

Social and Technological Network Analysis

Lecture 11: Epidemics Spreading

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In This Lecture



- In this lecture we introduce the process of spreading epidemics in networks.
 - This has been studied widely in various disciplines from different perspectives from theoretical models to simulations of real events.
 - But it also has important parallels and applications in information/idea diffusion in social and technological networks.

Epidemics vs Cascade Spreading



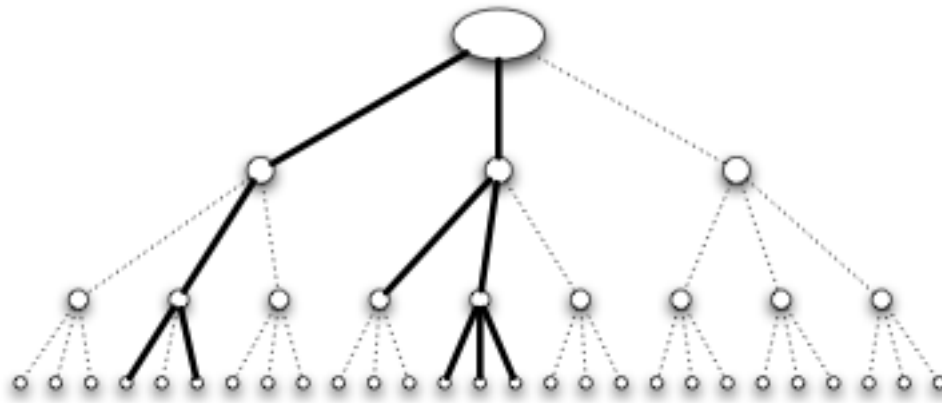
- In cascade spreading nodes make decisions based on pay-off benefits of adopting one strategy or the other.
- In epidemic spreading
 - Lack of decision making.
 - Process of contagion is complex and unobservable
 - In some cases it involves (or can be modeled as) randomness.



Branching Process

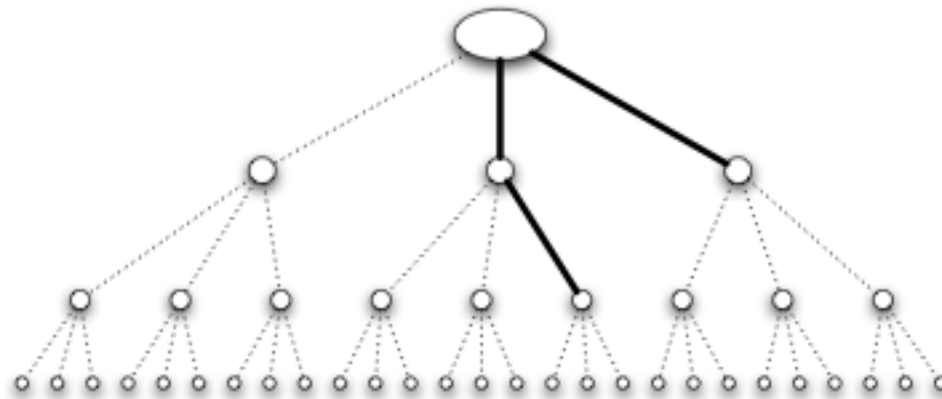
- Simple model.
- **First wave:** A person carrying a disease enters the population and transmits to all he meets with probability p . He meets k people: a portion of which will be infected.
- **Second wave:** each of the k people goes and meet k different people. So we have a second wave of $k*k=k^2$ people.
- **Subsequent waves:** same process.

Example with $k=3$



High contagion probability:
The disease spreads

Low contagion probability:
The disease dies out



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Basic Reproductive Number



- Basic Reproductive Number $R_0 = p * k$
 - It determines if the disease will spread or die out.
- In the branching process model, if $R_0 < 1$ the disease will die out after a finite number of waves. If $R_0 > 1$, with probability > 0 , the disease will persist by infecting at least one person in each wave.

Measures to Limit the Spreading



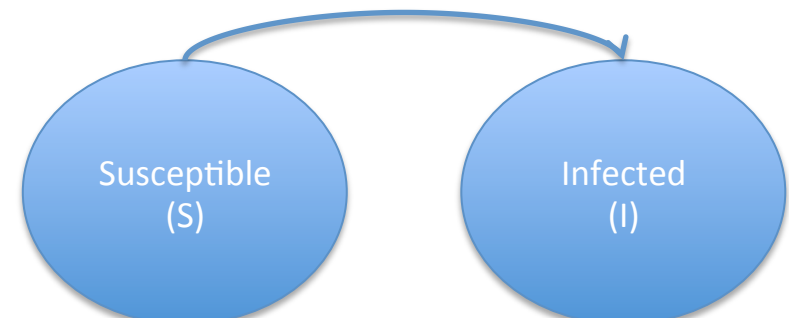
- When R_0 is close 1, slightly changing p or k can result in epidemics dying out or happening.
 - Quarantining people/nodes reduces k .
 - Encouraging better sanitary practices reduces germs spreading [reducing p].
- Limitations of this model:
 - No realistic contact networks: no triangles!
 - Nodes can infect only once.
 - No nodes recover.

Formal Epidemics Models

The SI Model



- S: susceptible individuals.
- I: infected individuals, when infected they can infect others continuously.
- n: total population.
- $\langle k \rangle$ average contacts per individual
- $\beta = \lambda \langle k \rangle$ is the infection rate per individual ($0 \leq \lambda \leq 1$)
- Susceptible contacts per unit of time $\beta S/n$.
- Overall rate of infection $I\beta S/n$.





SI Model

$$\frac{dI}{dt} = \beta \frac{SI}{n}$$

$$\frac{dS}{dt} = -\beta \frac{SI}{n}$$

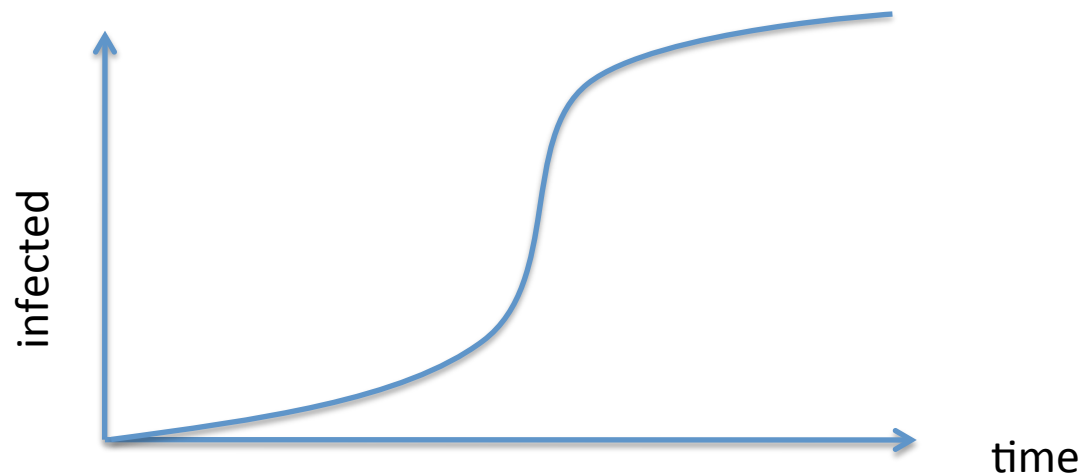
$$s = \frac{S}{n} \quad i = \frac{I}{n}$$

