

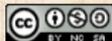
# Generalized Contagion

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Complex Networks | @networksvox  
CSYS/MATH 303, Spring, 2019

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Vermont Advanced Computing Core | University of Vermont



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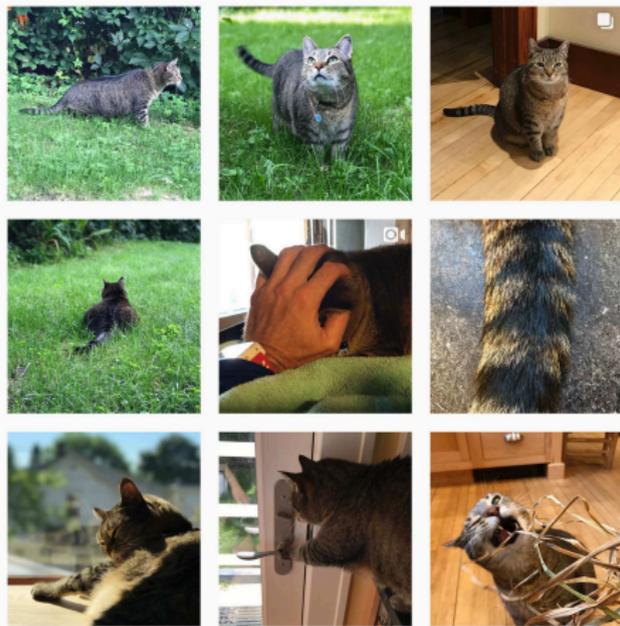


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## Special Guest Executive Producer



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 On Instagram at [pratchett\\_the\\_cat](https://www.instagram.com/pratchett_the_cat) 



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## “Universal Behavior in a Generalized Model of Contagion” ↗

Dodds and Watts,  
Phys. Rev. Lett., **92**, 218701, 2004. [5]



## “A generalized model of social and biological contagion” ↗

Dodds and Watts,  
J. Theor. Biol., **232**, 587–604, 2005. [6]

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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<sup>[11]</sup>

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The standard **SIR model** <sup>[11]</sup>

 = basic model of disease contagion

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The standard **SIR model** <sup>[11]</sup>

 = basic model of disease contagion

 Three states:

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The standard **SIR model** <sup>[11]</sup>

 = basic model of disease contagion

 Three states:

1.  $S$  = Susceptible

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The standard **SIR model** <sup>[11]</sup>

 = basic model of disease contagion

 Three states:

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

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The standard **SIR model** <sup>[11]</sup>

 = 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** <sup>[11]</sup>

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

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

 Presumes random interactions (mass-action principle)

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

 Presumes random interactions (mass-action principle)

 Interactions are independent (no memory)

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## The standard SIR model<sup>[11]</sup>

 = basic model of disease contagion

 Three states:

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3. R = Recovered or Removed or Refractory

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

 Presumes random interactions (mass-action principle)

 Interactions are independent (no memory)

 Discrete and continuous time versions

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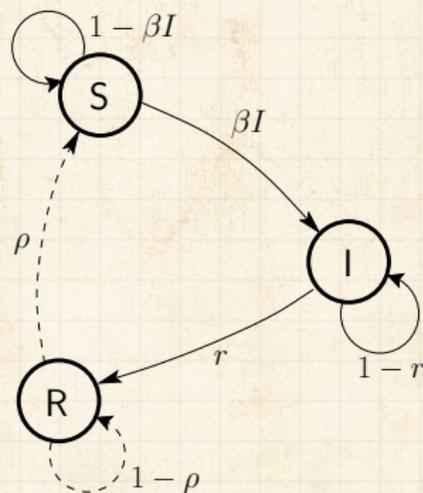


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Discrete time automata example:



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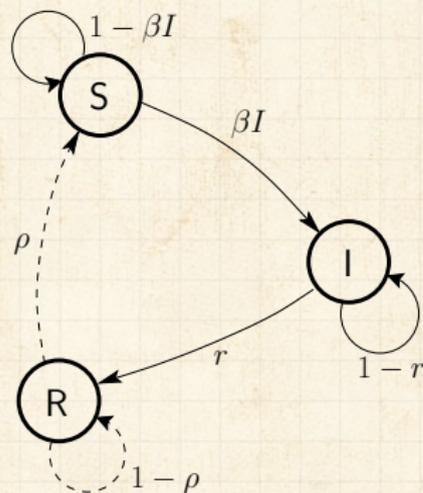


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Discrete time automata example:



Transition Probabilities:

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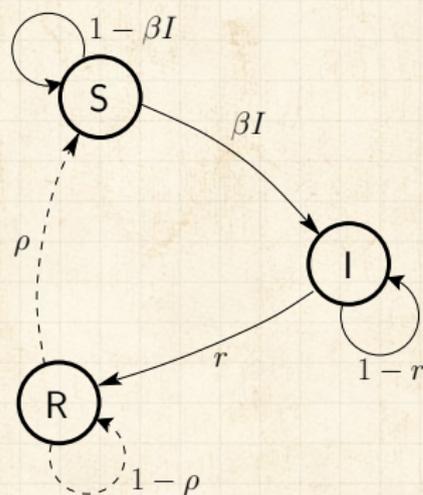


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Discrete time automata example:



Transition Probabilities:

$\beta$  for being infected given contact with infected

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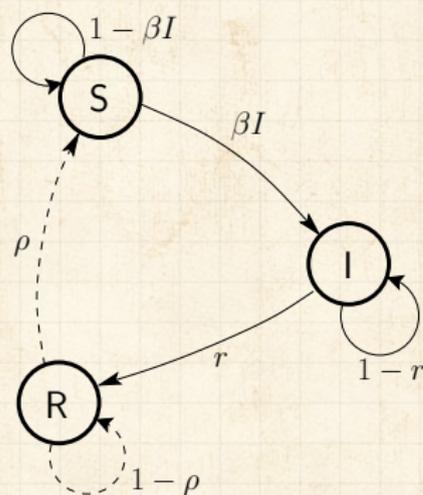


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Discrete time automata example:



Transition Probabilities:

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

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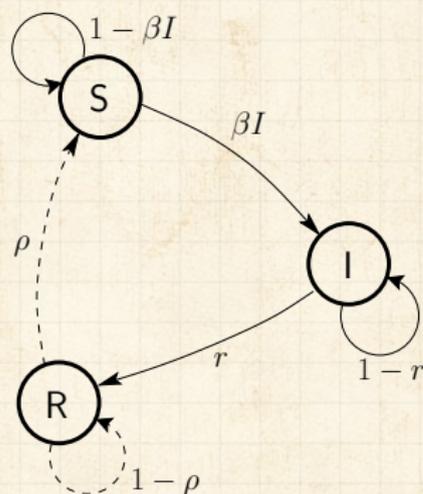
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## Discrete time automata example:



