# Introduction to the Mathematics of Epidemics on Graphs

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# **Epidemics with fixed infection times**

The action takes place on a graoh G in continuous time  $t \in [0, \infty)$ .

**SIR epidemic**. Susceptibles become infected at rate  $\lambda$  times the number of infected neighbors. An infection lasts for time 1, after which the individual is "Removed," no longer can be infected.

Since the infection time is fixed the events x infects  $y_i$  are independent for all the neighbors  $y_i$  of x.

Let T be exponential with rate  $\lambda$ , i.e.  $P(T > t) = e^{-\lambda t}$ . Let

$$\tau_f = P(T \le 1) = 1 - e^{-\lambda}$$

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# **Epidemic = Percolation**

Each edge will be S - I (or I - S) only once. Flip a coin with probability  $\tau_f$  of heads, and keep the edge if the coin shows heads, otherwise delete it. The individuals infected in an epidemic started by x = the connected component of the graph containing x.

**Erdös-Renyi graph.** There are *n* vertices. Each pair of vertices is indpendently connected by an edge with probability  $\mu/n$ . The thinned graph is  $ER(n, \mu\tau^f/n)$ .

A large epidemic occurs with positive probability if  $\mu \tau^f > 1$ . If  $z_0$  is the fixed point smaller than 1 of the generating function

$$G(z) = \exp(-\mu \tau^f (1-z)),$$

then  $1 - z_0$  gives both the limiting probability an infected individual will start a large epidemic, and the fraction of individuals infected when a large epidemic occurs.

# **Configuration model**

Let  $d_1, \ldots, d_n$  be independent and have  $P(d_i = k) = p_k$ . Since we want  $d_i$  to be the degree of vertex *i*, we condition on  $E_n = \{d_1 + \cdots + d_n \text{ is even}\}$ . To build the graph we think of  $d_i$  half-edges attached to *i* and then pair the half-edges at random.



#### Branching process viewpoint

Pick a vertex x to start from. Let  $Z_m$  be the number of vertices at distance m from x. In the Erdös-Renyi case, when m is small,  $Z_m$  is a branching process in which each individual has a Poisson( $\mu$ ) number of children.



# Branching process view of configuration model

First vertex chosen has j neighbors with probability  $p_j = P(D_i = j)$ .

Since we connect half-edges at random, a first generation vertex with degree k is k times as likely to be chosen as one with degree 1, so the distribution of the number of children of a first generation vertex is

$$q_{k-1} = rac{k p_k}{\mu}$$
 for  $k \geq 1$  where  $\mu = \sum_k k p_k$ 

The k-1 on the left-hand side comes from the fact that we used up one edge connecting to the vertex. If p has finite second moment, q has finite mean  $\nu = \sum_k k(k-1)p_k/\mu$ .

If  $\nu > 1$  then there is a component of order *n* with positive probability.

## **General infection times**

Suppose now that infections have duration S with density function  $f_S(s)$ . A common choice is exponential( $\gamma$ ) so that the set of infected sites is a Markov process.

Let  $\tau_c = P(T < S)$ . Infections from x to its neighbors  $y_i$  are no longer independent, but that does not matter, since our branching process will be supercritical if the mean number of children  $\nu \tau_c > 1$ .

$$\hat{G}_1(z) = \int_0^\infty ds \, f_{\mathsf{S}}(s) \sum_{j=0}^\infty z^j \sum_{k=j}^\infty q_k \binom{k}{j} (1 - e^{-\lambda s})^j (e^{-\lambda s})^{k-j}$$

If  $q_k \to p_k$  call this  $\hat{G}_0$ . When  $\nu > 1$  the probability of a large epidemic is  $1 - \hat{G}_0(\zeta)$  where  $\hat{G}_1(\zeta) = \zeta$  is the fixed point < 1.

# Probability of large epidemic $\neq$ final size.

For general infection times, the relevant graph for percolation has oriented edges x to neighbors y.

Probability of a giant component when we start at a randomly chosen vertex is percolation probability when we follow edges in the direction of their orientation. (Edges from x are dependent.)

Final size is probability of a giant component when we start at a randomly chosen vertex and follow edges in the direction OPPOSITE to their orientation. (Edges to x are independent.)

**Homework:** Compute the probability of a large epidemic and the final size on the complete graph when  $S = exponential(\gamma)$ . Out degree is shifted geometric, in degree is Poisson.

# Power law random graphs

 $p_k \sim Ck^{-\alpha}$ ,  $q_k \sim C'k^{1-\alpha}$ . If  $2 < \alpha < 3$  then  $\sum_k kp_k < \infty$  but  $\sum_k kq_k = \infty$ , i.e., the branching process has infinite mean. Internet (physical network of machines)  $\alpha = 2.16$  Faloutos<sup>3</sup> (1999) Movie actor network (connected by movies)  $\alpha = 2.3$ Liljeros et al (2001) Sexual network of 4,781 Swedes.  $\alpha_{male} = 3.3$ ,  $\alpha_{female} = 3.5$ . "Internet is robust yet fragile." Percolation probability for random attacks  $p_c = 0$  but is easily disconnected by targeting high degree nodes.