$$s = 1 - i$$

$$\frac{di}{dt} = \beta i(1 - i)$$

$$i(t) = \frac{i_0 e^{\beta t}}{1 - i_0 + i_0 e^{\beta t}}$$

Logistic Growth Equation





SIR Model

- Infected nodes recover at a rate γ .
- A node stays infected for τ time.
- Branching process is SIR with $\tau=1$.

$$\frac{ds}{dt} = -\beta si$$

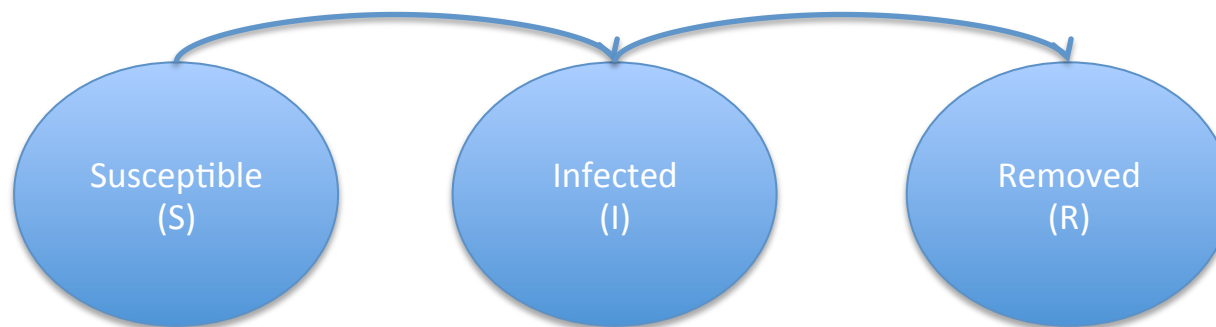
$$\frac{di}{dt} = \beta si - \gamma i$$

$$\frac{dr}{dt} = \gamma i$$

$$s + i + r = 1$$



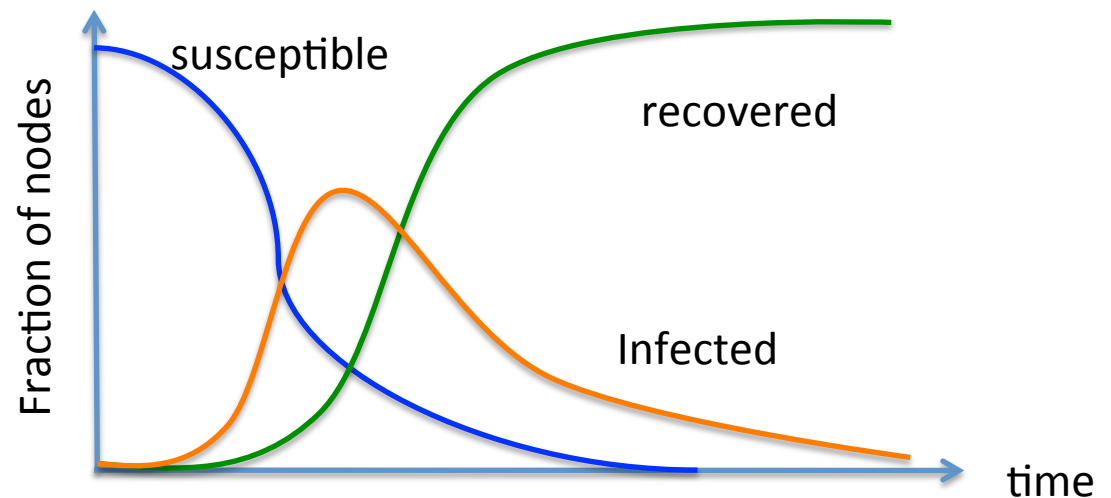
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Example



- Numerical examples of solution:
- $\beta=1$, $\gamma=0.4$, $s(\text{at start})=0.99$, $i(\text{at start})=0.01$, $r(\text{at start})=0$

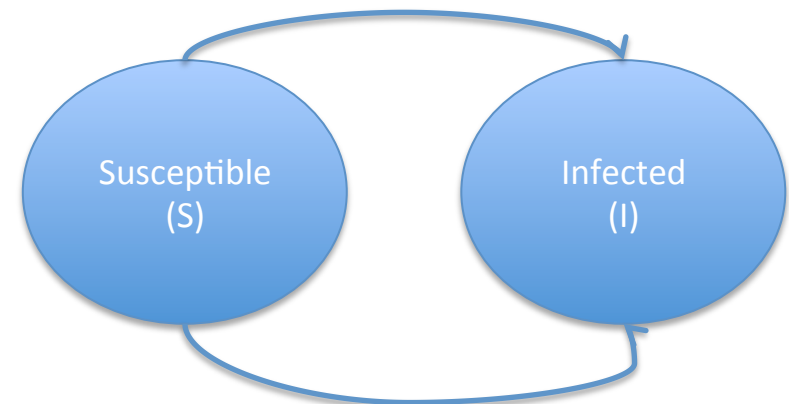




SIS Model

$$\frac{ds}{dt} = \gamma i - \beta si$$
$$\frac{di}{dt} = \beta si - \gamma i$$
$$s + i = 1$$

- If $\beta > \gamma$ growth curve like in SI but never reaching all population infected. The fraction of infected $\rightarrow 0$ as β approaches γ .
- If $\beta < \gamma$ the infection will die out exponentially.
- SIS has the same R_0 as SIR.



Epidemic Threshold



- When would the epidemic develop and when would it die out?
- It depends on the relationship of β and γ :
 - Basic Reproductive Number $R_0 = \beta/\gamma$
 - If the infection rate [per unit of time] is higher than the removal rate the infection will survive otherwise it will die out.
 - In SI, $\gamma=0$ so the epidemics always happen.

Limitations of SIR



- Contagion probability is uniform and “on-off”
- Extensions
 - Probability q of recovering in each step.
 - Infected state divided into intermediate states (early, middle and final infection times) with varying probability during each.
 - **We have assumed homogenous mixing** : assumes all nodes encounter each others with same probability: we could assume different probability per encounter.



