

Effect of initial seeding in epidemic process on different networks

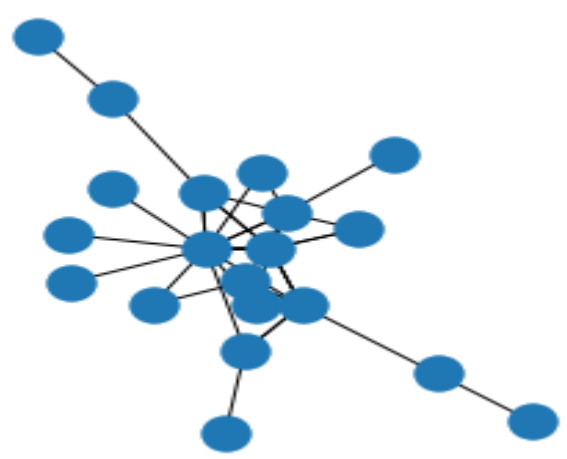
Research Question

Given the case count data for a certain period of time, when should one start the simulations? Are the simulations starting from day 1 with initial infections I similar to the simulations started from day T when initial infections are k times I .

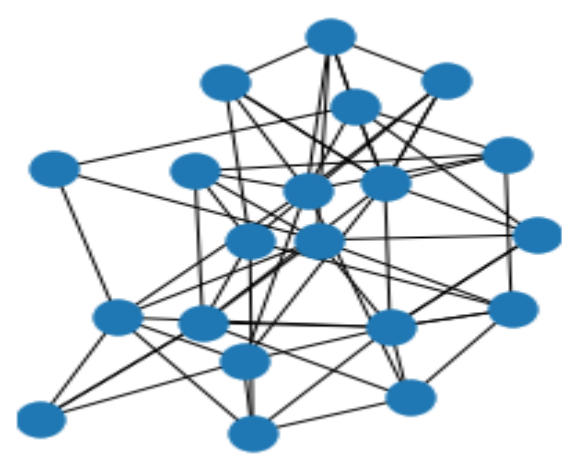
Background

Different networks can model different social situations, and three distinct networks are inspected

Truncated Power Law Network



Erdős-Rényi Network



Regular Network

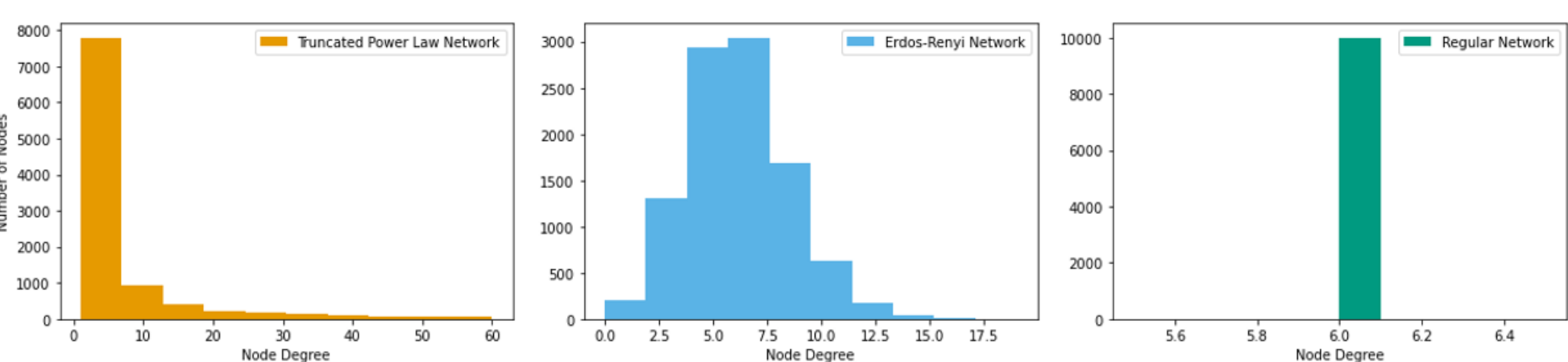
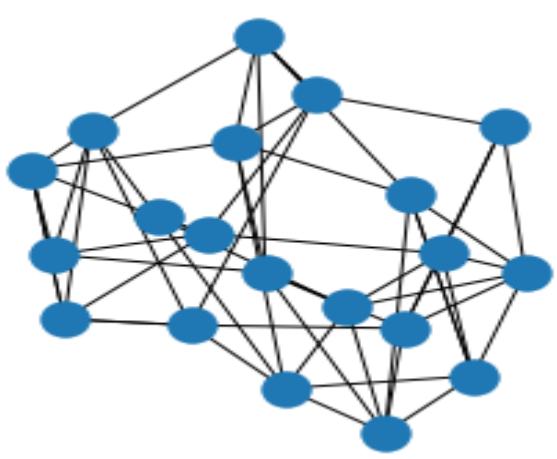
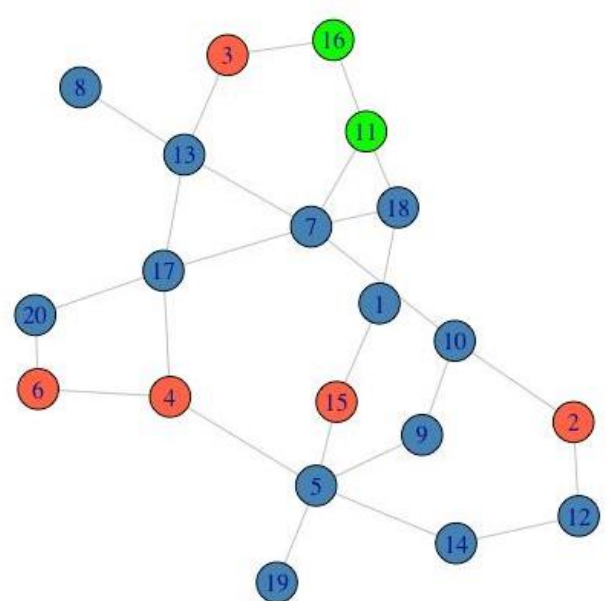


