



Social and Technological Network Analysis

Lecture 8: 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 biology.
 - But it also has important parallels in information/idea diffusion in 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).

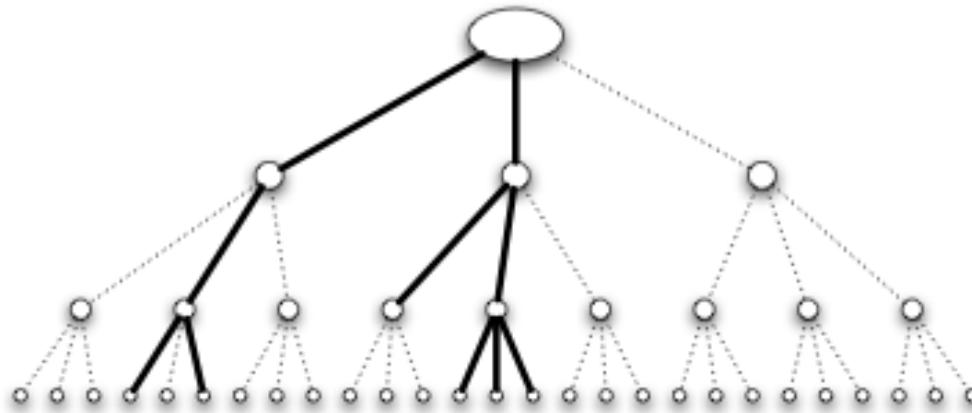


Branching Process

- Simple model.
- **First wave:** A person carrying a disease enters the population and transmit 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 $k \times k = k^2$ people.
- **Subsequent waves:** same process.

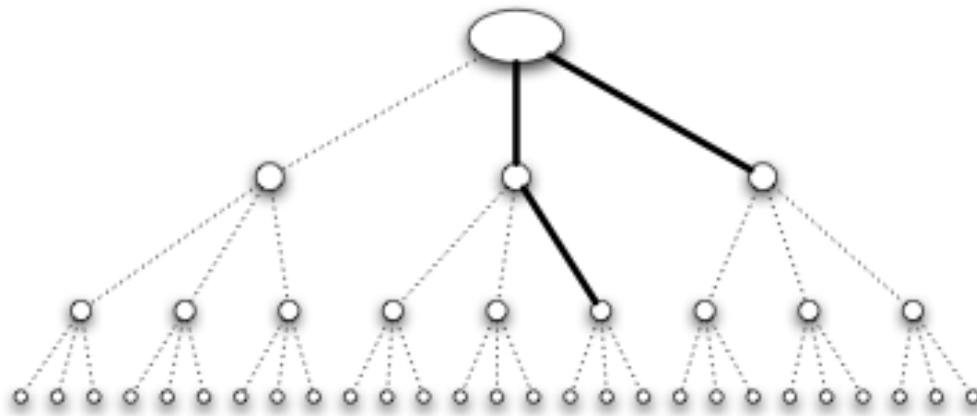


Example with $k=3$



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



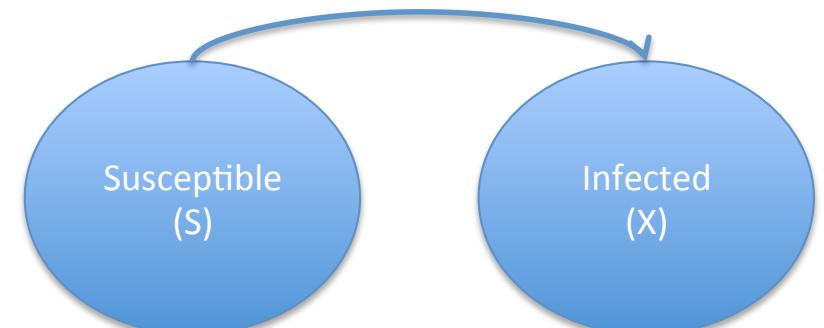
- When R_0 is close 1, slightly changing p or k can result in epidemics dying out or happening.
 - Quarantining people/nodes reduces k .
 - Encouraging better sanitary practices reduces germs spreading [reducing p].
- Limitations of this model:
 - No realistic contact networks: no triangles!
 - Nodes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

Formal Epidemics Models

The SI Model



- S: susceptible individuals.
- X: infected individuals, when infected they can infect others continuously (different from before).
- 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 $X\beta S/n$.





