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# Social and Technological Network Analysis

## Lecture 8: Epidemics Spreading

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# In This Lecture

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- 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

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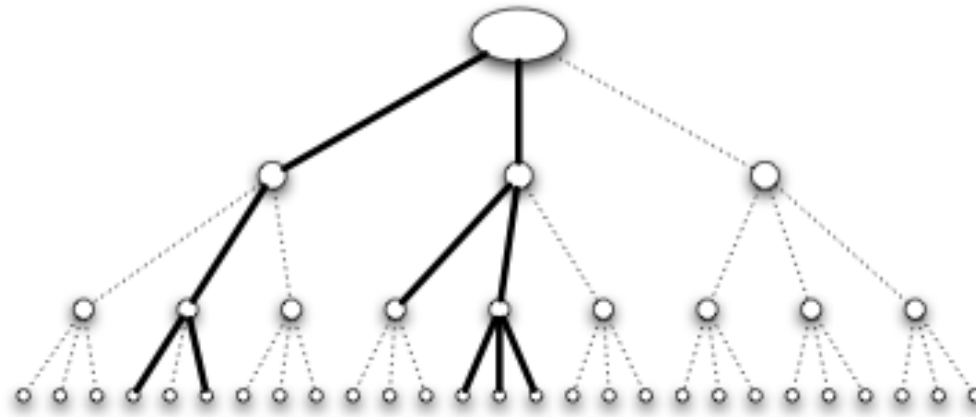
- In cascade spreading nodes make decisions based on pay-off benefits of adopting one strategy or the other.
- In epidemic spreading
  - Lack of decision making.
  - Process of contagion is complex and unobservable
    - In some cases it involves (or can be modeled as randomness).

# Branching Process



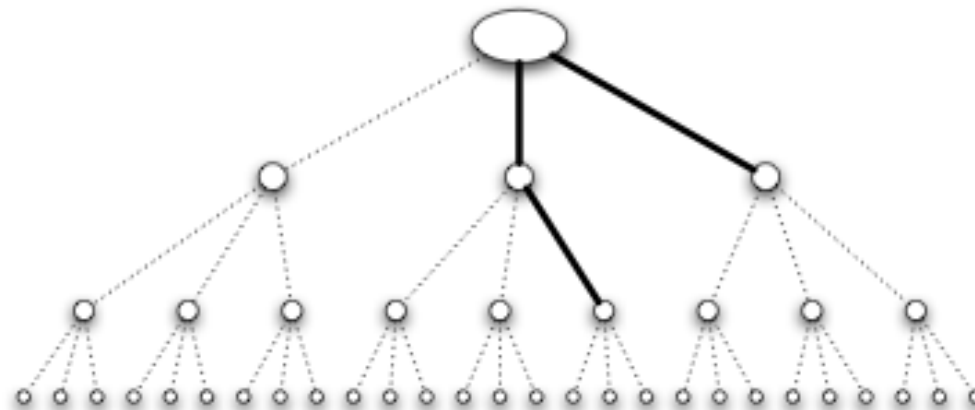
- Simple model.
- **First wave:** A person carrying a disease enters the population and transmit to all he meets with probability  $p$ . He meets  $k$  people: a portion of which will be infected.
- **Second wave:** each of the  $k$  people goes and meet  $k$  different people. So we have a second wave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



# Basic Reproductive Number

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- Basic Reproductive Number  $R_0 = p \cdot k$ 
  - 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.

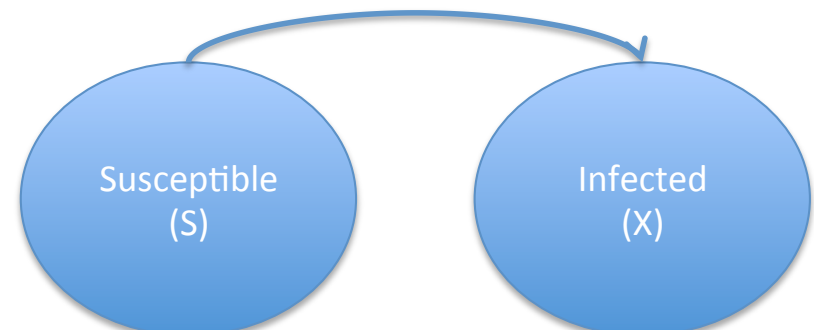
# Formal Epidemics Models

## The SI Model

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- 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

