

Epidemics on Networks

Introduction to Network Science

Instructor: Michele Starnini — <https://github.com/chattox/networks-science-course>



**Universitat
Pompeu Fabra**
Barcelona

Content

- **Degree-based solution of SIS model on SF networks**
- **Real-world network epidemiology**

Modeling epidemics

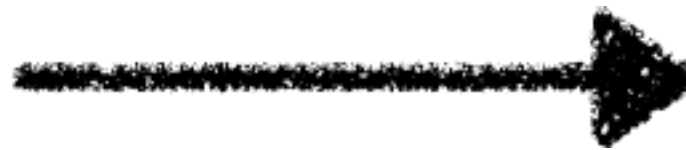
- **Modeling Epidemic process (dynamics):**
 - Branching process
 - SI model
 - SIR model
 - SIS model
- **Modeling underlying network substrate (static)**
 - Mean-field mixing (fully connected network)
 - Homogeneous networks (ER networks)
 - Heterogeneous networks (SF networks)

Modeling underlying network

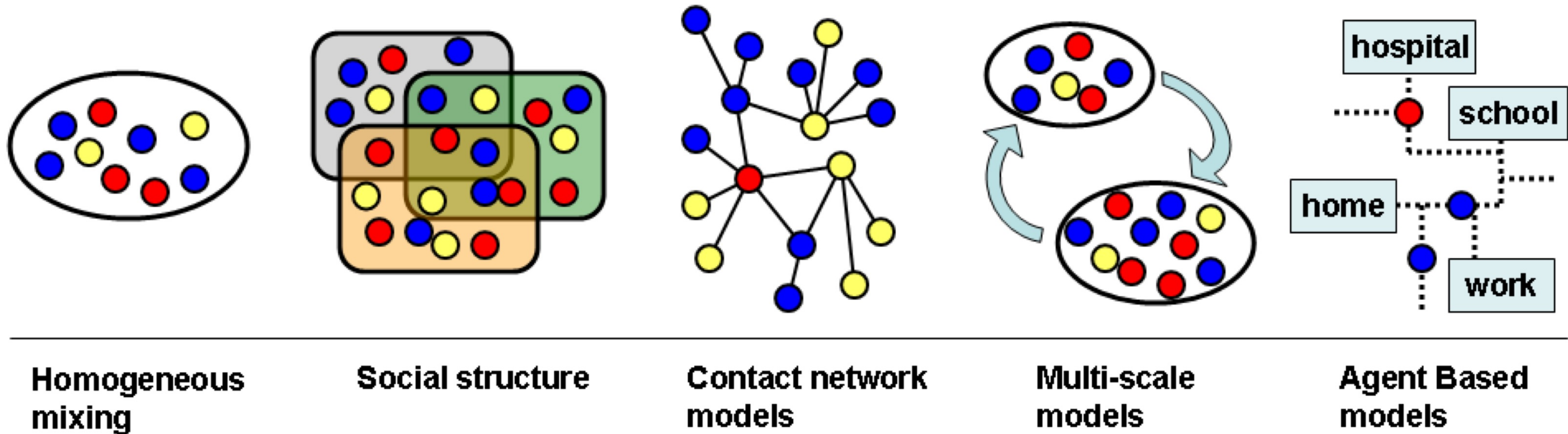


← reality

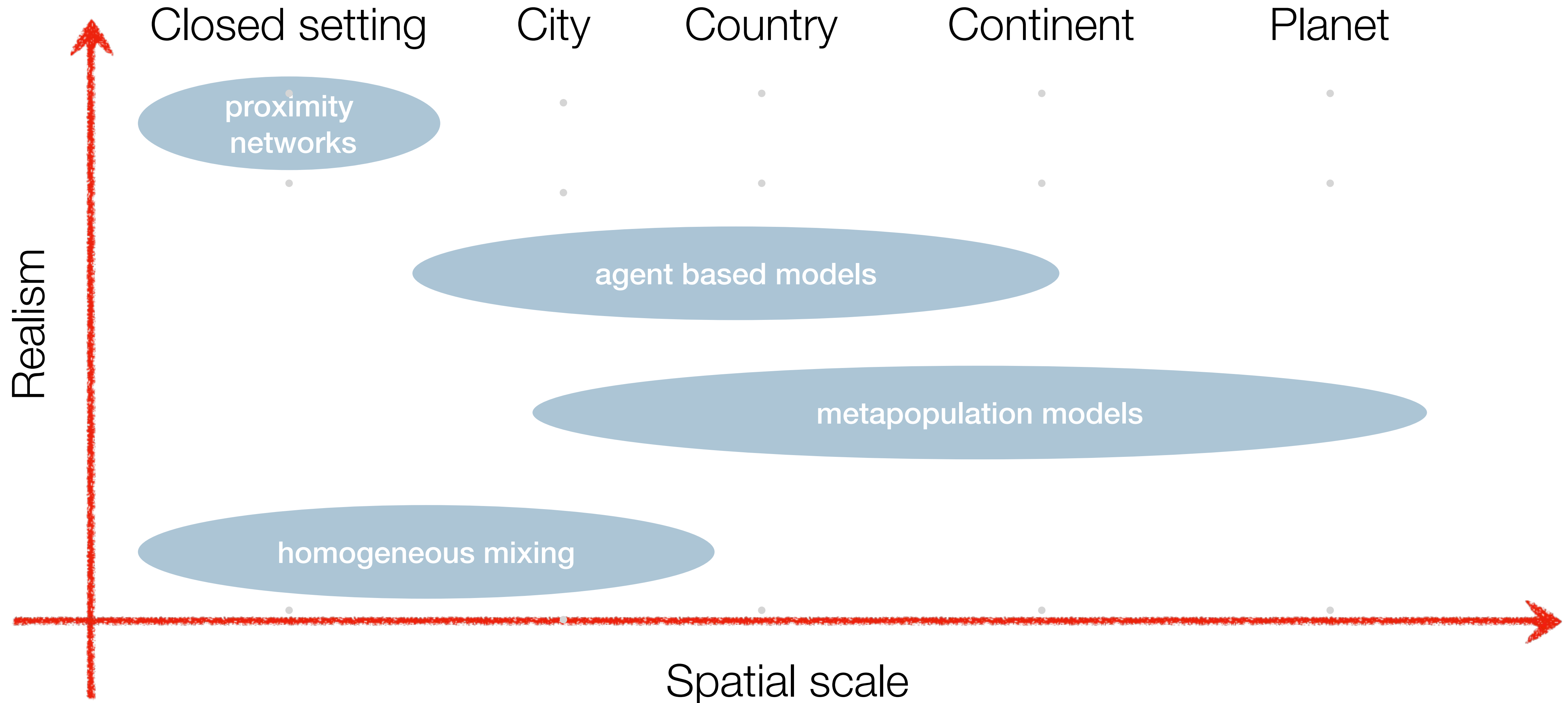
abstraction,
conceptualization



Modeling underlying networks

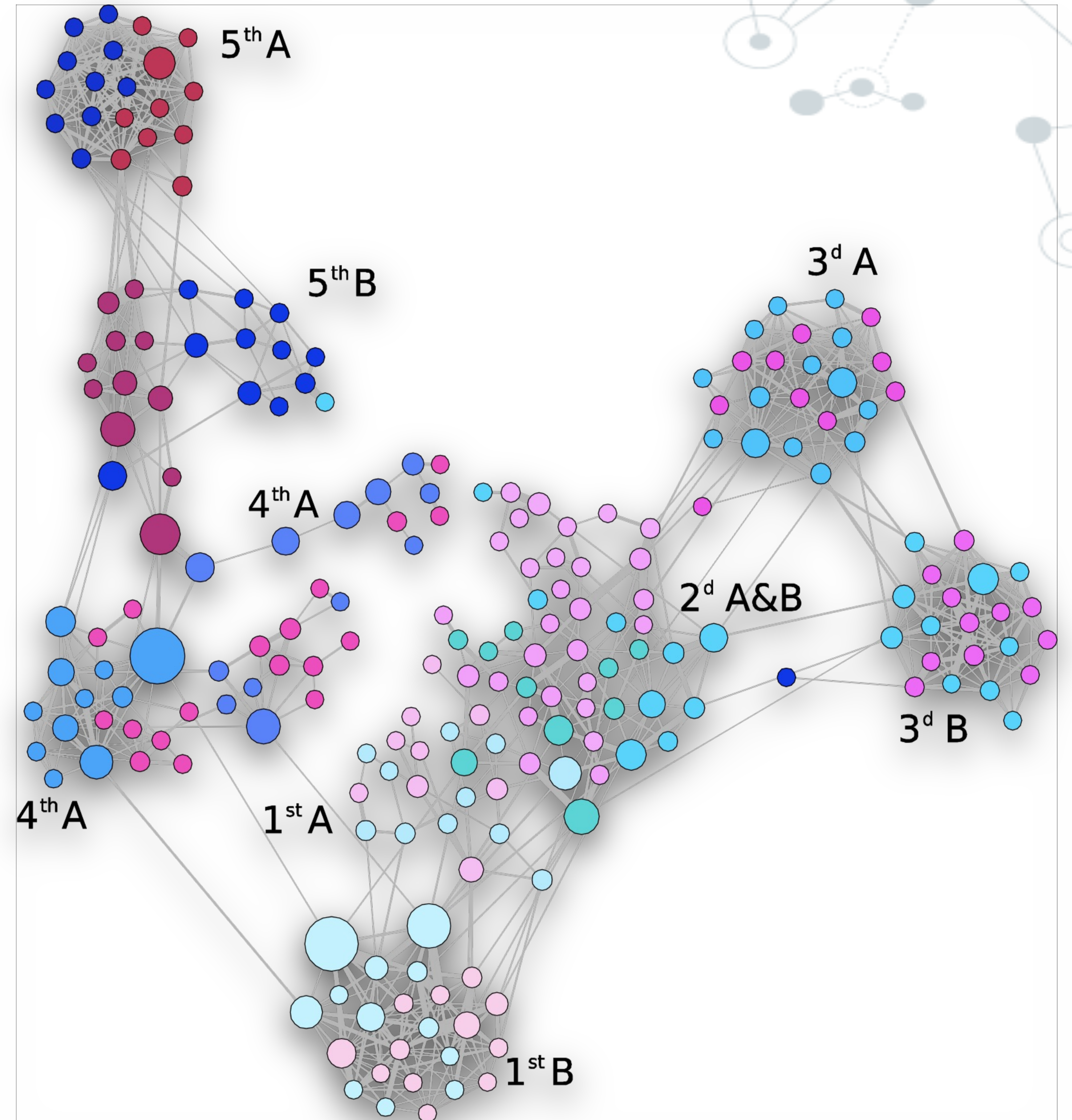


Spatial scales



Epidemics on networks

- ▶ Homogeneous mixing is not always realistic
- ▶ Contacts are not equal and not constant across groups.
- ▶ Real contact networks display high heterogeneities



Degree-based solution of the SIS model on scale-free networks



Epidemics on networks

- We consider a network of N nodes where each node can be in an epidemic state, S , I or R
- We define the density of nodes in a given state, as:

$$\rho^S(t) = \frac{S(t)}{N}, \rho^I(t) = \frac{I(t)}{N}, \rho^R(t) = \frac{R(t)}{N}$$

Degree-based mean field

- ▶ Nodes with the **same degree k** are considered as **statistically equivalent**
- ▶ Fraction of nodes in each compartment: ρ_k^α , $\alpha = S, I, R$
- ▶ These variables are not independent: $\sum_{\alpha} \rho_k^\alpha = 1$
- ▶ Fraction of individuals in compartment α at time t to $\rho^\alpha(t) = \sum_k P(k) \rho_k^\alpha(t)$

Degree-based mean field

- ▶ The network is considered in a mean-field perspective (**annealed network** approximation).
- ▶ The adjacency matrix is completely “destroyed”. Only the degree and the two-vertex correlations of each node are preserved.
- ▶ The adjacency matrix is replaced by its ensemble average:

$$\overline{A}_{ij} = \frac{k_j P(k_i | k_j)}{NP(k_i)}$$

The DBMF SIS model

$$\frac{d\rho_k^I(t)}{dt} = \beta k [1 - \rho_k^I(t)] \sum_{k'} P(k'|k) \rho_{k'}^I(t) - \mu \rho_k^I(t)$$

Sum over all possible k'

Number of nodes recovering

Transmission happens over k links

Prob of finding a node with degree k , susceptible

Probability that a node of degree k is connected to an infected node of degree k'

The DBMF SIS model

$$\frac{d\rho_k^I(t)}{dt} = \beta k [1 - \rho_k^I(t)] \sum_{k'} P(k'|k) \rho_{k'}^I(t) - \mu \rho_k^I(t)$$

If we assume the network to be **uncorrelated**: $P(k'|k) = \frac{k' P(k')}{\langle k \rangle}$

then
$$\frac{d\rho_k^I(t)}{dt} = \beta k [1 - \rho_k^I(t)] \Theta - \mu \rho_k^I(t)$$

where $\Theta = \sum_{k'} \frac{k' P(k')}{\langle k \rangle} \rho_{k'}^I(t)$ prob. of finding an infected node following a randomly chosen edge

Solution

Early stage approximation: $\rho_k^I(t) \ll 1$

then
$$\frac{d\Theta}{dt} = \left(\frac{\beta \langle k^2 \rangle}{\mu \langle k \rangle} - 1 \right) \Theta$$

which implies that Θ will grow only if:

$$\frac{\beta}{\mu} > \frac{\langle k \rangle}{\langle k^2 \rangle}$$

**Epidemic
threshold**

The DBMF threshold

$$\frac{\beta}{\mu} > \frac{\langle k \rangle}{\langle k^2 \rangle}$$

- ▶ In an infinite scale-free network, with $P(k) \sim k^{-\gamma}$, and $2 \leq \gamma \leq 3$, $\langle k^2 \rangle \rightarrow \infty$ which implies that **the epidemic threshold vanishes**.
- ▶ There is a finite prevalence for any value of the spreading parameters.

Homogeneous networks

