### **Generalized Contagion**

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Principles of Complex Systems, Vols. 1 & 2 CSYS/MATH 300 and 303, 2021–2022 | @pocsvox

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Computational Story Lab | Vermont Complex Systems Center Vermont Advanced Computing Core | University of Vermont























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# "Universal Behavior in a Generalized Model of Contagion"

Dodds and Watts, Phys. Rev. Lett., **92**, 218701, 2004. <sup>[5]</sup>



"A generalized model of social and biological contagion"

Dodds and Watts, J. Theor. Biol., **232**, 587–604, 2005. [6] Introduction
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Basic questions about contagion

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### Basic questions about contagion



How many types of contagion are there?

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### Basic questions about contagion



How many types of contagion are there?



How can we categorize real-world contagions?

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### Basic questions about contagion

How many types of contagion are there?

How can we categorize real-world contagions?

Can we connect models of disease-like and social contagion?

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### Basic questions about contagion

How many types of contagion are there?

How can we categorize real-world contagions?

Can we connect models of disease-like and social contagion?

& Focus: mean field models.

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The standard SIR model [11]

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The standard SIR model [11]



= basic model of disease contagion

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#### The standard SIR model [11]

🚓 = basic model of disease contagion

Three states:

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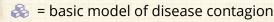
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#### The standard SIR model [11]



Three states:

1. S = Susceptible

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#### The standard SIR model [11]



= basic model of disease contagion



Three states:

- 1. S = Susceptible
- 2. I = Infective/Infectious

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#### The standard SIR model [11]



= basic model of disease contagion



Three states:

- 1. S = Susceptible
- 2. I = Infective/Infectious
- 3. R = Recovered

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#### The standard SIR model [11]



= basic model of disease contagion



Three states:

- 1. S = Susceptible
- 2. I = Infective/Infectious
- 3. R = Recovered or Removed or Refractory

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#### The standard SIR model [11]



= basic model of disease contagion



Three states:

- 1. S = Susceptible
- 2. I = Infective/Infectious
- 3. R = Recovered or Removed or Refractory

$$\Re S(t) + I(t) + R(t) = 1$$

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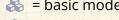
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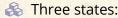
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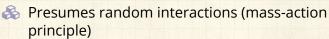
#### The standard SIR model [11]



= basic model of disease contagion



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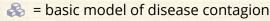
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#### The standard SIR model [11]



- Three states:
  - 1. S = Susceptible
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$$\Re S(t) + I(t) + R(t) = 1$$

Presumes random interactions (mass-action principle)

Interactions are independent (no memory)

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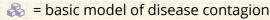
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#### The standard SIR model [11]



- Three states:
  - 1. S = Susceptible
  - 2. I = Infective/Infectious
  - 3. R = Recovered or Removed or Refractory

$$\Re S(t) + I(t) + R(t) = 1$$

- Presumes random interactions (mass-action principle)
- Interactions are independent (no memory)
- Discrete and continuous time versions

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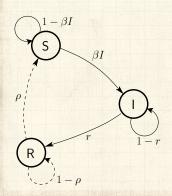
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### Discrete time automata example:



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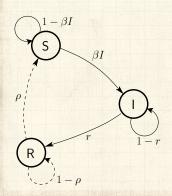
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### Discrete time automata example:



#### Transition Probabilities:

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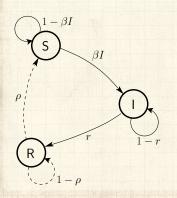
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### Discrete time automata example:



Transition Probabilities:

 $\beta$  for being infected given contact with infected

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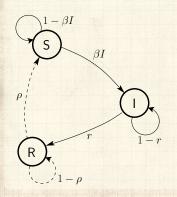
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### Discrete time automata example:



Transition Probabilities:

 $\beta$  for being infected given contact with infected r for recovery

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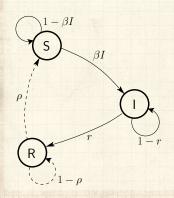
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### Discrete time automata example:



Transition Probabilities:

 $\beta$  for being infected given contact with infected r for recovery  $\rho$  for loss of immunity

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Original models attributed to

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### Original models attributed to



4 1920's: Reed and Frost

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### Original models attributed to

4 1920's: Reed and Frost

1920's/1930's: Kermack and McKendrick [8, 10, 9]

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### Original models attributed to



4 1920's: Reed and Frost



1920's/1930's: Kermack and McKendrick [8, 10, 9]



Coupled differential equations with a mass-action principle

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### Differential equations for continuous model

$$\frac{\mathrm{d}}{\mathrm{d}t}S = -\beta \underline{IS} + \rho R$$

$$\frac{\mathrm{d}}{\mathrm{d}t}I = \beta \underline{IS} - rI$$

$$\frac{\mathrm{d}}{\mathrm{d}t}R = rI - \rho R$$

 $\beta$ , r, and  $\rho$  are now rates.

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### Differential equations for continuous model

$$\frac{d}{dt}S = -\beta IS + \rho R$$

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 $\beta$ , r, and  $\rho$  are now rates.

Reproduction Number  $R_0$ :

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### Differential equations for continuous model

$$\frac{\mathrm{d}}{\mathrm{d}t}S = -\beta \underline{IS} + \rho R$$
$$\frac{\mathrm{d}}{\mathrm{d}t}I = \beta \underline{IS} - rI$$
$$\frac{\mathrm{d}}{\mathrm{d}t}R = rI - \rho R$$

 $\beta$ , r, and  $\rho$  are now rates.

### Reproduction Number $R_0$ :

 $R_0$  = expected number of infected individuals resulting from a single initial infective

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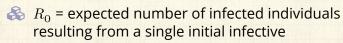


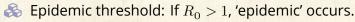
### Differential equations for continuous model

$$\frac{\mathrm{d}}{\mathrm{d}t}S = -\beta \underline{IS} + \rho R$$
 
$$\frac{\mathrm{d}}{\mathrm{d}t}I = \beta \underline{IS} - rI$$
 
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 $\beta$ , r, and  $\rho$  are now rates.

### Reproduction Number $R_0$ :





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## Reproduction Number $R_0$

#### Discrete version:



Set up: One Infective in a randomly mixing population of Susceptibles

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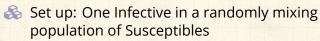
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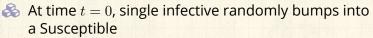
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## Reproduction Number $R_0$

#### Discrete version:





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#### Discrete version:

- Set up: One Infective in a randomly mixing population of Susceptibles
- At time t=0, single infective randomly bumps into a Susceptible
- $\ensuremath{\mathfrak{S}}$  Probability of transmission =  $\beta$

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#### Discrete version:

- Set up: One Infective in a randomly mixing population of Susceptibles
- At time t=0, single infective randomly bumps into a Susceptible
- $\ensuremath{\mathfrak{S}}$  Probability of transmission =  $\beta$
- $\clubsuit$  At time t=1, single Infective remains infected with probability 1-r

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#### Discrete version:

- Set up: One Infective in a randomly mixing population of Susceptibles
- $\clubsuit$  At time t=0, single infective randomly bumps into a Susceptible
- $\ensuremath{\mathfrak{S}}$  Probability of transmission =  $\beta$
- At time t=1, single Infective remains infected with probability 1-r
- At time t=k, single Infective remains infected with probability  $(1-r)^k$

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#### Discrete version:



Expected number infected by original Infective:

$$R_0 = \beta + (1 - r)\beta + (1 - r)^2\beta + (1 - r)^3\beta + \dots$$

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#### Discrete version:



Expected number infected by original Infective:

$$R_0 = \beta + (1-r)\beta + (1-r)^2\beta + (1-r)^3\beta + \dots$$

$$=\beta \left(1+(1-r)+(1-r)^2+(1-r)^3+...\right)$$

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#### Discrete version:



Expected number infected by original Infective:

$$R_0 = \beta + (1-r)\beta + (1-r)^2\beta + (1-r)^3\beta + \dots$$

$$= \beta \left( 1 + (1-r) + (1-r)^2 + (1-r)^3 + \dots \right)$$

$$=\beta \frac{1}{1-(1-r)}$$

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#### Discrete version:



Expected number infected by original Infective:

$$R_0 = \beta + (1-r)\beta + (1-r)^2\beta + (1-r)^3\beta + \dots$$

$$=\beta \left( 1+(1-r)+(1-r)^2+(1-r)^3+\ldots \right)$$

$$=\beta \frac{1}{1-(1-r)} = \beta/r$$

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#### Discrete version:



Expected number infected by original Infective:

$$R_0 = \beta + (1-r)\beta + (1-r)^2\beta + (1-r)^3\beta + \dots$$

$$=\beta \left(1+(1-r)+(1-r)^2+(1-r)^3+...\right)$$

$$=\beta \frac{1}{1-(1-r)} = \beta/r$$



Similar story for continuous model.