For proofs, Bollobás and Riordan (2004) Internet Math 1, 1-35

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# SIS model

Nodes become infected at rate  $\lambda$ . At rate 1, infected nodes become healthy (and again susceptible to disease)

If infected  $\rightarrow$  occupied, and susceptible  $\rightarrow$  vacant this is the contact process invented by Ted Harris in 1974 (on  $\mathbb{Z}^d$ ).

 $\lambda_c$  is threshold for survival,  $heta(\lambda) =$  survival probability,  $heta(\lambda) \sim C(\lambda - \lambda_c)^{eta}$ 

According to degree-based mean field calculations

If 
$$\alpha \leq 3$$
,  $\lambda_c = 0$ ,  $\beta = 1/(3 - \alpha)$   
If  $3 < \alpha < 4$ ,  $\lambda_c > 0$  but  $\beta = 1/(\alpha - 3) > 1$   
If  $\alpha \geq 4$ ,  $\lambda_c > 0$ .  $\beta = 1$ 

Epidemics in Complex Networks Rev. Modern Phys. 87, 925-979

# Berger, Borgs, Chayes and Saberi (2005)

Considered the contact process on Barabási-Albert preferential attachment graph, has a power law degree distribution with  $\alpha = 3$ .

**Theorem.** (a) The probability that the process will survive from a randomly chosen vertex is  $\lambda^{\Theta(1)}$  and hence  $\lambda_c = 0$ . (b) With probability  $1 - O(\lambda^2)$  the survival probability is

 $\lambda^{\Theta[\log(1/\lambda)/\log\log(1/\lambda)]}$ 

Idea: If the infection can reach a vertex with degree  $C\lambda^{-2}$  there is survival **Lemma.** On the star graph with k leaves, if  $k\lambda^2 \to \infty$ 

$$P\left(\xi_{\exp(k\lambda^2/10)}^0 
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# (Shirshendu) Chatterjee and Durrett (2009)

Suppose  $\alpha > 3$ ,  $P(d_i \le 2) = 0$ , and  $G_n$  has no self-loops or parallel edges.

**Theorem.** Let  $\xi_t^1$ ,  $t \ge 0$  denote the contact process on the random graph  $G_n$  starting from all sites occupied. Then for any  $\lambda > 0$ , there is a positive constant  $p(\lambda)$  so that for any  $\delta > 0$ 

$$\inf_{t \leq \exp(n^{1-\delta})} P\left(\frac{|\xi_t^1|}{n} \geq p(\lambda)\right) \to 1 \quad \text{as } n \to \infty.$$

**Main idea:** Look at all the vertices with  $d_i \ge n^{\epsilon}$ . The infection survives for a long time on their stars. The graph has diameter  $O(\log n)$  so if a star becomes healthy it is easy to push infection from another star to it.

Bounds were also proved on the critical exponent  $\beta$ .

# Mountford and friends (2013, 2016)

Mountford, T., Valesin, D., and Yao, Q. (2013) extended the result to cover  $2 < \alpha \leq 3$  and proved upper and lower bounds on survival probability, that are the same up to a constant.

$$\theta(\lambda) = \begin{cases} \lambda^{1/(3-\alpha)} & 2 < a \le 5/2\\ \lambda^{2\alpha-3} \log^{2-\alpha}(1/\lambda) & 5/2 < \alpha \le 3\\ \lambda^{2\alpha-3} \log^{4-2\alpha}(1/\lambda) & 3 < \alpha \end{cases}$$

Mountford, T., Mourrat, J-C, Valesin, D., and Yao, Q. (2016) showed that survival holds for times  $t \leq e^{cn}$ .

# **Physics versus Rigorous Critical Exponents**



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#### Contact process on Galton-Watson trees.

It is easier to work on the infinite object because there is survival for all time rather than just metastability. On trees there are two critical values: if  $\lambda > \lambda_1$  the process does not die out. If  $\lambda > \lambda_2$  the root is occupied infinitely often.

**Theorem.** Pemantle (1992) If the offspring distribution in the Galton-Watson tree is a stretched exponential  $p_k = c_{\gamma} \exp(-k^{\gamma})$  with  $\gamma < 1$  and has mean  $\mu > 1$  then  $\lambda_2 = 0$ .

**Theorem** Zoe Huang and Durrett (arXiv:1810.06040) If the offspring distribution  $p_k$  for a Galton-Watson tree is subexponential

$$\limsup_{k\to\infty}(1/k)\log p_k=0.$$

and has mean  $\mu > 1$  then  $\lambda_2 = 0$ .

**Theorem.** Consider the contact process on the Galton-Watson tree with offspring distribution  $\zeta$  with  $E(\exp(c\zeta)) < \infty$  for some c > 0, then  $\lambda_1 > 0$ .

Proof is based on a very simple recursive equation. They also get bounds on the critical value and on survival times. To lower bound survival times they use subtrees with good expansion properties.

**Problem.** Is  $\lambda_1 < \lambda_2$ ?

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# DOMath project, Duke Summer 2020

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# **Two Topics**

Duke is unusual because students are required to live on campus 3 out of 4 years, so 80% of students live on campus. How much would spread in the dormitories be reduced if all students had single rooms?

