Homework 1

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Reminders:

- 1. Out of 100 points. 4 Questions. Contains 8 pages.
- 2. If you use Late days, mark how many you are using (out of maximum 4 available) at the top of your answer PDF.
- 3. There could be more than one correct answer. We shall accept them all.
- 4. Whenever you are making an assumption, please state it clearly.
- 5. You will submit a solution pdf LASTNAME.pdf containing your answers and the plots as well as a tar-ball LASTNAME.tgz that contains your code and any output files.
- 6. Please type your answers either in LATEX document or in a separate file like a Word document and then convert it into a pdf file. Typed answers are strongly encouraged. Illegible handwriting may get no points, at the discretion of the grader. Only drawings may be hand-drawn, as long as they are neat and legible.
- 7. Additionally, you will submit one tar-ball LASTNAME. tgz that contains your code and any results files. Code and results for each question should be contained in a separate sub-directory (Eg: Q1) and there should be a README. txt file for each sub-directory explaining any packages to install, command to run the code files and location of the expected output. Please follow the naming convention strictly.
- 8. If a question asks you to submit code please enter the file path (Eg: Q1/Q-1.3.1.py) in the solution pdf.
- 9. You can download all the datasets needed for this homework from canvas files, you can check the information about the datasets in the README.txt file.

1. (50 points) SIR Model

Q 1.1 (5 points) Everyone loves DS for Epi. It's the beginning of Fall 2022 semester and initially only 2% of the student population loves doing Data Science for Epidemiology (DSE). However, it is very contagious: once someone becomes interested in DSE they try to spread their interest to any uninterested people for a short time before they can no longer do so. Interest for DSE is also very persistent: those who develop interested in DSE stay that way. We model the spread of interest in DSE using an SIR model: Susceptible people are uninterested, infected students recently caught interest and can spread it to others, "Recovered" students are interested for life but can't spread anymore.

We observe that at the end of a very long semester $(t \to \infty)$ 60% of the student population are interested in DSE. What is the expected fraction of fellow students an interested (infected) student spreads their interest to?

Hint: You don't need to derive the ODE solution. You may use the results from the slides.

Solution:

Source: https://services.math.duke.edu/education/postcalc/sir/sir5.html We need to find the average number of individuals infected by a particular individual. This will become β/δ .

From the equations of s and i in the terms of t, $ds/dt = -\beta * s * i$ $di/dt = \beta * s * i - \delta * i$ Using both the equations applying chain rule we get, $di/ds = -1 + \delta/\beta * s$ For the simplicity of the equation, we replace β/δ with a variable γ . $di/ds = -1 + 1/\gamma * s$ Integrating the above equation, we get $i = -s + \ln(s)/\gamma + c$ At t = 0 and $t = \infty$, infected population i will be 0, $0.02 = -0.98 + \ln(0.98)/\gamma + c$ $0 = -0.40 + \ln(0.40)/\gamma + c$ Using them both, $0.6 = \frac{\ln(0.98/0.40)}{\gamma}$ $\gamma = \frac{\ln(0.98/0.40)}{0.6} = 1.493$

- Q 1.2 (13 points) **ODE** \Leftrightarrow **Networks** Let us try to derive the SIR ODE model from the SIR network model. Assume that the network is a clique of size N (i.e. everyone is connected to everyone else with a total population of N individuals). Further assume the model evolves in discrete time steps. At each step:
 - 1. each susceptible individual u picks a person v from the population, uniformly randomly
 - 2. if v is infected, then u gets infected with probability β .
 - 3. each infected individual goes into the R state with probability γ

Let X(t) be the number of individuals who are in the S state at a time t. Similarly, Y(t) is the number of individuals who are in state I and Z(t) be the number of individuals in state R.

Q 1.2.1 (2 points) What is the probability P(t) of a single susceptible individual getting infected at time t? Clearly the total number of expected new infections at time t+1 will be $P(t)\times X(t)$.