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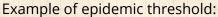
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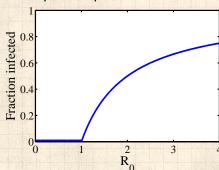
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### Independent Interaction models





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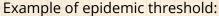
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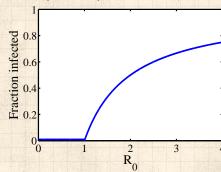
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### Independent Interaction models





Continuous phase transition.

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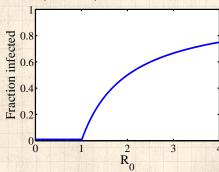
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### Independent Interaction models

### Example of epidemic threshold:



Continuous phase transition.

Fine idea from a simple model.

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Valiant attempts to use SIR and co. elsewhere:

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### Valiant attempts to use SIR and co. elsewhere:



Adoption of ideas/beliefs (Goffman & Newell, 1964)[7]

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### Valiant attempts to use SIR and co. elsewhere:

Adoption of ideas/beliefs (Goffman & Newell, 1964) [7]

Spread of rumors (Daley & Kendall, 1964, 1965) [3, 4]

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#### Valiant attempts to use SIR and co. elsewhere:

- Adoption of ideas/beliefs (Goffman & Newell, 1964) [7]
- Spread of rumors (Daley & Kendall, 1964, 1965) [3, 4]
- A Diffusion of innovations (Bass, 1969)[1]

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### Valiant attempts to use SIR and co. elsewhere:

- Adoption of ideas/beliefs (Goffman & Newell, 1964) [7]
- Spread of rumors (Daley & Kendall, 1964, 1965) [3, 4]
- Diffusion of innovations (Bass, 1969)
- Spread of fanatical behavior (Castillo-Chávez & Song, 2003) [2]

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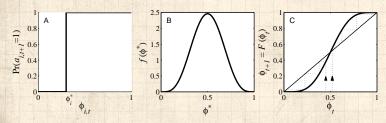
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### Granovetter's model (recap of recap)

Action based on perceived behavior of others.



Two states: S and I.

Recovery now possible (SIS).

 $\Leftrightarrow \phi$  = fraction of contacts 'on' (e.g., rioting).

Discrete time, synchronous update.

This is a Critical mass model.

Interdependent interaction model.

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Disease models assume independence of infectious events.

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Disease models assume independence of infectious events.

Threshold models only involve proportions:  $3/10 \equiv 30/100$ .

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- Disease models assume independence of infectious events.
- Threshold models only involve proportions:  $3/10 \equiv 30/100$ .
- Threshold models ignore exact sequence of influences

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- Disease models assume independence of infectious events.
- Threshold models only involve proportions:  $3/10 \equiv 30/100$ .
- Threshold models ignore exact sequence of influences
- Threshold models assume immediate polling.

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- Disease models assume independence of infectious events.
- Threshold models only involve proportions:  $3/10 \equiv 30/100$ .
- Threshold models ignore exact sequence of influences
- Threshold models assume immediate polling.
- Mean-field models neglect network structure

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- Disease models assume independence of infectious events.
- Threshold models only involve proportions:  $3/10 \equiv 30/100$ .
- Threshold models ignore exact sequence of influences
- Threshold models assume immediate polling.
- Mean-field models neglect network structure
- Network effects only part of story: media, advertising, direct marketing.

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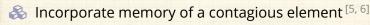
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### Basic ingredients:



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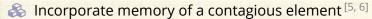
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### Basic ingredients:



 $\red{\$}$  Population of N individuals, each in state S, I, or R.

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### Basic ingredients:

Incorporate memory of a contagious element [5, 6]

 $\ensuremath{\mathfrak{S}}$  Population of N individuals, each in state S, I, or R.

Each individual randomly contacts another at each time step. The PoCSverse Generalized Contagion 17 of 65

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### Basic ingredients:

- Incorporate memory of a contagious element [5, 6]
- $\ensuremath{\mathfrak{S}}$  Population of N individuals, each in state S, I, or R.
- Each individual randomly contacts another at each time step.
- $\phi_t$  = fraction infected at time t = probability of <u>contact</u> with infected individual

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### Basic ingredients:

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- $\ensuremath{\mathfrak{S}}$  With probability p, contact with infective leads to an exposure.

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### Basic ingredients:

- Incorporate memory of a contagious element [5, 6]
- $\ensuremath{\mathfrak{S}}$  Population of N individuals, each in state S, I, or R.
- Each individual randomly contacts another at each time step.
- $\phi_t$  = fraction infected at time t = probability of <u>contact</u> with infected individual
- & With probability p, contact with infective leads to an exposure.
- $\Re$  If exposed, individual receives a dose of size d drawn from distribution f. Otherwise d=0.

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#### Generalized Model

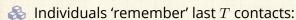
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$$D_{t,i} = \sum_{t'=t-T+1}^t d_i(t')$$

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 $\clubsuit$  Individuals 'remember' last T contacts:

$$D_{t,i} = \sum_{t'=t-T+1}^t d_i(t')$$

Infection occurs if individual i's 'threshold' is exceeded:

$$D_{t,i} \geq d_i^*$$

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brace Individuals 'remember' last T contacts:

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Infection occurs if individual i's 'threshold' is exceeded:

$$D_{t,i} \ge d_i^*$$

 $\Leftrightarrow$  Threshold  $d_i^*$  drawn from arbitrary distribution g at t=0.

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When  $D_{t,i} < d_i^*$ , individual i recovers to state R with probability r.

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When  $D_{t,i} < d_i^*$ , individual i recovers to state R with probability r.



Once in state R, individuals become susceptible again with probability  $\rho$ .

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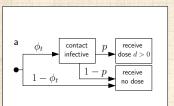
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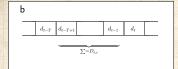
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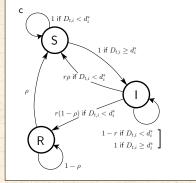
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### A visual explanation







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Study SIS-type contagion first:

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### Study SIS-type contagion first:

Recovered individuals are immediately susceptible again:

 $\rho = 1$ .

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#### Study SIS-type contagion first:

Recovered individuals are immediately susceptible again:

$$\rho = 1$$
.

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### Study SIS-type contagion first:

Recovered individuals are immediately susceptible again:

$$\rho = 1$$
.

- & Look for steady-state behavior as a function of exposure probability p.
- & Denote fixed points by  $\phi^*$ .

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#### Study SIS-type contagion first:

Recovered individuals are immediately susceptible again:

$$\rho = 1$$
.

- Solution Look for steady-state behavior as a function of exposure probability p.
- $\ensuremath{\&}$  Denote fixed points by  $\phi^*$ .

Homogeneous version:

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#### Study SIS-type contagion first:

Recovered individuals are immediately susceptible again:

$$\rho = 1$$
.

- & Look for steady-state behavior as a function of exposure probability p.
- $\ensuremath{\&}$  Denote fixed points by  $\phi^*$ .

#### Homogeneous version:

 $\clubsuit$  All individuals have threshold  $d^*$ 

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### Study SIS-type contagion first:

Recovered individuals are immediately susceptible again:

$$\rho = 1$$
.

- & Look for steady-state behavior as a function of exposure probability p.
- $\red{\$}$  Denote fixed points by  $\phi^*$ .

#### Homogeneous version:

 $\triangle$  All individuals have threshold  $d^*$ 

 $\clubsuit$  All dose sizes are equal: d=1

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

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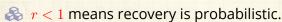
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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



r < 1 means recovery is probabilistic.

T = 1 means individuals forget past interactions.

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



r < 1 means recovery is probabilistic.



T = 1 means individuals forget past interactions.