### 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



1920's: Reed and Frost



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Original models attributed to



1920's: Reed and Frost



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



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## Original models attributed to

-  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

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

$$\frac{d}{dt}I = \beta IS - rI$$

$$\frac{d}{dt}R = rI - \rho R$$

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

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## Differential equations for continuous model

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Reproduction Number  $R_0$ :

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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 $R_0$ :

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

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## Differential equations for continuous model

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

$$\frac{d}{dt}I = \beta IS - rI$$

$$\frac{d}{dt}R = rI - \rho R$$

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

## Reproduction Number $R_0$ :

-   $R_0$  = expected number of infected individuals resulting from a single initial infective
-  Epidemic threshold: If  $R_0 > 1$ , 'epidemic' occurs.

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## Discrete version:

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

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# Reproduction Number $R_0$

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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

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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
-  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
-  Probability of transmission =  $\beta$
-  At time  $t = 1$ , single Infective remains infected with probability  $1 - r$

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# Reproduction Number $R_0$

## Discrete version:

- Set up: One Infective in a randomly mixing population of Susceptibles
- At time  $t = 0$ , single infective randomly bumps into a Susceptible
- 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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# Reproduction Number $R_0$

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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:

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

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

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Discrete version:

 Expected number infected by original Infective:

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

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

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

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 Expected number infected by original Infective:

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

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

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# Reproduction Number $R_0$

## Discrete version:

 Expected number infected by original Infective:

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

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

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

 Similar story for continuous model.

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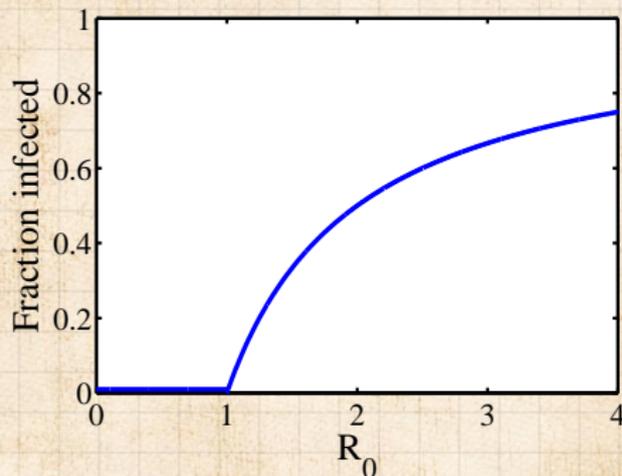


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Example of epidemic threshold:



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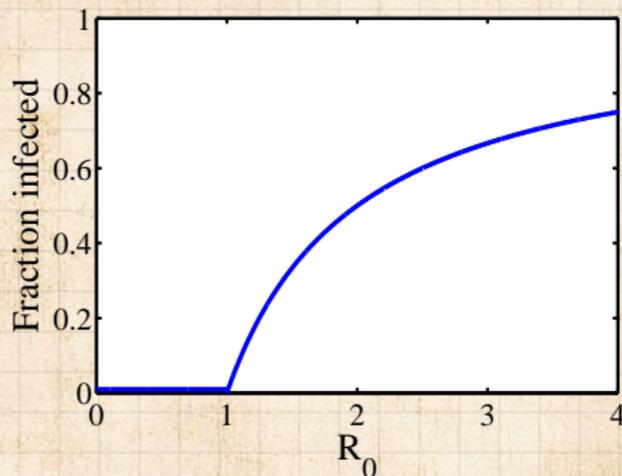


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Example of epidemic threshold:



Continuous phase transition.

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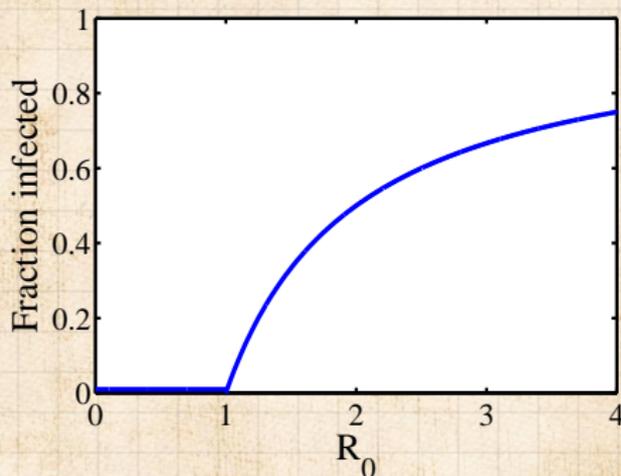


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Example of epidemic threshold:



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)<sup>[7]</sup>

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

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

 Spread of rumors (Daley & Kendall, 1964, 1965)<sup>[3, 4]</sup>

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

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

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

-  Adoption of ideas/beliefs (Goffman & Newell, 1964)<sup>[7]</sup>
-  Spread of rumors (Daley & Kendall, 1964, 1965)<sup>[3, 4]</sup>
-  Diffusion of innovations (Bass, 1969)<sup>[1]</sup>
-  Spread of fanatical behavior (Castillo-Chávez & Song, 2003)<sup>[2]</sup>

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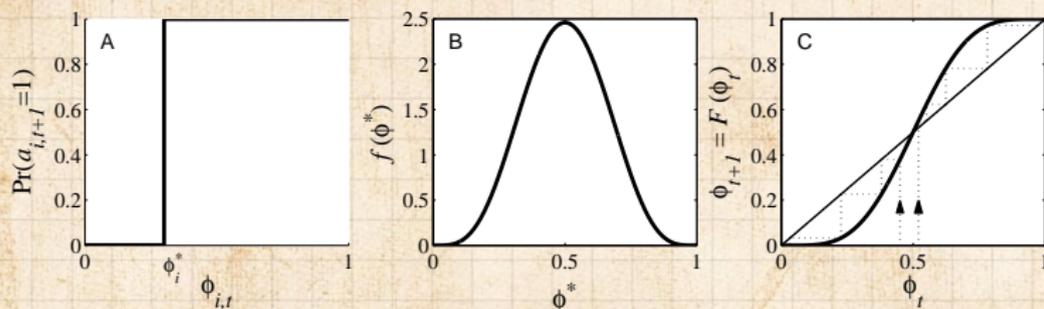
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# Granovetter's model (recap of recap)

 Action based on perceived behavior of others.