SI Model

$$\frac{dX}{dt} = \beta \frac{SX}{n}$$

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

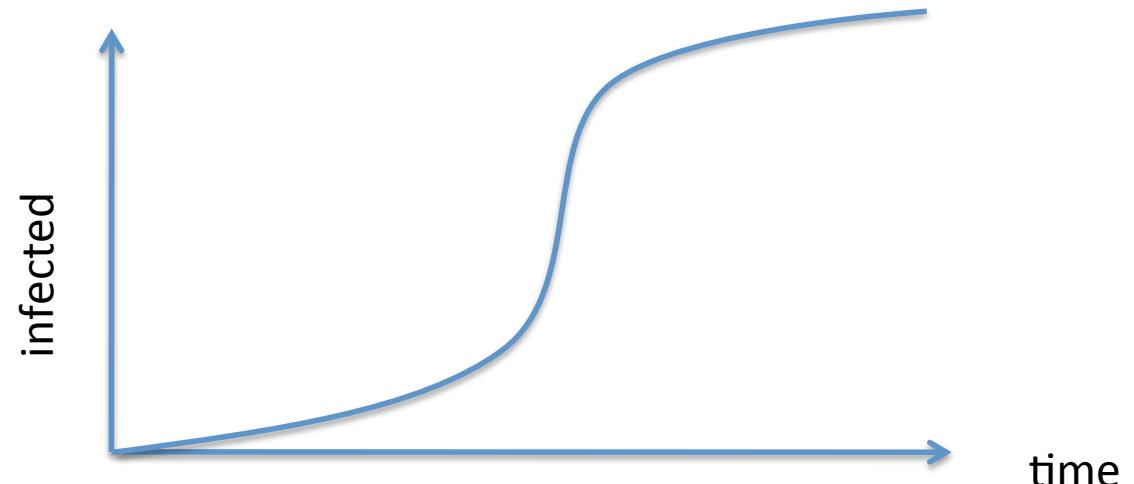
$$S = \frac{S}{n} \qquad X = \frac{X}{n}$$

$$s = 1 - x$$

$$\frac{dx}{dt} = \beta x(1 - x)$$

$$x(t) = \frac{x_0 e^{\beta t}}{1 - x_0 + x_0 e^{\beta t}}$$

Logistic Growth Equation





Microscopic explanation

- Probability for a node to not be infected from each infected contact: $1 - \lambda$
- Number of infected contacts per node: $\langle k \rangle x$
- Probability for a susceptible node to avoid infection from all its $\langle k \rangle x$ infected contacts: $p = (1 - \lambda)^{\langle k \rangle x}$
- Probability to be infected from at least one infected contact: $1 - p = 1 - (1 - \lambda)^{\langle k \rangle x}$
- If $\lambda \ll 1$ then $p \approx \lambda \langle k \rangle x = \beta x$
- Therefore, the increase of infected nodes reads:

$$\frac{dx}{dt} = \beta x s = \beta x(1 - x)$$



SIR Model

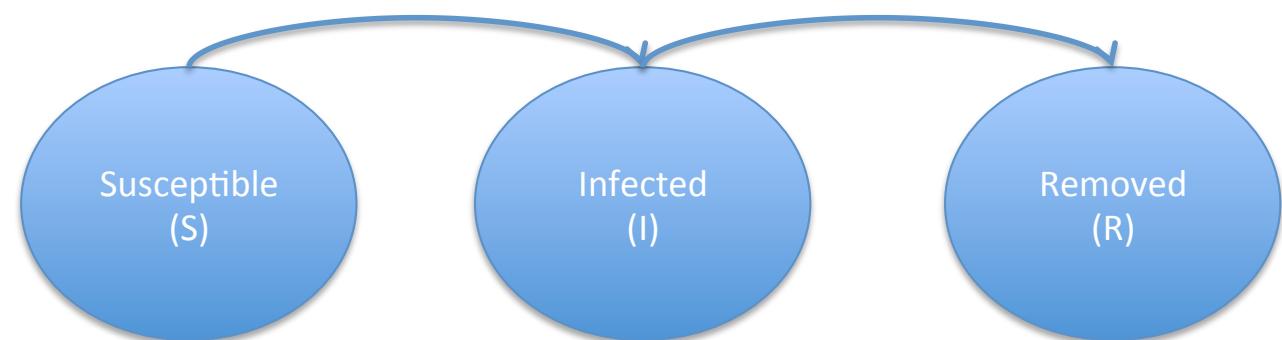
- Infected nodes recover at a rate γ .
- A node stays infected for τ time.
- Branching process is SIR with $\tau=1$.