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

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

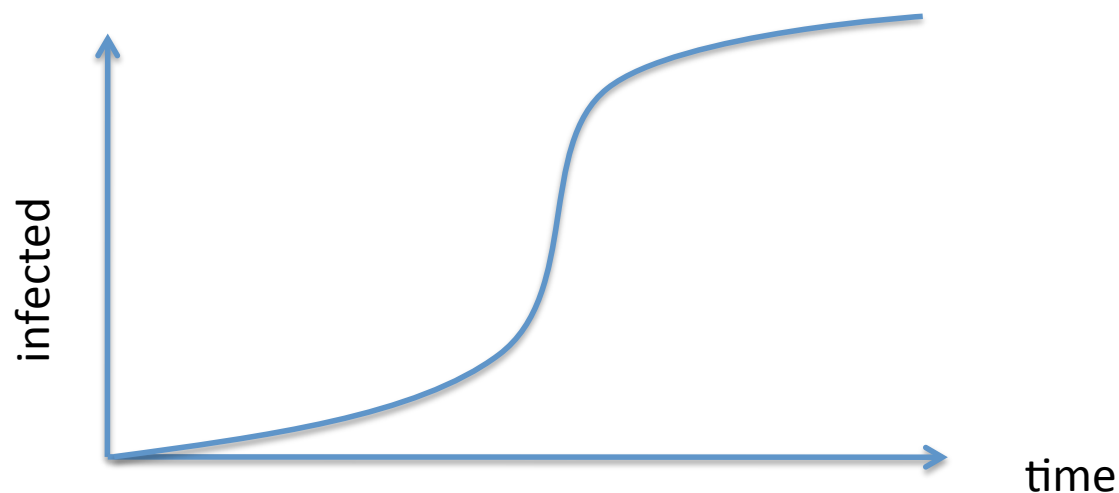
$$s = \frac{S}{n} \quad x = \frac{X}{n}$$

$$s = 1 - x$$

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

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

**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

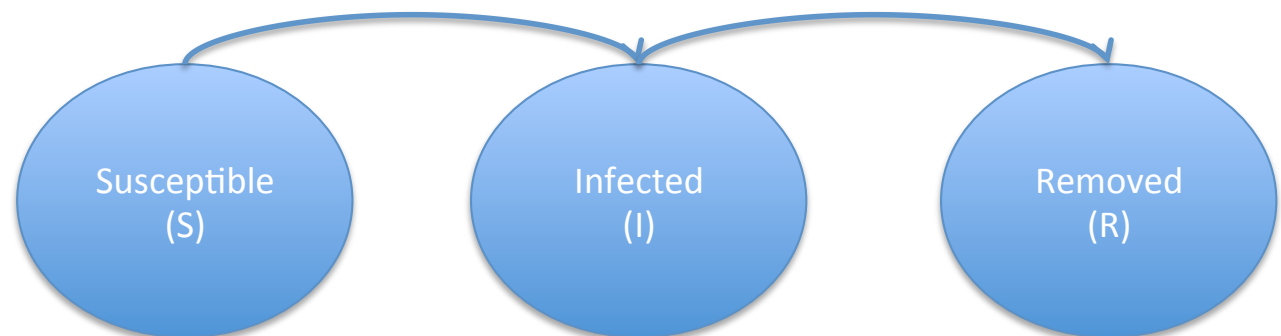
- Infected nodes recover at a rate  $\gamma$ .
- A node stays infected for  $\tau$  time.
- Branching process is SIR with  $\tau=1$ .

