have degree $d_1 = 5$, N/2 nodes have degree $d_2 = 35$.

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Pairwise: dashed, Compact pairwise: continuous red, Simulation (average of 200 runs): grey thick curve

A graph with N nodes is given

The nodes can be susceptible (S) or infected (I)

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AIM: Derive a simple system of differential equations for the expected number of infected nodes [I](t).

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Models:

- Mean-field: no network structure
- Pairwise: regular random graph
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Analysis of the ODE models

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Degree-based mean-field model at the level of singles:

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$$\langle d \rangle = \frac{1}{N} \sum_{k=1}^{K} d_k N_k$$
 and $\langle d^2 \rangle = \frac{1}{N} \sum_{k=1}^{K} d_k^2 N_k$.

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If $\tau(\langle d^2 \rangle - \langle d \rangle) > \gamma \langle d \rangle$, then the endemic steady state is stable.

Models:

- Mean-field: number of nodes in different states
- Pairwise: number of nodes and edges in different states
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Adaptive network: cutting SI links, creating SS links

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The nodes can be susceptible (S) or infected (I)

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Transitions:

- infection, rate: $k\tau$, k is the number of I neighbours.
- recovery, rate: γ
- SS link creation, rate α
- SI link deletion, rate ω

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SIS EPIDEMIC ON AN ADAPTIVE NETWORK



MATHEMATICAL MODEL

$$\begin{aligned} &[\dot{I}] &= \tau[SI] - \gamma[I], \\ &[\dot{S}I] &= \gamma([II] - [SI]) + \tau([SSI] - [ISI] - [SI]) \\ &[\dot{I}I] &= -2\gamma[II] + 2\tau([ISI] + [SI]), \\ &[\dot{S}S] &= 2\gamma[SI] - 2\tau[SSI] \end{aligned}$$

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$$\begin{split} &[\dot{I}] &= \tau[SI] - \gamma[I], \\ &[\dot{S}I] &= \gamma([II] - [SI]) + \tau([SSI] - [ISI] - [SI]) - \omega[SI], \\ &[\dot{I}I] &= -2\gamma[II] + 2\tau([ISI] + [SI]), \\ &[\dot{S}S] &= 2\gamma[SI] - 2\tau[SSI] + \alpha([S]([S] - 1) - [SS]). \end{split}$$

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POSSIBLE MODEL OUTCOMES



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 $\tau = 0.1, \, \gamma = 1, \, \textit{N} = 1000$

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The infection rate is different in each layer: $\tau_{home} = 1$, $\tau_{wp} = 1/2$, $\tau_{sch} = 1/2$, $\tau_{geom} = 1/10$, $\tau_{store} = 1/20$.

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Network of a model city: weighted graph with four layers.

Total population N = 10000, is divided into households of different sizes.

Household structure: number of children, adults and elderly people.

Geometry: adjacency matrix of the households.

Work places and schools: complete graphs with given sizes and randomly chosen members.

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The network is not a random graph, hence ODE approximations are more difficult to derive.

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This is a joint work with Ágnes Backhausz and Bence Bolgár.

Results of a Gillespie simulation on the above network with open (red) and closed (black) schools.

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 R_{∞} : final epidemic size, i.e. proportion of the population having immunity when the epidemic is over. ($N = 10000, R_0 \approx 2.$)

The results for epidemic processes are summarized in our book:

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Thank you for your attention!

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