Solution: New infections at time t+1 should be equal to $\beta \times Y(t) \times X(t)$ as per the SIR Model.

Probability of single individual being infected at time t+1 should be $P(t) = \beta \times \frac{Y(t)}{N}$

Q 1.2.2 (4 points) What is the number of expected susceptible individuals at time t+1 i.e. what is $\mathbb{E}[X(t+1)]$? Write it down first in terms of P(t) and X(t) and then substitute P(t) from Q 1.2.1. Similarly, write down $\mathbb{E}[Y(t+1)]$ and $\mathbb{E}[Z(t+1)]$.

Solution:

The equation of updated susceptible population would have the original

susceptible and the change of susceptible population or new people getting infected. New people who get infected are $P(t) \times X(t)$.

$$\mathbb{E}[X(t+1] = X(t) - P(t) \times X(t)]$$

From Q 1.2.1, we substitute P(t) with $\beta \times \frac{Y(t)}{N}$.

$$\mathbb{E}[X(t+1] = X(t) - \beta \times \frac{Y(t)}{N} \times X(t)$$

Similarly,

$$\mathbb{E}[Y(t+1)] = Y(t) + \beta \times \frac{Y(t)}{N} \times X(t) - \gamma \times Y(t)$$

$$\mathbb{E}[(Z(t+1)] = Z(t) + \gamma \times Y(t)$$

Q 1.2.3 (2 points) Assuming $\mathbb{E}[X(t)] \approx X(t)^1$, write down the expression for X(t+1)-X(t)? Similarly, write down Y(t+1)-Y(t) and Z(t+1)-Z(t).

Solution:

With $\mathbb{E}[X(t)] \approx X(t)$, we substitute $\mathbb{E}[X(t)]$ with X(t+1).

$$X(t+1) - X(t) = -\beta \times \frac{Y(t)}{N} \times X(t)$$

Similarly,

$$Y(t+1) - Y(t) = \beta \times \frac{Y(t)}{N} \times X(t) - \gamma \times Y(t)$$

$$Z(t+1) - Z(t) = \gamma \times Y(t)$$

The steps above assumed 1 unit of time. Let us change the unit of time to a small Δt instead. Hence, β should now become $\beta \Delta t$ (ditto for γ).

Q 1.2.4 (2 points) Write down $X(t+\Delta t)-X(t), Y(t+\Delta t)-Y(t), \text{ and } Z(t+\Delta t)-Z(t), \text{ using your answers in Q 1.2.3 and the change mentioned above.}$

Solution:

Now, the previous values of X(t), Y(t), Z(t) would be unchanged and wouldn't change with the change in the decrease of the time step. But β and γ would change with the change in the time step and become a fraction of itself as they are measured over a single time step. Considering a uniformity across the time-step, β would become $\beta \Delta t$ and the same for γ also.

The equations start looking like this,

$$X(t + \Delta t) - X(t) = -\beta \Delta t \times \frac{Y(t)}{N} \times X(t)$$

$$Y(t+\Delta t) - Y(t) = \beta \Delta t \times \frac{Y(t)}{N} \times X(t) - \gamma \Delta t \times Y(t)$$

¹This is an example of the 'mean field' approximation, frequently used in the analysis of non-linear models.

$$Z(t + \Delta t) - Z(t) = \gamma \Delta t \times Y(t)$$

Q 1.2.5 (3 points) Starting from your answers in Q 1.2.4, derive the standard SIR ODE equations you have seen in class. Make sure you show all the steps.