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

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

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

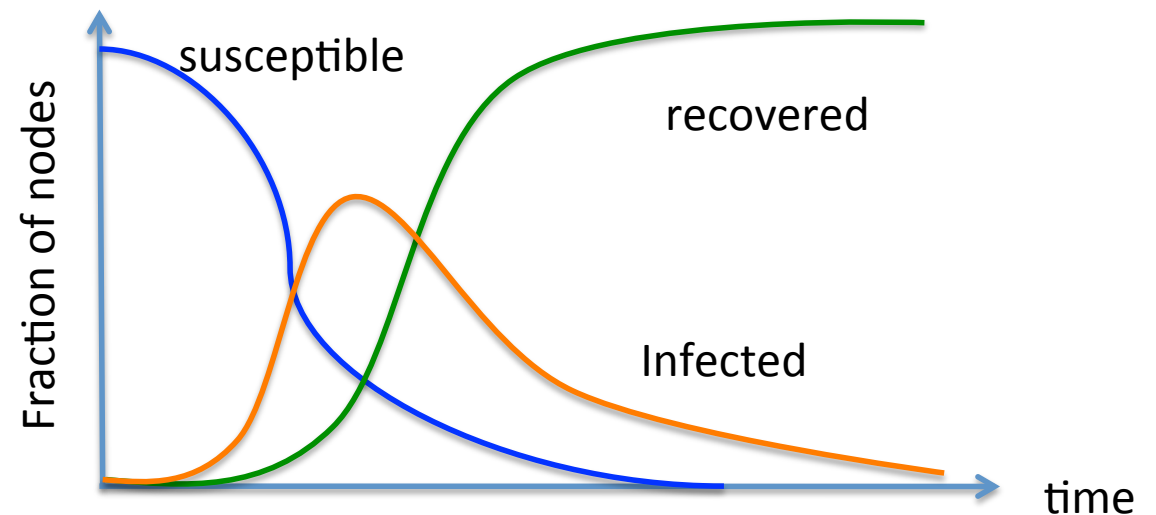
$$s + x + r = 1$$



# 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  $\beta$  and  $\gamma$ :
  - 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

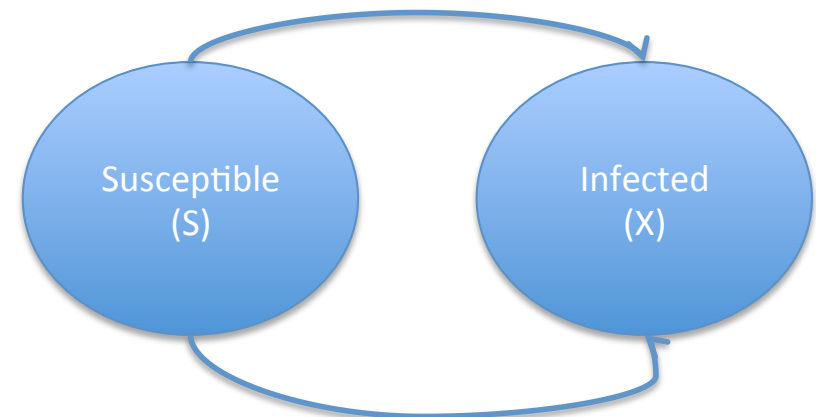
$$\frac{ds}{dt} = \gamma x - \beta s x$$

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

$$s + x = 1$$

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

- If  $\beta > \gamma$  growth curve like in SI but never reaching all population infected. The fraction of infected  $\rightarrow 0$  as  $\beta$  approaches  $\gamma$ .
- 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!

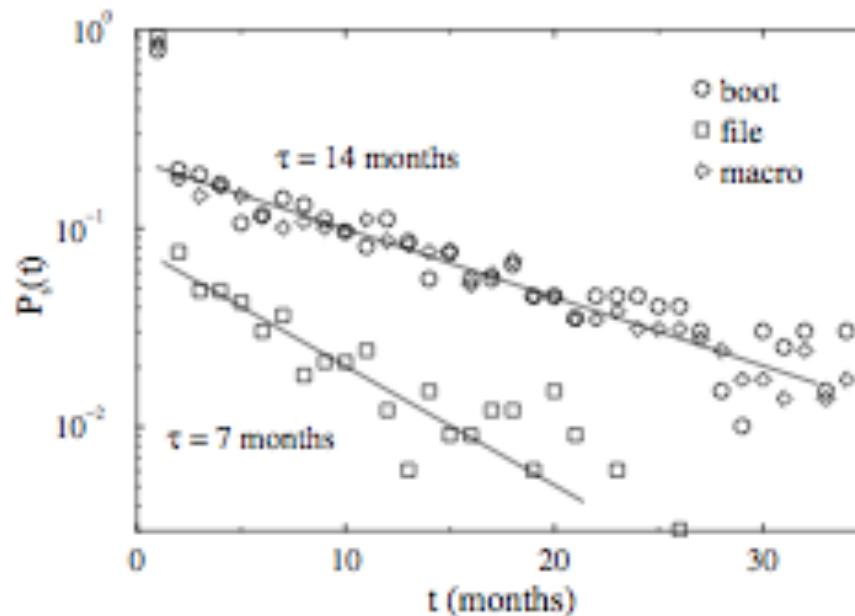
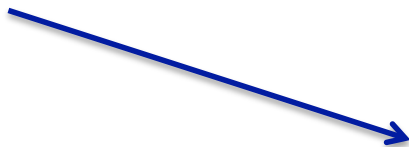