 Two states: S and I.

 Recovery now possible (SIS).

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

 Discrete time, synchronous update.

 This is a **Critical mass model**.

 **Inter**dependent 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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# Some (of many) issues

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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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# Some (of many) issues

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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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# Generalized model

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Basic ingredients:

 Incorporate memory of a contagious element [5, 6]

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## Basic ingredients:

-  Incorporate memory of a contagious element [5, 6]
-  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]
-  Population of  $N$  individuals, each in state  $S$ ,  $I$ , or  $R$ .
-  Each individual randomly contacts another at each time step.

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# Generalized model

## Basic ingredients:

-  Incorporate memory of a contagious element [5, 6]
-  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 contact with infected individual

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## Basic ingredients:

-  Incorporate memory of a contagious element [5, 6]
-  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 contact with infected individual
-  With probability  $p$ , contact with infective leads to an exposure.

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## Basic ingredients:

-  Incorporate memory of a contagious element [5, 6]
-  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 contact with infected individual
-  With probability  $p$ , contact with infective leads to an exposure.
-  If exposed, individual receives a dose of size  $d$  drawn from distribution  $f$ . Otherwise  $d = 0$ .

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$$S \Rightarrow I$$

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S  $\Rightarrow$  I

 Individuals 'remember' last  $T$  contacts:

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

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S  $\Rightarrow$  I

 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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S  $\Rightarrow$  I

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:

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

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

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I  $\Rightarrow$  R

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

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I  $\Rightarrow$  R

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

R  $\Rightarrow$  S

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

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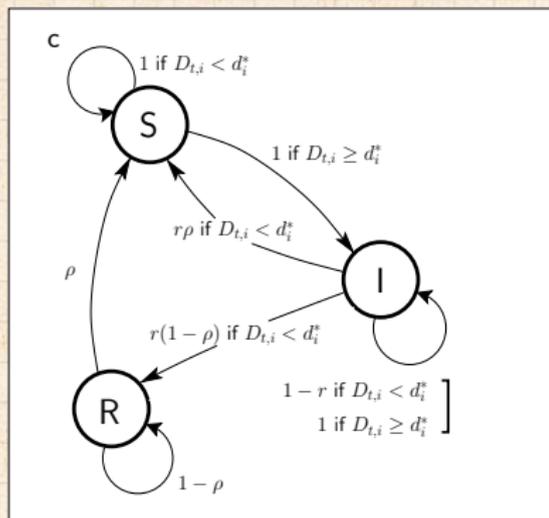
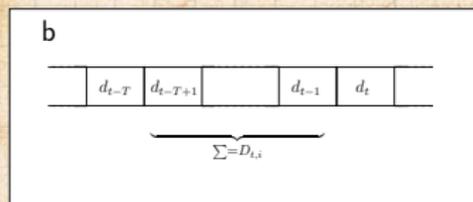
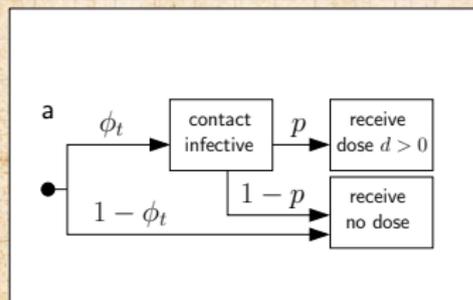
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# A visual explanation



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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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# Generalized mean-field model

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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  $p$ .

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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:

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 Look for steady-state behavior as a function of exposure probability  $p$ .

 Denote fixed points by  $\phi^*$ .

Homogeneous version:

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Study SIS-type contagion first:

 Recovered individuals are immediately susceptible again:

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 Look for steady-state behavior as a function of exposure probability  $p$ .

 Denote fixed points by  $\phi^*$ .

Homogeneous version:

 All individuals have threshold  $d^*$

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# Generalized mean-field model

## 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  $\phi^*$ .

## Homogeneous version:

 All individuals have threshold  $d^*$

 All dose sizes are equal:  $d = 1$

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# Homogeneous, one hit models:

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# Homogeneous, one hit models:

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

  $r < 1$  means recovery is probabilistic.

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# Homogeneous, one hit models:

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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# Homogeneous, one hit models:

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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# Homogeneous, one hit models:

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:

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

# Homogeneous, one hit models:

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:

$$\phi_{t+1} = p \underbrace{\phi_t}_a$$

- a: Fraction infected between  $t$  and  $t + 1$ , independent of past state or recovery.

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# Homogeneous, one hit models:

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:

$$\phi_{t+1} = \underbrace{p\phi_t}_a + \underbrace{\phi_t(1 - p\phi_t)}_b$$

- 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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# Homogeneous, one hit models:

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:

$$\phi_{t+1} = \underbrace{p\phi_t}_a + \underbrace{\phi_t(1-p\phi_t)}_b \underbrace{(1-r)}_c.$$

- 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  $T = 1$ :

 Set  $\phi_t = \phi^*$ :

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# Homogeneous, one hit models:

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

 Set  $\phi_t = \phi^*$ :

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

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# Homogeneous, one hit models:

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

 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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# Homogeneous, one hit models:

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 Set  $\phi_t = \phi^*$ :

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

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

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# Homogeneous, one hit models:

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

 Set  $\phi_t = \phi^*$ :

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 Critical point at  $p = p_c = r$ .

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# Homogeneous, one hit models:

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

 Set  $\phi_t = \phi^*$ :

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

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

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

 Critical point at  $p = p_c = r$ .

 Spreading takes off if  $p/r > 1$

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# Homogeneous, one hit models:

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

 Set  $\phi_t = \phi^*$ :

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

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

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

 Critical point at  $p = p_c = r$ .

 Spreading takes off if  $p/r > 1$

 Find continuous phase transition as for SIR model.

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# Homogeneous, one hit models:

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

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 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  $T > 1$

  $r = 1$  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  $T > 1$

-   $r = 1$  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.

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

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-  Call  $\phi^*$  the steady state level of infection.

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-   $\text{Pr}(\text{infected}) = 1 - \text{Pr}(\text{uninfected})$ :

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# Simple homogeneous examples

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

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

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# Homogeneous, one hit models:

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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# Homogeneous, one hit models:

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  $p_c$ .

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

 Closed form expression for  $\phi^*$ :

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 As  $\phi^* \rightarrow 0$ , we see

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

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# Homogeneous, one hit models:

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 Again find continuous phase transition ...



# Homogeneous, one hit models:

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  $p_c$ .

 As  $\phi^* \rightarrow 0$ , we see

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

 Again find continuous phase transition ...