 $d^* = 1$  means one positive interaction will infect an individual.

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



r < 1 means recovery is probabilistic.



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Evolution of infection level:

$$\phi_{t+1} =$$

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



r < 1 means recovery is probabilistic.



T = 1 means individuals forget past interactions.



 $d^* = 1$  means one positive interaction will infect an individual.



Evolution of infection level:

$$\phi_{t+1} = \underbrace{p\phi_t}_{\mathsf{a}}$$

a: Fraction infected between t and t+1, independent of past state or recovery. The PoCSverse Generalized Contagion 23 of 65

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



r < 1 means recovery is probabilistic.



T = 1 means individuals forget past interactions.



 $d^* = 1$  means one positive interaction will infect an individual.



Evolution of infection level:

$$\phi_{t+1} = \underbrace{p\phi_t}_{\mathsf{a}} + \underbrace{\phi_t(1 - p\phi_t)}_{\mathsf{b}}$$

- a: Fraction infected between t and t+1, independent of past state or recovery.
- b: Probability of being infected and not being reinfected.

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:



r < 1 means recovery is probabilistic.



T = 1 means individuals forget past interactions.



 $d^* = 1$  means one positive interaction will infect an individual.



Evolution of infection level:

$$\phi_{t+1} = \underbrace{p\phi_t}_{\mathsf{a}} + \underbrace{\phi_t(1-p\phi_t)}_{\mathsf{b}} \underbrace{(1-r)}_{\mathsf{C}}.$$

- a: Fraction infected between t and t+1, independent of past state or recovery.
- b: Probability of being infected and not being reinfected.
- c: Probability of not recovering.

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

$$\Leftrightarrow$$
 Set  $\phi_t = \phi^*$ :

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

$$\phi^*=p\phi^*+(1-p\phi^*)\phi^*(1-r)$$

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

$$\clubsuit$$
 Set  $\phi_t = \phi^*$ :

$$\phi^* = p\phi^* + (1 - p\phi^*)\phi^*(1 - r)$$

$$\Rightarrow 1=p+(1-p\phi^*)(1-r), \quad \phi^*\neq 0,$$

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

$$\phi^* = p\phi^* + (1 - p\phi^*)\phi^*(1 - r)$$

$$\Rightarrow 1=p+(1-p\phi^*)(1-r), \quad \phi^*\neq 0,$$

$$\Rightarrow \phi^* = \frac{1 - r/p}{1 - r}$$
 and  $\phi^* = 0$ .

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

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 $\red {\red {\Bbb S}}$  Critical point at  $p=p_c=r$ .

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

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 and  $\phi^* = 0$ .

 $\red{solution}$  Critical point at  $p=p_c=r$ .

Spreading takes off if p/r > 1

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Fixed points for r < 1,  $d^* = 1$ , and T = 1:

$$\clubsuit$$
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$$\phi^* = p\phi^* + (1 - p\phi^*)\phi^*(1 - r)$$

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$$\Rightarrow \phi^* = rac{1 - r/p}{1 - r}$$
 and  $\phi^* = 0$ .

$$\red{shift}$$
 Critical point at  $p=p_c=r$ .

$$\clubsuit$$
 Spreading takes off if  $p/r > 1$ 

Find continuous phase transition as for SIR model.

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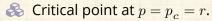
Fixed points for r < 1,  $d^* = 1$ , and T = 1:

$$\clubsuit$$
 Set  $\phi_t = \phi^*$ :

$$\phi^*=p\phi^*+(1-p\phi^*)\phi^*(1-r)$$

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 $\clubsuit$  Spreading takes off if p/r > 1

Find continuous phase transition as for SIR model.

& Goodness: Matches  $R_o = \beta/\gamma > 1$  condition.

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Fixed points for r=1,  $d^*=1$ , and T>1

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Fixed points for r = 1,  $d^* = 1$ , and T > 1



r = 1 means recovery is immediate.

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Fixed points for r = 1,  $d^* = 1$ , and T > 1



r = 1 means recovery is immediate.



T > 1 means individuals remember at least 2 interactions.

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#### Fixed points for r = 1, $d^* = 1$ , and T > 1



r = 1 means recovery is immediate.



T > 1 means individuals remember at least 2 interactions.



 $d^* = 1$  means only one positive interaction in past T interactions will infect individual.

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#### Fixed points for r = 1, $d^* = 1$ , and T > 1

- $rac{1}{4}$  means recovery is immediate.
- T > 1 means individuals remember at least 2 interactions.
- $d^* = 1$  means only one positive interaction in past T interactions will infect individual.
- Effect of individual interactions is independent from effect of others.

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#### Fixed points for r = 1, $d^* = 1$ , and T > 1

- r = 1 means recovery is immediate.
- T > 1 means individuals remember at least 2 interactions.
- $d^* = 1$  means only one positive interaction in past T interactions will infect individual.
- Effect of individual interactions is independent from effect of others.
- $\Leftrightarrow$  Call  $\phi^*$  the steady state level of infection.

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#### Fixed points for r = 1, $d^* = 1$ , and T > 1

- $rac{1}{4}$  means recovery is immediate.
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- $\Leftrightarrow$  Call  $\phi^*$  the steady state level of infection.
- Pr(infected) = 1 Pr(uninfected):

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#### Fixed points for r = 1, $d^* = 1$ , and T > 1

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- $\Leftrightarrow$  Call  $\phi^*$  the steady state level of infection.
- Pr(infected) = 1 Pr(uninfected):

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

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Fixed points for r = 1,  $d^* = 1$ , and T > 1



& Closed form expression for  $\phi^*$ :

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

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Fixed points for r = 1,  $d^* = 1$ , and T > 1

& Closed form expression for  $\phi^*$ :

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

 $\red {\Bbb R}$  Look for critical infection probability  $p_c.$ 

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Fixed points for r = 1,  $d^* = 1$ , and T > 1

& Closed form expression for  $\phi^*$ :

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

& Look for critical infection probability  $p_c$ .

 $\clubsuit$  As  $\phi^* \to 0$ , we see

$$\phi^* \simeq pT\phi^*$$

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Fixed points for r = 1,  $d^* = 1$ , and T > 1

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$$\phi^* \simeq pT\phi^* \Rightarrow p_c = 1/T.$$

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Fixed points for r = 1,  $d^* = 1$ , and T > 1

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🙈 Again find continuous phase transition ...

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Fixed points for r = 1,  $d^* = 1$ , and T > 1

& Closed form expression for  $\phi^*$ :

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

- & Look for critical infection probability  $p_c$ .
- $\clubsuit$  As  $\phi^* \to 0$ , we see

$$\phi^* \simeq pT\phi^* \Rightarrow p_c = 1/T.$$

- Again find continuous phase transition ...
- $\red{ }$  Note: we can solve for p but not  $\phi^*$ :

$$p = (\phi^*)^{-1}[1 - (1 - \phi^*)^{1/T}].$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

 $\clubsuit$  Start with r=1,  $d^*=1$ , and  $T\geq 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Start with r=1,  $d^*=1$ , and  $T\geq 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

 $\clubsuit$  For r < 1, add to right hand side fraction who:

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Start with r=1,  $d^*=1$ , and  $T\geq 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

For r < 1, add to right hand side fraction who: 1. Did not receive any infections in last T time steps, The PoCSverse Generalized Contagion 27 of 65

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Start with r=1,  $d^*=1$ , and  $T\geq 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

- $\clubsuit$  For r < 1, add to right hand side fraction who:
  - 1. Did not receive any infections in last T time steps,
  - 2. And did not recover from a previous infection.

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Start with r=1,  $d^*=1$ , and  $T\geq 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

- $\clubsuit$  For r < 1, add to right hand side fraction who:
  - 1. Did not receive any infections in last T time steps,
  - 2. And did not recover from a previous infection.
- Define corresponding dose histories. Example:

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Start with r=1,  $d^*=1$ , and  $T\geq 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

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  - 1. Did not receive any infections in last T time steps,
  - 2. And did not recover from a previous infection.
- Define corresponding dose histories. Example:

$$H_1 = \{\dots, d_{t-T-2}, d_{t-T-1}, 1, \underbrace{0, 0, \dots, 0, 0}_{T \text{ 0's}}\},$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Start with r = 1,  $d^* = 1$ , and  $T \ge 1$  case we have just examined:

$$\phi^* = 1 - (1 - p\phi^*)^T.$$

- $\clubsuit$  For r < 1, add to right hand side fraction who:
  - 1. Did not receive any infections in last T time steps,
  - 2. And did not recover from a previous infection.
- Define corresponding dose histories. Example:

$$H_1 = \{\dots, d_{t-T-2}, d_{t-T-1}, 1, \underbrace{0, 0, \dots, 0, 0}_{T \text{ 0's}}\},\$$

With history  $H_1$ , probability of being infected (not recovering in one time step) is 1-r.