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

$$\frac{dx}{dt} = \beta sx - \gamma x$$

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

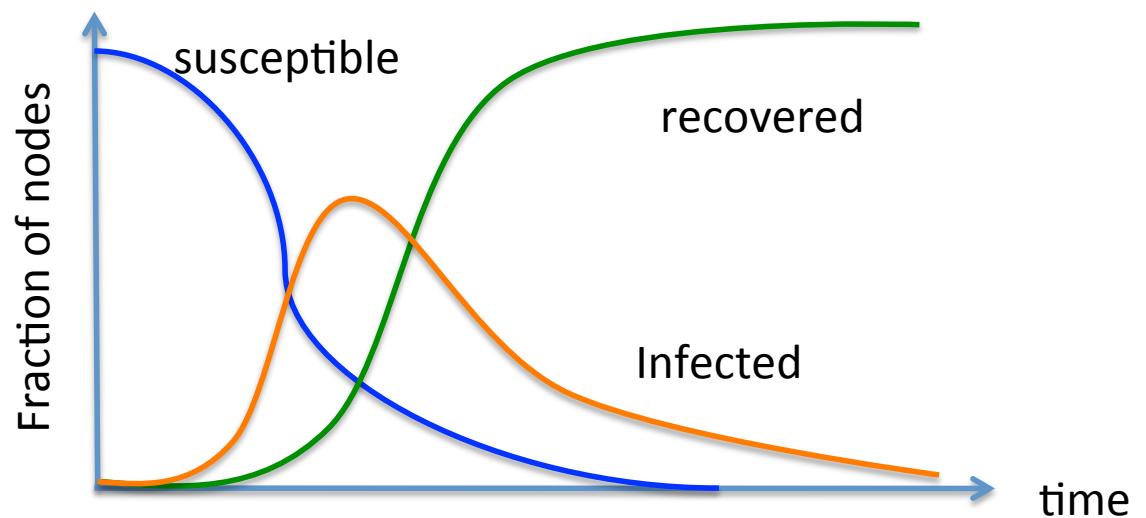
$$s + x + r = 1$$





Example

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



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.



SIS Model

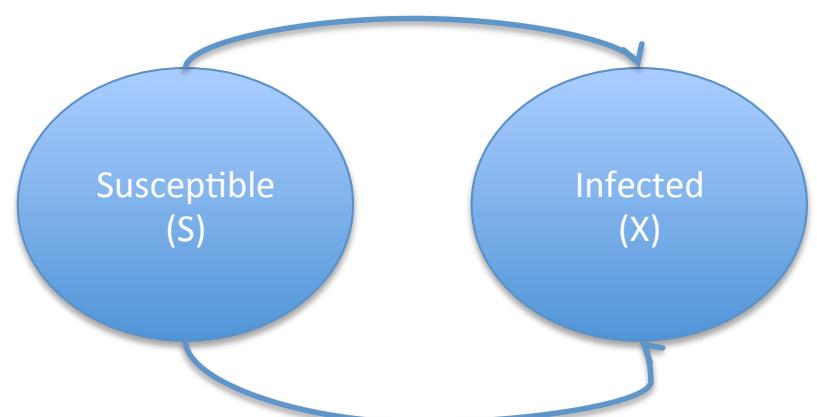
$$\frac{ds}{dt} = \gamma x - \beta s x$$

$$\frac{dx}{dt} = \beta s x - \gamma x$$

$$s + x = 1$$

$$\frac{dx}{dt} = (\beta - \gamma - \beta x)x$$

- 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.



