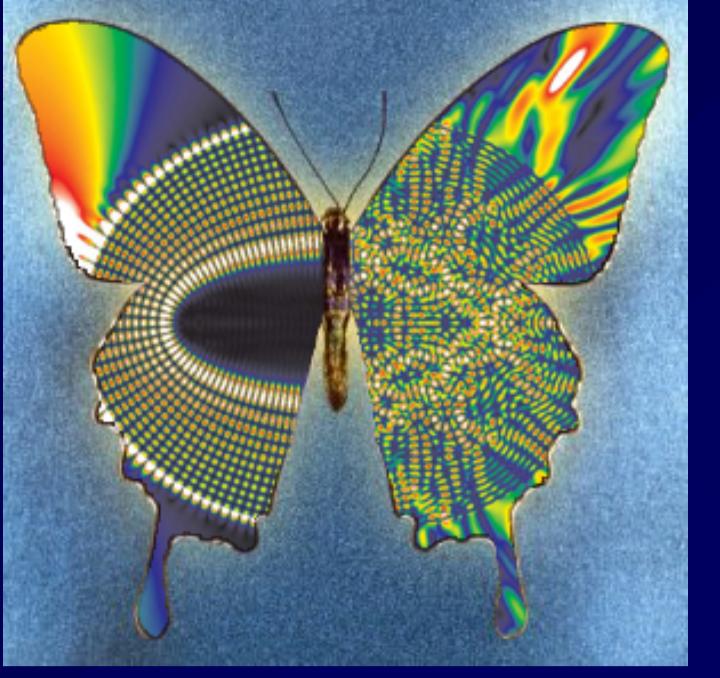


Time Evolution of Epidemics on Complex Networks

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Motivation

Efficient tools are required by decision makers in matter of disease propagation to:

- **predict** and **quantify** risks;
- **optimize** prevention methods; and
- **evaluate** intervention scenario efficacy.

Numerical simulations? Part of the solution...

Pros

- Generality } Describe the dynamics,
- Availability } program what you have described!

Cons

- Computation time (some simulations “impossible”).
- Lack of insight (compared to analytical solutions).

Our goal is to provide **analytical tools** that are as general and as close to the real-world system as possible.

Only for diseases?
Not at all!
Almost anything that spread from neighbour to neighbour can be simulated with such formalisms.
For example: computer viruses, rumors, email chains...

Fundamentals

At a given time, each individual is in a specific **state**.

Susceptible individuals do not have the disease but can get infected by contact with infectious.

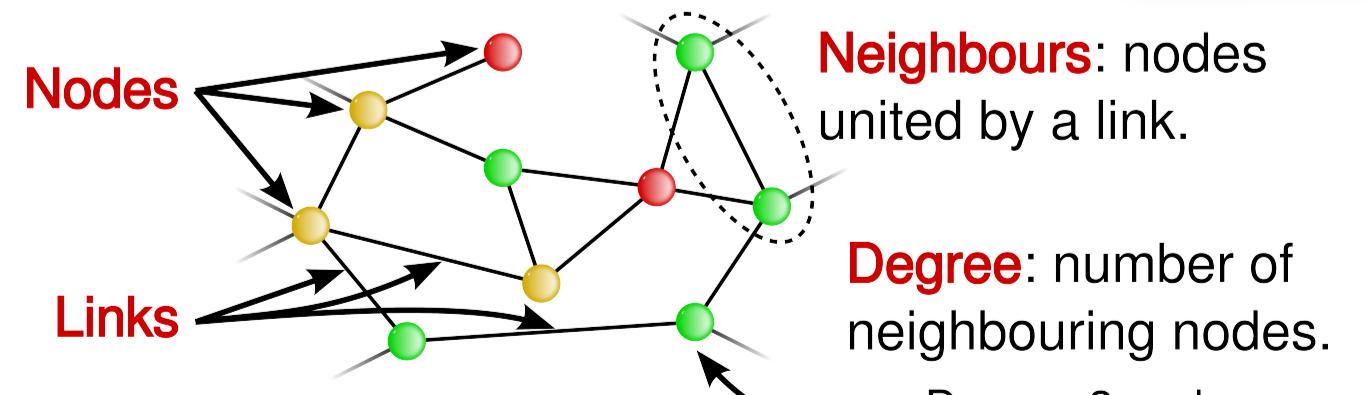
Infectious individuals have the disease and can transmit it to susceptibles through contacts.