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

In general, relevant dose histories are:

$$H_{m+1} = \{\dots, d_{t-T-m-1}, 1, \underbrace{0, 0, \dots, 0, 0}_{m \text{ 0's}}, \underbrace{0, 0, \dots, 0, 0}_{T \text{ 0's}}\}.$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

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Overall probabilities for dose histories occurring:

$$P(H_1) = p\phi^*(1 - p\phi^*)^T(1 - r),$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

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Overall probabilities for dose histories occurring:

$$P(H_1) = p \phi^* (1 - p \phi^*)^T (1 - r),$$

$$P(H_{m+1}) = \underbrace{p\phi^*}_a$$

a: Pr(infection T + m + 1 time steps ago)

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

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Overall probabilities for dose histories occurring:

$$P(H_1) = p \phi^* (1 - p \phi^*)^T (1 - r),$$

$$P(H_{m+1}) = \underbrace{p\phi^*}_{a} \underbrace{(1 - p\phi^*)^{T+m}}_{b}$$

a: Pr(infection T + m + 1 time steps ago)

b: Pr(no doses received in T+m time steps since)

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

In general, relevant dose histories are:

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$$P(H_{m+1}) = \underbrace{p\phi^*}_{a} \underbrace{(1 - p\phi^*)^{T+m}}_{b} \underbrace{(1 - r)^{m+1}}_{c}.$$

a: Pr(infection T + m + 1 time steps ago)

b: Pr(no doses received in T + m time steps since)

c:  $Pr(no\ recovery\ in\ m\ chances)$ 

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

 $\Re$  Pr(recovery) = Pr(seeing no doses for at least T time steps and recovering)

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

 $\ref{Pr}$  Pr(recovery) = Pr(seeing no doses for at least T time steps and recovering)

$$= \mathop{r}\limits_{m=0}^{\infty} P(H_{T+m})$$

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 $\ref{Pr}$  Pr(recovery) = Pr(seeing no doses for at least T time steps and recovering)

$$= \mathop{r} \sum_{m=0}^{\infty} P(H_{T+m}) = \mathop{r} \sum_{m=0}^{\infty} p \phi^* (1 - p \phi^*)^{T+m} (1 - r)^m$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Pr(recovery) = Pr(seeing no doses for at least T time steps and recovering)

$$\begin{split} &= \mathop{r} \sum_{m=0}^{\infty} P(H_{T+m}) = \mathop{r} \sum_{m=0}^{\infty} p \phi^* (1 - p \phi^*)^{T+m} (1 - r)^m \\ &= \mathop{r} \frac{p \phi^* (1 - p \phi^*)^T}{1 - (1 - p \phi^*) (1 - r)}. \end{split}$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

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Using the probability of not recovering, we end up with a fixed point equation:

$$\phi^* = 1 - \frac{r(1 - p\phi^*)^T}{1 - (1 - p\phi^*)(1 - r)}.$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 



Fixed point equation (again):

$$\phi^* = 1 - \frac{r(1 - p\phi^*)^T}{1 - (1 - p\phi^*)(1 - r)}.$$

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Fixed point equation (again):

$$\phi^* = 1 - \frac{r(1 - p\phi^*)^T}{1 - (1 - p\phi^*)(1 - r)}.$$

 $\Longrightarrow$  Find critical exposure probability by examining above as  $\phi^* \to 0$ .

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

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$$\Rightarrow \quad p_c = \frac{1}{T + 1/r - 1} = \frac{1}{T + \tau}.$$

where  $\tau$  = mean recovery time for simple relaxation process.

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Fixed points for  $r \le 1$ ,  $d^* = 1$ , and  $T \ge 1$ 

Fixed point equation (again):

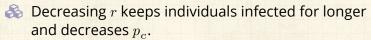
$$\phi^* = 1 - \frac{r(1 - p\phi^*)^T}{1 - (1 - p\phi^*)(1 - r)}.$$

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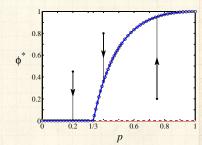
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### Epidemic threshold:

Fixed points for  $d^* = 1$ ,  $r \le 1$ , and  $T \ge 1$ 





 $\clubsuit$  Example details:  $T=2 \& r=1/2 \Rightarrow p_c=1/3$ .



Blue = stable, red = unstable, fixed points.



 $\Rightarrow \tau = 1/r - 1$  = characteristic recovery time = 1.



 $T + \tau \simeq \text{average memory in system} = 3.$ 

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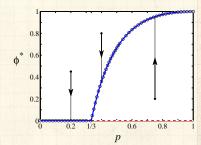
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### Epidemic threshold:

Fixed points for  $d^* = 1$ ,  $r \le 1$ , and  $T \ge 1$ 





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Blue = stable, red = unstable, fixed points.



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 $T + \tau \simeq$  average memory in system = 3.

Phase transition can be seen as a transcritical bifurcation. [12]

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All right:  $d^* = 1$  models correspond to simple disease spreading models.

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All right:  $d^* = 1$  models correspond to simple disease spreading models.

 $\clubsuit$  What if we allow  $d^* \geq 2$ ?

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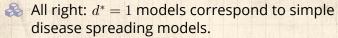
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 $\clubsuit$  What if we allow  $d^* \geq 2$ ?

Again first consider SIS with immediate recovery (r = 1)

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All right:  $d^* = 1$  models correspond to simple disease spreading models.

Again first consider SIS with immediate recovery (r = 1)

Also continue to assume unit dose sizes  $(f(d) = \delta(d-1))$ .

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 $\ref{Solution}$  To be infected, must have at least  $d^*$  exposures in last T time steps.

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Again first consider SIS with immediate recovery (r = 1)

Also continue to assume unit dose sizes  $(f(d) = \delta(d-1))$ .

 $lap{3}{\ }$  To be infected, must have at least  $d^*$  exposures in last T time steps.

Fixed point equation:

$$\phi^* = \sum_{i=d^*}^T \binom{T}{i} (p\phi^*)^i (1 - p\phi^*)^{T-i}.$$

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Fixed point equation:

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 $\clubsuit$  As always,  $\phi^* = 0$  works too.

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Fixed points for r=1,  $d^*>1$ , and  $T\geq 1$ 

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Fixed points for r = 1,  $d^* > 1$ , and  $T \ge 1$ 



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Fixed points for r = 1,  $d^* > 1$ , and T > 1



& e.g., for  $d^* = 2$ , T = 3:

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Fixed points for r = 1,  $d^* > 1$ , and T > 1



& Exactly solvable for small T.

\$ e.g., for  $d^* = 2$ , T = 3:



Fixed point equation:

$$\begin{array}{l} \phi^* = \\ 3p^2 {\phi^*}^2 (1-p\phi^*) + p^3 {\phi^*}^3 \end{array}$$

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Fixed points for r = 1,  $d^* > 1$ , and T > 1



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Fixed point equation:

$$\phi^* = 3p^2 \phi^{*2} (1 - p\phi^*) + p^3 \phi^{*3}$$

See new structure: a saddle node bifurcation [12] appears as p increases.

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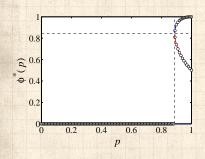


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Fixed point equation:

$$\phi^* = 3p^2 \phi^{*2} (1 - p\phi^*) + p^3 \phi^{*3}$$

as p increases.

$$(p_b, \phi^*) = (8/9, 27/32).$$

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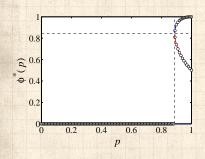
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Fixed points for r=1,  $d^*>1$ , and  $T\geq 1$ 

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8

Fixed point equation:  $\phi^* = \frac{3n^2 \phi^{*2} (1 - n\phi^*) + n^3 \phi^*}{1 + n^3 \phi^*}$ 

$$3p^2\phi^{*2}(1-p\phi^*)+p^3\phi^{*3}$$

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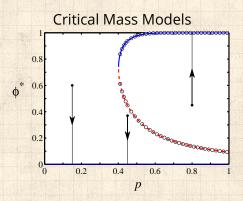
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Behavior akin to output of Granovetter's threshold model.