Solution:

Taking $X(t + \Delta t) - X(t)$ equation first,

$$X(t + \Delta t) - X(t) = -\beta \Delta t \times \frac{Y(t)}{N} \times X(t)$$

$$\frac{X(t + \Delta t) - X(t)}{\Delta t} = -\beta \times \frac{Y(t)}{N} \times X(t)$$

To derive standard ODE we reduce the Δt time step to the smallest possible value tending to 0. Therefore we will put the limit to 0.

$$\lim_{\Delta t \rightarrow 0} \frac{X(t+\Delta t) - X(t)}{\Delta t} = -\beta \times \frac{Y(t)}{N} \times X(t)$$

This becomes the differential which will then look like this,

$$\frac{dX(t)}{dt} = -\beta \times \frac{Y(t)}{N} \times X(t)$$

Similarly, for Y(t) and Z(t),

$$\frac{dY(t)}{dt} = \beta \times \frac{Y(t)}{N} \times X(t) - \gamma \times Y(t)$$

$$\frac{dZ(t)}{dt} = \gamma \times Y(t)$$

- Q 1.3 (32 points) **Implementation and Calibration.** In this question you will implement standard SIR ODE and SIR network models.
 - Q 1.3.1 (6 points) Implement the standard SIR ODE model (lecture 3, slide 16-21) in your favorite language. Your code should take in the β and γ values, and the initial fractions S(0), I(0), R(0), and max_time as input, and give the fraction of infections, susceptible and recovered population (i.e. list of tuples (t, S(t), I(t), R(t)) for $t = [0, \ldots, max_time]$) at each time-step till max_time. Submit the code.

Set the values of parameters as $\beta = 0.1$, $\gamma = 0.05$ and initialize S(0) = 0.95, I(0) = 0.05, R(0) = 0. Show SIR curves (plot of S(t), I(t), R(t) over time) for up to T = 200.

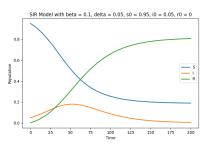
Now set $\beta = 0.05$, $\gamma = 0.1$ and plot the SIR curves. Do you notice any difference in nature of the curves? Explain.

(You may use your favourite programming language like Python, R, Matlab, C++ and Julia. Instructors are familiar with Python, R and Julia and may provide help with those. You may use basic arithmetic functions from numpy and math Python packages and packages like scipy package for integration. As a rule of thumb, make

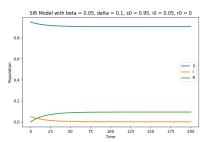
sure the packages you use only help with arithmetic operations or integration and don't solve the problem for you. If in doubt, please ask the instructors.)

Solution:

The plot with initial parameters of $\beta = 0.1$, $\delta = 0.05$, S(0) = 0.95, I(0) = 0.05 and R(0) = 0, looks like this



The plot with initial parameters of $\beta = 0.05$, $\delta = 0.1$, S(0) = 0.95, I(0) = 0.05 and R(0) = 0, looks like this



With the $\beta < \delta$, the infection dies out before even starting as the infected folks start to recover before getting a chance to infect the susceptible population.

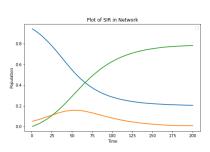
Q 1.3.2 (10 points) Implement the discrete-time SIR Network model in your favorite language. Your code should take a graph-edge-list, the total number of nodes in the graph N, the β value, the γ value, the initial fractions S(0), I(0), R(0), and the max_time as input, and give the fraction of infected nodes I(t), susceptible nodes S(t) and recovered nodes R(t) at each time-step till max_time. You can set any random set of nodes to I/R/S states at time t=0 to reach the desired fraction. Submit the code.

Also provide SIR plots for the network simulation with N=1000 given the initialization parameters from previous question Q 1.3.1.

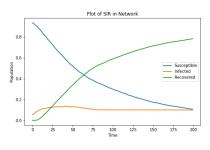
(As the SIR network model is stochastic, make sure you run the simulation 50 times, and take the average to get I(t), S(t), R(t). Further you can use packages to construct graphs like the networkx Python package or SimpleGraph, LightGraph Julia packages.)

Solution:

Code and graph in the tarball archive. Network when taking N=1000 cliques



Network when using the example network



Q 1.3.3 (8 points) Calibrate your SIR ODE model using the COVID mortality time-series for GA from July 2021 - Nov 2021 (download from canvas). The dataset contains cumulative fraction of cases and deaths observed each week. You will be using deaths as ground truth $R_{observed}(0) \dots, R_{observed}(T)$. The objective function to minimize is

$$\mathcal{L} = \sum_{t=1}^{T} (R(t) - R_{observed}(t))^{2}$$

as given in slide 8, lecture 4 (assume the R state corresponds to mortality).