Removed individuals neither have nor can get the disease. This include death, quarantine and recovery by immunization.

In traditional compartmental models, anyone can have contact with anyone (the fully-mixed approximation) and the dynamics is governed by “mass-action” ODE ($S + I + R = N = \text{constant}$).

$$\begin{aligned}\dot{S} &= -\beta SI \\ \dot{I} &= \beta SI - \mu I \\ \dot{R} &= \mu I\end{aligned}$$

In network models, contacts are restricted by the **network structure**.



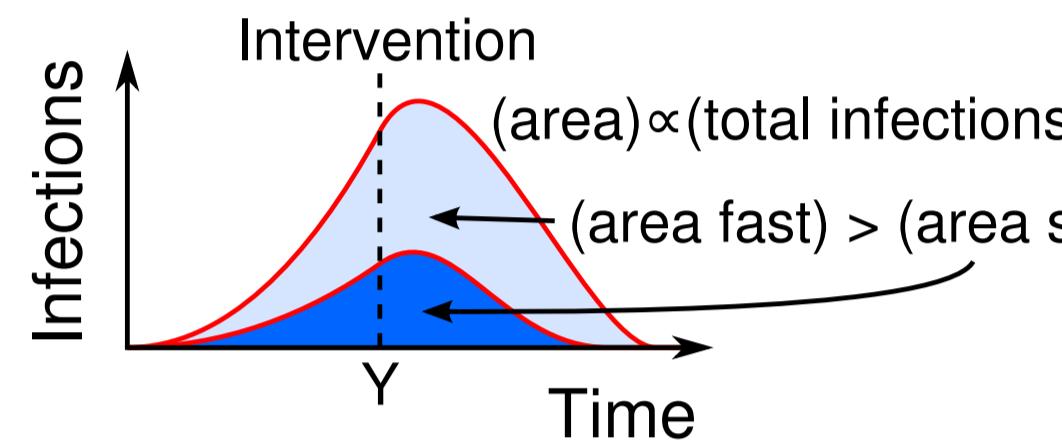
Many real-world networks are highly **heterogeneous in degree** (e.g. STIs). The formalisms presented here hold for random networks with arbitrary degree distribution.

ODE?
Ordinary Differential Equation.
STI?
Sexually Transmitted Infections.
PGF?
Probability Generating Function.