Another example:



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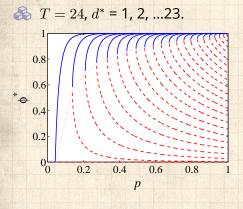
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$$r = 1, d^* = 3, T = 12$$

# Fixed points for r=1, $d^*>1$ , and $T\geq 1$



 $d^* = 1 \rightarrow d^* > 1$ : jump between continuous phase transition and pure critical

&

Unstable curve for  $d^* = 2$  does not hit  $\phi^* = 0$ .

mass model.

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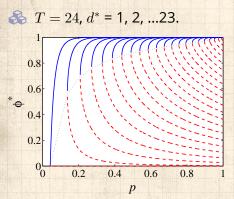
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### Fixed points for r=1, $d^*>1$ , and $T\geq 1$



 $d^* = 1 \rightarrow d^* > 1$ : jump between continuous phase transition and pure critical mass model.

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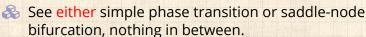
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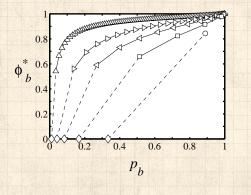




# Fixed points for r=1, $d^*>1$ , and $T\geq 1$



 $\Longrightarrow$  Bifurcation points for example fixed T, varying  $d^*$ :



$$3 T = 96$$
 ( ).

$$\Rightarrow T = 24 (\triangleright),$$

$$3$$
  $T = 12 (<),$ 

$$3$$
  $T=6 (\square),$ 

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 For r < 1, need to determine probability of recovering as a function of time since dose load last dropped below threshold.

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For r < 1, need to determine probability of recovering as a function of time since dose load last dropped below threshold.

Partially summed random walks:

$$D_i(t) = \sum_{t'=t-T+1}^t d_i(t')$$

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**Solution** Example for T = 24,  $d^* = 14$ :

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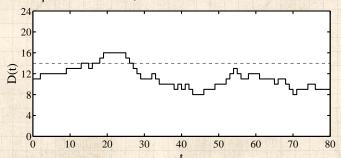


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 $\bigotimes$  Define  $\gamma_m$  as fraction of individuals for whom D(t)last equaled, and has since been below, their threshold m time steps ago,

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 $\begin{align*}{l} \end{align*} \begin{align*}{l} \end{align*$ 

Fraction of individuals below threshold but not recovered:

$$\Gamma(p,\phi^*;r) = \sum_{m=1}^{\infty} (1-r)^m \gamma_m(p,\phi^*).$$

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 $\begin{align*}{l} \end{align*} \begin{align*}{l} \end{align*$ 

Fraction of individuals below threshold but not recovered:

$$\Gamma(p,\phi^*;r) = \sum_{m=1}^{\infty} (1-r)^m \gamma_m(p,\phi^*).$$

Fixed point equation:

$$\phi^* = \Gamma(p, \phi^*; r) + \sum_{i=d^*}^T \binom{T}{i} (p\phi^*)^i (1 - p\phi^*)^{T-i}.$$

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Example: T = 3,  $d^* = 2$ 

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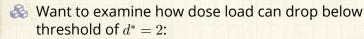
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Example: T = 3,  $d^* = 2$ 



$$D_n=2\Rightarrow D_{n+1}=1$$

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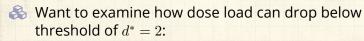
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Example: T = 3,  $d^* = 2$ 



$$D_n = 2 \Rightarrow D_{n+1} = 1$$

Two subsequences do this:

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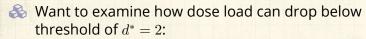
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Example: T = 3,  $d^* = 2$ 



$$D_n = 2 \Rightarrow D_{n+1} = 1$$

Two subsequences do this:

$$\{d_{n-2}, d_{n-1}, d_n, d_{n+1}\} = \{1, 1, 0, {\color{red}0}\}$$

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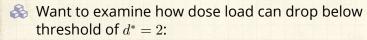
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Example: T = 3,  $d^* = 2$ 



$$D_n = 2 \Rightarrow D_{n+1} = 1$$

Two subsequences do this:

$$\begin{aligned} \{d_{n-2}, d_{n-1}, d_n, d_{n+1}\} &= \{1, 1, 0, \textcolor{red}{0}\} \\ \text{and } \{d_{n-2}, d_{n-1}, d_n, d_{n+1}, d_{n+2}\} &= \{1, 0, 1, \textcolor{red}{0}, \textcolor{red}{0}\}. \end{aligned}$$

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Example: T = 3,  $d^* = 2$ 

Want to examine how dose load can drop below threshold of  $d^* = 2$ :

$$D_n = 2 \Rightarrow D_{n+1} = 1$$

Two subsequences do this:

$$\begin{aligned} \{d_{n-2},d_{n-1},d_n,d_{n+1}\} &= \{1,1,0,\textcolor{red}{\mathbf{0}}\}\\ \text{and } \{d_{n-2},d_{n-1},d_n,d_{n+1},d_{n+2}\} &= \{1,0,1,\textcolor{red}{\mathbf{0}},\textcolor{red}{\mathbf{0}}\}. \end{aligned}$$

Note: second sequence includes an extra 0 since this is necessary to stay below  $d^* = 2$ .

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Example: T = 3,  $d^* = 2$ 

Want to examine how dose load can drop below threshold of  $d^* = 2$ :

$$D_n = 2 \Rightarrow D_{n+1} = 1$$

Two subsequences do this:

$$\begin{aligned} \{d_{n-2}, d_{n-1}, d_n, d_{n+1}\} &= \{1, 1, 0, \textcolor{red}{0}\} \\ \text{and } \{d_{n-2}, d_{n-1}, d_n, d_{n+1}, d_{n+2}\} &= \{1, 0, 1, \textcolor{red}{0}, \textcolor{red}{0}\}. \end{aligned}$$

- Note: second sequence includes an extra 0 since this is necessary to stay below  $d^* = 2$ .
- To stay below threshold, observe acceptable following sequences may be composed of any combination of two subsequences:

$$a = \{0\}$$
 and  $b = \{1, 0, 0\}.$ 

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keep dose load below  $d^* = 2$ .

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ightharpoonup Determine number of sequences of length m that keep dose load below  $d^*=2$ .

 $N_a$  = number of  $a = \{0\}$  subsequences.

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ightharpoonup Determine number of sequences of length m that keep dose load below  $d^*=2$ .

 $\begin{cases} \&\ N_a = \mbox{number of } a = \{0\} \mbox{ subsequences.} \end{cases}$ 

 $\aleph$   $N_b$  = number of  $b = \{1, 0, 0\}$  subsequences.

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 $\red$  Determine number of sequences of length m that keep dose load below  $d^*=2$ .

 $\begin{tabular}{ll} \& N_a = \mbox{number of } a = \{0\} \mbox{ subsequences.} \end{tabular}$ 

 $\aleph$   $N_b$  = number of  $b = \{1, 0, 0\}$  subsequences.

$$m = N_a \cdot 1 + N_b \cdot 3$$

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 $lap{Rel}$  Determine number of sequences of length m that keep dose load below  $d^*=2$ .

 $\begin{tabular}{ll} \& N_a = \mbox{number of } a = \{0\} \mbox{ subsequences.} \end{tabular}$ 

 $\Re N_b$  = number of  $b = \{1, 0, 0\}$  subsequences.

$$m = N_a \cdot 1 + N_b \cdot 3$$

Possible values for  $N_b$ :

$$0,1,2,\ldots,\left\lfloor \frac{m}{3} \right\rfloor.$$

where | | means floor.

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 $lap{Rel}$  Determine number of sequences of length m that keep dose load below  $d^*=2$ .