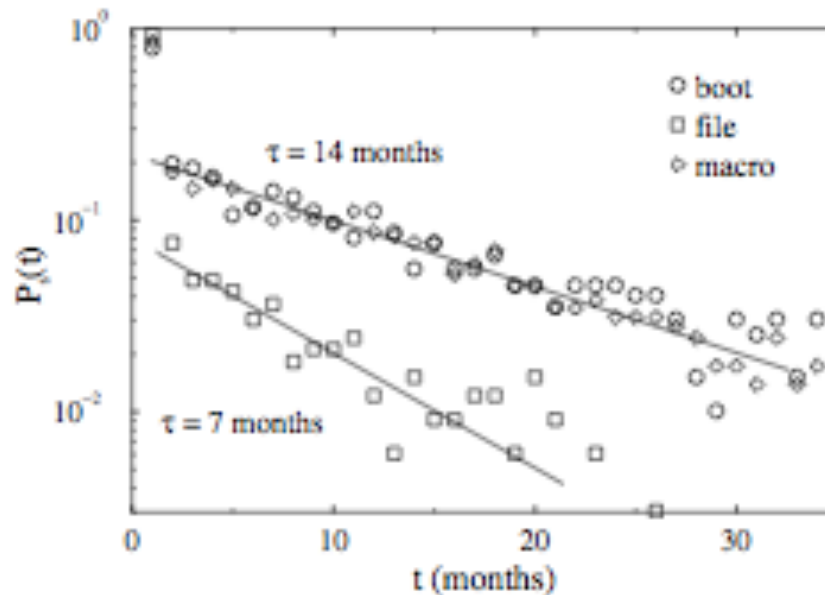
Relaxing Assumptions

- Homogeneous Mixing: a node connects to the same average number of other nodes as any other.
- Most real networks are not Erdos-Renyi random networks (for which the homogeneous mixing assumption holds).
- Most networks have heterogeneous degree distributions.
 - Scale free networks!

Would the Model Apply to SF?

- Pastor-Satorras and Vespignani [2001] have considered the life of computer viruses over time on the Internet:

Surviving probability of virus



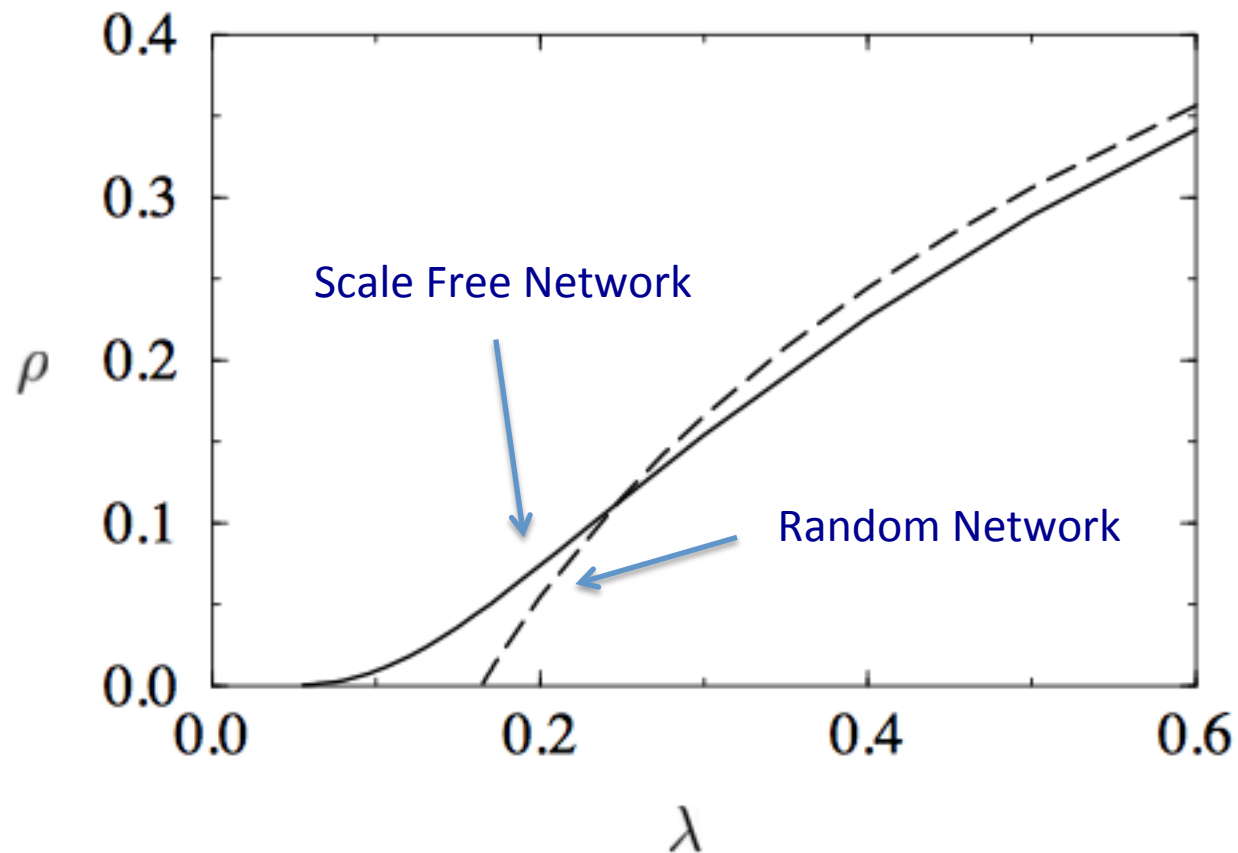
Virus survived on average 6-9/14 months depending on type

How to Justify This Survival Time?



- The virus survival time is considerably high with respect to the results of epidemic models of spreading/recovering:
 - Something wrong with the epidemic threshold!
- Experiment: SIS over a generated Scale Free network (exponent -3).

No Epidemic Threshold for SF!



Network Immunisation



- Random network can be immunized with some sort of uniform immunization process [oblivious of the characteristics of nodes].
- **Random immunisation does not work in SF networks** no matter how many nodes are immunized [unless it is all of them].
- Targeted immunization in SF must be used instead.
 - Keeping into account degree!