 Note: we can solve for  $p$  but not  $\phi^*$ :

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

# Homogeneous, one hit models:

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

 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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# Homogeneous, one hit models:

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

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:

# Homogeneous, one hit models:

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

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1. Did not receive any infections in last T time steps,

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

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

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 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.

# Homogeneous, one hit models:

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

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

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- 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:

# Homogeneous, one hit models:

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

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

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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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# Homogeneous, one hit models:

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

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 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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# Homogeneous, one hit models:

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

 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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# Homogeneous, one hit models:

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

 In general, relevant dose histories are:

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

 Overall probabilities for dose histories occurring:

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

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# Homogeneous, one hit models:

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

 In general, relevant dose histories are:

$$H_{m+1} = \{\dots, d_{t-T-m-1}, 1, \underbrace{0, 0, \dots, 0, 0}_m, \underbrace{0, 0, \dots, 0, 0}_T\}.$$

 Overall probabilities for dose histories occurring:

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

$$P(H_{m+1}) =$$

# Homogeneous, one hit models:

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

 In general, relevant dose histories are:

$$H_{m+1} = \{\dots, d_{t-T-m-1}, 1, \underbrace{0, 0, \dots, 0, 0}_m, \underbrace{0, 0, \dots, 0, 0}_T\}.$$

 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)



# Homogeneous, one hit models:

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

 In general, relevant dose histories are:

$$H_{m+1} = \{\dots, d_{t-T-m-1}, 1, \underbrace{0, 0, \dots, 0, 0}_m, \underbrace{0, 0, \dots, 0, 0}_T\}.$$

 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)

# Homogeneous, one hit models:

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

 In general, relevant dose histories are:

$$H_{m+1} = \{\dots, d_{t-T-m-1}, 1, \underbrace{0, 0, \dots, 0, 0}_m, \underbrace{0, 0, \dots, 0, 0}_T\}.$$

 Overall probabilities for dose histories occurring:

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

$$P(H_{m+1}) = \underbrace{p\phi^*}_a \underbrace{(1 - p\phi^*)^{T+m}}_b \underbrace{(1 - r)^{m+1}}_c.$$

- 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)

# Homogeneous, one hit models:

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

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

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# Homogeneous, one hit models:

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

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

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

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# Homogeneous, one hit models:

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

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

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

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

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# Homogeneous, one hit models:

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

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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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# Homogeneous, one hit models:

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

 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  $r \leq 1$ ,  $d^* = 1$ , and  $T \geq 1$

 Fixed point equation (again):

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

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

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# Homogeneous, one hit models:

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

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

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# Homogeneous, one hit models:

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 Fixed point equation (again):

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

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

 Decreasing  $r$  keeps individuals infected for longer and decreases  $p_c$ .

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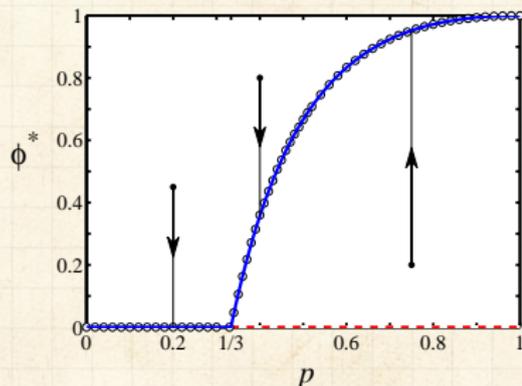
# Epidemic threshold:

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

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

  $\phi^* = 0$

  $p_c = 1/(T + \tau)$



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

 Blue = stable, red = unstable, fixed points.

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

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

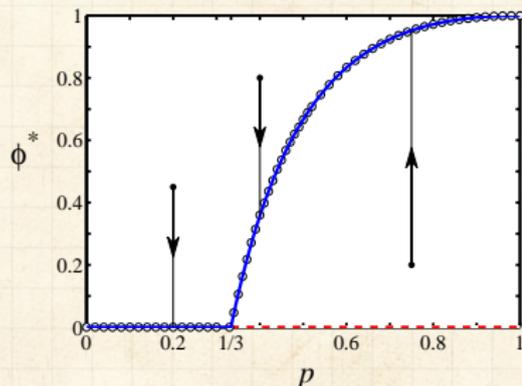
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  $\phi^* = 0$

  $p_c = 1/(T + \tau)$



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

 Blue = stable, red = unstable, fixed points.

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

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

 Phase transition can be seen as a **transcritical bifurcation**.<sup>[12]</sup>

# Homogeneous, multi-hit models:



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

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# Homogeneous, multi-hit models:

-  All right:  $d^* = 1$  models correspond to simple disease spreading models.
-  What if we allow  $d^* \geq 2$ ?

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# Homogeneous, multi-hit models:

- ☰ All right:  $d^* = 1$  models correspond to simple disease spreading models.
- ☰ What if we allow  $d^* \geq 2$ ?
- ☰ Again first consider SIS with immediate recovery ( $r = 1$ )

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# Homogeneous, multi-hit models:

-  All right:  $d^* = 1$  models correspond to simple disease spreading models.
-  What if we allow  $d^* \geq 2$ ?
-  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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# Homogeneous, multi-hit models:

- All right:  $d^* = 1$  models correspond to simple disease spreading models.
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$$\phi^* = \sum_{i=d^*}^T \binom{T}{i} (p\phi^*)^i (1 - p\phi^*)^{T-i}.$$

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



# Homogeneous, multi-hit models:

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

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# Homogeneous, multi-hit models:

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

 Exactly solvable for small  $T$ .

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# Homogeneous, multi-hit models:

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

 Exactly solvable for small  $T$ .

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

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# Homogeneous, multi-hit models:

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

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

 Fixed point equation:

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

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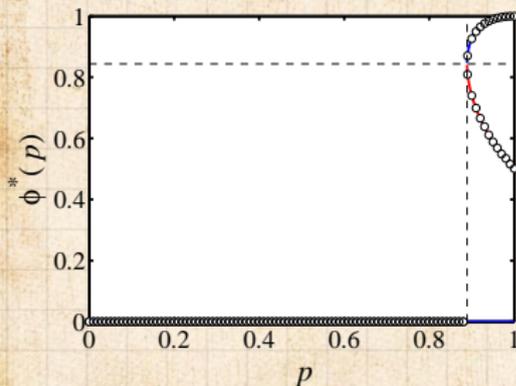


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  $(p_b, \phi^*) = (8/9, 27/32)$ .