# Would the model apply to SF?



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

Surviving probability of virus



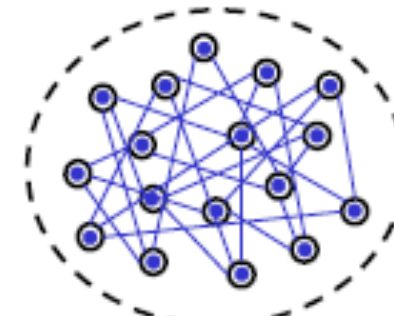
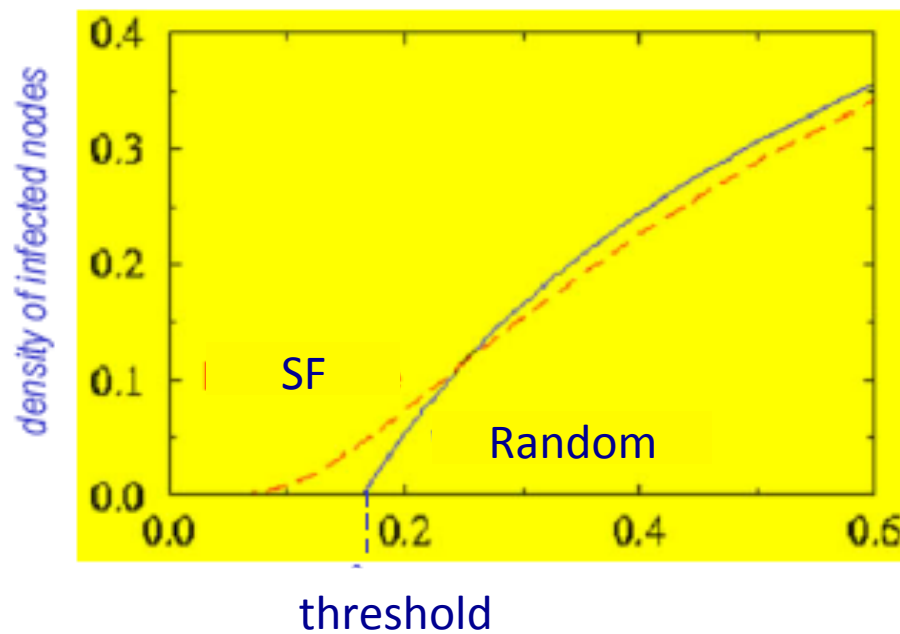
Virus survived on average 6-9/14 months depending on type

# How to justify this survival time?

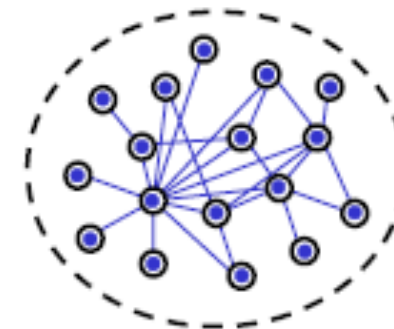


- The virus survival time is considerably high with respect to the results of epidemic models of spreading/recovering:
  - Something wrong with the epidemic threshold!
- Experiment: SIS over a generated Scale Free network (exponent -3).

# No Epidemic Threshold for SF!



Random Network



Scale Free Network

Infections proliferate in SF networks independently of their spreading rates!

