## BNFO 262 2024

**Homework 2:** single cell RNA-seq and Networks (Robert Morey and Hannah Carter)

## Instructions

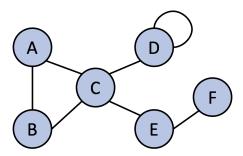
Answer the following questions in your own words and upload a PDF of your answers to Gradescope. Make sure to write your name and PID at the start of your answers. This assignment is due **2/15/24** at **9:00AM**.

## Part 1: Single cell RNA-seq

- 1. Describe the experimental differences between droplet based methods and physical separation methods. Start by explaining each. Then discuss when it might be better to use one or the other. Limit 5 sentences. (3 points)
- 2. When analyzing scRNA-seq data, how do you know which cell a read comes from? Limit 5 sentences. (1 point)
- 3. Describe what a doublet is and how it is generated. Why do you not want doublets in your dataset when you perform analysis? Limit 5 sentences.(2 points)
- When preprocessing scRNA-seq data, there are multiple metrics to perform QC on.
  Please list three QC metrics and explain what each controls for? Limit 5 sentences. (3 points)
- 5. Come up with an experiment in which you'd prefer to use scRNA-seq rather than RNA-seq. Justify why you believe this assay would be preferable for this experiment. Limit 5 sentences. (2 points)

## Part 2: Networks

6. Answer the following questions according to the graph (3 points)



- a. What is the adjacency matrix of this graph? (1 point)
- b. List the degree of each node in this graph (1 point)
- c. Which node has the highest degree of centrality? Why? (1 point)
- 7. Give an example of a biological network. Describe the node and edge. Explain what is the biological meaning of a node with high centrality in this network. Limit 5 sentences. (3 points)