In the case of a **homogeneous network** with a regular (Poisson) degree distribution:

$$\langle k^2 \rangle = \langle k \rangle^2 + \langle k \rangle$$

$$\langle k^2 \rangle / \langle k \rangle \simeq \langle k \rangle$$

The epidemic threshold then becomes:


$$\frac{\beta}{\mu} \gtrsim \frac{1}{\langle k \rangle}$$

which is finite and it does only depend on the average connectivity of the network.

Immunization

A decorative network diagram in the top right corner of the slide. It consists of several interconnected nodes, represented by circles of varying sizes and colors (some solid grey, some hollow white with a grey outline). The nodes are connected by thin grey lines, forming a complex, branching structure that resembles a molecular or biological network.

In the case of complex networks, we can consider three different immunization strategies:

- uniform immunization
 - proportional immunization
 - targeted immunization
- 
- A decorative network diagram in the bottom left corner of the slide. It features a cluster of nodes connected by lines, similar in style to the diagram in the top right. The nodes are represented by circles, some solid and some hollow, connected by thin lines, creating a complex, interconnected network structure.

Uniform immunization

In the case of uniform immunization, **individuals are randomly chosen to be vaccinated** with a density of immune nodes g .

This corresponds to an effective rescaling of the spreading rate:

$$\beta \rightarrow \beta(1 - g)$$

The threshold is affected in a uniform way:

$$\frac{\beta}{\mu}(1 - g) > \frac{\langle k \rangle}{\langle k^2 \rangle}$$

Uniform immunization

$$\frac{\beta}{\mu}(1 - g) > \frac{\langle k \rangle}{\langle k^2 \rangle}$$

In infinite scale-free network, with $P(k) \sim k^{-\gamma}$, and $2 \leq \gamma \leq 3$, $\langle k^2 \rangle \rightarrow \infty$

which implies that the **uniform immunization is not effective** unless we

immunize all the network: $g = 1$

Proportional immunization

We can find a better solution through a **proportional immunization**.

Let us define the fraction of immune individuals with connectivity k : g_k

If we impose the condition:

$$\tilde{\beta} \equiv \beta k (1 - g_k) = \text{const.}$$

The system equation becomes:

$$\frac{d\rho_k^I(t)}{dt} = \tilde{\beta} [1 - \rho_k^I(t)]^\Theta - \mu \rho_k^I(t)$$

Proportional immunization

In the case of early stage approximation and low density of infectious individuals, we recover an epidemic threshold:

$$\beta k(1 - g_k) - \mu > 0$$

which defines a threshold on density of immunized individuals:

$$g_k > 1 - \frac{\mu}{\beta k}$$

for every class of degree k , to stop the epidemic.

Targeted immunization

Optimum approach: immunize a fraction of all nodes with the largest degree.

This way we introduce a cut-off in the degree distribution.

We need to immunize a fraction of nodes g such that:

$$\frac{\beta}{\mu} < \frac{\langle k \rangle_g}{\langle k^2 \rangle_g}$$

In the case of the BA network, it is possible to show that: $g_c \simeq e^{-\frac{2\mu}{m\beta}}$

The fraction of nodes to immunize is exponentially small with β

How do we find the hubs?

- ▶ Targeted immunisation is very hard to achieve in practice, the full network structure is not known
- ▶ We need a strategy to find hubs based on a **local knowledge** of the network
- ▶ In scale-free networks, this can be done efficiently with the **acquaintance immunisation** (Cohen et al. Phys. Rev. Lett. 2003)
- ▶ Instead of immunizing nodes at random, we pick random nodes and for each we immunise one of their neighbours at random.