Figure #1: Degree Distribution for Truncated Power Law Network (Left), Erdős-Rényi Network (Center), and Regular Network (Right)

SIR Dynamics on Networks

Compartmental modeling is a method to simulate disease transmission

- Susceptible, Infectious, Recovered (SIR) model utilized in this experiment
- This type of modeling can also be utilized on contact networks, where the nodes represent people and edges represent contacts between people



$$\begin{aligned}\dot{S}(t) &= -\tau I(t)S(t) \\ \dot{I}(t) &= \tau I(t)S(t) - \gamma I(t) \\ \dot{R}(t) &= \gamma I(t)\end{aligned}$$

Figure #3: Susceptible (Blue), Infectious (Red), Recovered (Green) model on a network

Future Work

Explore scaling on different networks, specifically networks that better represent contact networks for disease transmission

- Truncated power law attempts to model the social network of disease transmission, but other networks can also model this network

What happens when the proportion of the population infected, ρ , is held constant and the size of the population, N , increases?

- Is there a scalar limit, k , at which a base population of size N fails to display the same disease dynamics as a population of size kN ?

References

- Kiss IZ, Miller JC, Simon P. (Book) Mathematics of epidemics on networks: from exact to approximate models. Springer; 2017.
- Okabe, Y., Shudo, A. Spread of variants of epidemic disease based on the microscopic numerical simulations on networks. Sci Rep 12, 523 (2022). <https://doi.org/10.1038/s41598-021-04520-0>
- Hethcote, H. W. (2000). The Mathematics of Infectious Diseases. SIAM Review, 42(4), 599–653. <http://www.jstor.org/stable/2653135>

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Current Work

Exploration of how network structure affect disease dynamics:

- After generating networks, running SIR models on those networks to simulate a contagion with a fixed number of initial infections distributed uniformly on the network
- Examined how the network structure and initial number of infections affects disease dynamics of the SIR simulation
 - Fixed Seeding - Determined how the initial seeding of infected nodes (I_0) within a specific network affects disease dynamics
 - Fixed Network - Compared disease dynamics with initial infections (I_0) and scaled initial infections ($k I_0$)