Relaxing Assumptions

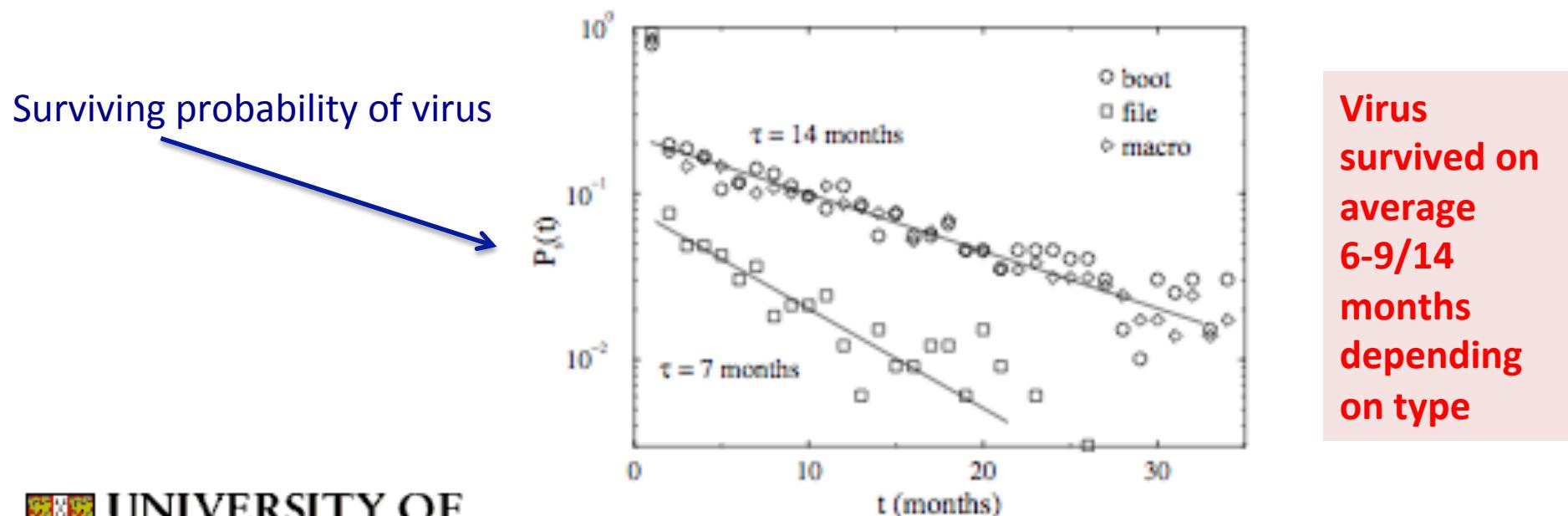


- Homogeneous Mixing: a node connects to the same average number of other nodes as any other.
- Most real networks are not Erdos-Renyi random networks (for which the homogeneous mixing assumption holds).
- Most networks have heterogeneous degree distributions.
 - Scale free networks!



Would the model apply to SF?

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

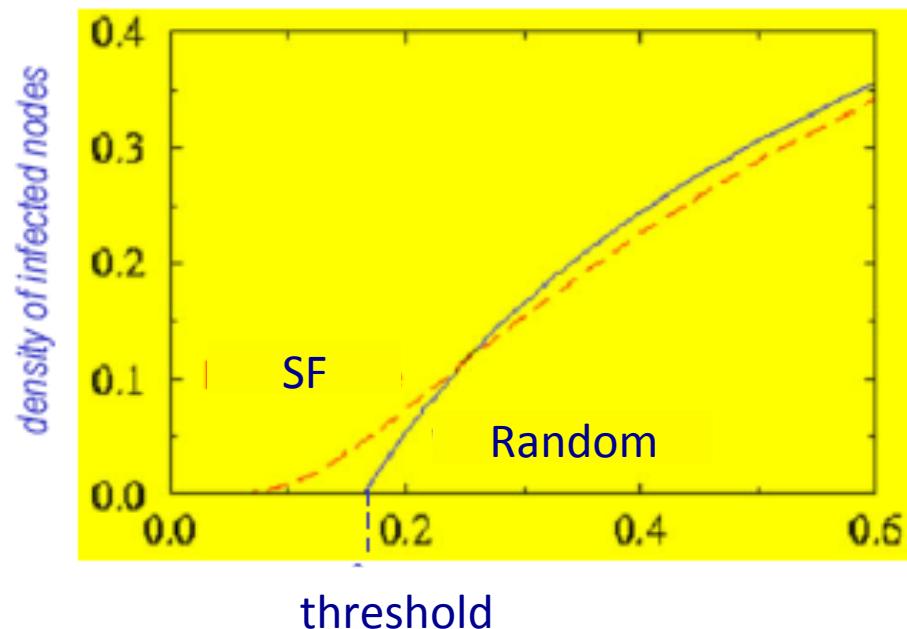


How to justify this survival time?

