



- 1) Use a graphical or flowchart format to describe the steps of purification starting with a single colony growing on an agarose plate to the final dialyzed purified protein.
- 2) How do you know it is "OK" to switch from wash to elution?
- 3) How is your protein binding to the column? What is in the elution buffer to remove the protein from the resin/chromatography medium.
- 4) Give a detailed description on what you will do to elute and capture your protein.