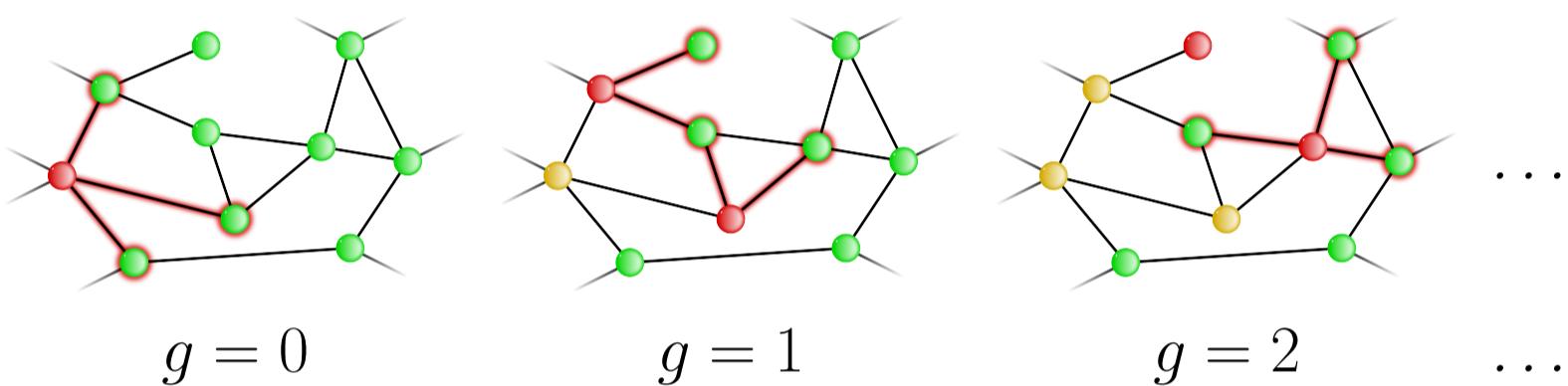
Generations and finite-size

[Noël et al. 2009]

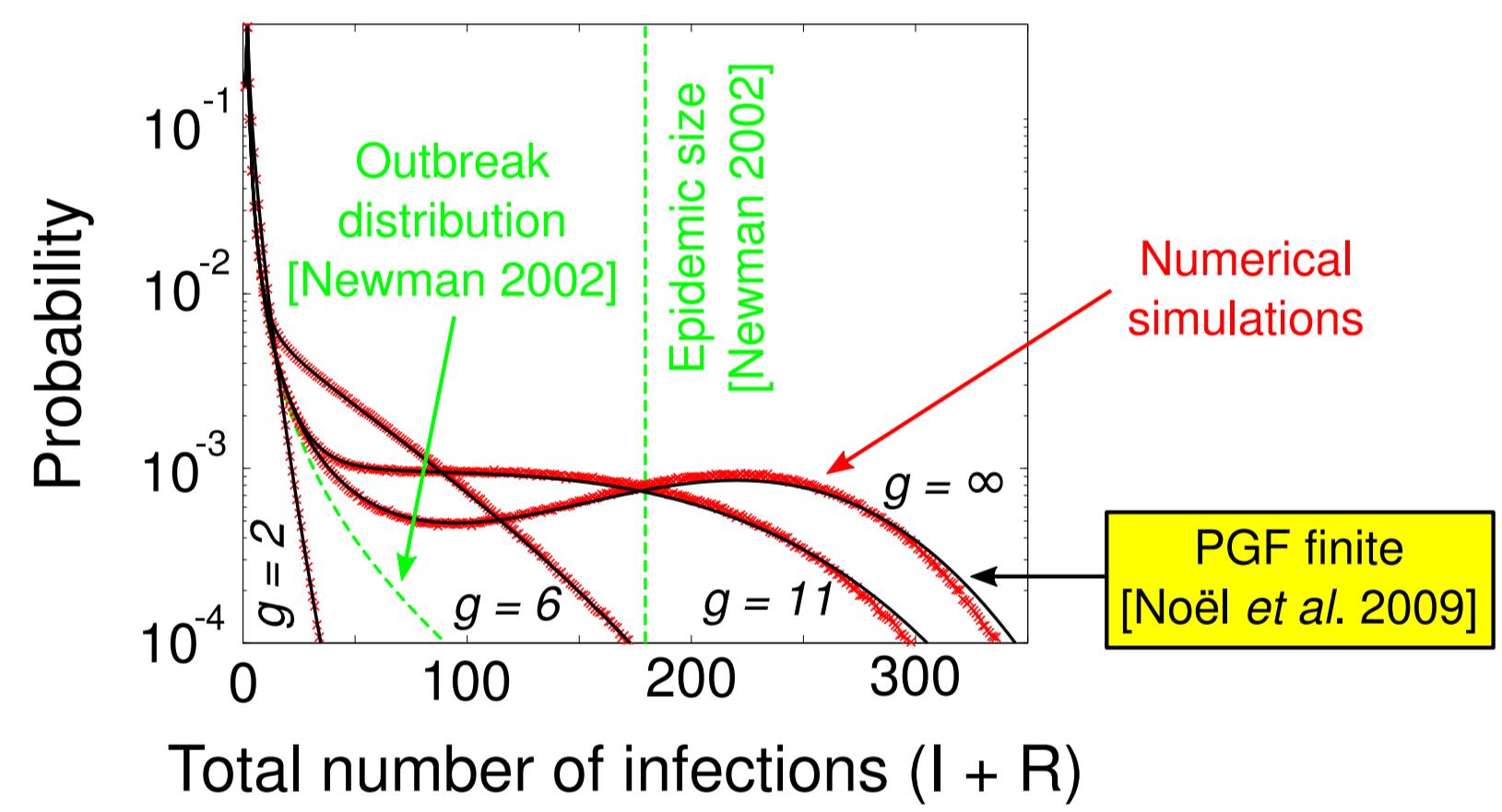
Time is **crucial** for intervention scenarios. For example, what if a $X\%$ efficient vaccine is discovered after Y months?