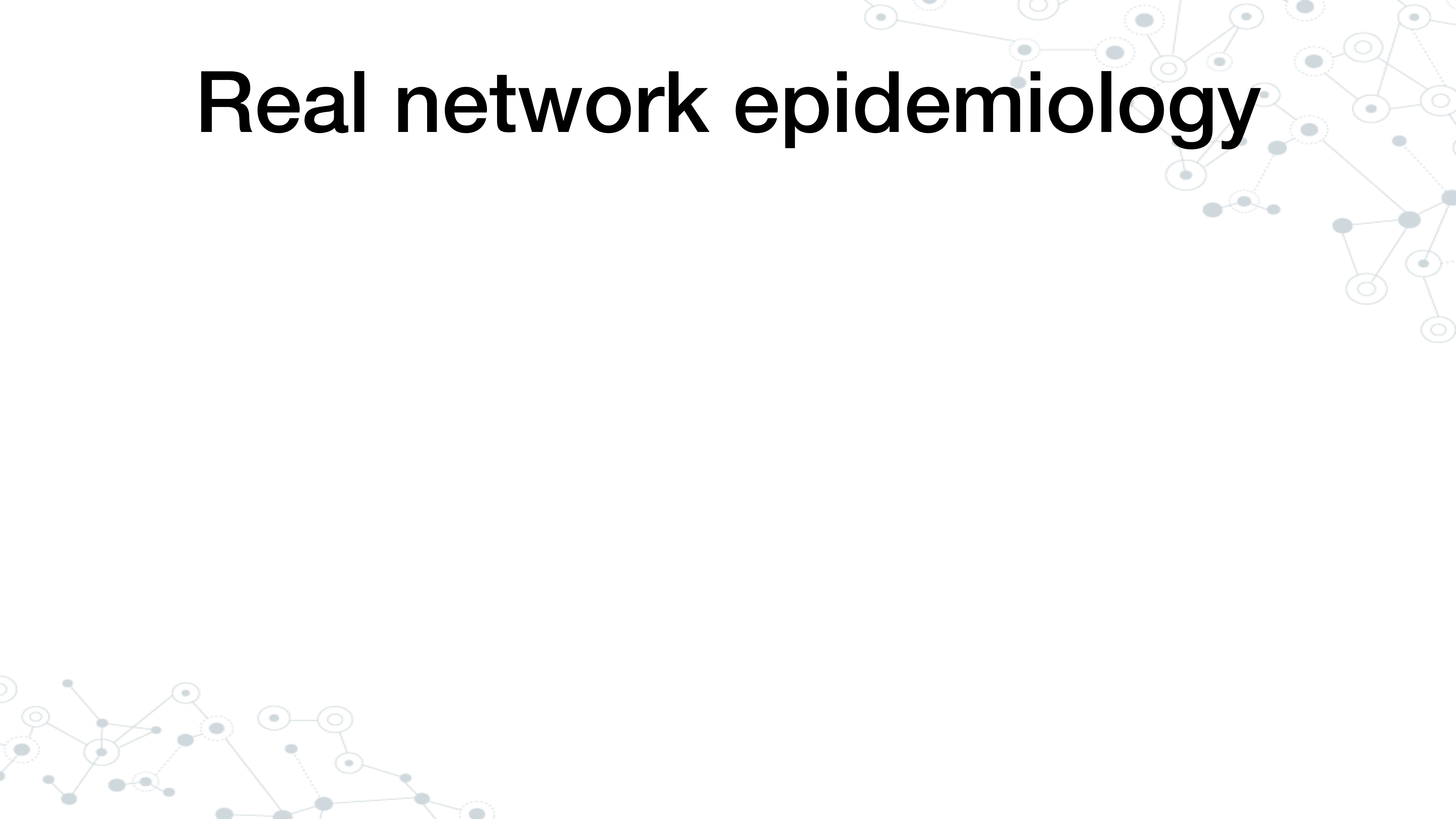
How do we find the hubs?

- ▶ Instead of immunizing nodes at random, we pick random nodes and for each we immunise one of their neighbours at random.

$$k_{nn}^{\text{unc}} = \frac{\langle k^2 \rangle}{\langle k \rangle}$$

- ▶ My neighbours are more probably hubs than myself! This is also known as the **friendship paradox**

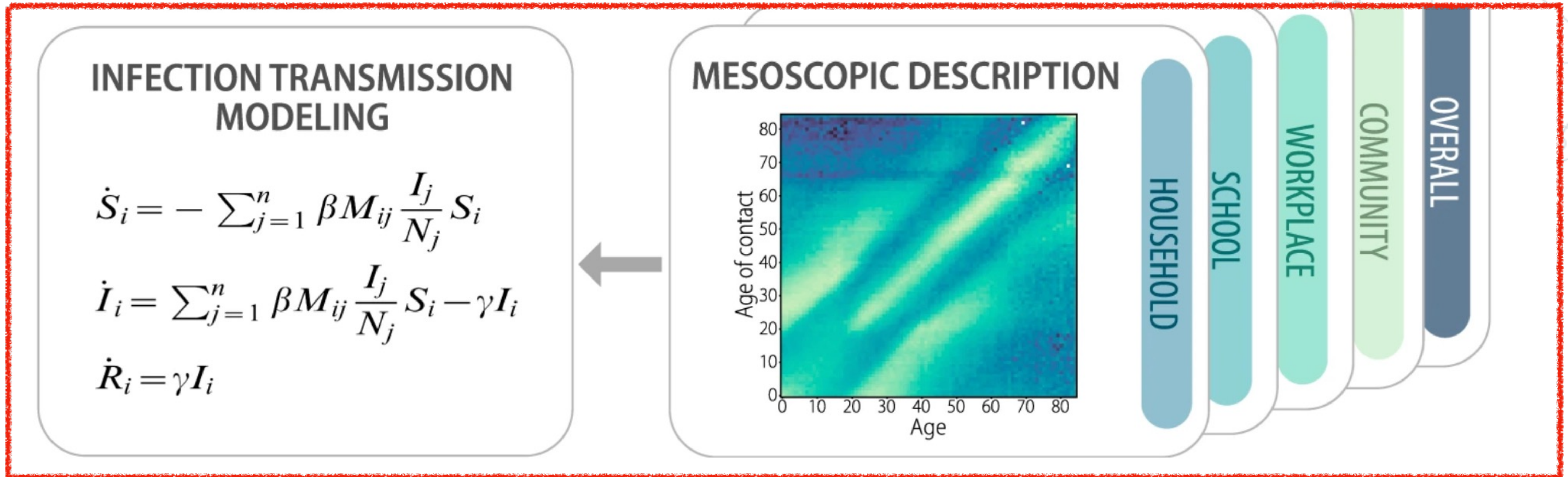
Real network epidemiology



Real network epidemiology

- More sophisticated compartmental models (incubation period, hospitalization)
- Age-structured population
- Estimation of real contact matrices
- Mobility
- Lots of numerical simulations (no nice analytical solutions!)

Age-structured models



- Compartments are structured into n age classes
- M_{ij} represents the average contact rate between individuals of age i and j

Contact matrices

- Contact matrices can be estimated in different ways
- Through **empirical surveys**, which are more accurate but require significant resources (Mossong et al. 2008).
- By the creation of **synthetic populations** (Fumanelli et al. 2012).