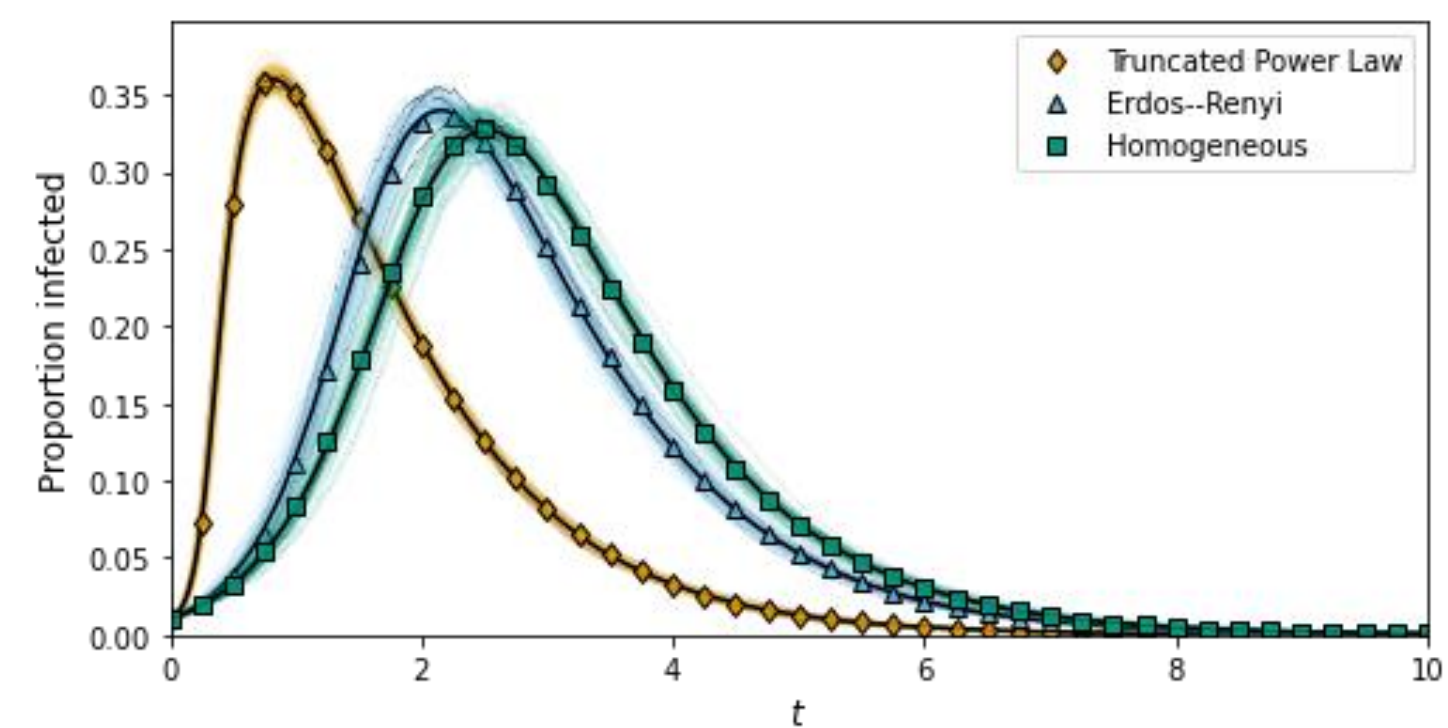


Figure #2: Mean epi curve for 50 SIR simulations on a network of size 10,000, $\tau = 0.5$, $\gamma = 1$

Design

- Generated previously described networks and ran SIR simulations with uniformly selected initial infected nodes (I_0)
- Altered the number of infected nodes (I_0) by a scalar, k , and the transmission rate, τ

Experiment

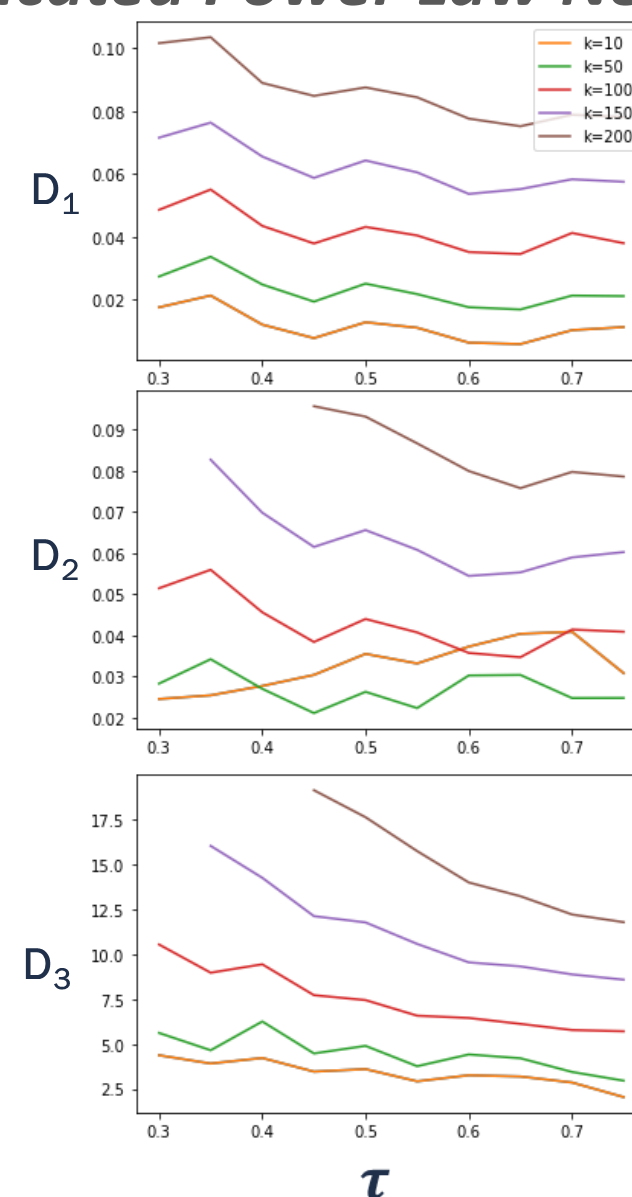
1. Generate truncated power law network and Erdős-Rényi Network
2. Run SIR simulations with I_0 and $k I_0$ infected nodes
3. Inspected the mean infection curves for different k and τ ,
4. Determined the shift (Δ_k) at which the unit epi curve ($k = 1$) reaches $k I_0$