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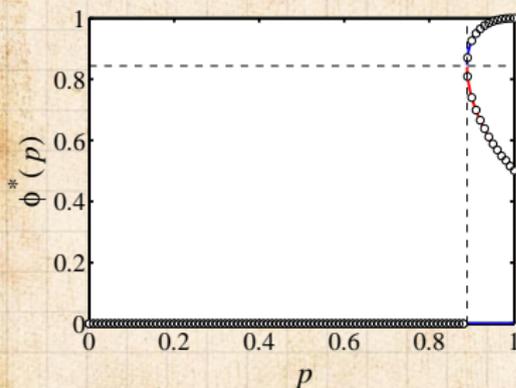


# Homogeneous, multi-hit models:

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

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 See new structure: a **saddle node bifurcation** <sup>[12]</sup> appears as  $p$  increases.

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

 Behavior akin to output of Granovetter's threshold model.

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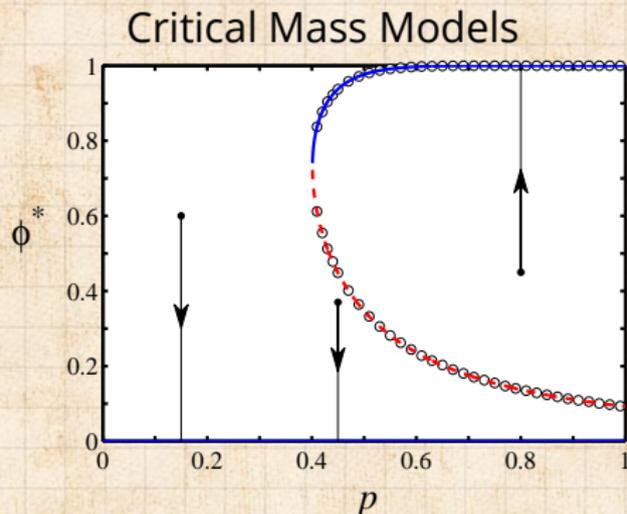


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 Another example:



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

Saddle-node bifurcation.

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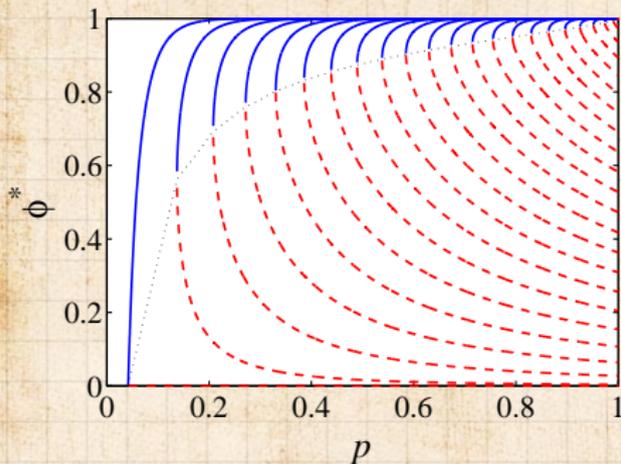
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# Fixed points for $r = 1$ , $d^* > 1$ , and $T \geq 1$

  $T = 24$ ,  $d^* = 1, 2, \dots, 23$ .



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

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

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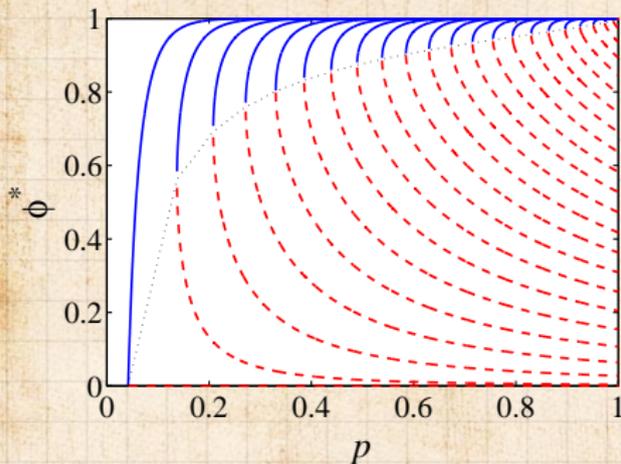
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# Fixed points for $r = 1$ , $d^* > 1$ , and $T \geq 1$

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jump between  
continuous  
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mass model.

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

 See **either** simple phase transition or saddle-node bifurcation, nothing in between.

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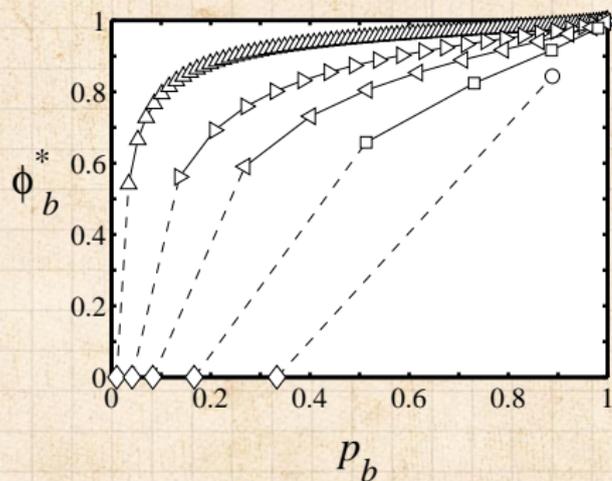
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# Fixed points for $r = 1$ , $d^* > 1$ , and $T \geq 1$

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



  $T = 96$  (.),

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

  $T = 12$  ( $\triangleleft$ ),

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

  $T = 3$  ( $\circ$ ),

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

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

 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:

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



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

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

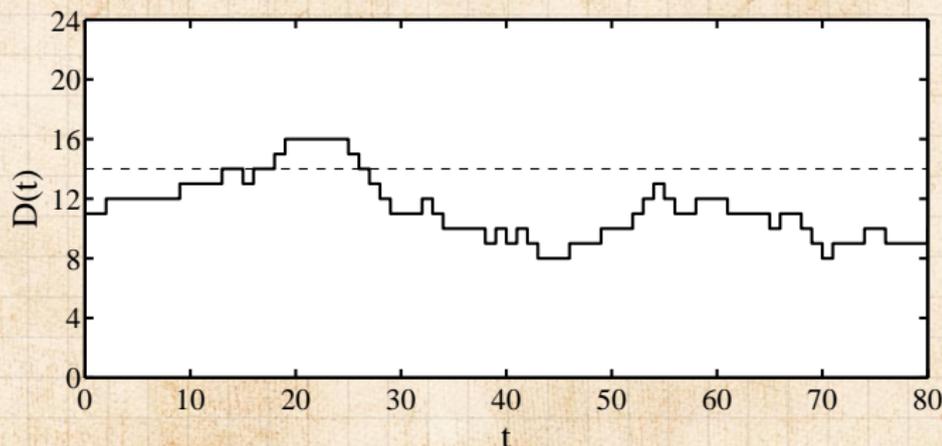
# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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



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

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

- Define  $\gamma_m$  as fraction of individuals for whom  $D(t)$  last equaled, and has since been below, their threshold  $m$  time steps ago,
- 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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- 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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# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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

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

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

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

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



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

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 Two subsequences do this:  
 $\{d_{n-2}, d_{n-1}, d_n, d_{n+1}\} = \{1, 1, 0, 0\}$



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

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- 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:

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

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

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

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

- ☰ Determine number of sequences of length  $m$  that keep dose load below  $d^* = 2$ .
- ☰  $N_a$  = number of  $a = \{0\}$  subsequences.