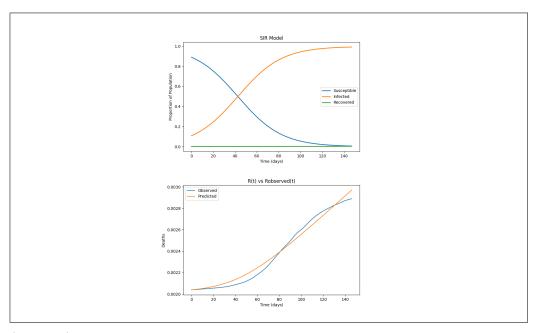
To initialize the boundary conditions, we set the initial values of I(0) as the fraction of cases and R(0) as fraction of deaths on the first day from the dataset. Set the initial value of S(0) as S(0) = 1 - R(0) - I(0). Now calibrate the SIR model to find the best set of parameters β, γ that minimizes the objective function \mathcal{L} .

Write down the values of β , γ you find after calibration. Plot S(t), I(t), R(t) vs time (just like the plots we saw in slide 19, lecture 3). Also plot another figure showing $R_{observed}(t)$ you used to calibrate and the resultant R(t) after calibration. Submit the plots.

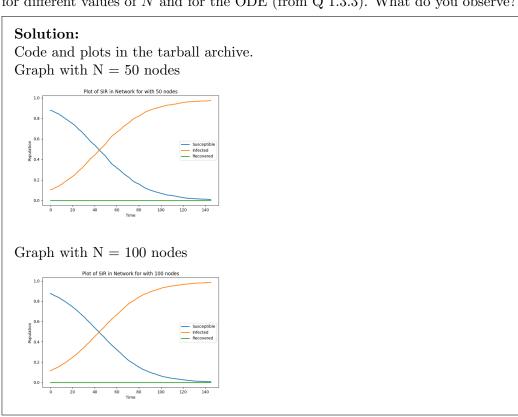
(Hint: You may use pre-built optimization functions/packages. For Python you may consider using the scipy.optimize.minimize function from scipy.optimize import minimize. Similarly, you may use DifferentialEquations Julia library.)

Solution:

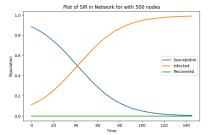
Attaching the plots here as well. Code in the tarball. Optimal parameters of β and γ are 0.0497 and 9.1278 × 10⁻⁶ respectively.



Q 1.3.4 (8 points) Run the SIR Network model on a clique of size N=100. Set the β and γ values as the ones you get in Q 1.3.3. Set the max_time as 900. For the simulation, set the initial number of infected individuals $I0 = \lfloor I(0) * N \rfloor \rfloor$ (where I(0) is value you set in Q1.3.3), initial number of susceptibles $S0 = \lfloor (S(0) * N) \rfloor$, and the remaining nodes as R0 (i.e. $R0 = N - \lfloor I(0) * N \rfloor - \lfloor S(0) * N \rfloor$). Note that $\lfloor x \rfloor$ is the function which just rounds x to closest integer less than x. Plot the S(t), I(t), R(t) vs t on a plot as in Q1.3.3 (submit the plot). Repeat the same steps for N=1000 and N=10,000. Submit the plots. Compare the plots for different values of N and for the ODE (from Q 1.3.3). What do you observe?



Graph with N = 500 nodes



With the different values of N and the ODE model plots, it's quite evident that with the increase of nodes the discreteness of the plots reduce and the plots become more smoother due to the presence of more data-points to plot.

2. (5 points) Cayley Trees

Consider a symmetric k-regular tree in which each vertex is connected to the same number k of others, till we reach the leaves. This is also sometimes called a 'Cayley' Tree.