 $\begin{cases} \&\ N_a = \mbox{number of } a = \{0\} \mbox{ subsequences.} \end{cases}$ 

 $\Re N_b$  = number of  $b = \{1, 0, 0\}$  subsequences.

$$m = N_a \cdot 1 + N_b \cdot 3$$

Possible values for  $N_b$ :

$$0,1,2,\ldots,\left\lfloor \frac{m}{3} \right\rfloor.$$

where | | means floor.

& Corresponding possible values for  $N_a$ :

$$m, m-3, m-6, \ldots, m-3 \left\lfloor \frac{m}{3} \right\rfloor$$
.

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 $\Longrightarrow$  How many ways to arrange  $N_a$  a's and  $N_b$  b's?

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 $\Re$  How many ways to arrange  $N_a$  a's and  $N_b$  b's? Think of overall sequence in terms of subsequences:

$$\{Z_1,Z_2,\dots,Z_{N_a+N_b}\}$$

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 $\Re$  How many ways to arrange  $N_a$  a's and  $N_b$  b's?



Think of overall sequence in terms of subsequences:

$$\{Z_1,Z_2,\dots,Z_{N_a+N_b}\}$$



 $N_a + N_b$  slots for subsequences.

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 $\red{\$}$  How many ways to arrange  $N_a~a'$ s and  $N_b~b'$ s?

Think of overall sequence in terms of subsequences:

$$\{Z_1,Z_2,\dots,Z_{N_a+N_b}\}$$

 $N_a + N_b$  slots for subsequences.

& Choose positions of either a's or b's:

$$\binom{N_a+N_b}{N_a}=\binom{N_a+N_b}{N_b}.$$

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3 Total number of allowable sequences of length m:

$$\sum_{N_b=0}^{\lfloor m/3\rfloor} \binom{N_b+N_a}{N_b} = \sum_{k=0}^{\lfloor m/3\rfloor} \binom{m-2k}{k}$$

where  $k = N_h$  and we have used  $m = N_a + 3N_h$ .

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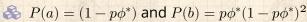




 $\clubsuit$  Total number of allowable sequences of length m:

$$\sum_{N_b=0}^{\lfloor m/3\rfloor} \binom{N_b+N_a}{N_b} = \sum_{k=0}^{\lfloor m/3\rfloor} \binom{m-2k}{k}$$

where  $k = N_b$  and we have used  $m = N_a + 3N_b$ .



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 $\mathfrak{S}$  Total number of allowable sequences of length m:

$$\sum_{N_b=0}^{\lfloor m/3\rfloor} \binom{N_b+N_a}{N_b} = \sum_{k=0}^{\lfloor m/3\rfloor} \binom{m-2k}{k}$$

where  $k = N_b$  and we have used  $m = N_a + 3N_b$ .

$$P(a) = (1 - p\phi^*) \text{ and } P(b) = p\phi^*(1 - p\phi^*)^2$$

 $\Re$  Total probability of allowable sequences of length m:

$$\chi_m(p,\phi^*) = \sum_{k=0}^{\lfloor m/3 \rfloor} \binom{m-2k}{k} (1-p\phi^*)^{m-k} (p\phi^*)^k.$$

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 $\mathfrak{S}$  Total number of allowable sequences of length m:

$$\sum_{N_b=0}^{\lfloor m/3\rfloor} \binom{N_b+N_a}{N_b} = \sum_{k=0}^{\lfloor m/3\rfloor} \binom{m-2k}{k}$$

where  $k=N_b$  and we have used  $m=N_a+3N_b$ .

$$Rrac{1}{8} P(a) = (1 - p\phi^*) \text{ and } P(b) = p\phi^*(1 - p\phi^*)^2$$

 $\Re$  Total probability of allowable sequences of length m:

$$\chi_m(p,\phi^*) = \sum_{k=0}^{\lfloor m/3 \rfloor} \binom{m-2k}{k} (1-p\phi^*)^{m-k} (p\phi^*)^k.$$

Notation: Write a randomly chosen sequence of a's and b's of length m as  $D_m^{a,b}$ .

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- Nearly there ...must account for details of sequence endings.
- Three endings ⇒ Six possible sequences:

$$D_1 = \{1, 1, 0, 0, D_{m-1}^{a,b}\}$$

$$D_2 = \{1, 1, 0, 0, D_{m-2}^{a,b}, 1\}$$

$$D_3 = \{1, 1, 0, 0, D_{m-3}^{a,b}, 1, 0\}$$

$$D_4 = \{1, 0, 1, 0, 0, D_{m-2}^{a,b}\}$$

$$D_5 = \{1, 0, 1, 0, 0, D_{m-3}^{a,b}, 1\}$$

$$D_6 = \{1, 0, 1, 0, 0, D_{m-4}^{a,b}, 1, 0\}$$

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- Nearly there ...must account for details of sequence endings.
- Three endings ⇒ Six possible sequences:

$$D_1 = \{1, 1, 0, 0, D_{m-1}^{a,b}\}$$

$$D_2 = \{1, 1, 0, 0, D_{m-2}^{a,b}, 1\}$$

$$D_3 = \{1, 1, 0, 0, D_{m-3}^{a,b}, 1, 0\}$$

$$D_4 = \{1, 0, 1, 0, 0, D_{m-2}^{a,b}\}$$

$$D_5 = \{1, 0, 1, 0, 0, D_{m-3}^{a,b}, 1\}$$

$$D_6 = \{1, 0, 1, 0, 0, D_{m-4}^{a,b}, 1, 0\}$$

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- Nearly there ...must account for details of sequence endings.

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$$\begin{split} D_2 &= \{1,1,0,0,D_{m-2}^{a,b},1\} \\ D_3 &= \{1,1,0,0,D_{m-3}^{a,b},1,0\} \end{split} \qquad P_2 = (p\phi)^3(1-p\phi)^2\chi_{m-2}(p,\phi) \end{split}$$

$$P_3 = (p\phi)^3 (1 - p\phi)^3 \chi_{m-3}(p,\phi)$$

$$(p\phi)^3(1-p\phi)$$

 $P_1 = (p\phi)^2 (1 - p\phi)^2 \chi_{m-1}(p, \phi)$ 

$$P_4 = (p\phi)^2 (1-p\phi)^3 \chi_{m-2}(p,\phi) \qquad \qquad \text{PoCS}$$

$$P_5 = (p\phi)^3 (1 - p\phi)^3$$

$$P_5 = (p\phi)^3 (1-p\phi)^3 \chi_{m-3}(p,\phi)$$

$$D_6 = \{1, 0, 1, 0, 0, D_{m-4}^{a,b}, 1, 0\}$$

 $D_5 = \{1, 0, 1, 0, 0, D_{m-3}^{a,b}, 1\}$ 

 $D_4 = \{1, 0, 1, 0, 0, D_{m-2}^{a,b}\}$ 

 $D_1 = \{1, 1, 0, 0, D_{m-1}^{a,b}\}$ 

$$P_6 = (p\phi)^3 (1-p\phi)^4 \chi_{m-4}(p,\phi)$$

$$\text{F.P. Eq: } \phi^* = \Gamma(p,\phi^*;r) + \sum_{i=d^*}^T \binom{T}{i} (p\phi^*)^i (1-p\phi^*)^{T-i}.$$

where  $\Gamma(p, \phi^*; r) =$ 

$$(1-r)(p\phi)^{2}(1-p\phi)^{2} + \sum_{m=1}^{\infty} (1-r)^{m}(p\phi)^{2}(1-p\phi)^{2} \times$$

$$\left[\chi_{m-1} + \chi_{m-2} + 2p\phi(1-p\phi)\chi_{m-3} + p\phi(1-p\phi)^2\chi_{m-4}\right]$$

and

$$\chi_m(p,\phi^*) = \sum_{k=0}^{\lfloor m/3 \rfloor} \binom{m-2k}{k} (1-p\phi^*)^{m-k} (p\phi^*)^k.$$

Note:  $(1-r)(p\phi)^2(1-p\phi)^2$  accounts for  $\{1,0,1,0\}$  sequence.