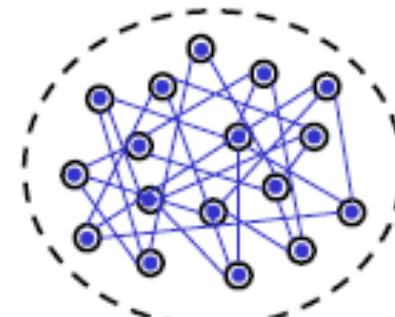


- The virus survival time is considerably high with respect to the results of epidemic models of spreading/recovering:
 - Something wrong with the epidemic threshold!
- Experiment: SIS over a generated Scale Free network (exponent -3).

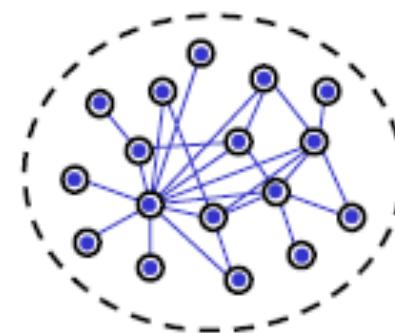
No Epidemic Threshold for SF!



Infections proliferate in SF networks
independently of their spreading rates!



Random Network



Scale Free Network



The vanishing threshold

- Percentage of infected of degree k :

$$x_k(t) = i_0(1 + ck(e^{t/\tau} - 1))$$

$$\tau = \frac{\langle k \rangle}{\beta(\langle k^2 \rangle - \langle k \rangle)}$$

- Larger degree nodes are infected with higher probability
- In scale-free networks with $2 < \gamma \leq 3$ $\langle k^2 \rangle$ diverges as $N \rightarrow \infty$, so the epidemic spreads very fast