PLOS COMPUTATIONAL BIOLOGY

OPEN ACCESS PEER-REVIEWED

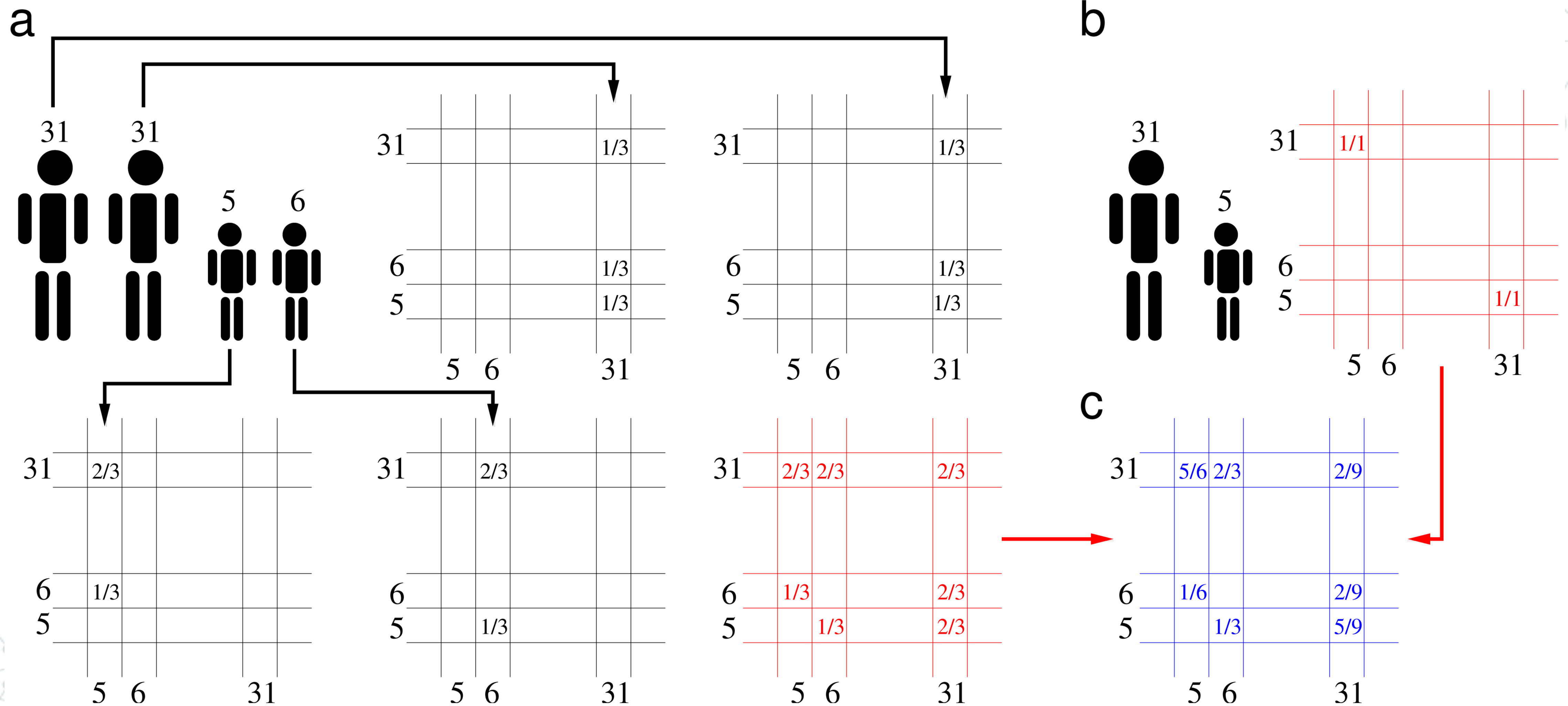
RESEARCH ARTICLE

Inferring the Structure of Social Contacts from Demographic Data in the Analysis of Infectious Diseases Spread