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

-  Determine number of sequences of length  $m$  that keep dose load below  $d^* = 2$ .
-   $N_a$  = number of  $a = \{0\}$  subsequences.
-   $N_b$  = number of  $b = \{1, 0, 0\}$  subsequences.

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

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

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

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

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

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

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

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

Possible values for  $N_b$ :

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

where  $\lfloor \cdot \rfloor$  means floor.

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

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

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Possible values for  $N_b$ :

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

where  $\lfloor \cdot \rfloor$  means floor.

 Corresponding possible values for  $N_a$ :

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

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

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

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

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 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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  $N_a + N_b$  slots for subsequences.

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 Think of overall sequence in terms of subsequences:

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  $N_a + N_b$  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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# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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

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

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

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

 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  $D_m^{a,b}$ .

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

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

 Three endings  $\Rightarrow$  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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# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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

 Three endings  $\Rightarrow$  Six possible sequences:

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

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

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

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

$$P_2 = (p\phi)^3(1-p\phi)^2\chi_{m-2}(p, \phi)$$

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

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

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

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

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# Fixed points for $r < 1$ , $d^* = 2$ , and $T = 3$

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

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

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

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

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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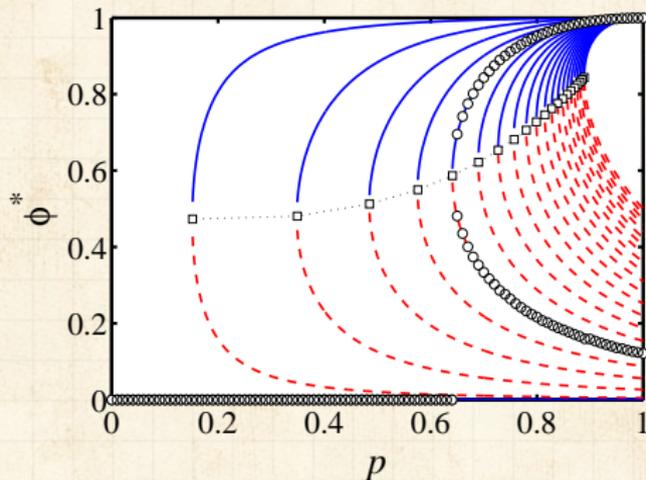


# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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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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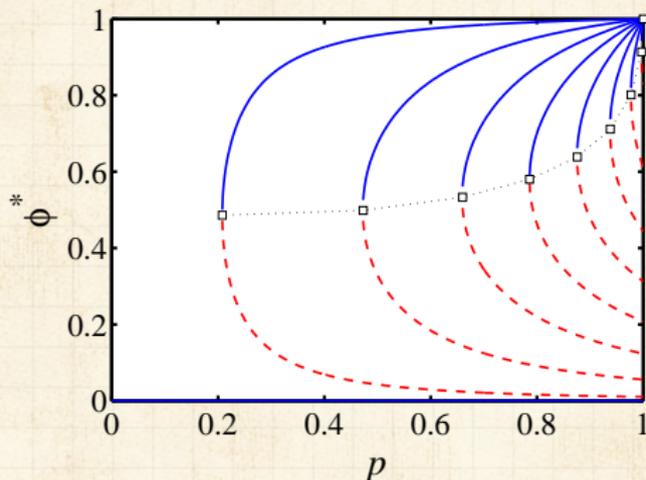


# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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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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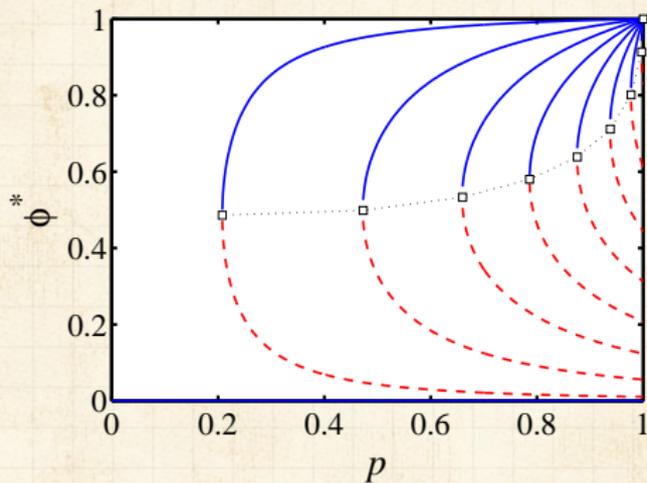


# Fixed points for $r < 1$ , $d^* > 1$ , and $T \geq 1$

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



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

 No spreading for  $r \gtrsim 0.382.$

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 Two kinds of contagion processes:

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# What we have now:

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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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# Generalized model

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

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# Generalized model

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

 Key quantities:

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

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# Generalized model

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

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

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

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# Generalized model

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

 Key quantities:

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

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

 e.g.,  
 $P_1$  = Probability that one dose will exceed the threshold of a random individual  
= Fraction of most vulnerable individuals.

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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:

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

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

$$pP_1T \geq 1$$

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# Generalized model—heterogeneity, $r = 1$

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 Fixed point equation:

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

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

$$pP_1T \geq 1$$

or

$$\Rightarrow p_c = 1/(TP_1)$$

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# Generalized model—heterogeneity, $r = 1$

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

$$pP_1T \geq 1$$

or

$$\Rightarrow p_c = 1/(TP_1)$$

 Very good:

1.  $P_1T$  is the expected number of vulnerables the initial infected individual meets before recovering.