Following result on Immunization



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

Immunization on SF Networks

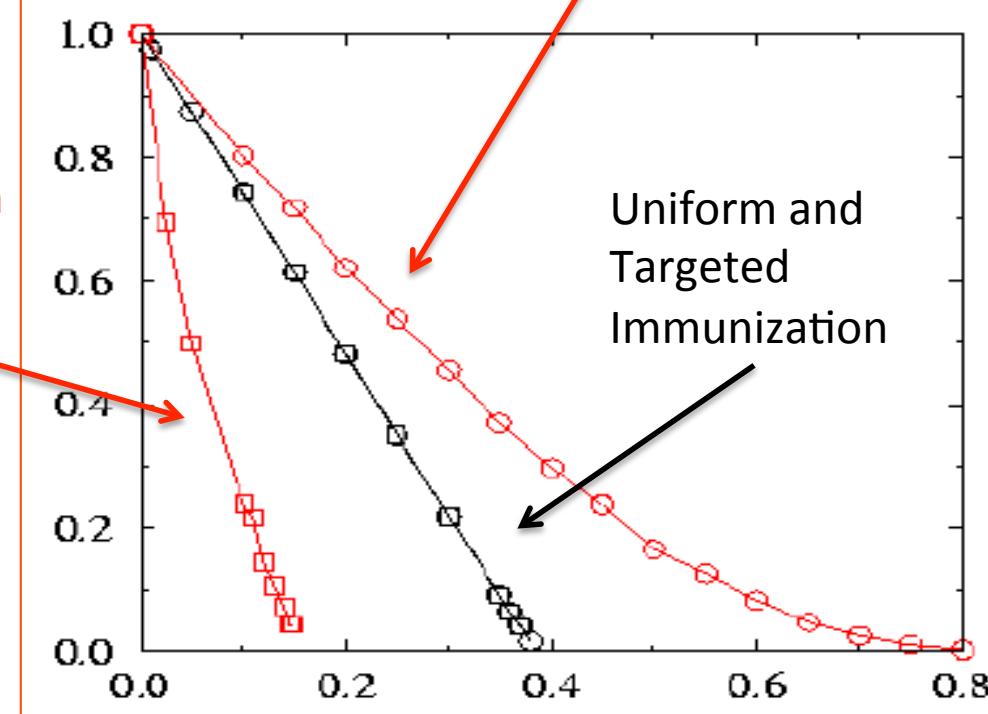


- Red=SF
- Black= Random

Targeted Immunization

Uniform Immunization

Uniform and
Targeted
Immunization



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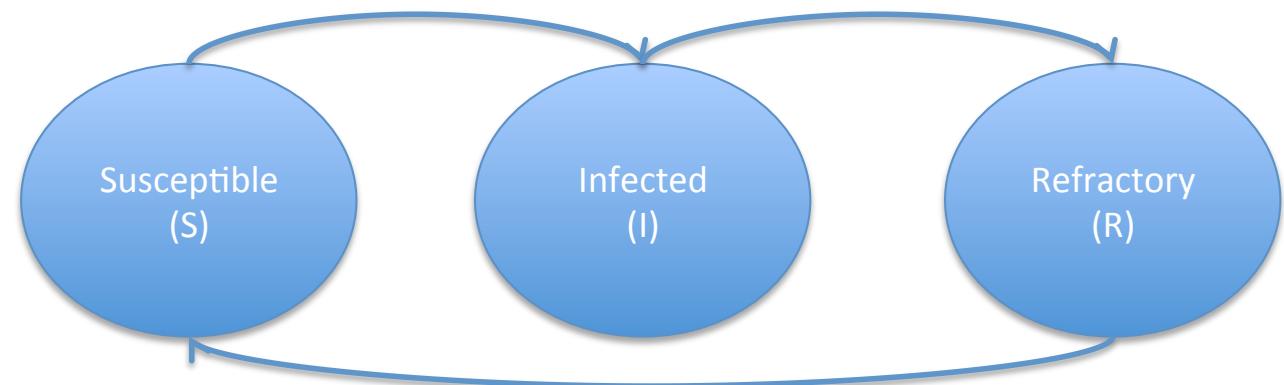
Local Immunization

- Global knowledge on the network structure is rarely (or never) available
- 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!)

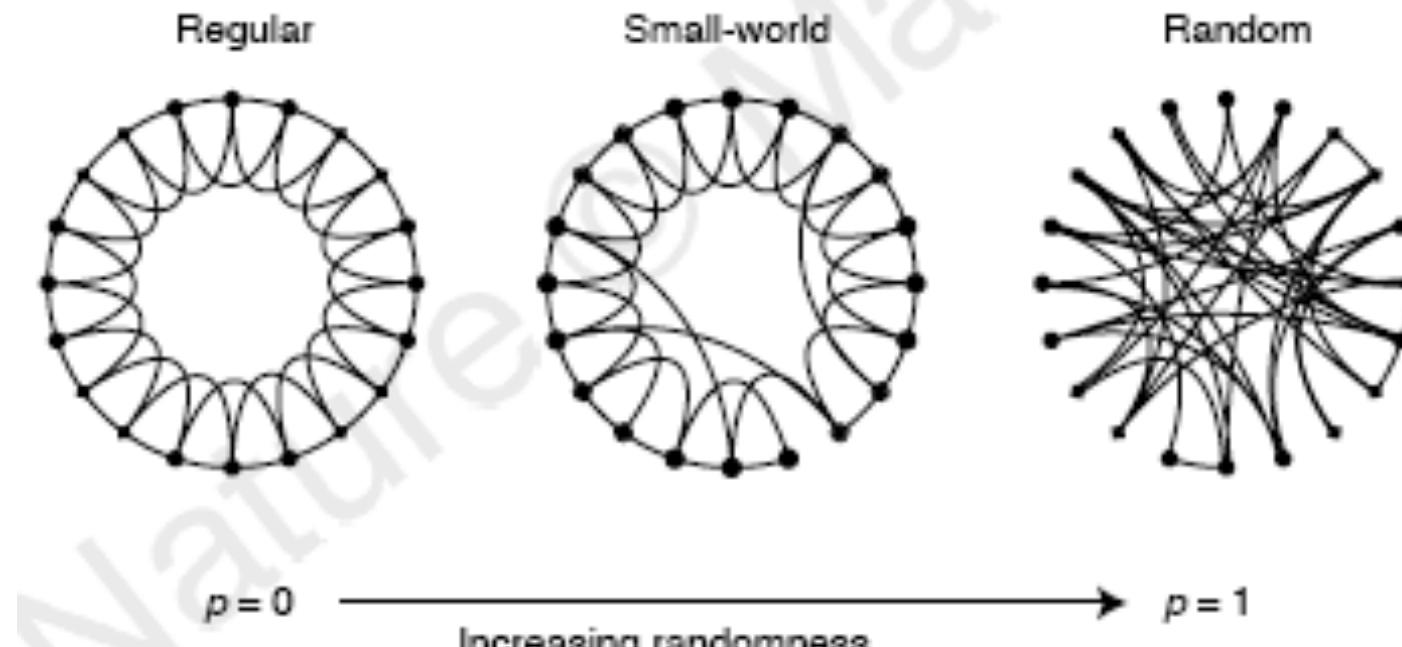


SIRS Model

- SIR but after some time an R node can become susceptible again.
- A number of epidemics spread in this manner (remaining latent for a while and having bursts).