Immunization on SF Networks

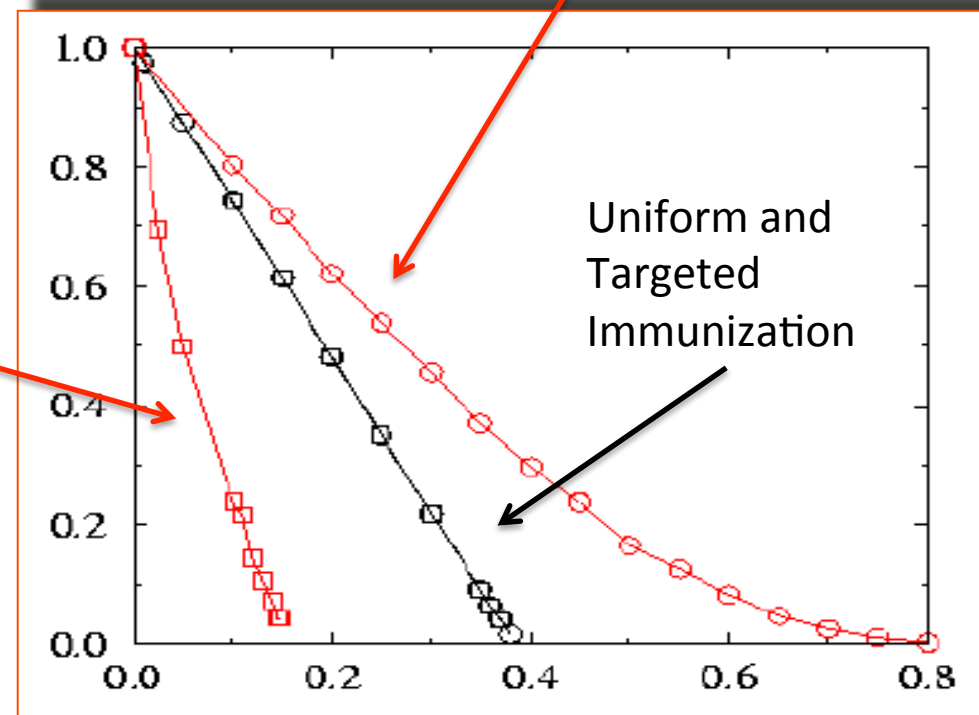


- Red=SF
- Black= Random

Targeted Immunization

Uniform Immunization

Density of
Infected



Fraction of immunized nodes



Local Immunization

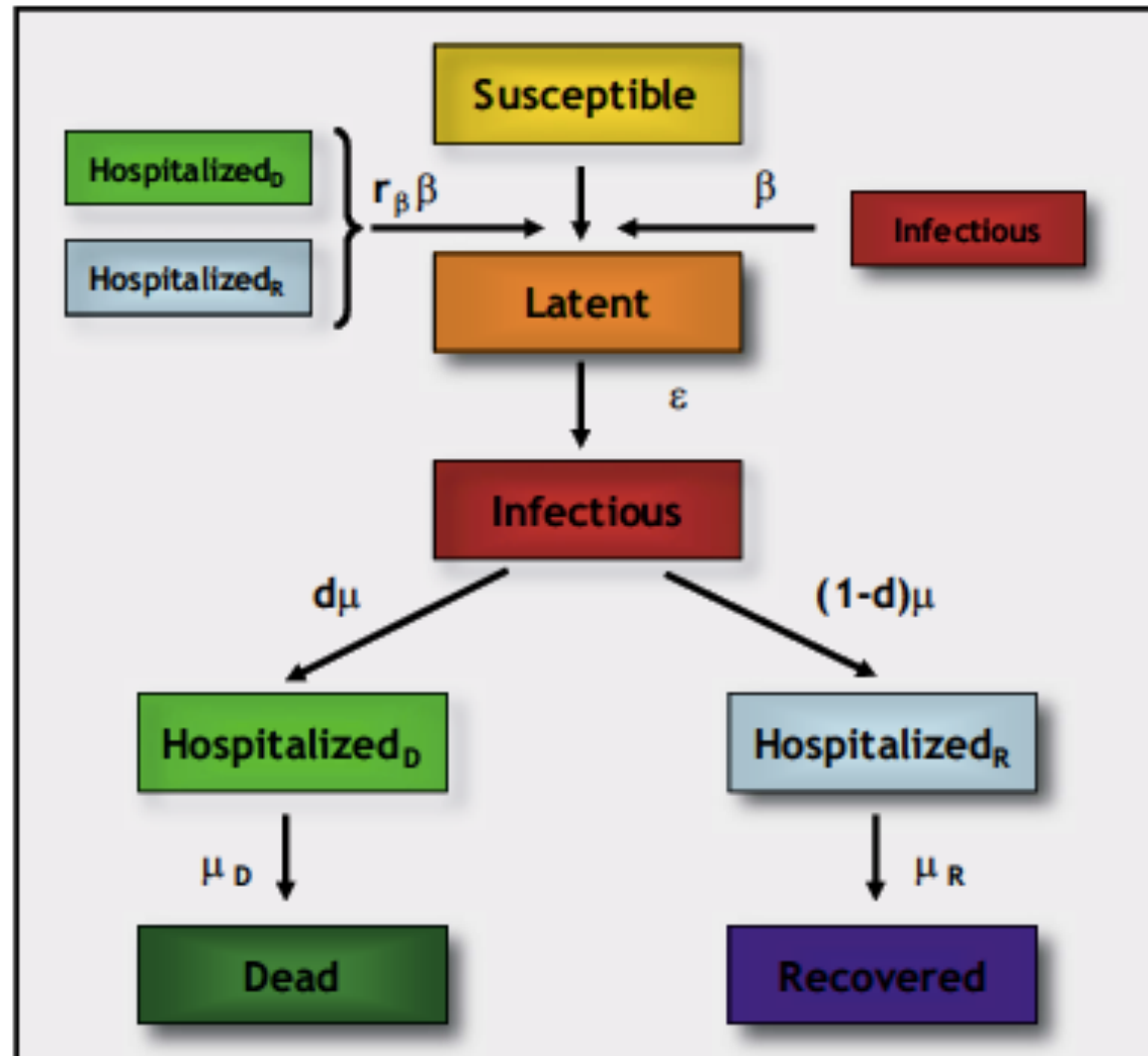
- Global knowledge on the network structure is rarely available (more on this later)
- Local immunization strategy:
 - Select g nodes at random
 - Ask to each of them to pass over the vaccine to one of their neighbors
 - As a result, a node with degree k is immunized with a probability $kP(k)$ (**hubs are immunized with higher probability!**)

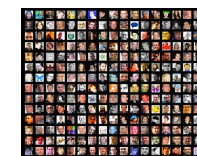
Modelling SARS Spreading



- SARS: severe acute respiratory syndrome
- SIR like model with more parameters and homogenous mixing
- Travel data and census data
- WHO data about SARS spreading to evaluate the model
- Outbreak starts in Hong Kong

Epidemics Flow





The Parameters of the Model

- Parameters used:

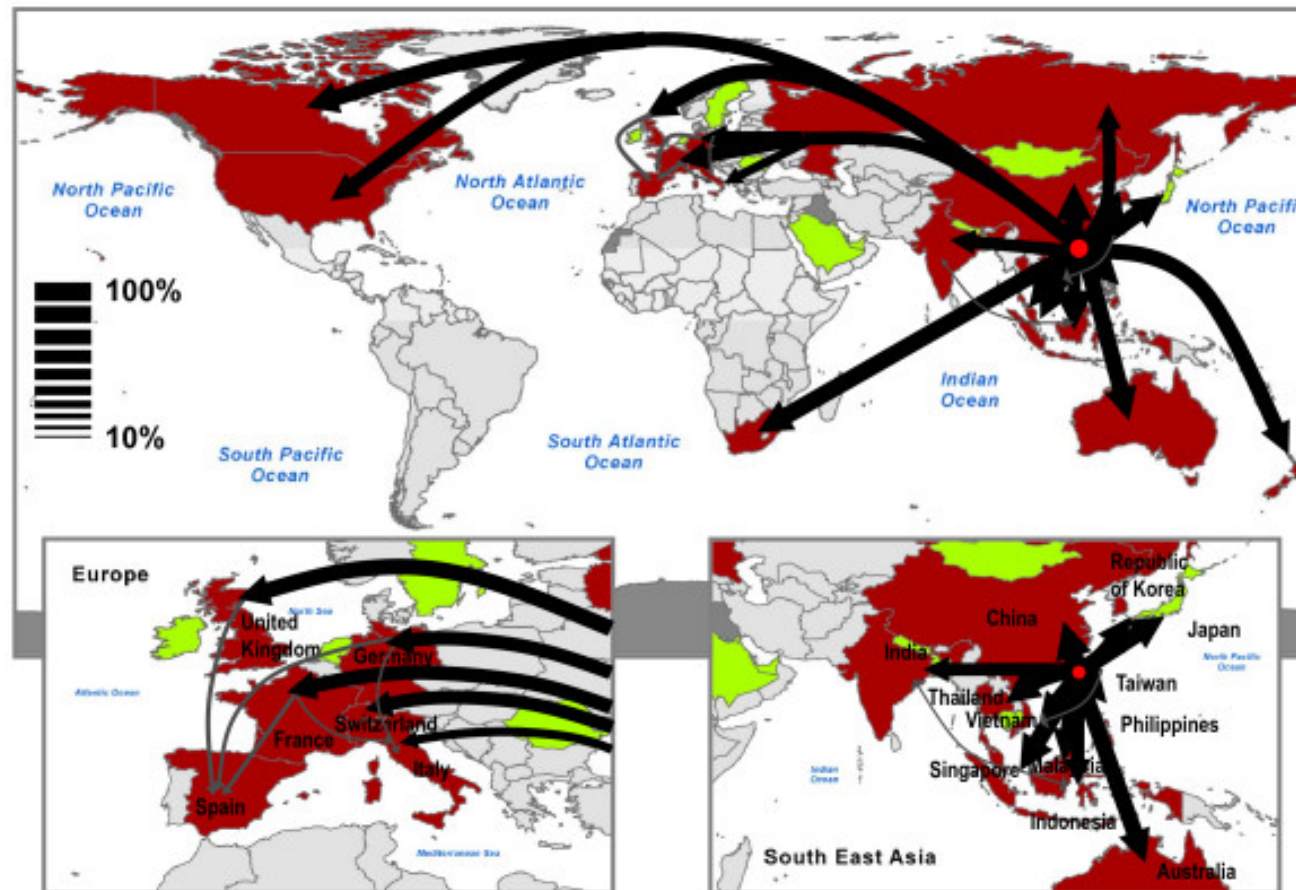
Parameter	Description	Baseline value	
T_0	Initial offset from 21 February (days)	3*	
β	Rate of transmission	0.57*	
$L(t = 0)$	Number of initial latent individuals	10*	
$s_f(t)$	Scaling factor for the rate of transmission	21 February + T_0 -20 March	1.00
		21 March – 9 April	0.37
		10 April – 11 July	0.06
r_β	Relative infectiousness of patients at the hospital	0.2	
ε^{-1}	Average latency period (days)	4.6	
$\mu^{-1}(t)$	Average period from onset of symptoms to admission (days)	21 February + T_0 -25 March	4.84
		25 March – 1 April	3.83
		2 April – 11 July	3.67
μ_R^{-1}	Average period from admission to recovery (days)	23.5	
μ_D^{-1}	Average period from admission to death (days)	35.9	
d	Case fatality rate	0.2	

