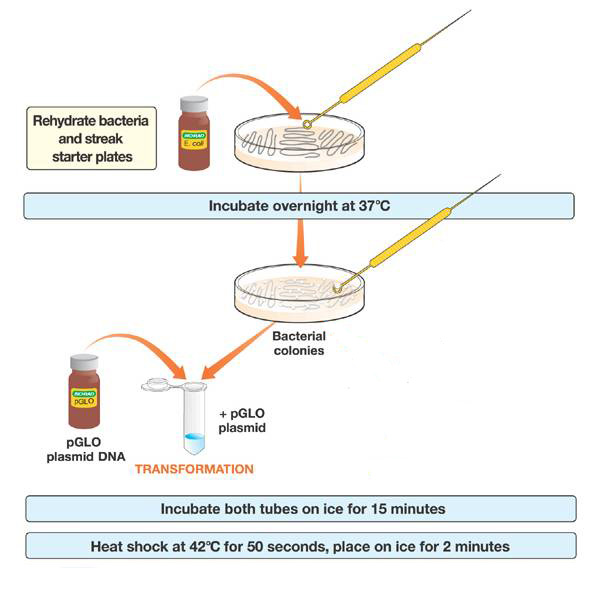
Name: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

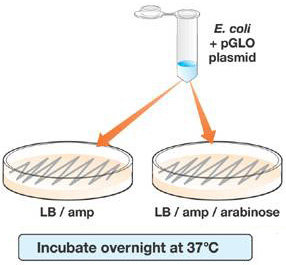
Going Over Lab 4.1 pGlo Bacteria Transformation & Recombinant DNA



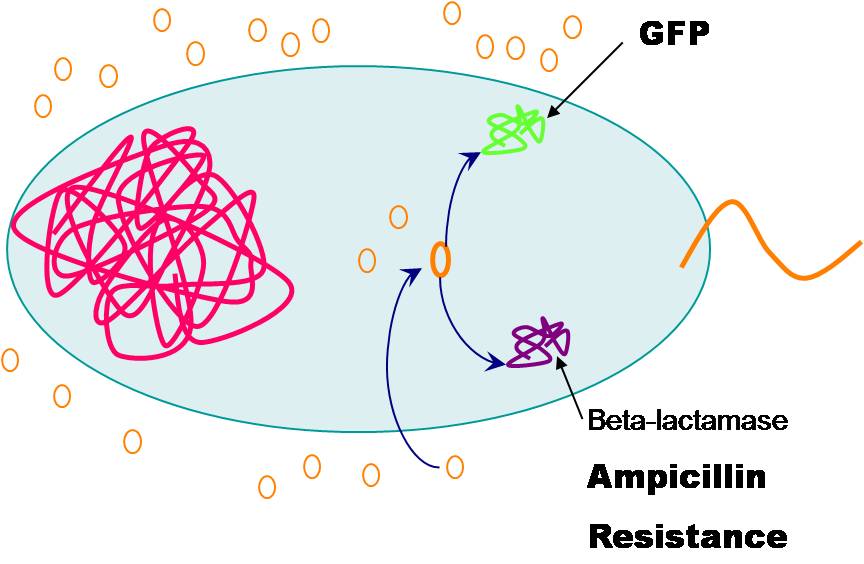
Summarize what we did on each of the days of the lab:

Previous to lab (teacher prep):

Lab Day:



1. The GFP (Glow Fluourescence Protein) gene had already been spliced into the bacterial plasmid. Explain how we inserted the plasmid into the bacteria on lab day:

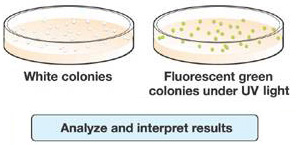


Summarize:

Day 2 - Analysis:

No Sugar

Sugar

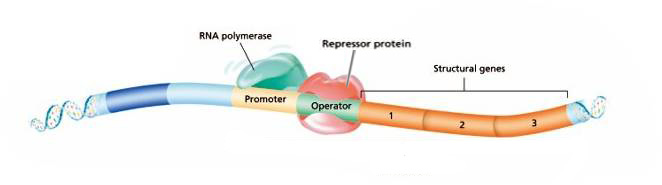


2. Explain what actually glows?

pGLO_results.jpg                                               00016E3F
KSD Server                     B471509A:

3. Bacteria grew on each of the new plates, and all of the bacteria had the plasmid inserted, so why didn’t the bacteria grown on the plate without sugar added glow? Inversely, why did the plate with sugar added allow for the bacteria to glow?

Explain how the GFP gene could be *expressed* or *not* *expressed*?



GFP Gene