For example, if k = 3 the network will look like Figure 1.

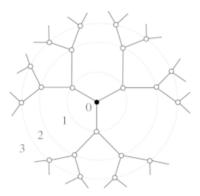


Figure 1: k-regular Cayley tree for k=3

Q 2.1 (2 points) In a k-Cayley tree, what is the number of nodes reachable in d steps from the central node? (e.g., the number of nodes reachable in d=1 steps in the figure above is equal to 3). Show your derivation.

Note: A node u is reachable from the center node in d steps if there is a path of exactly d steps from center node to node u.

Solution:

Considering a k-Cayley tree with each k-1 children nodes, the first step would be encountering k paths or nodes. For the next layer it will be k-1. And for the next as well. This indicates an exponential relation with number of nodes reachable is proportional to the d exponential of each number of children nodes.

This indicates the series $k * (k-1) * (k-1) * (k-1) * \dots$ uptil d terms which will eventually become, $k \cdot (k-1)^{(d-1)}$ for a node reachable in d steps.

Q 2.2 (3 points) What is the diameter of such a network (in terms of k and n)? The diameter of a graph is defined as the longest shortest path in the graph. Let the length of the shortest path between a pair of vertices be called the **distance** between the vertices. The maximum distance between any pair of vertices is called the diameter of the graph. Please show your steps. Here n is the total number of nodes in the graph.

Solution:

The diameter of the k-Cayley tree will be 2d as it takes d steps to reach from central node to exterior node.

Now we know the total number of nodes is n which is equivalent to the sum of all nodes present from the first layer to uptill the d layer. This becomes

$$n = k.(1 + (k-1) + (k-1)^2 + (k-1)^3 + \dots + (k-1)^{(d-1)}) + 1$$

Being a normal geometric progression expression we can find the sum.

$$n = k * \frac{((k-1)^d - 1)}{(k-1)-1} + 1$$

Upon simplification, we get

$$\frac{n(k-2)+2}{k} = (k-1)^d$$

Getting the expression in terms of $2d=2.\frac{\log(n(k-2)+2)-\log(k)}{\log(k-1)}$

Q 2.3 (6 points) [Bonus] Suppose we take our original k-Cayley tree with a depth d and add a node to it. Then, we add an edge from the added node to any other node in the tree. What is the expected value of the diameter of the resulting graph?

If instead of adding a node we add an edge between a pair of vertices that are not previously connected, what is the maximum length of resulting cycle?

Solution:

The diameter of our k-Cayley tree only increases if the node added is on the boundary nodes or extremities or leaves of the network.

This will become

 $\mathbb{E}[D] = 2.d + P(\text{node attached to the leaves of the network})$

Substituting the values,

$$\mathbb{E}[D] = 2.d + \frac{k.(k-1)^{(d-1)}}{k*\frac{((k-1)^d-1}{(k-1)-1})+1}$$

3. (20 points) Calibrating IC Models

In this question we will try out a simple way to calibrate the IC model on a graph, given

some cascade traces. Recall that the IC model is just an SIR model on a network such that any infected node cures itself in 1 time-step, and each edge (v, u) may have a different constant beta probability p_{vu} . Remember that p_{vu} may not be equal to p_{uv} .

We will work with the following two datasets (download from canvas):

- 1. Network: network.txt ² Note that the graph is directed. Edge $v \to u$ is denoted as (v, u).
- 2. Cascades/Action log: Ratings.timed.csv ³.

This file contains an action on each line. An action is a user rating a movie. That is, if a user v rates "The King's Speech", and then later v's friend, say u, does the same, we consider the action of rating "The King's Speech" as having propagated (cascaded) from v to u. In epidemiological terms you can think of v 'infected' u.

In this setting, the goal of the calibration procedure is to find out the probability p_{vu} over each edge (v, u).