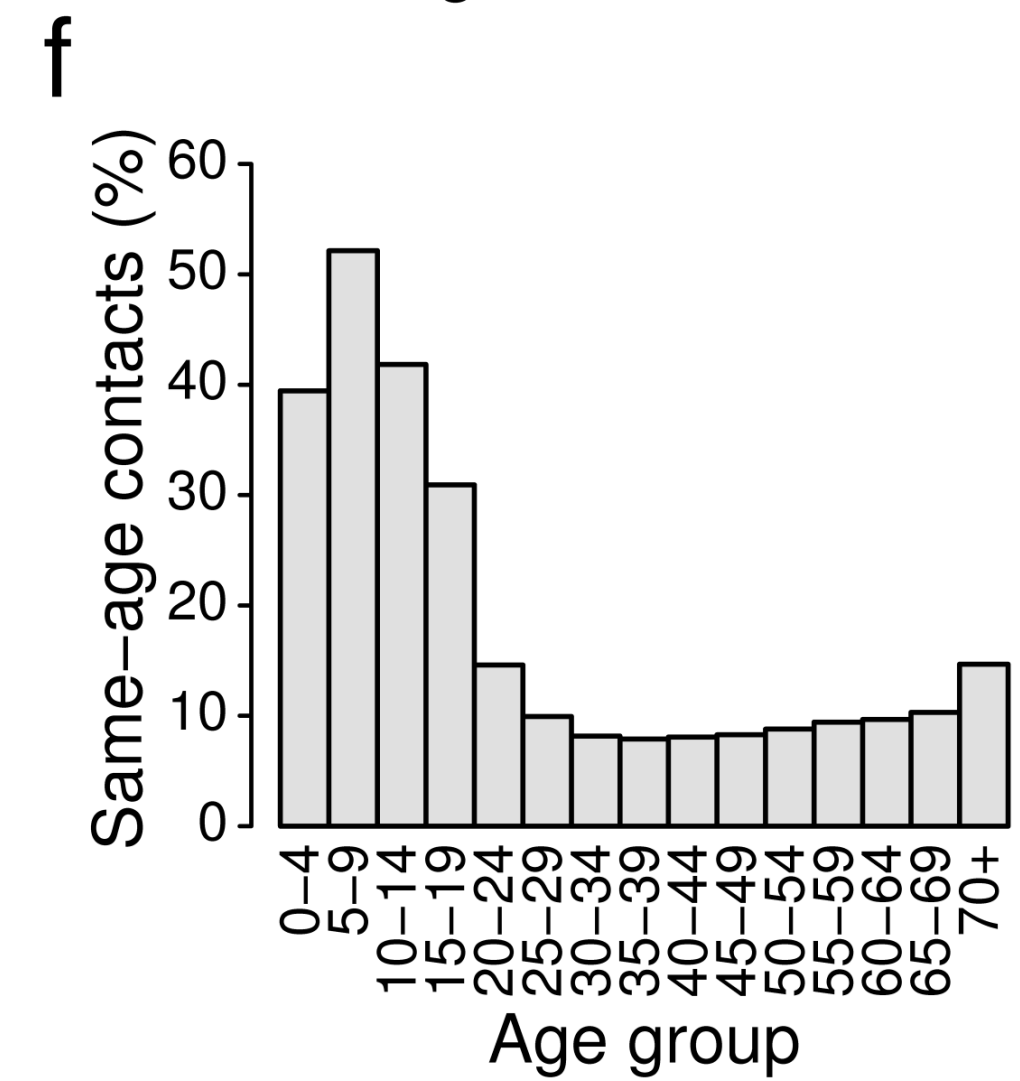
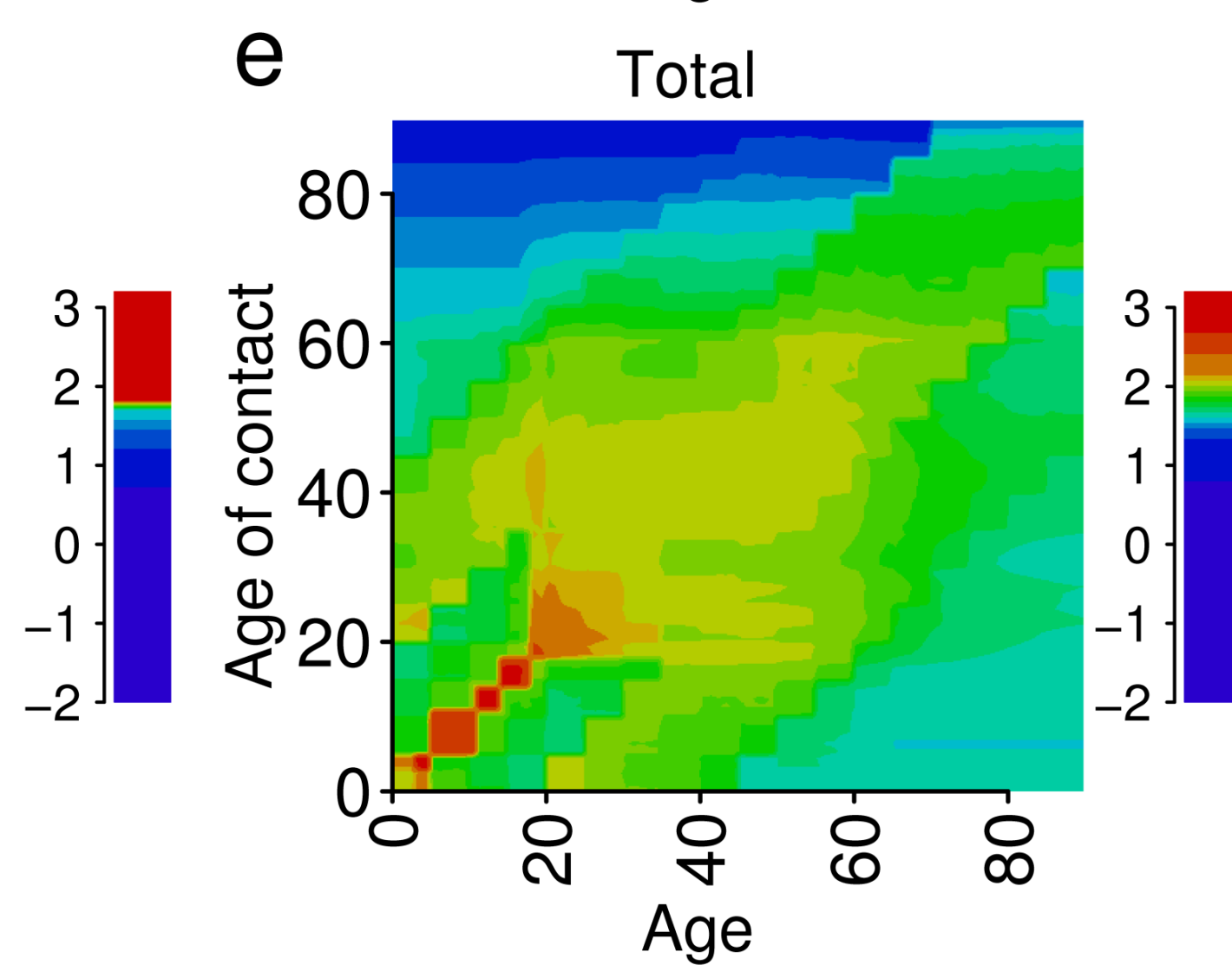
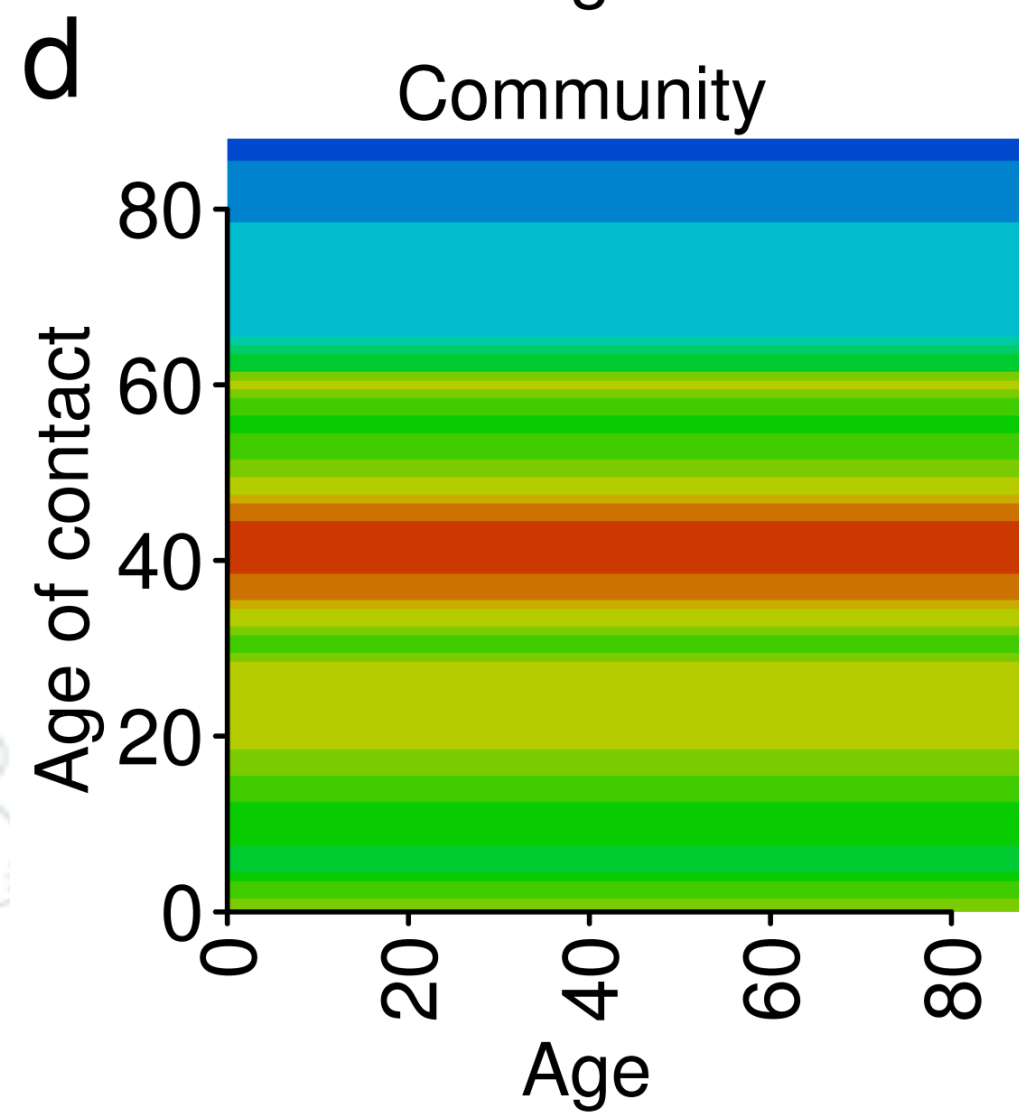
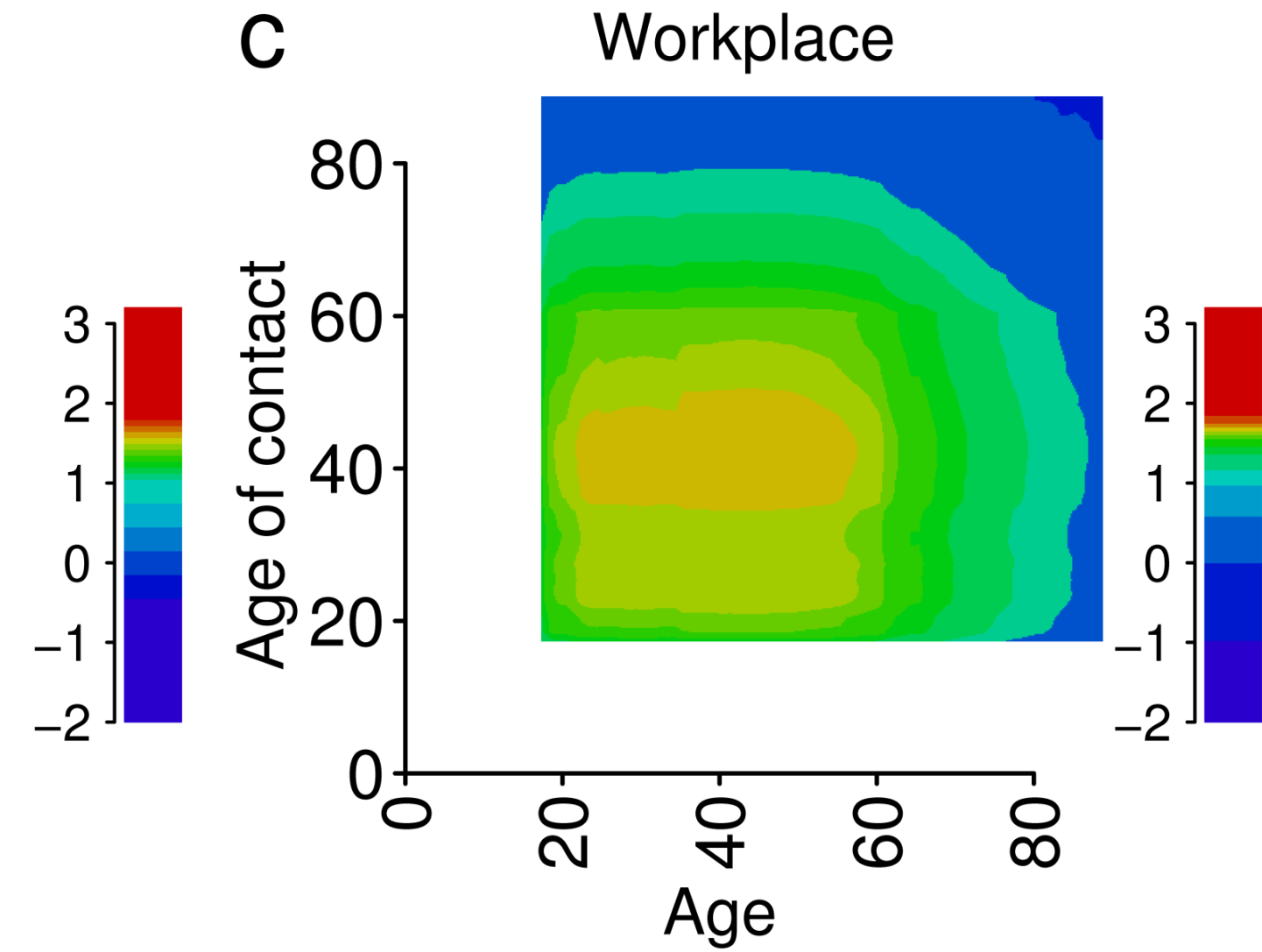