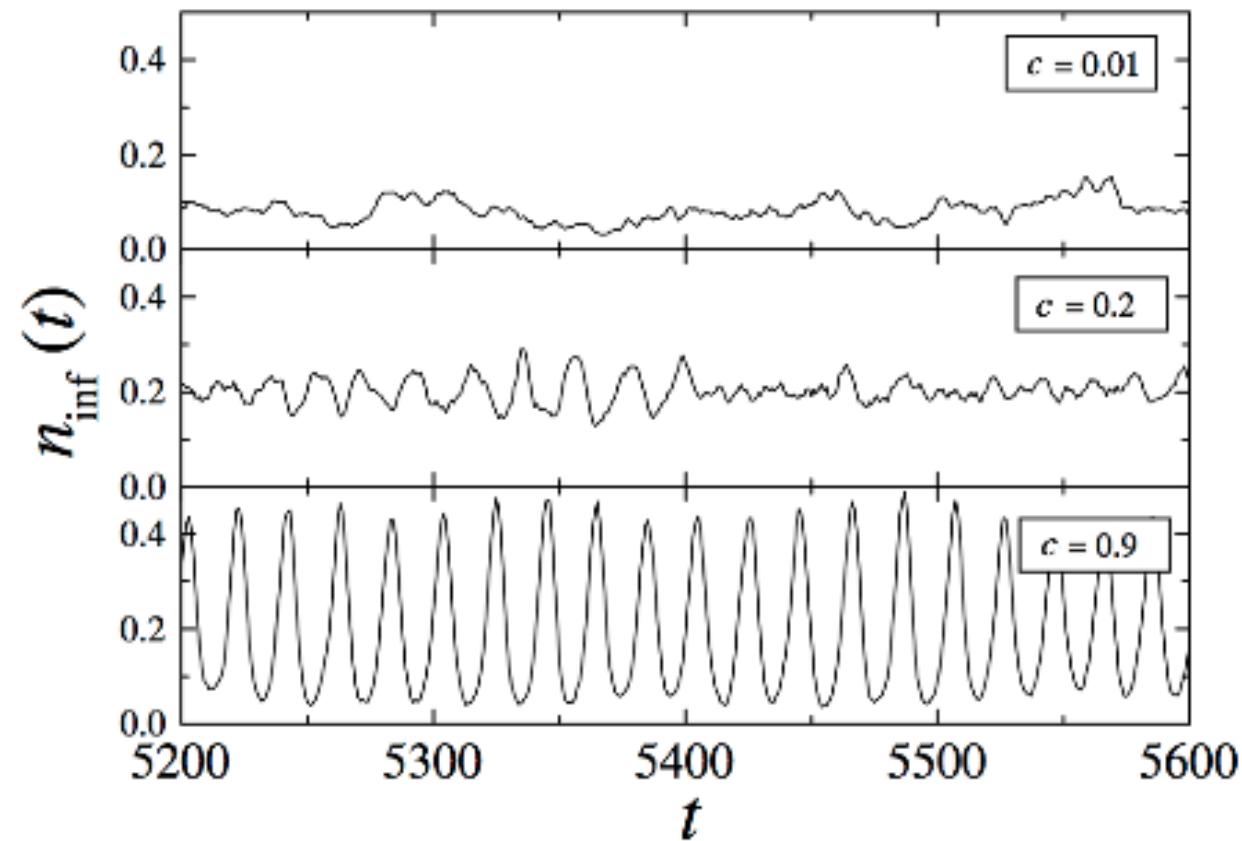
Application of SIRS to Small World Models





Numerical Results

- c is the rewiring probability



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.



References

- Chapter 21
- Pastor-Satorras, R. and Vespignani, A. Epidemic Spreading in Scale-Free Networks. *Phys. Rev. Lett.* (86), n.14. Pages = 3200--3203. 2001.
- Pastor-Satorras, R. and Vespignani, A. Immunization of Complex Networks. *Physical Review E* 65. 2002.
- Marcelo Kuperman and Guillermo Abramson. Small world effect in an epidemiological model. *Physical Review Letters*, 86(13):2909–2912, March 2001.