- Q 3.1 (5 points) Let A_{v2u} be the total number of actions of node v which cascaded to node u later. An action cascades from v to u if both the following conditions satisfy:
 - 1. v must have taken the action before u.
 - 2. Edge (v, u) exists. In other words, v is an in-neighbor of u.

Similarly, let A_v be the total number of actions of node v in the database. Then you can think of node v making A_v tries to infect node u, of which only A_{v2u} tries succeeded. Write down the probability L of A_{v2u} tries succeeding out of A_v tries, assuming the probability of success is p_{vu} . L is also called the likelihood of the data.

Solution:

 $Source: \ https://stats.stackexchange.com/questions/181035/how-to-derive-the-likelihood-function-for-binomial-distribution-for-parameter-es$

The general log likelihood function where its derivative is zero should look like $ln(nC_x * p^x * (1-p)^(n-x)) = ln(nC_x) + x * ln(p) + (n-x)ln(1-p)$

Taking a partial derivative w.r.t. the probability p and setting it to 0.

$$\frac{d}{dp} * ln(nC_x) + xln(p) + (n-x)ln(1-p) = 0$$

$$\frac{x}{p} - \frac{n-x}{1-p} = 0$$

$$p = \frac{x}{n}$$

This becomes when we substitute successful infection x for A_{v2u} out of possible infections n for A_v . Thus,

$$p_{vu}^* = \frac{A_v 2u}{A_v}$$

²The original much larger raw file we constructed this from is here: http://konect.cc/networks/flixster/

³The original much larger raw file we constructed this from is here: https://sites.google.com/view/
mohsenjamali/flixter-data-se

We find the Maximum Likelihood Estimator p_{vv}^* . i.e

$$p_{vu}^* = \arg\max_{p_{vu}} L = \frac{A_{v2u}}{A_v}.$$

(Try to derive it on you own.)

- Q 3.2 (15 points) Now we calibrate the IC model to estimate $p*_{vu}$ for each edge (v,u). Using the two datasets given above, write code to calculate the p^*_{vu} for each edge (v,u) of the network. The code should take in the two datasets as input, and print " v,u,p^*_{vu} " (without quotes) on each line for each edge (v,u). Submit both the code and the output you get after running it on the two datasets.
 - *Note 1:* If there two actions which happen on the same date in the cascades/action log, you can consider the one which appears earlier in the file to be temporally earlier (i.e. use line numbers to break the ties).
 - Note 2: There are many users u who have zero actions. In such cases you may set $p_{vu}^* = 0$.
 - Note 3: There are multiple ways to do this efficiently. We will not grade on efficiency of your code. However, it might be interesting for you to hear that you can do this in < 2 scans of the action/cascades log.

Solution:

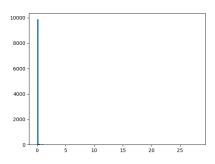
The code is in the tarball with the results in a file called Q-3.2-Output_file.txt.

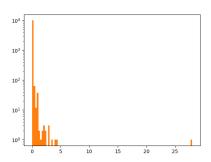
Q 3.3 (5 points) [Bonus] Let $p_u^* = \sum_v p_{uv}^*$ i.e. if we consider the p_{uv}^* 's as the weights on the network, then p^*u is just the weighted out-degree degree of each node u. Plot this weighted out-degree distribution. Make the y-axis log-scale. Submit the plot. What do you observe?

Hint: You can use the numpy function histogram() for this.

Solution:

The weighted out-degree distribution plots attached below.





With the frequency of the degrees being substantially higher, the log scale provides a clearer picture of the weight distribution.

4. (20 points) Zombie Apocalypse

There is a new disease outbreak in a town that turns people into zombies.

- Q 4.1 (5 points) Stopping the outbreak using branching process. In the initial stage of the outbreak, the zombies don't harm humans, they only infect them with the disease. The town has set aside 2 million dollars to deal with this outbreak which is currently in its infancy. The public health officials have two measures to slow down the outbreak:
 - 1. Controlling how many people a zombie comes in contact with: If the officials spend x dollars, they expect a zombie to come in contact with $40 \frac{x}{200,000}$ people.
 - 2. Controlling the probability of transmission: If the officials spend y dollars, they expect the probability of a zombie transmitting the disease to another person it comes in contact with is $0.04 \frac{y}{100.000.000}$ dollars.