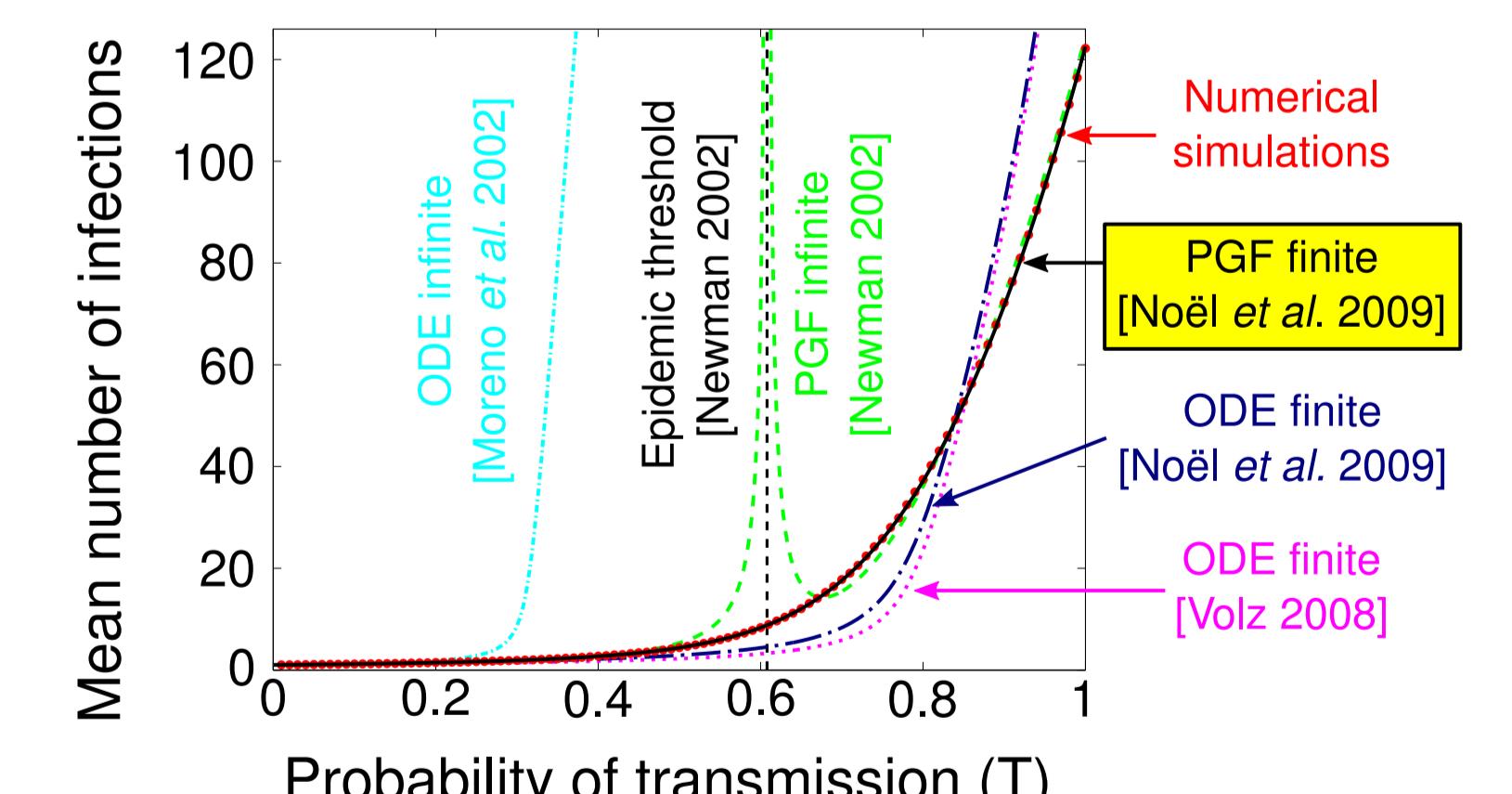
Discrete time approach: **generations**.