# 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

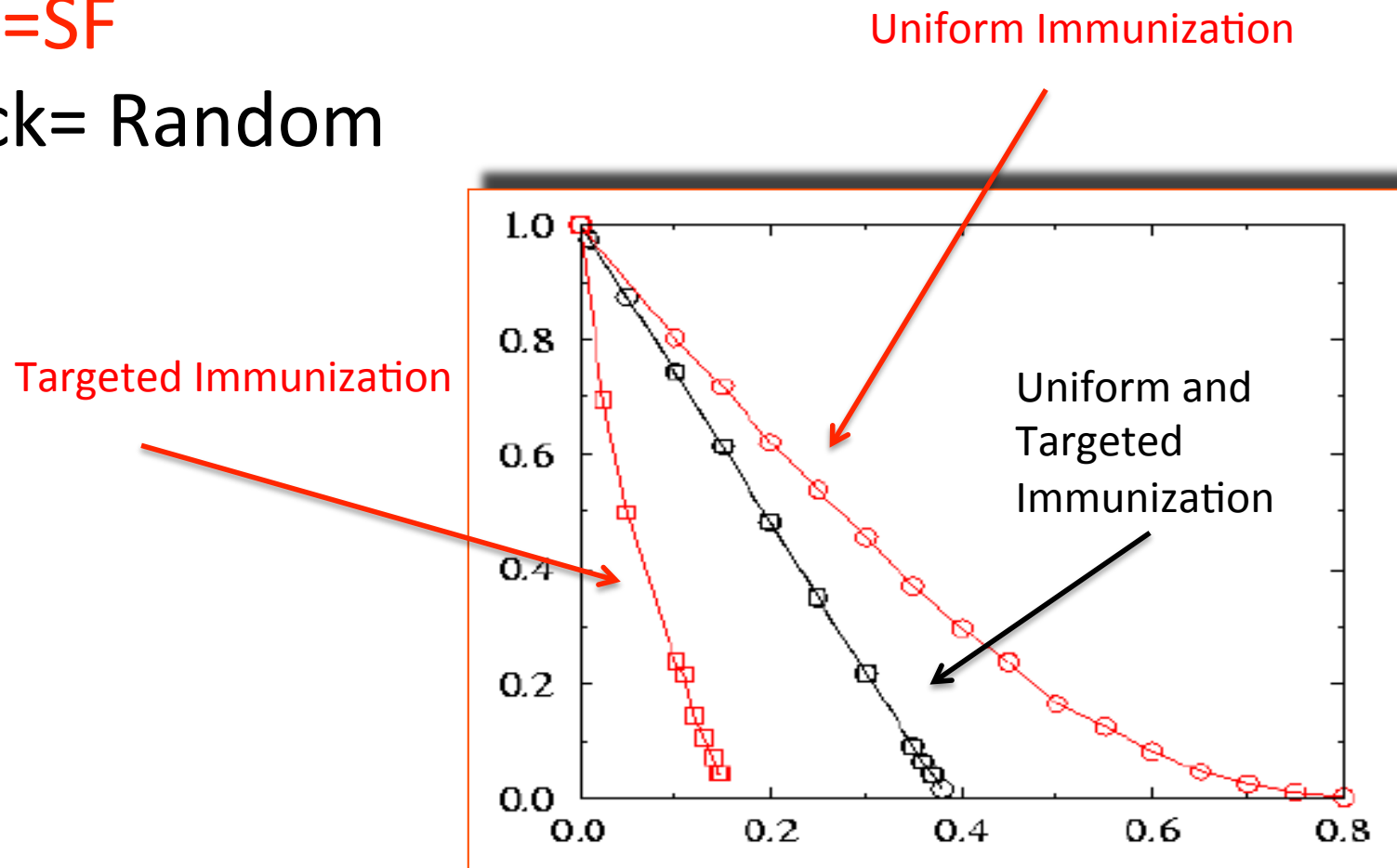


- Random network can be immunized with some sort of uniform immunization process [oblivious of the characteristics of nodes].
- **This does not work in SF networks** no matter how many nodes are immunized [unless it is all of them].
- Targeted immunization needs to be applied
  - Keeping into account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