Results

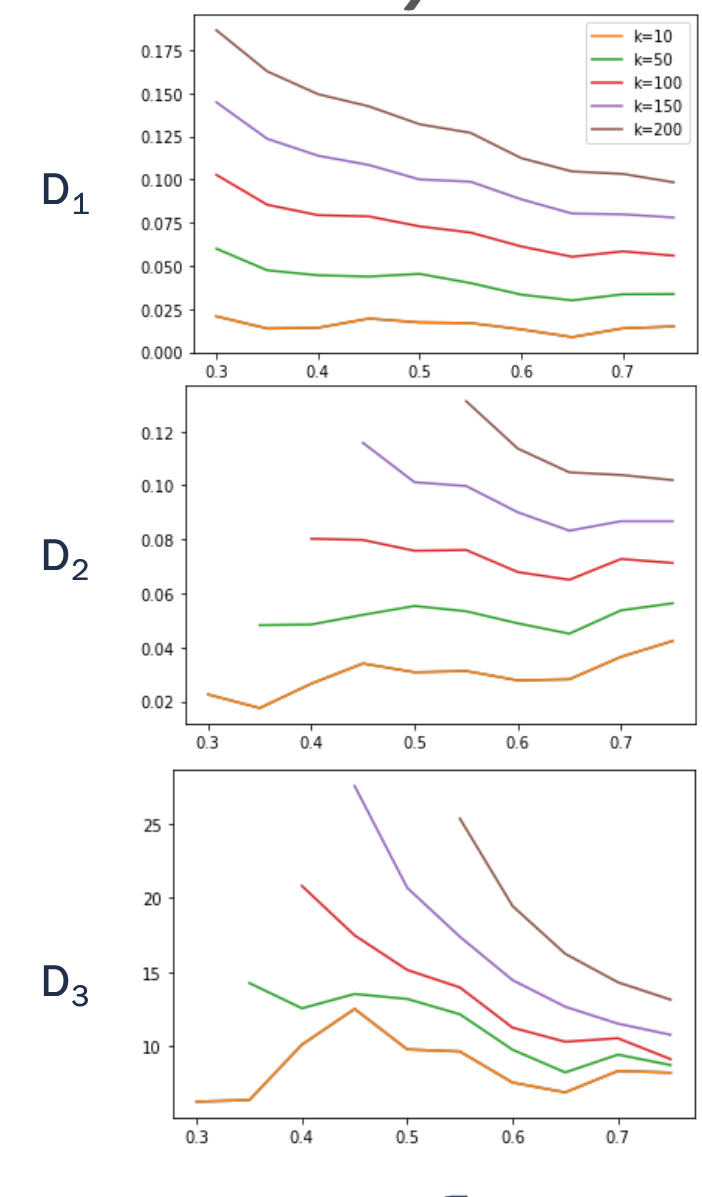
Three distances to represent how similar the scaled epi curve is to the unit epi curve:

- Difference in Total Proportion Infected (D_1)
 - $D_1(i_k, i_1) = |\sum_{\Delta_k < t} i_k(t + \Delta_k) - \sum_{\Delta_k < t} i_1(t)|$
- Maximum Distance between Curves (D_2)
 - $D_2(i_k, i_1) = \max_{\Delta_k < t} |i_k(t + \Delta_k) - i_1(t)|$
- Area between Curves (D_3)
 - $D_3(i_k, i_1) = \sum_{\Delta_k < t} |i_k(t + \Delta_k) - i_1(t)|$

Truncated Power Law Network



Erdős-Rényi Network



Conclusion

- For both networks, as τ increases, the distance between the unit epi curve and the scaled epi curve decreases. Therefore, for higher values of τ the scaled disease can be more accurately predicted.
- Additionally, for small scale factors ($k < 50$) the scaled base curve is a close estimate of the true curve.