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# Generalized model—heterogeneity, $r = 1$

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or

$$\Rightarrow p_c = 1/(TP_1)$$

 Very good:

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

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# Generalized model—heterogeneity, $r = 1$

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

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or

$$\Rightarrow p_c = 1/(TP_1)$$

 Very good:

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

 Observe:  $p_c$  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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# Heterogeneous case

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

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

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# Heterogeneous case

- 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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# Heterogeneous case

- Next: Determine slope of fixed point curve at critical point  $p_c$ .
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- Find slope depends on  $(P_1 - P_2/2)$  [6] (see Appendix).
- Behavior near fixed point depends on whether this slope is

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# Heterogeneous case

-  **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).
-  Behavior near fixed point depends on whether this slope is
  1. positive:  $P_1 > P_2/2$  (continuous phase transition)

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# Heterogeneous case

-  **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).
-  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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# Heterogeneous case

-  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).
-  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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# Heterogeneous case

## Example configuration:

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

 Memory span:  $T = 10$ .

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# Heterogeneous case

## Example configuration:

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

 Memory span:  $T = 10$ .

 Thresholds are uniformly set at

1.  $d_* = 0.5$
2.  $d_* = 1.6$
3.  $d_* = 3$

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# Heterogeneous case

## Example configuration:

-  Dose sizes are lognormally distributed with mean 1 and variance 0.433.
-  Memory span:  $T = 10$ .
-  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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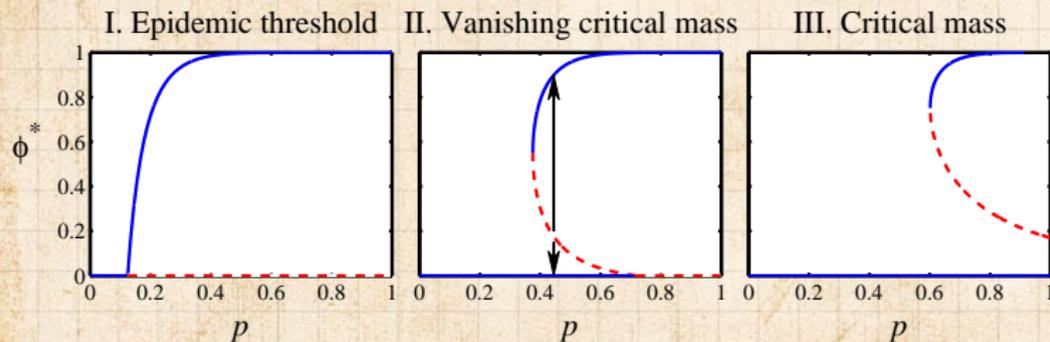
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# Three universal classes

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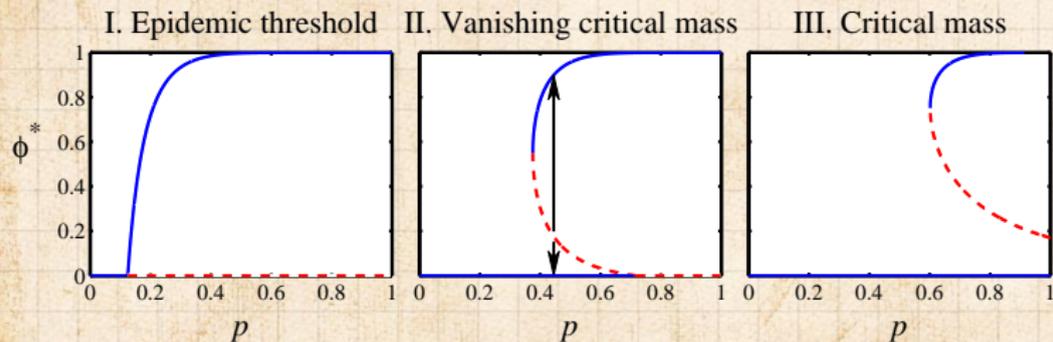
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# Three universal classes

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

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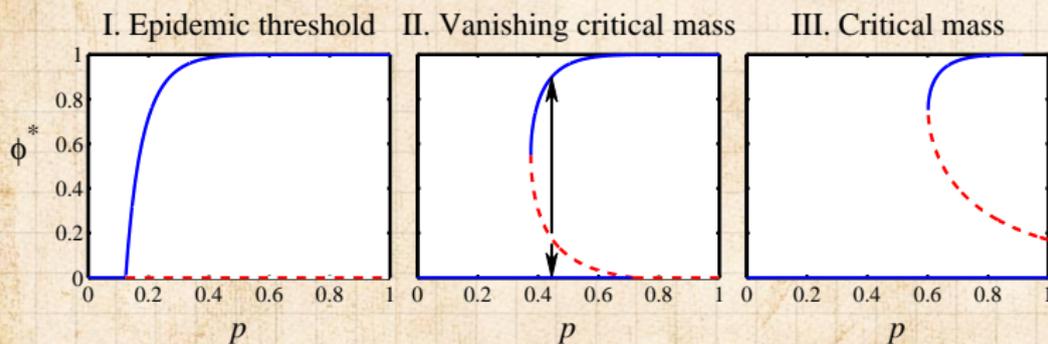
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# Three universal classes