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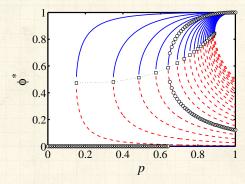
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$$T=3, d^*=2$$



 $r = 0.01, 0.05, 0.10, 0.15, 0.20, \dots, 1.00.$ 

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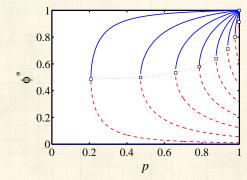
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$$T=2, d^*=2$$



 $r = 0.01, 0.05, 0.10, \dots, 0.3820 \pm 0.0001.$ 

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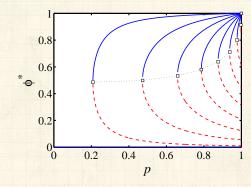
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$$T=2, d^*=2$$



 $r = 0.01, 0.05, 0.10, \dots, 0.3820 \pm 0.0001.$ 

Arr No spreading for  $r \gtrsim 0.382$ .

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Two kinds of contagion processes:

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Two kinds of contagion processes:

1. Continuous phase transition: SIR-like.

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#### Two kinds of contagion processes:

- 1. Continuous phase transition: SIR-like.
- 2. Saddle-node bifurcation: threshold model-like.

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Two kinds of contagion processes:

- 1. Continuous phase transition: SIR-like.
- 2. Saddle-node bifurcation: threshold model-like.



 $d^* = 1$ : spreading from small seeds possible.

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Two kinds of contagion processes:

- 1. Continuous phase transition: SIR-like.
- 2. Saddle-node bifurcation: threshold model-like.



 $d^* = 1$ : spreading from small seeds possible.



 $d^* > 1$ : critical mass model.

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Two kinds of contagion processes:

- 1. Continuous phase transition: SIR-like.
- 2. Saddle-node bifurcation: threshold model-like.



 $d^* = 1$ : spreading from small seeds possible.

 $d^* > 1$ : critical mass model.

Are other behaviors possible?

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 $\aleph$  Now allow for general dose distributions (f) and threshold distributions (*q*).

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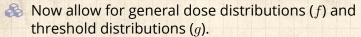
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& Key quantities:

$$P_k = \int_0^\infty \mathrm{d} d^* \, g(d^*) P\left(\sum_{j=1}^k d_j \geq d^*\right) \text{ where } 1 \leq k \leq T.$$

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Now allow for general dose distributions (f) and threshold distributions (a).

Key quantities:

$$P_k = \int_0^\infty \mathrm{d} d^* \, g(d^*) P\left(\sum_{j=1}^k d_j \geq d^*\right) \text{ where } 1 \leq k \leq T.$$

 $P_k$  = Probability that the threshold of a randomly selected individual will be exceeded by k doses.

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Now allow for general dose distributions (f) and threshold distributions (g).

Key quantities:

$$P_k = \int_0^\infty \mathrm{d} d^* \, g(d^*) P\left(\sum_{j=1}^k d_j \geq d^*\right) \text{ where } 1 \leq k \leq T.$$

 $P_k$  = Probability that the threshold of a randomly selected individual will be exceeded by k doses.

ቆ e.g.,

 P<sub>1</sub> = Probability that <u>one dose</u> will exceed the threshold of a random individual
 = Fraction of most vulnerable individuals. The PoCSverse Generalized Contagion 49 of 65

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### Generalized model—heterogeneity, r = 1



Fixed point equation:

$$\phi^* = \sum_{k=1}^T \binom{T}{k} (p\phi^*)^k (1 - p\phi^*)^{T-k} \underline{P_k}$$

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## Generalized model—heterogeneity, r = 1

Fixed point equation:

$$\phi^* = \sum_{k=1}^T \binom{T}{k} (p\phi^*)^k (1 - p\phi^*)^{T-k} \underline{P_k}$$

 $\clubsuit$  Expand around  $\phi^* = 0$  to find when spread from single seed is possible:

$$pP_1T \ge 1$$

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### Generalized model—heterogeneity, r=1

Fixed point equation:

$$\phi^* = \sum_{k=1}^T \binom{T}{k} (p\phi^*)^k (1 - p\phi^*)^{T-k} \underline{P_k}$$

 $\clubsuit$  Expand around  $\phi^* = 0$  to find when spread from single seed is possible:

$$\boxed{pP_1T\geq 1} \qquad \text{or} \qquad \boxed{\Rightarrow p_c=1/(TP_1)}$$

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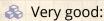
### Generalized model—heterogeneity, r = 1

Fixed point equation:

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 $\clubsuit$  Expand around  $\phi^* = 0$  to find when spread from single seed is possible:

$$pP_1T \ge 1$$
 or  $\Rightarrow p_c = 1/(TP_1)$ 



1.  $P_1T$  is the expected number of vulnerables the initial infected individual meets before recovering. The PoCSverse Generalized Contagion 50 of 65

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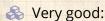
## Generalized model—heterogeneity, r=1

Fixed point equation:

$$\phi^* = \sum_{k=1}^T \binom{T}{k} (p\phi^*)^k (1 - p\phi^*)^{T-k} \underline{P_k}$$

 $\clubsuit$  Expand around  $\phi^* = 0$  to find when spread from single seed is possible:

$$pP_1T \ge 1$$
 or  $\Rightarrow p_c = 1/(TP_1)$ 



- 1.  $P_1T$  is the expected number of vulnerables the initial infected individual meets before recovering.
- 2.  $pP_1T$  is : the expected number of successful infections (equivalent to  $R_0$ ).

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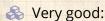
## Generalized model—heterogeneity, r = 1

Fixed point equation:

$$\phi^* = \sum_{k=1}^T \binom{T}{k} (p\phi^*)^k (1 - p\phi^*)^{T-k} \underline{P_k}$$

 $\clubsuit$  Expand around  $\phi^* = 0$  to find when spread from single seed is possible:

$$pP_1T \ge 1$$
 or  $\Rightarrow p_c = 1/(TP_1)$ 



- 1.  $P_1T$  is the expected number of vulnerables the initial infected individual meets before recovering.
- 2.  $pP_1T$  is : the expected number of successful infections (equivalent to  $R_0$ ).
- Observe: p<sub>c</sub> may exceed 1 meaning no spreading from a small seed.

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Next: Determine slope of fixed point curve at critical point  $p_c$ .

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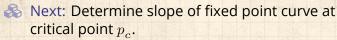
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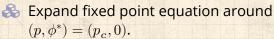
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Next: Determine slope of fixed point curve at critical point  $p_c$ .

& Expand fixed point equation around  $(p, \phi^*) = (p_c, 0)$ .

Find slope depends on  $(P_1 - P_2/2)^{[6]}$  (see Appendix).

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 $\ref{Next:}$  Determine slope of fixed point curve at critical point  $p_c$ .

& Expand fixed point equation around  $(p, \phi^*) = (p_c, 0)$ .

 $\ref{eq:property}$  Find slope depends on  $(P_1-P_2/2)^{[6]}$  (see Appendix).

Behavior near fixed point depends on whether this slope is The PoCSverse Generalized Contagion 51 of 65

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Next: Determine slope of fixed point curve at critical point  $p_c$ .

 $\Leftrightarrow$  Expand fixed point equation around  $(p,\phi^*)=(p_c,0).$ 

 $\ref{Find slope depends on } (P_1-P_2/2)^{[6]}$  (see Appendix).

Behavior near fixed point depends on whether this slope is

1. positive:  $P_1 > P_2/2$  (continuous phase transition)

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 $lap{Next:}$  Determine slope of fixed point curve at critical point  $p_c$ .

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Behavior near fixed point depends on whether this slope is

1. positive:  $P_1 > P_2/2$  (continuous phase transition)

2. negative:  $P_1 < P_2/2$  (discontinuous phase transition)

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 $lap{Next:}$  Determine slope of fixed point curve at critical point  $p_c$ .

 $\Leftrightarrow$  Expand fixed point equation around  $(p,\phi^*)=(p_c,0).$ 

 $\ref{Find slope depends on } (P_1-P_2/2)^{[6]}$  (see Appendix).

Behavior near fixed point depends on whether this slope is

- 1. positive:  $P_1 > P_2/2$  (continuous phase transition)
- 2. negative:  $P_1 < P_2/2$  (discontinuous phase transition)
- Now find three basic universal classes of contagion models ...

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#### Example configuration:



Dose sizes are lognormally distributed with mean 1 and variance 0.433.

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#### Example configuration:

Dose sizes are lognormally distributed with mean 1 and variance 0.433.