What is the best way to allocate the 2 million dollar budget?

(Assume that most people are uninfected. Therefore, number of people an infected zombie comes in contact with is assumed to be constant.)

Hint: Refer to Lecture 4 Slides 36-41.

Solution:

From the hint given and the model being branching processes, the probability that there is an infected node at depth h is

$$p_h = 1 - (1 - q.p_{h-1})^d$$

For the infection to stop we try to reduce p_h so that it tends to zero. For this condition to be true, q.d < 1.

Thus, to stop the spread of the infection, the equation becomes,

$$(40 - \frac{x}{200000})(0.04 - \frac{y}{100000000}) < 1$$

With the budget constraints this essentially becomes a quadratic equation. Substituting y for 2000000-x

$$(40 - \frac{x}{200000})(0.04 - \frac{2000000 - x}{100000000}) < 1$$

Solving the roots, we get an idea that x has to be in the range of [0,763932) dollars.

- Q 4.2 (15 points) You failed to stop the outbreak. Now the zombie disease has turned sinister. The infected zombies don't merely turn humans to zombies, instead they consume
 - Q 4.2.1 (6 points) Assume x(t) is the population that is human and y(t) the population that are zombies at time t. We use the Lotka-Volterra Model (LVM) to model the dynamics of the populations:

$$\frac{dx}{dt} = \alpha x - \beta xy
\frac{dy}{dt} = \gamma xy - \delta y$$
(1)

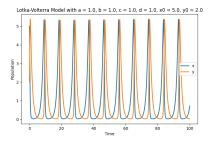
$$\frac{dy}{dt} = \gamma xy - \delta y \tag{2}$$

Similar to Q 1.3.1, implement the ODE model above that takes in parameters $\alpha, \beta, \gamma, \delta$ and initial states x(0), y(0) and returns fraction of populations x(t), y(t)till time-step max_time. Submit the code for the model.

Set $\alpha = \beta = \gamma = \delta = 1$. Now set the values of initial population x(t) = 5, y(t) =2 and plot the values of x(t), y(t) vs. t till T = 100. Submit the plot. Now experiment with two other initializations of x(0), y(0) for which the values of x and y don't change over time.

Solution:

Code and plots in the tarball archive.



Only two other initializations exist

where x and y do not change is when both and x(0) and y(0) initialize at (0,0) and (1,1).

Q 4.2.2 (3 points) The zombie disease has mutated further! Now you realize that there are actually two variants of zombies that are consuming humans. Assume x(t) is the population that is human and y(t) the the population that are zombie variant 1 and z(t) the the population that are zombie variant 2 at time t.

We model the population dynamics as a variant of LVM called LVM2:

$$\frac{dx}{dt} = \alpha x - \beta xy - \phi xz \tag{3}$$

$$\frac{dx}{dt} = \alpha x - \beta xy - \phi xz \tag{3}$$

$$\frac{dy}{dt} = \gamma xy - \delta y \tag{4}$$

$$\frac{dz}{dt} = \rho xz - \epsilon z \tag{5}$$

$$\frac{dz}{dt} = \rho xz - \epsilon z \tag{5}$$

What are the fixed points of this LVM2 model?

Solution:

To find the fixed points of these equations, we equate these equations to 0 to get the fixed points of x, y and z.

The equations then start looking like this,

$$\alpha x - \beta xy - \phi xz = 0$$

$$\gamma xy - \delta y = 0$$

$$\rho xz - \epsilon z = 0$$

Solving them we get,

keeping x = 0,

we get the fixed point at origin (0,0,0).

keeping y = 0,

we get fixed point at $(\frac{\epsilon}{a}, 0, \frac{\alpha}{b})$

keeping z = 0,

we get fixed point at $(\frac{\delta}{\gamma}, \frac{\alpha}{\beta}, 0)$

Q 4.2.3 (5 points) Implement the model in your favourite language that takes in parameters $\alpha, \beta, \gamma, \delta, \phi, \rho, \epsilon$ and initial states x(0), y(0), z(0) and returns fraction of populations x(t), y(t), z(t) till time-step max_time. Submit the code.