Adding Travel and Geography

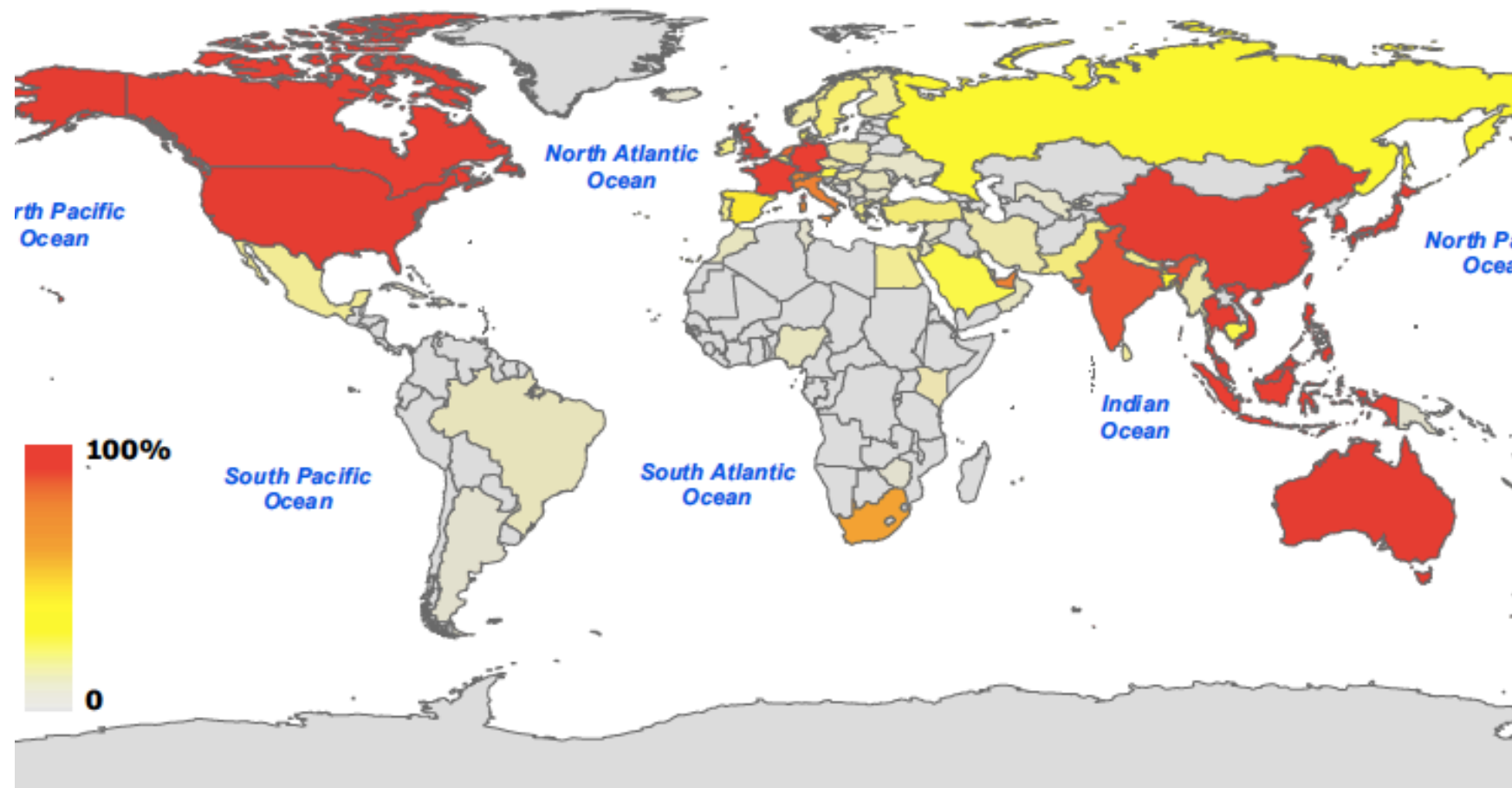


- A model per city (“meta-population model”)
 - Each compartmental model describes the epidemics in a given city.
 - Models per city are coupled using information from air travel
 - Possible simplification of the reality but effective abstraction.
- Stochastic model
 - Probabilities of an individual of moving from one city to the next follows proportions of traffic observed in the air travel data.

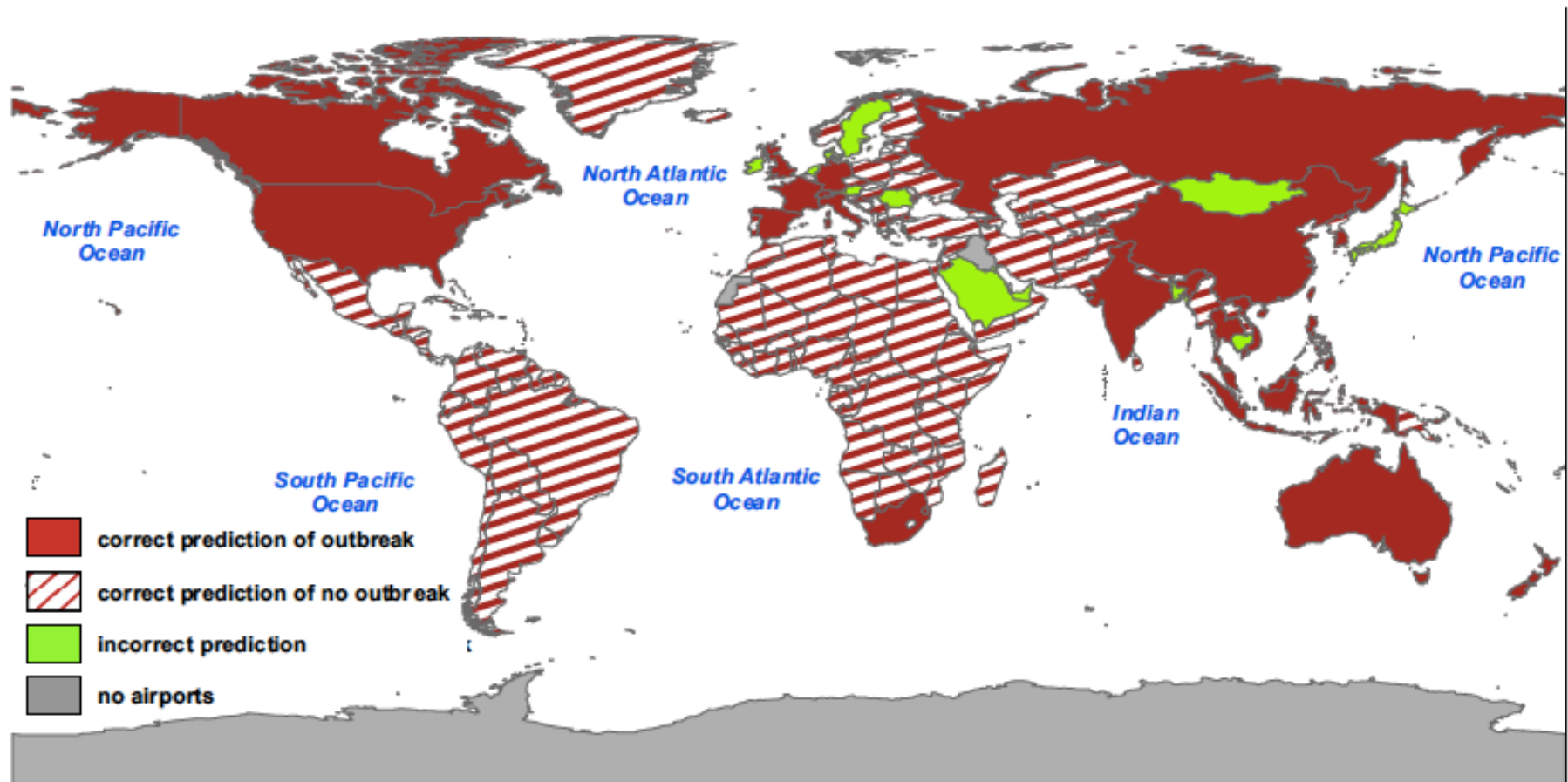
Epidemic Pathways



Predicted Outbreak Likelihood



Comparison with Data



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Epidemic Spreading Models and Real Data