We first use an infinite-size network approximation based on a **PGF** formalism [Newman 2002] adapted to include generations [Marder 2007] then add **corrections** for the **finite size** of the network.



The results for the final state ($g = \infty$) of this finite-size algorithm also compares favourably with those of other formalisms.



Contribution A

Formalism for **discrete time evolution** on **finite-size** networks with **heterogeneous degree distribution**.

Information about the simulations

Network of $N = 1000$ nodes. Each node has probability $p_k \propto k^{-2} e^{-k/5}$ to be of degree k . In the first figure, the transmission probability is $T = 0.8$.

Types of nodes

[Allard et al. 2009]

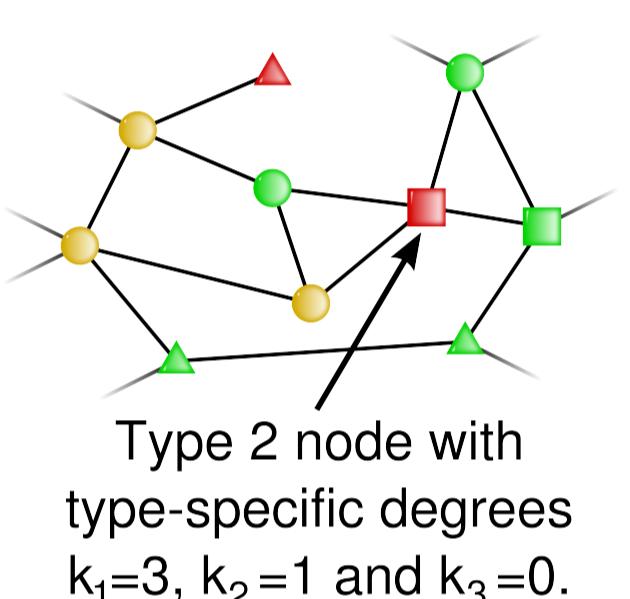
Individuals differ from one another (e.g. age, gender, sociocultural group, ...). This affects the dynamics in two important ways by:

- introducing **correlations** on how the nodes are linked to one another (**network structure**) and
- modifying the **probability of transmission** between connected nodes.

We use a **PGF** formalism [Newman 2002] generalized for many different **types** of nodes.

type 1: , type 2: , type 3: , ...