# Immunization on SF Networks



- Red=SF
- Black= Random



# Local Immunization

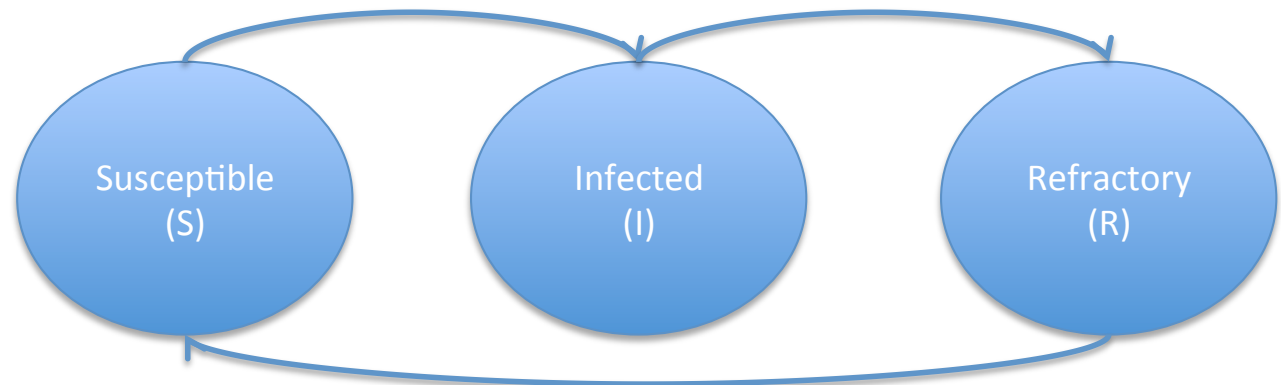


- Global knowledge on the network structure is rarely (or never) available
- Local immunization strategy:
  - Select  $g$  nodes at random
  - Ask to each of them to pass over the vaccine to one of their neighbors
  - As a result, a node with degree  $k$  is immunized with a probability  $kP(k)$  (hubs are immunized with higher probability!)



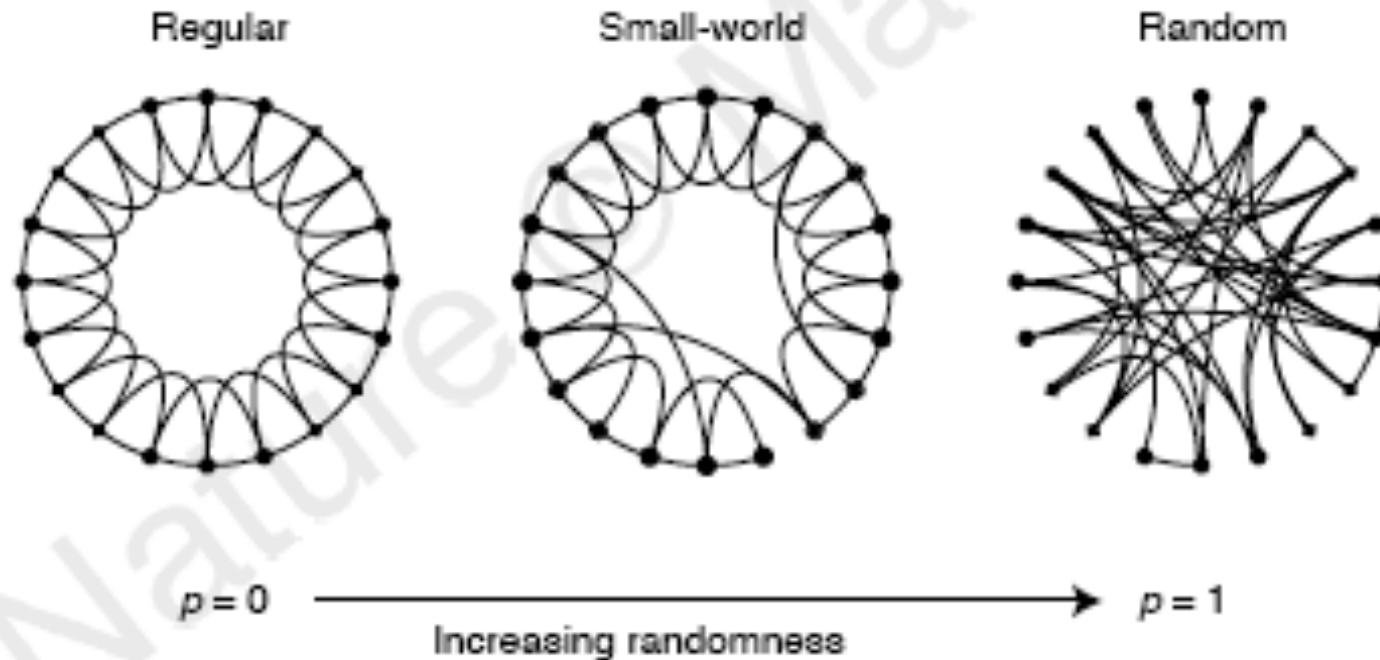
# SIRS Model

- 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).