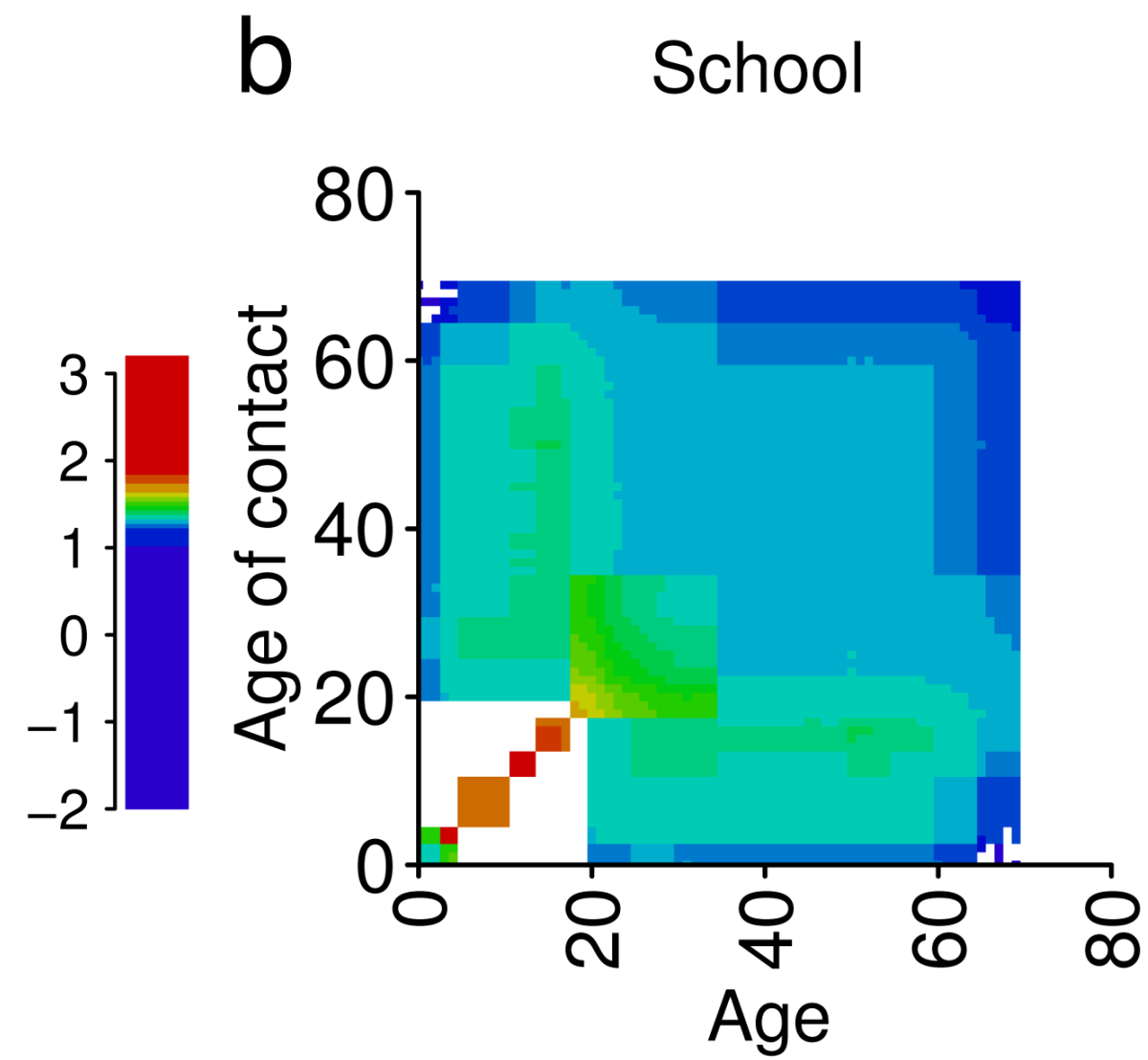
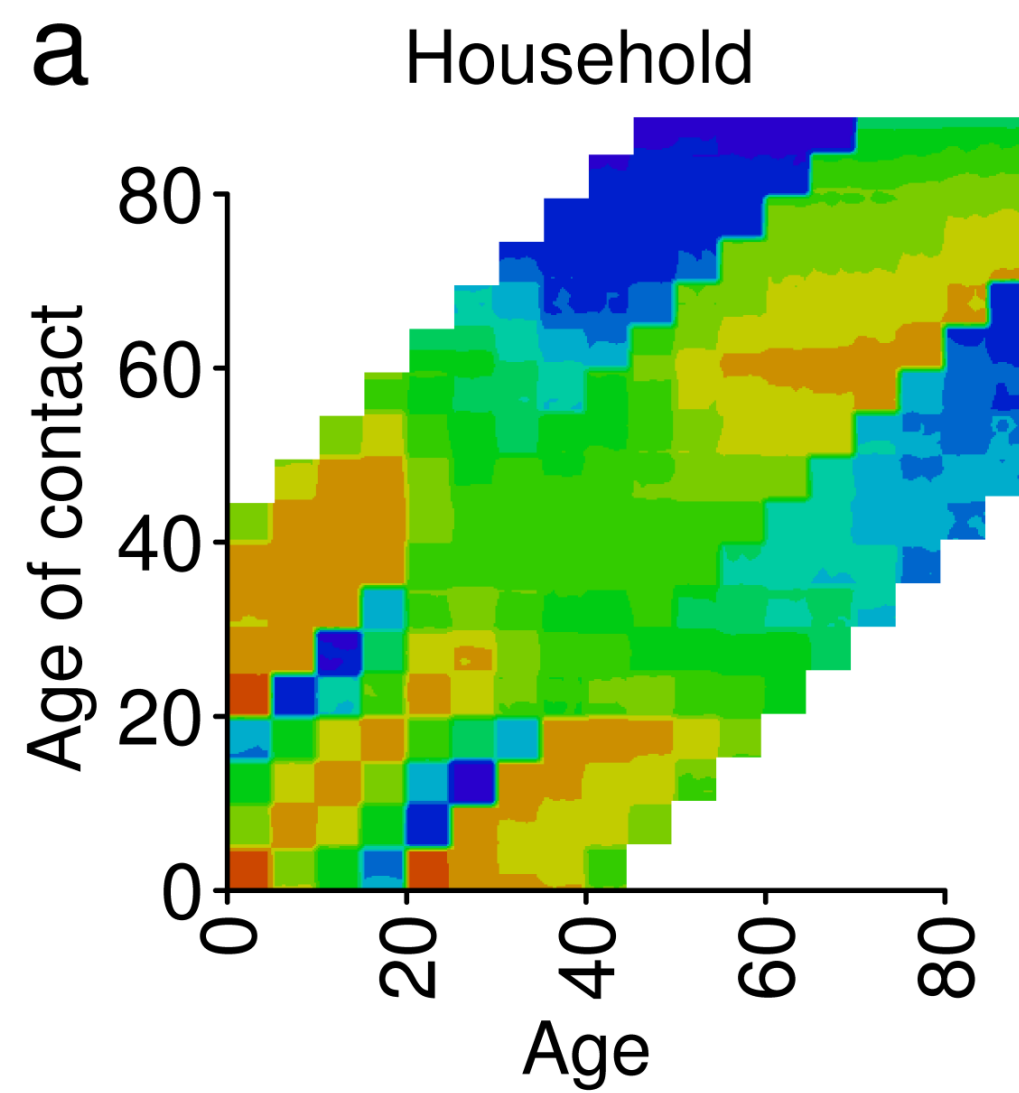
Laura Fumanelli , Marco Ajelli, Piero Manfredi, Alessandro Vespignani, Stefano Merler