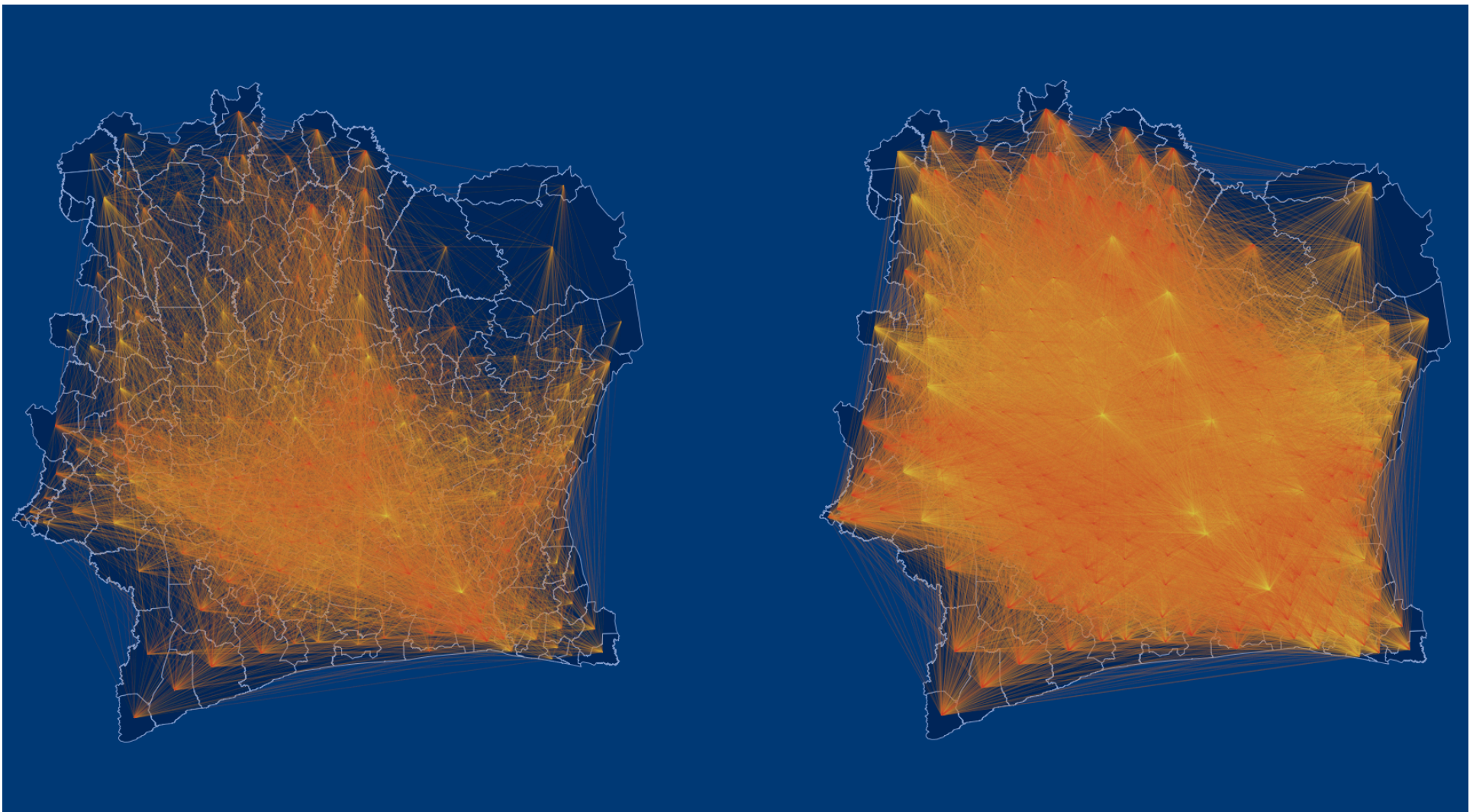


- A key problem is how to extract information for modelling the spreading the disease from real data.
- One possibility is to use information coming from the cellular network:
 - Transitions between base-stations for modelling the mobility;
 - Phone calls graph for modelling the underlying social networks between callers and callees.

Epidemic Spreading Models and Real Data



- In order to study possible strategies of containment of epidemics, an important aspect is how to model information campaigns for example for mass vaccination.
- Vaccination can happen through the population by exploiting the “strong” ties between them (family ties or friendships, etc.)
- Presence of two concurrent processes:
 - Epidemic spreading
 - Information diffusion (spreading of “immunising information”)



Mobility

Calls

Data for Development Challenge



- Data mining competition using a data set containing movement and call data of 5 million individuals in Ivory Coast in order to help to address society development questions in novel way
- Information extracted from the Call Data Records (CDRs) of the Orange network in Ivory Coast



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Call Data Records



- A Call Data Record (CDR) entry contains information about a specific phone call (usually called “metadata”), including:
 - The phone number of the subscriber originating the call;
 - The phone number of the called party;
 - The identification of the equipment writing the record (base station).
 - From this information it is possible to extract the geographic location of the caller.

Mobility Matrix



- Movement data extracted from the registration patterns to the cellular infrastructure (i.e., CDRs) are used to evaluate the influence of human mobility on the spreading of the disease in a given geographic area.
- From the data it is possible to extract the probability of transitions between different areas, in this case “sub-prefectures” (counties) of Ivory Coast.
- Using this information, we build a mobility matrix representing movement in the country as a Markov process.

Mobility Matrix



Number of times a user u moves
from the sub-prefecture i to the
sub-prefecture j

$$m_{ij} = \frac{\sum_u \mathcal{M}_{ij}^u}{\sum_k \sum_u \mathcal{M}_{ik}^u}$$

Probability that an individual moves
from the sub-prefecture i to the
sub-prefecture j

Number of times a user u moves
from the sub-prefecture i to the
sub-prefecture k

Call Graph



Number of phone calls initiated
from the sub-prefecture i and
directed to the sub-prefecture j