Gressman and Peck (arXiv:2006.03175) have used a stochastic agent-based model of a university to argue that to reduce spread all classes larger than 30 should be online only. We will use a much simpler models to show that the answer depends on how contagious the disease is.

# Household model

This well-studied model is one step more realistic than homogeneously mixing. See e.g., Ball, Mollison, and Scalia-Tomba (1997) Ann. Prob.

- $p_H = P($ infect a given person within the house)
- $p_G = (G \text{ is for glabal}) P(\text{infect a given person outside the house})$
- $G_H(z) = g.f.$  of the size of the epidemic within the house started by one person
- $G_{\nu}(z) = \exp(Np_G(z-1) \text{ g.f. of number of long distance infections})$ caused by one person, distribution Poisson( $Np_G$ )

Probability of a large epidemic is the positive solution of

$$1-\zeta = G_H(G_\nu(\zeta))$$

## Dorm with all single rooms

- $G_D^1(z) = g.f.$  of the size of the epidemic within the dorm started by one person
- G<sub>ν</sub>(z) = exp(Np<sub>G</sub>(z 1)) g.f. of number of long distance infections caused by one person, distribution Poisson(Np<sub>G</sub>)

Spread of the epidemic in the dorm is an branching process with progeny Poisson( $\lambda_1$ ),  $\lambda_1 = np_D$ . Let  $G_b^1$  be the g.f. of total progeny

Mean number infected in dorm:  $(1 - \lambda_1)^{-1}$ 

$$G_D^1$$
 solves  $G_D^1(z) = zG_b^1(G_D^1(z))$ 

Solve by iteration. Probability of a large epidemic solves

$$1-\zeta = G_D^1(G_
u(\zeta))$$
 where  $G_
u(z) = \exp(Np_G(z-1))$ 

#### Dorm with all double rooms

 $p_L$  = probability you infect your roommate Epidemic in the dorm is an branching process with progeny  $p_I \operatorname{Poisson}(2 \cdot 2n_1 p_D) + (1 - p_I) \operatorname{Poisson}(2n_1 p_D)$ Let  $G_{h}^{2}$  be the generating function of total progeny Let  $\lambda_2 = (1 + p_I) 2n_1 p_D$ . Mean number infected:  $(1 - \lambda_2)^{-1}$ g.f. of size of epidemic within the dorm solves  $G_D^2(z) = zG_b^2(G_D^2(z))$ Probability of a large epidemic solves

$$1-\zeta = G_D^2(G_
u(\zeta))$$
 where  $G_
u(z) = \exp(Np_G(z-1))$ 

# Comparison of epidemic probabilities

 $p_L = 0.7$ ,  $n = 2n_1$  so same number of students per dorm.



Figure: When  $2n_1p_D = 0.7$  or 0.9 there is an epidemic even when  $p_G = 0$ . When  $2n_1p_D = 0.3$  or 0.5 there is a positive threshold for  $p_G$  but it is 50% larger in the single room situation.

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# **Class model**

We consider some simplifieid models for a university where each student takes three classes. The situation can be described by a graph in which there are *n* students and *m* classes with sizes  $c_i \ 1 \le i \le m$ , and there is an edge from each student to the three classes they are enrolled in. Each classroom is assumed to be homogeneously mixing.

An individual that is infected in class i at time t, adds his two alter egos (his presence in other classes) to the infected population at time t + 1/2 and then infections at time t + 1 are produced. None of the time t + 1/2 infecteds are present at time t + 1. Two time steps = one week.

**Scenario 1.** 100 classes of size 30. Each student takes 3 classes so this means 1000 students total. An individual infected in class 1 at time 0 will produce a Poisson(87p) infecteds in class 1 at time 1.

#### Scenario 2

25 classes of size 60  $(1 \le i \le 25)$  and 75 of size 20  $(26 \le i \le 100)$ , so again 1000 students. m(i,j) expected number of infections in class j caused by one infected in in class i.

$$m_{i,j} = \begin{cases} (c_i - 1)p & j = i \\ 2 \cdot \frac{c_j}{3000 - c_i} \cdot (c_j - 1)p & j \neq i \end{cases}$$

Since maximum eigenvalue has  $x_i = a$  for  $1 \le i \le 25$  and  $x_i = b$  for  $26 \le i \le 100$ , the eigenvalue problem for the  $100 \times 100$  matrix can be reduced to one for a  $2 \times 2$  matrix and we have

$$R_0 = 129.38p$$

which is 48% larger than all classes of size 30.

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### Scenario 3

Consider a university with 111 classes, one each of sizes ranging from 10, 11... 120. Total enrollment in classes is 10 + 11 + ... + 120 = 7215, or 2405 students. m(i,j) is the expected number of infections in class j caused by one infected student in i.

$$m_{i,j} = \begin{cases} (j+8)p & i=j\\ \frac{2(j+9)}{7215-(i+9)} \cdot (j+8)p & i\neq j \end{cases}$$

The maximum eigenvalue  $R_0 = 251.5p$ 

If all classes of size > k are moved online, m(i, j) = 0 for j > k.

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