Published: September 13, 2012 • <https://doi.org/10.1371/journal.pcbi.1002673>

Synthetic populations



Synthetic populations

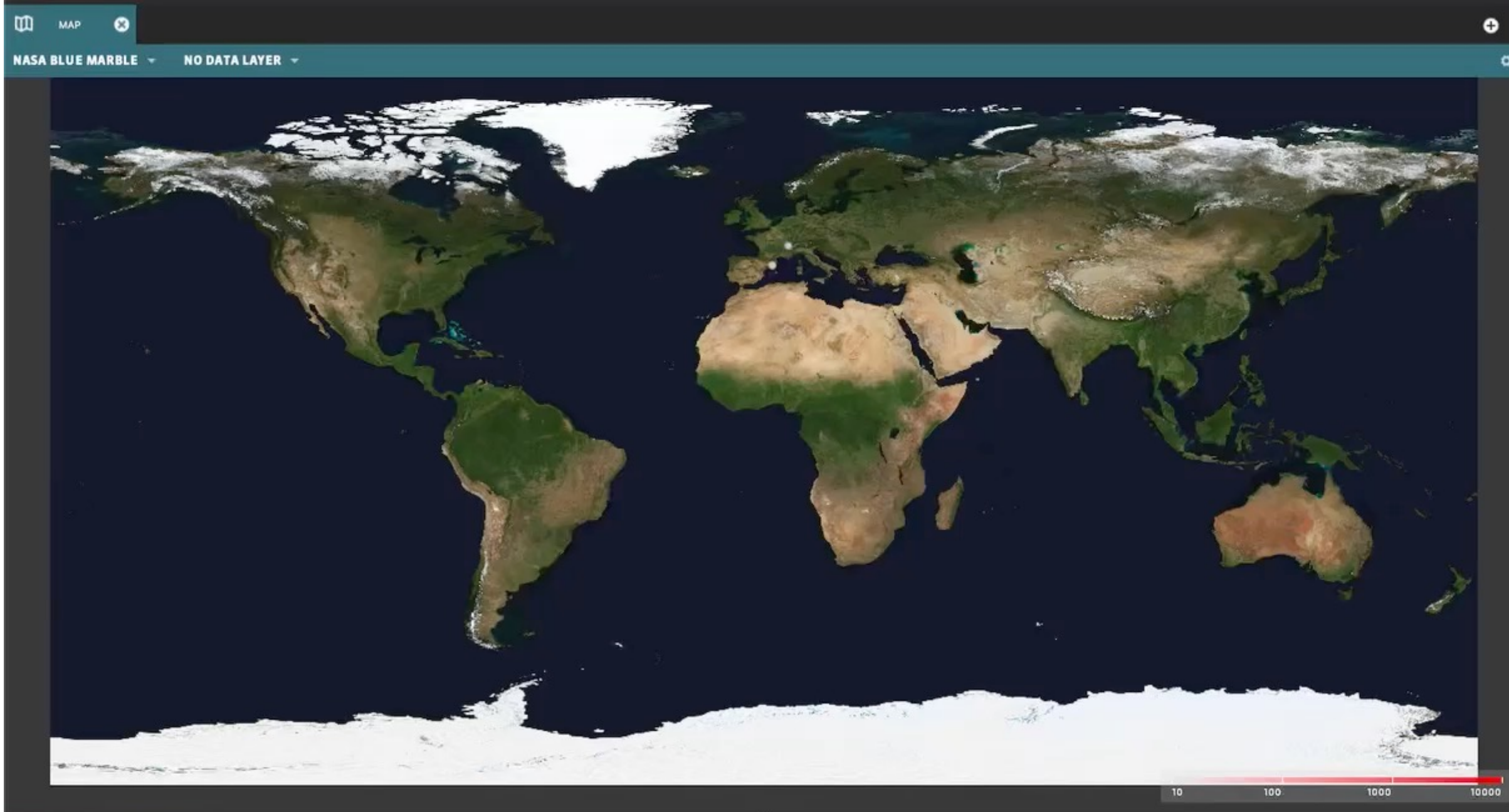




Using big data and computational modeling to fight infectious diseases

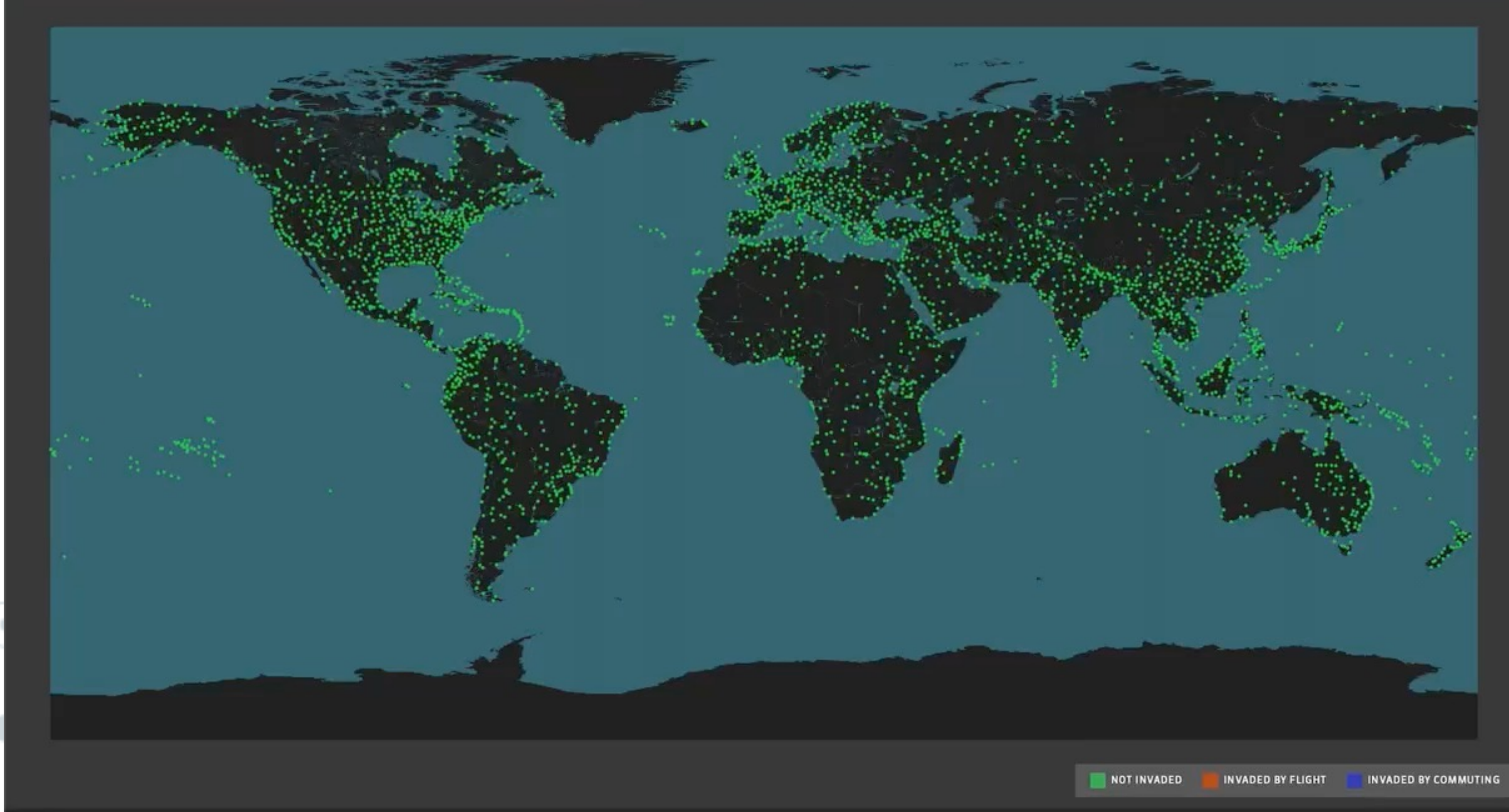
COVID-19 Research

Global Epidemic and Mobility project
<https://www.youtube.com/watch?v=YstB9VWDUqE>