$$c_{ij} = \frac{C_{ij}}{\sum_k C_{ik}}$$

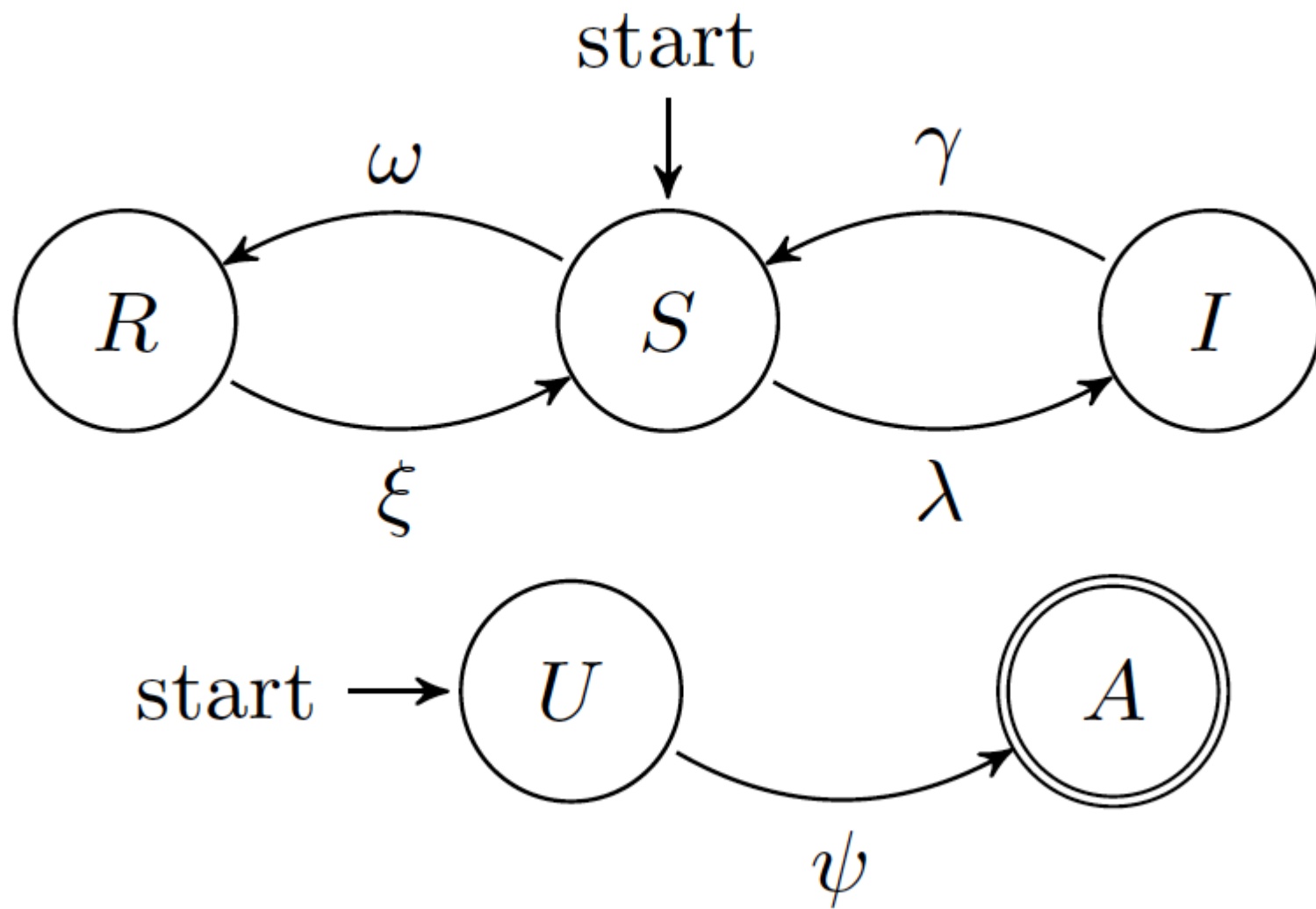
Probability of a call being
established between sub-
prefectures i and j

Number of phone calls initiated
from the sub-prefecture i and
directed to the sub-prefecture k

States



- Disease spreading process:
 - S: Susceptible
 - I: Infected
 - R: Resistant/recovered
- Information spreading process:
 - U: unaware
 - A: aware
- $N(t) = S(t) + I(t) + R(t) = U(t) + A(t)$





$$\begin{aligned} I_i[t+1] &= \sum_{j=1}^n m_{ji} \left[I_j[t] + \lambda \frac{S_j[t]}{N_j[t]} I_j[t] - \gamma I_j[t] \right] \\ S_i[t+1] &= \sum_{j=1}^n m_{ji} \left[S_j[t] - \lambda \frac{S_j[t]}{N_j[t]} I_j[t] + \gamma I_j[t] + \xi R_j[t] + \right. \\ &\quad \left. - \omega S_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \\ R_i[t+1] &= \sum_{j=1}^n m_{ji} \left[R_j[t] - \xi R_j[t] + \omega S_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \\ A_i[t+1] &= \sum_{j=1}^n m_{ji} \left[A_j[t] + \psi U_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \\ U_i[t+1] &= \sum_{j=1}^n m_{ji} \left[U_j[t] - \psi U_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \end{aligned} \quad (3)$$

Mobility Matrix

$$\begin{aligned} I_i[t+1] &= \sum_{j=1}^n m_{ji} \left[I_j[t] + \lambda \frac{S_j[t]}{N_j[t]} I_j[t] - \gamma I_j[t] \right] \\ S_i[t+1] &= \sum_{j=1}^n m_{ji} \left[S_j[t] - \lambda \frac{S_j[t]}{N_j[t]} I_j[t] + \gamma I_j[t] + \xi R_j[t] + \right. \\ &\quad \left. - \omega S_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \\ R_i[t+1] &= \sum_{j=1}^n m_{ji} \left[R_j[t] - \xi R_j[t] + \omega S_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \\ A_i[t+1] &= \sum_{j=1}^n m_{ji} \left[A_j[t] + \psi U_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \\ U_i[t+1] &= \sum_{j=1}^n m_{ji} \left[U_j[t] - \psi U_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \end{aligned} \quad (3)$$



Mobility Matrix

$$I_i[t + 1] = \sum_{j=1}^n m_{ji} \left[I_j[t] + \lambda \frac{S_j[t]}{N_j[t]} I_j[t] - \gamma I_j[t] \right]$$

$$S_i[t + 1] = \sum_{j=1}^n m_{ji} \left[S_j[t] - \lambda \frac{S_j[t]}{N_j[t]} I_j[t] + \gamma I_j[t] + \xi R_j[t] + \right. \\ \left. - \omega S_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right]$$

Call Matrix

$$R_i[t + 1] = \sum_{j=1}^n m_{ji} \left[R_j[t] - \xi R_j[t] + \omega S_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right]$$

$$A_i[t + 1] = \sum_{j=1}^n m_{ji} \left[A_j[t] + \psi U_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right]$$

$$U_i[t + 1] = \sum_{j=1}^n m_{ji} \left[U_j[t] - \psi U_j[t] \frac{\sum_{k=1}^n c_{kj} A_k[t]}{\sum_{k=1}^n c_{kj} N_k[t]} \right] \quad (3)$$





Real-time Predictive Modelling

- By using a model like this one, real-time predictive modelling might be possible
- Policy-makers might extract the parameters of the model, such as the mobility matrix, in real-time.
- Transmission models have been widely used for assessing potential strategies for containing diseases such as influenza.
- “Mobile big data” might help in developing more accurate models.



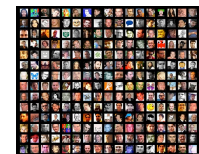
Real-time Predictive Modelling

- One of the key problems is understanding the uncertainty associated to the model.
- It is difficult to understand the contribution of many factors such as:
 - Climatic factors
 - Transmission seasonality
 - Long-term immunity of a population
 - ...
- Scenario-based modelling is routinely used in order to predict future evolution of epidemics.



Summary

- Epidemics are very complex processes.
- Existing models have been increasingly capable of capturing their essence.
- However there are still a number of open issues related to the modelling of real disease spreading or information dissemination.
- Mixing geographic factors with epidemics model is very relevant and important.
- The availability of “big data” might help in building more realistic and possibly real-time epidemic models.



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