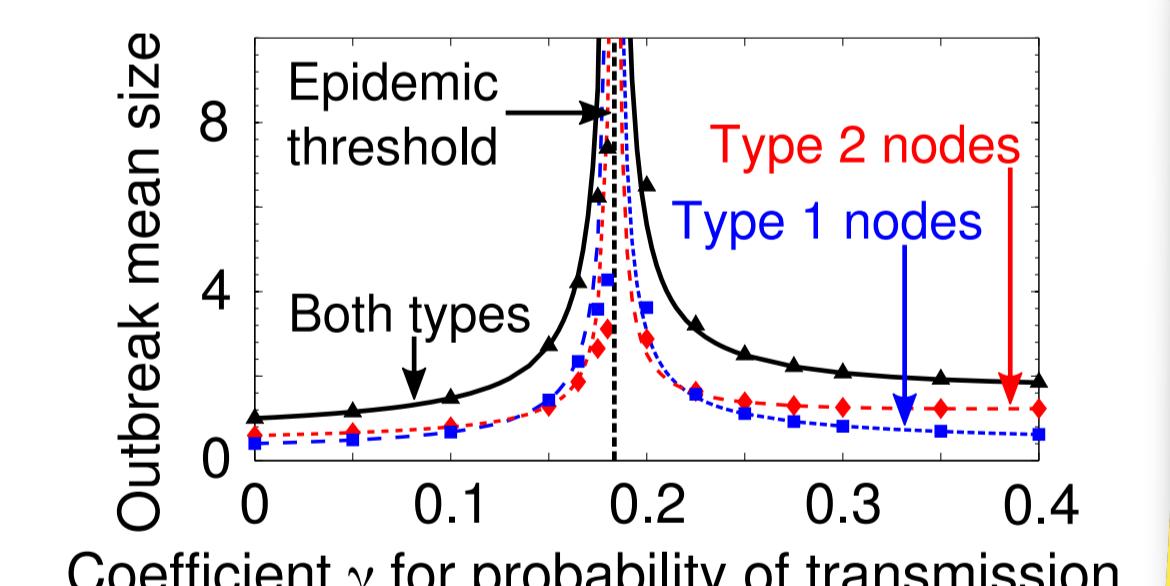
This allows for new **network structures** where the degree k_j of a node must be specified for each possible type j .



	$T_{11} T_{12} T_{13} \dots$
	$T_{21} T_{22} T_{23} \dots$
	$T_{31} T_{32} T_{33} \dots$
\vdots	$\vdots \vdots \ddots$

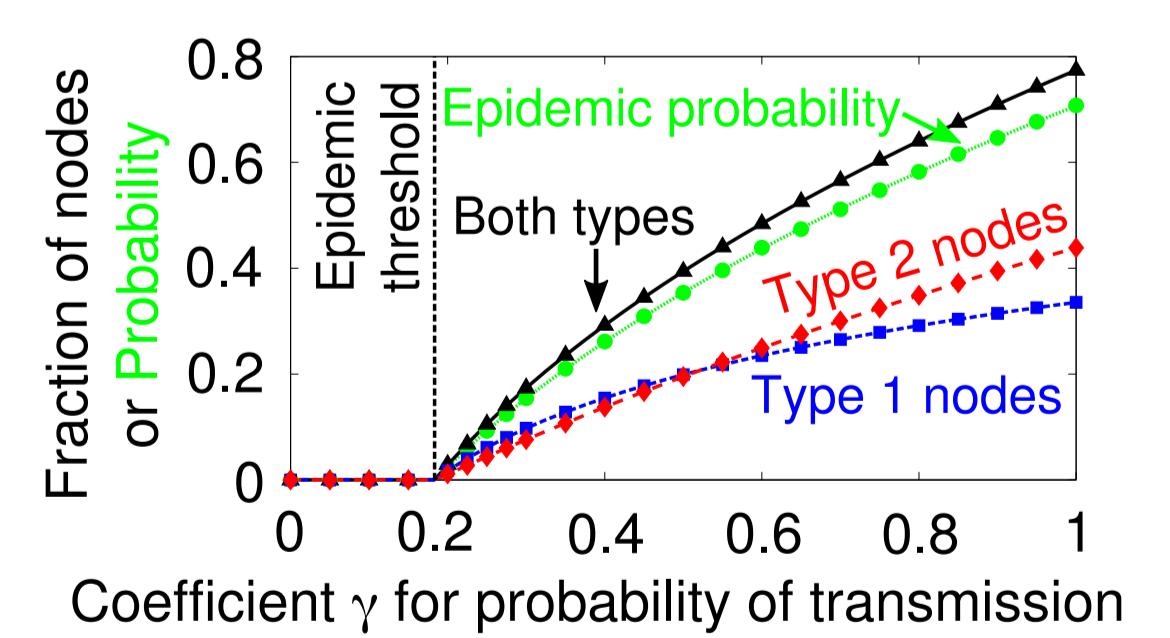
Different **probability of transmission** are considered for each ordered pair of infectious and susceptible node types.