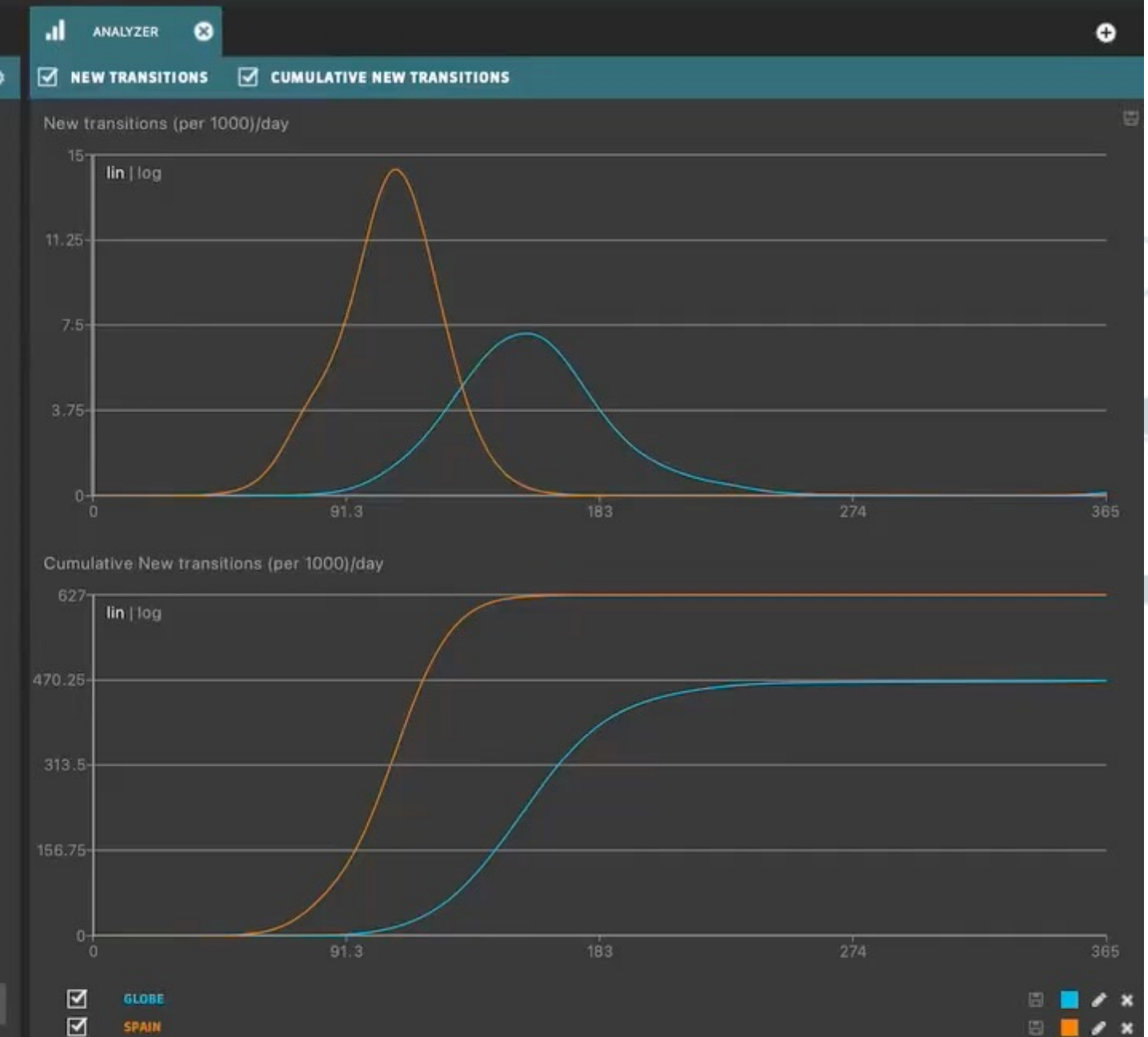
INVASIONTREE

TIME INTERVAL 1 365 BASIC DARK CITY



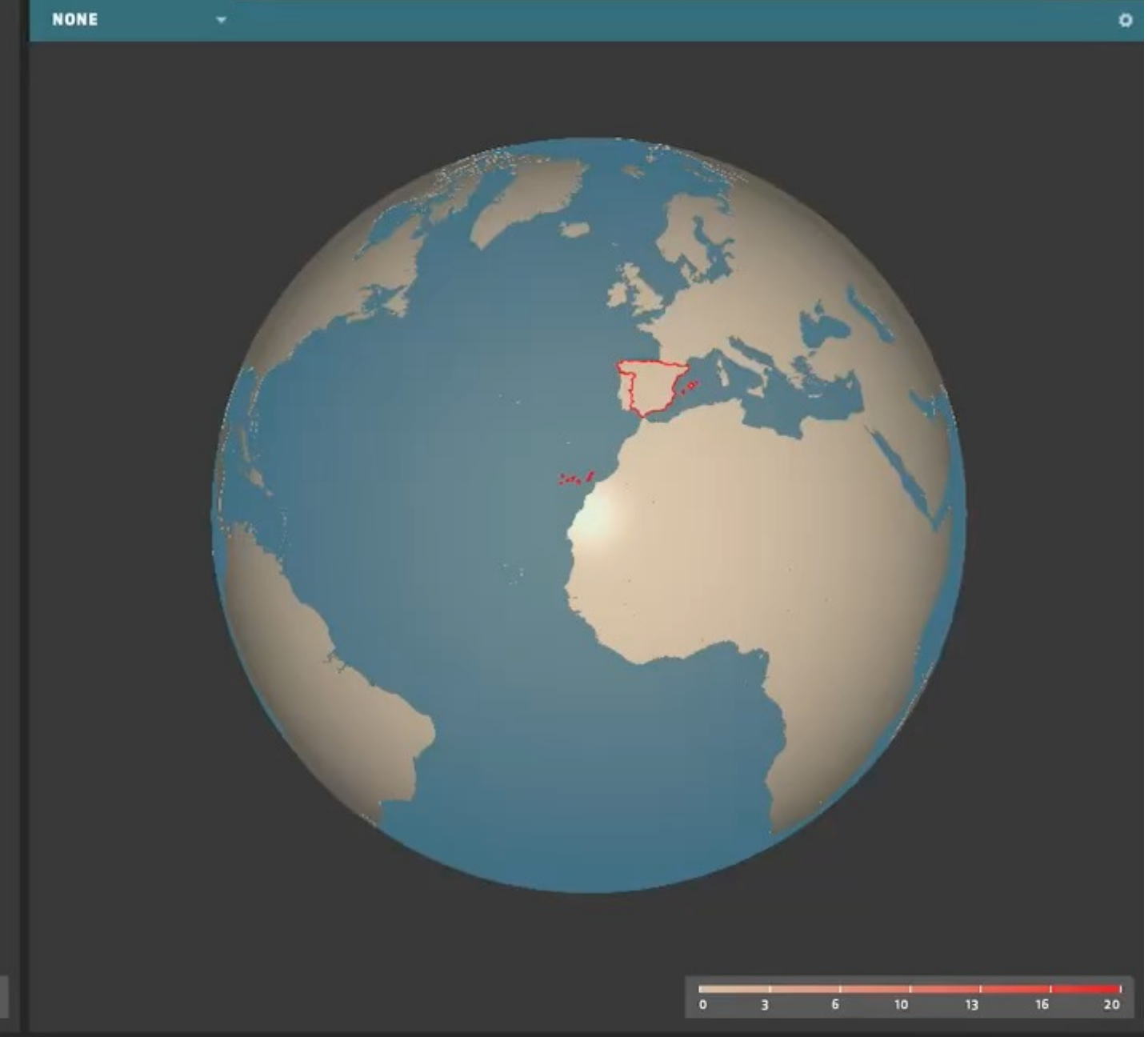
H1N1 Flu - Barcelona

3 SELECTED COMPARTMENTS NEW INDIVIDUALS DAILY



X3D GLOBE

NONE

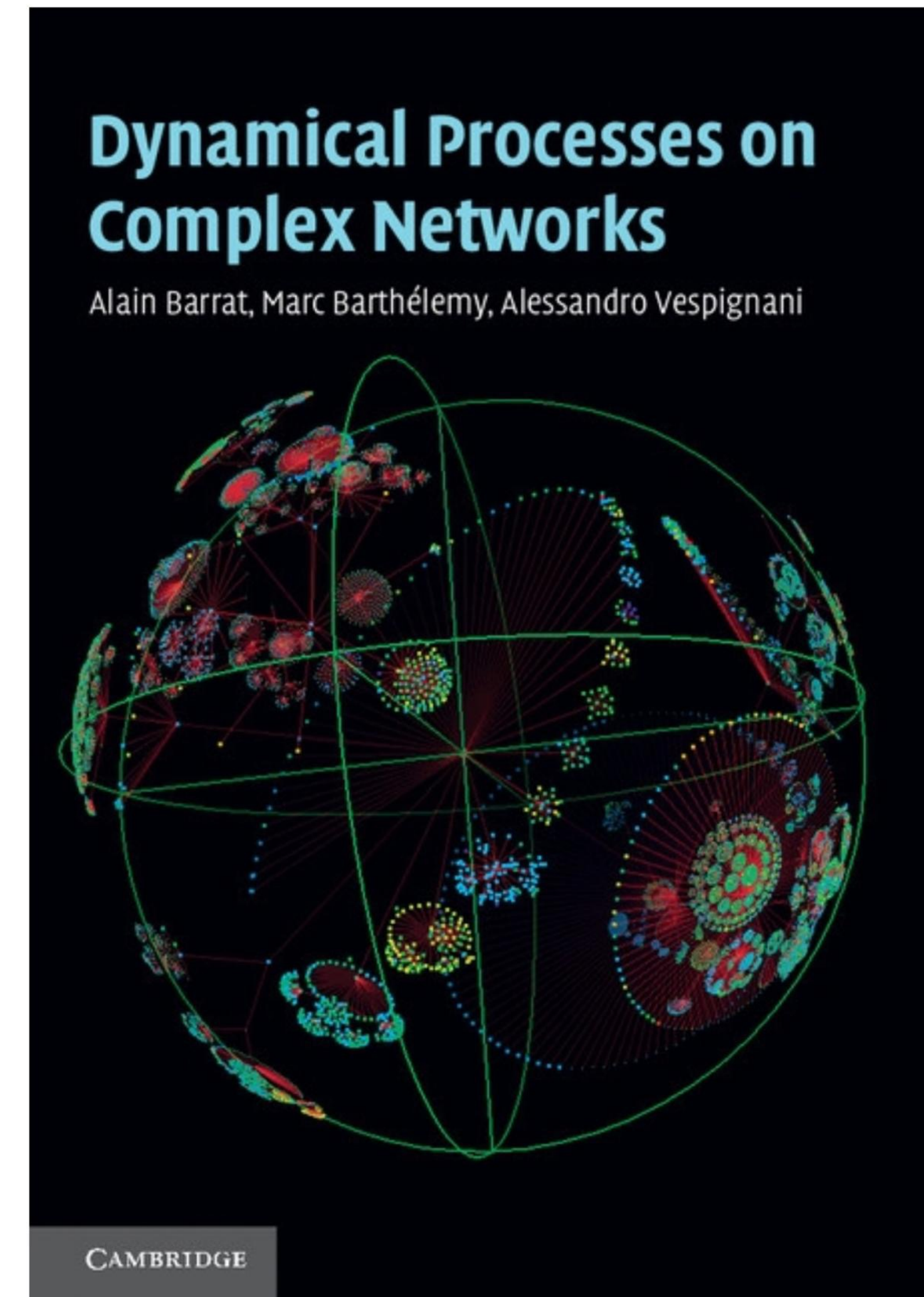


DAY 0 15/11/2024



Sources

- ▶ Pastor-Satorras et al. Epidemic processes in complex networks. Rev. Mod. Phys. 87, 925 (2015)
- ▶ Pastor-Satorras, and Vespignani. Epidemic spreading in scale-free networks. Phys. Rev. Lett. 86, 14 (2000)
- ▶ Barrat, Barthelemy, Vespignani. Dynamical processes on complex networks. Cambridge University Press



Thank you!

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