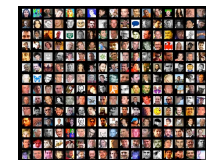




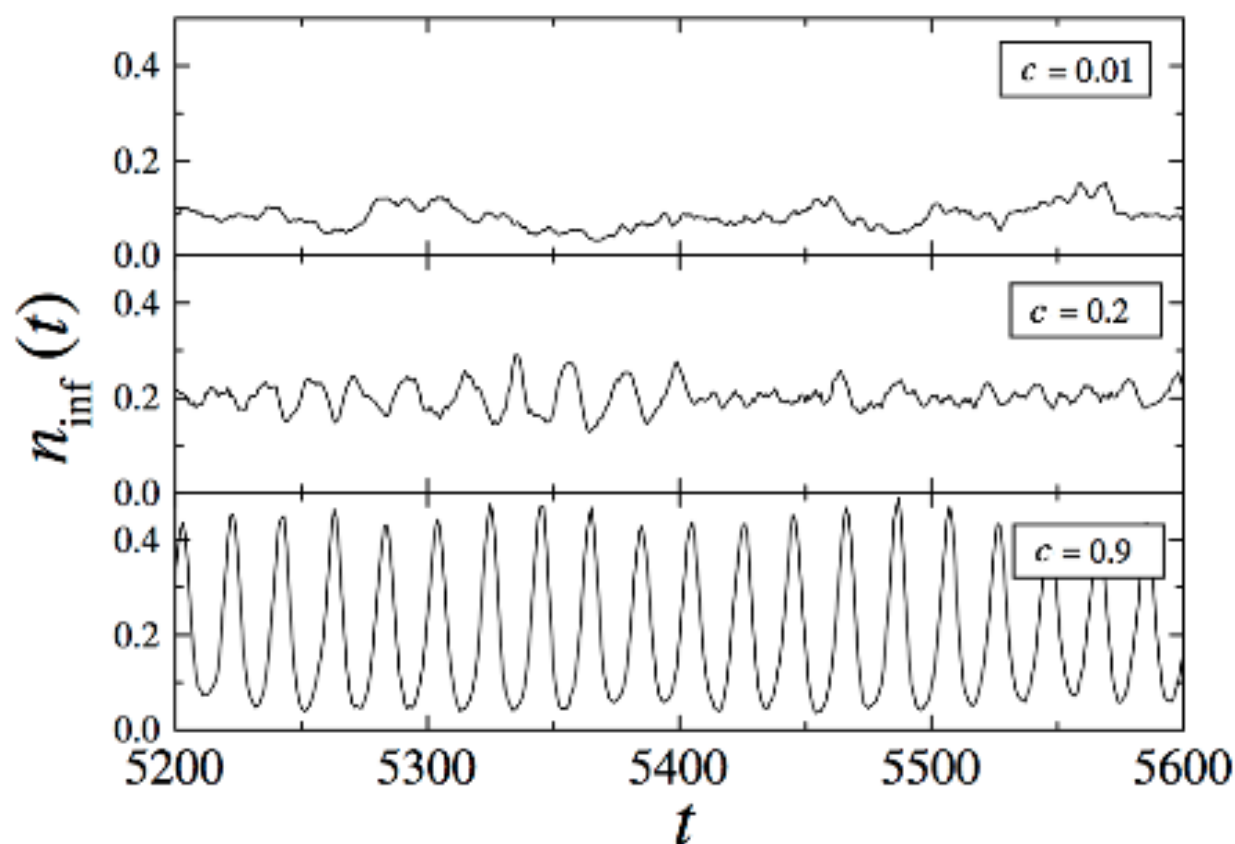
# Application of SIRS to Small World Models



# Numerical Results



- $c$  is the rewiring probability



# Summary

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- 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.