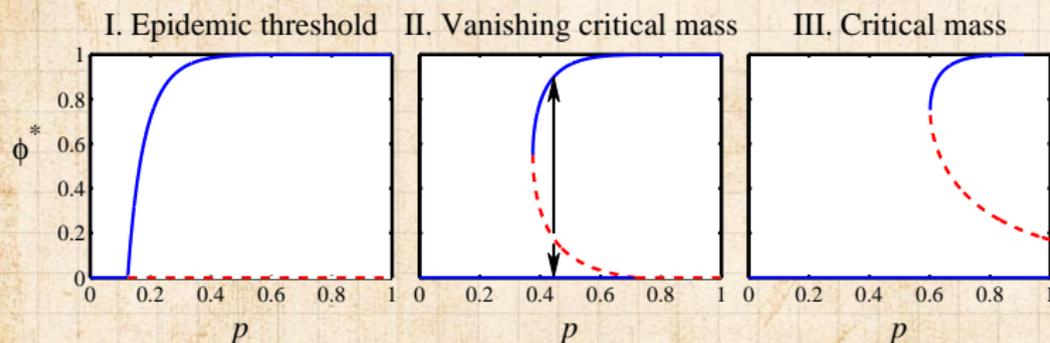


 Epidemic threshold:  $P_1 > P_2/2, p_c = 1/(TP_1) < 1$

 Vanishing critical mass:  $P_1 < P_2/2, p_c = 1/(TP_1) < 1$



# Three universal classes



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

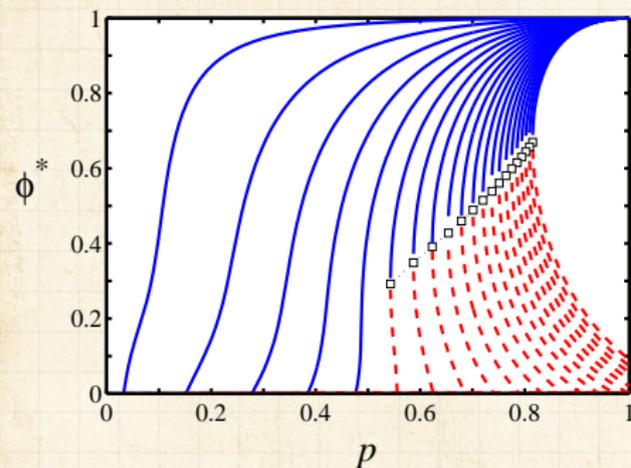
 Vanishing critical mass:  $P_1 < P_2/2, p_c = 1/(TP_1) < 1$

 Pure critical mass:  $P_1 < P_2/2, p_c = 1/(TP_1) > 1$



# 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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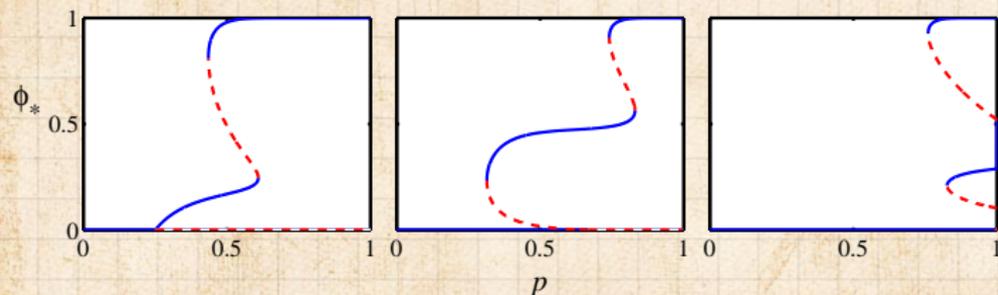
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# More complicated models

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- 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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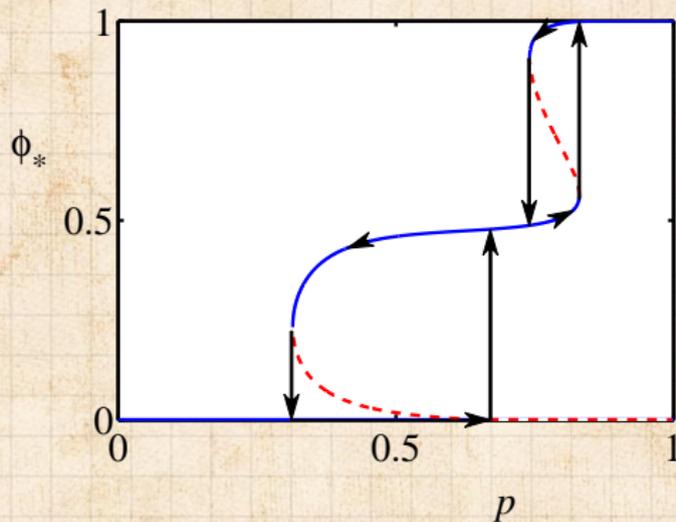
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# Hysteresis in vanishing critical mass models

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 Memory is a natural ingredient.

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Memory is a natural ingredient.



Three universal classes of contagion processes:

1. I. Epidemic Threshold
2. II. Vanishing Critical Mass
3. III. Critical Mass

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Memory is a natural ingredient.



Three universal classes of contagion processes:

1. I. Epidemic Threshold
2. II. Vanishing Critical Mass
3. III. Critical Mass



Dramatic changes in behavior possible.

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# Nutshell (one half)

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-  Memory is a natural ingredient.
-  Three universal classes of contagion processes:
  1. I. Epidemic Threshold
  2. II. Vanishing Critical Mass
  3. III. Critical Mass
-  Dramatic changes in behavior possible.
-  To change kind of model: 'adjust' memory, recovery, fraction of vulnerable individuals ( $T$ ,  $r$ ,  $\rho$ ,  $P_1$ , and/or  $P_2$ ).

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-  Memory is a natural ingredient.
-  Three universal classes of contagion processes:
  1. I. Epidemic Threshold
  2. II. Vanishing Critical Mass
  3. III. Critical Mass
-  Dramatic changes in behavior possible.
-  To change kind of model: 'adjust' memory, recovery, fraction of vulnerable individuals ( $T$ ,  $r$ ,  $\rho$ ,  $P_1$ , and/or  $P_2$ ).
-  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  $pP_1(T + \tau) \geq 1$ .

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

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

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- Single seed infects others if  $pP_1(T + \tau) \geq 1$ .
- Key quantity:  $p_c = 1/[P_1(T + \tau)]$
- If  $p_c < 1 \Rightarrow$  contagion can spread from single seed.

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-  Single seed infects others if  $pP_1(T + \tau) \geq 1$ .
-  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 ( $P_1$ ).

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- Depends only on:
  - System Memory ( $T + \tau$ ).
  - Fraction of highly vulnerable individuals ( $P_1$ ).
- Details unimportant: Many threshold and dose distributions give same  $P_k$ .

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# Nutshell (other half)

- Single seed infects others if  $pP_1(T + \tau) \geq 1$ .
- Key quantity:  $p_c = 1/[P_1(T + \tau)]$
- If  $p_c < 1 \Rightarrow$  contagion can spread from single seed.
- Depends only on:
  - System Memory ( $T + \tau$ ).
  - Fraction of highly vulnerable individuals ( $P_1$ ).
- 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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# Appendix: Details for Class I-II transition:



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

# Appendix: Details for Class I-II transition:

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

since

$$\begin{aligned} \binom{T}{k} \binom{T-k}{m-k} &= \frac{T!}{k!(T-k)!} \frac{(T-k)!}{(m-k)!(T-m)!} \\ &= \frac{T!}{m!(T-m)!} \frac{m!}{k!(m-k)!} \\ &= \binom{T}{m} \binom{m}{k}. \end{aligned}$$

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# Appendix: Details for Class I-II transition:

## Linearization gives

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

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

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# Appendix: Details for Class I-II transition:

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

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

## 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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# Appendix: Details for Class I-II transition:

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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  
 $C_2 = \binom{T}{2}(-2P_1 + P_2)$ .

 Using  $p_c = 1/(TP_1)$ :

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

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

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