 $\clubsuit$  Memory span: T=10.

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#### Example configuration:

Dose sizes are lognormally distributed with mean 1 and variance 0.433.

 $\clubsuit$  Memory span: T=10.

Thresholds are uniformly set at

1.  $d_* = 0.5$ 

2.  $d_* = 1.6$ 

3.  $d_{*}^{*} = 3$ 

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#### Example configuration:

- Dose sizes are lognormally distributed with mean 1 and variance 0.433.
- Thresholds are uniformly set at
  - 1.  $d_* = 0.5$
  - 2.  $d_* = 1.6$
  - 3.  $d_* = 3$
- Spread of dose sizes matters, details are not important.

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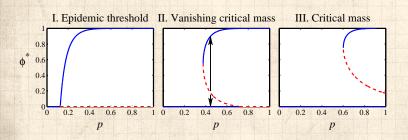
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#### Three universal classes



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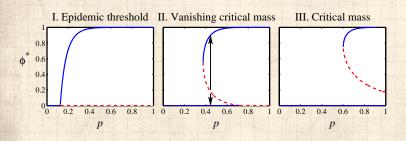
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### Three universal classes

Epidemic threshold:



 $P_1 > P_2/2$ ,  $p_c = 1/(TP_1) < 1$ 

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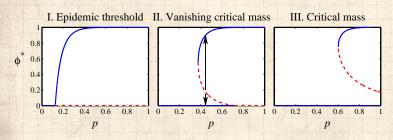
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### Three universal classes



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Epidemic threshold:

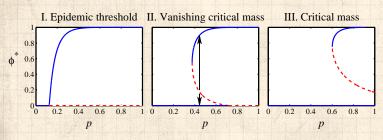
$$P_1 > P_2/2 \text{, } p_c = 1/(TP_1) < 1$$

 $P_1 < P_2/2$ 

Vanishing critical mass:

 $p_c = 1/(TP_1) < 1$ 

### Three universal classes



Epidemic threshold:

$$P_1 > P_2/2 \text{, } p_c = 1/(TP_1) < 1$$

Vanishing critical mass:  $p_c = 1/(TP_1) < 1$ 

$$P_1 < P_2/2$$
,

Pure critical mass:

$$P_1 < P_2/2 \text{, } p_c = 1/(TP_1) > 1$$

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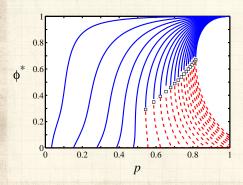
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# Heterogeneous case

### Now allow r < 1:



 II-III transition generalizes:  $p_c=1/[P_1(T+\tau)]$  where  $\tau=1/r-1=$  expected recovery time

🚴 I-II transition less pleasant analytically.

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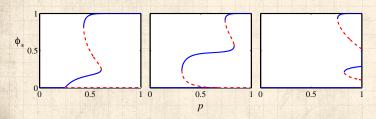
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# More complicated models



🙈 Due to heterogeneity in individual thresholds.

Three classes based on behavior for small seeds.

🙈 Same model classification holds: I, II, and III.

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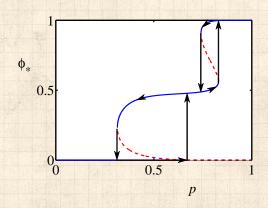
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# Hysteresis in vanishing critical mass models



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Memory is a natural ingredient.

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Memory is a natural ingredient.



Three universal classes of contagion processes:

I. Epidemic Threshold II. Vanishing Critical Mass III. Critical Mass

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Memory is a natural ingredient.



Three universal classes of contagion processes:

I. Epidemic Threshold II. Vanishing Critical Mass III. Critical Mass



Dramatic changes in behavior possible.

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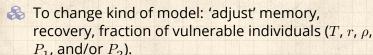
Memory is a natural ingredient.

Three universal classes of contagion processes:

I. Epidemic Threshold II. Vanishing Critical Mass III. Critical Mass



Dramatic changes in behavior possible.



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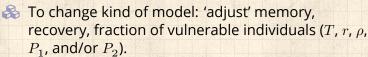
Memory is a natural ingredient.

Three universal classes of contagion processes:

I. Epidemic Threshold II. Vanishing Critical Mass III. Critical Mass



Dramatic changes in behavior possible.



To change behavior given model: 'adjust' probability of exposure (p) and/or initial number infected ( $\phi_0$ ).

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Single seed infects others if  $pP_1(T+\tau) \ge 1$ .

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Single seed infects others if  $pP_1(T+\tau) \geq 1$ .

 $\Re$  Key quantity:  $p_c = 1/[P_1(T+\tau)]$ 

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Single seed infects others if  $pP_1(T+\tau) \geq 1$ .



 $\Re$  Key quantity:  $p_c = 1/[P_1(T+\tau)]$ 



 $\Re$  If  $p_c < 1 \Rightarrow$  contagion can spread from single seed.

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Single seed infects others if  $pP_1(T+\tau) \geq 1$ .



 $\Leftrightarrow$  Key quantity:  $p_c = 1/[P_1(T+\tau)]$ 



If  $p_c < 1 \Rightarrow$  contagion can spread from single seed.



Depends only on:

1. System Memory  $(T + \tau)$ . 2. Fraction of highly vulnerable individuals  $(P_1)$ . The PoCSverse Generalized Contagion 58 of 65

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Single seed infects others if  $pP_1(T+\tau) > 1$ .

 $\Leftrightarrow$  Key quantity:  $p_c = 1/[P_1(T+\tau)]$ 

If  $p_c < 1 \Rightarrow$  contagion can spread from single seed.

Depends only on:

1. System Memory  $(T + \tau)$ .

2. Fraction of highly vulnerable individuals  $(P_1)$ .

Details unimportant: Many threshold and dose distributions give same  $P_{h}$ .

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Single seed infects others if  $pP_1(T+\tau) \ge 1$ .

 $\clubsuit$  If  $p_c < 1 \Rightarrow$  contagion can spread from single seed.

Depends only on:

1. System Memory ( $T + \tau$ ).

2. Fraction of highly vulnerable individuals ( $P_1$ ).

Arr Details unimportant: Many threshold and dose distributions give same  $P_k$ .

Another example of a model where vulnerable/gullible population may be more important than a small group of super-spreaders or influentials.

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$$\begin{split} \phi^* &= \sum_{k=1}^T \binom{T}{k} P_k (p\phi^*)^k (1-p\phi^*)^{T-k}, \\ &= \sum_{k=1}^T \binom{T}{k} P_k (p\phi^*)^k \sum_{j=0}^{T-k} \binom{T-k}{j} (-p\phi^*)^j, \\ &= \sum_{k=1}^T \sum_{j=0}^{T-k} \binom{T}{k} \binom{T-k}{j} P_k (-1)^j (p\phi^*)^{k+j}, \\ &= \sum_{m=1}^T \sum_{k=1}^m \binom{T}{k} \binom{T-k}{m-k} P_k (-1)^{m-k} (p\phi^*)^m, \\ &= \sum_{m=1}^T C_m (p\phi^*)^m \end{split}$$

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$$C_m = (-1)^m \binom{T}{m} \sum_{k=1}^m (-1)^k \binom{m}{k} P_k,$$

since

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Linearization gives

$$\phi^* \simeq C_1 p \phi^* + C_2 p_c^2 {\phi^*}^2.$$

where 
$$C_1=TP_1(=1/p_c)$$
 and  $C_2={T\choose 2}(-2P_1+P_2).$ 

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Linearization gives

$$\phi^* \simeq C_1 p \phi^* + C_2 p_c^2 {\phi^*}^2.$$

where 
$$C_1=TP_1(=1/p_c)$$
 and  $C_2={T\choose 2}(-2P_1+P_2).$ 



 $\Leftrightarrow$  Using  $p_c = 1/(TP_1)$ :

$$\phi^* \simeq \frac{C_1}{C_2 p_c^2} (p-p_c) = \frac{T^2 P_1^3}{(T-1)(P_1-P_2/2)} (p-p_c).$$

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Linearization gives

$$\phi^* \simeq C_1 p \phi^* + C_2 p_c^2 {\phi^*}^2.$$

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Sign of derivative governed by  $P_1 - P_2/2$ .

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