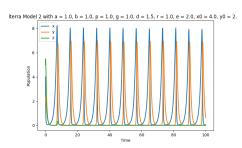
Assume $\alpha = \beta = \phi = \gamma = \rho = 1$, $\delta = 1.5$ and $\epsilon = 2$. For x(0) = 4, y(0) = 2, z(0) = 5 plot x(t), y(t), z(t) vs. t till T = 100. Also plot for other initializations of x(0), y(0), z(0) for which they are fixed points and show that the population doesn't change over time. Submit the plots.

Solution:

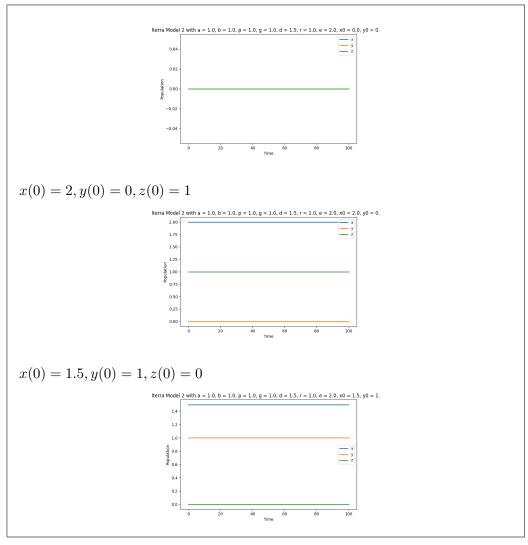
Code and plots are in the tarball archive.

Plot for the following initialization

$$x(0) = 4, y(0) = 2, z(0) = 5$$



$$x(0) = 0, y(0) = 0, z(0) = 0$$



Q 4.2.4 (1 point) Are the LVM2 plots periodic?

Solution:

For the entirety they are not periodic, but become periodic once the z population dies out.

5. (5 points) Closing Triangles

Imagine a social network consisting of the students in this class (if x knows/is friends with y, then there is an undirected edge between x and y). Let us create a subgraph centered around you. First, list down all the people from the class you know. These students are your neighboring nodes in the graph. Now, for each of your neighbors, ask them to list down the people they know from the class. Connect these people to your subgraph as well. These are your 'friends-of-friends' (f-o-f).

Naturally some of your f-o-f will also be your friends. These create 'triangles' centered around you in the graph.

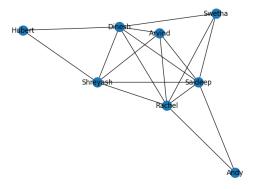
1. Draw the constructed subgraph with names of your friends or f-o-f on the nodes. (You can use any plotting package or submit a handwritten graph.)

- 2. How many triangles are you already part of?
- 3. How many <u>incomplete</u> triangles are you part of? These are exactly the people who are friends of your friends but not (yet) yours aka 'potential' friends:-)

Note: You have to name people in the constructed graph. Feel free to use any method to get this answer: Face-Face meetings, Phone calls, Social Media, Zoom/Teams chats.

Solution:

1. Constructed subgraph



- 2. Shreyash is part of 7 triangles. (H&D, D&Ar, D&Sa, D&R, Ar&Sa, Ar&R, Sa&R)
- 3. Shreyash has 2 friends-of-friends who aren't his friends yet.

 Shreyash isn't a friend of Swetha and has 3 incomplete triangles with her through Dinesh, Rachel and Saideep.

 Shreyash isn't a friend of Andy and has 2 incomplete triangles with him through Saideep and Rachel.

The total incomplete triangles that Shreyash is part of are 5.