Using a 2 types network, simulations (symbols) confirm analytical results (lines) for outbreaks...



Outbreak?
In arbitrarily large networks, an outbreak's size (number of individuals affected) does not scale with the network size (e.g. unchanged by increasing network size by a factor 10). Outbreaks can occur both below and above the epidemic threshold.

... as well as for large-scale epidemics!



Epidemics?
Epidemics affect a fraction of the network; the total number affected scale with network size (e.g. 10 times more affected in a 10 times larger network). In addition to the fraction affected, epidemics are also characterized by their probability of occurrence.

Work in progress

Mapping generations to continuous time

This is an add-on to **Contribution A** aiming at mapping this **discrete-time generation based formalism** to **observables varying continuously in time**.

Basically, we get the infected proportion of each generations at any given time then translate this to the desired observable. Complications occur when one introduces finite-size.

Genuine continuous-time formalism

This is an alternative to **Contribution A** aiming for a true **continuous time evolution** formalism on **finite-size** networks with **heterogeneous degree distribution**.

We have designed a high-dimensional ODE system that fulfil these requirements. Some properties of the formalism indicate that an analytical solution might be possible for important special cases.

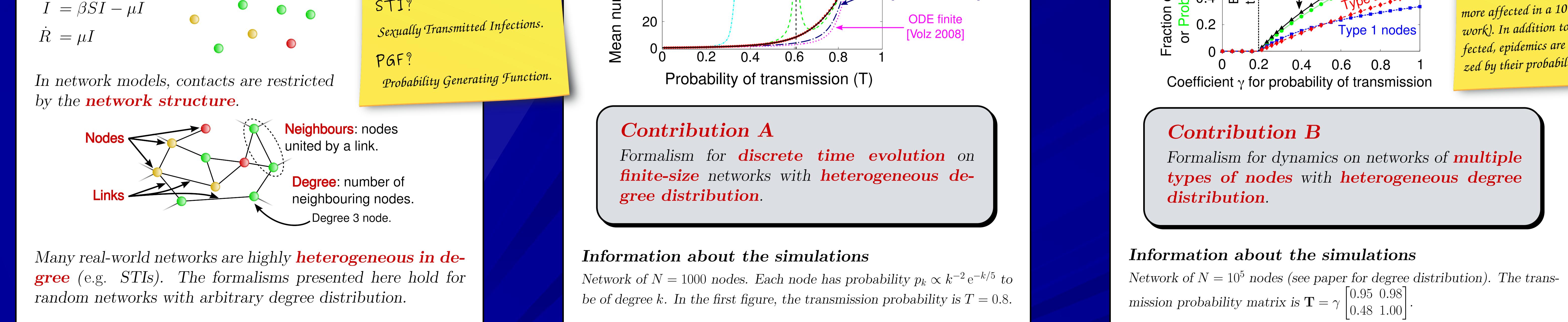
Types of links

We extend the **PGF** formalism of **Contribution B** in order to include different **types of links**.

Similarly to the “type of nodes” approach of Contribution B that needs a list of degrees for every node type, this approach requires a list of degrees for every link type (types of nodes are a special case of this formalism). In addition to allow new network structures, this could make the formalism particularly well adapted for survey data.

Future directions

- Inclusion of clustering.
- Dynamical networks that evolve in time.
- Co-evolution of networks and dynamics (retroaction of process dynamics on network topology).
- Co-infection (interaction of more than one disease).
- Networks of networks.



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