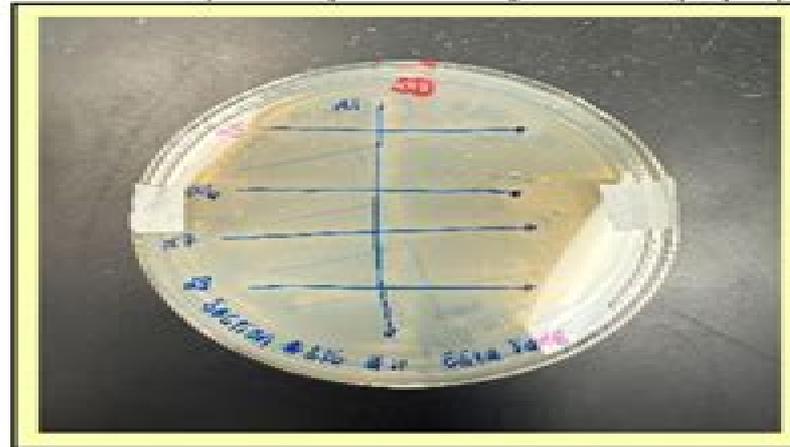


| | | | | | |
|-----------|--|------------|--|---------------|--|
| Last Name | | First Name | | Lab Section # | |
|-----------|--|------------|--|---------------|--|

1) (0.5 pt) Show an image of the bottom of your SD plate showing that it was properly labeled.



2) (0.5 pt) The protocol for cleaning the PCR sample includes a binding step, a washing step and an elution step. Accidentally using the elution buffer to "wash" the column would result in the DNA being lost in the "wash" step. Why?

Using the elution buffer would lead to DNA's binding affinity to the silica in the spin column decreasing because of the elution buffer's low salt concentration. Then during centrifugation, the DNA will get spun down the column because of its lack of affinity to the spin column and be discarded because it will be collected in the flow-through of the tube.

3) (1 pt) You set up 4 sequencing reactions using 30 ng of your PCR product for each. Include an image of the gel you used **and show the calculations** you performed to determine how much volume of your PCR product you had to use to obtain 30 ng of DNA. (If you did not get any products, then use the positive control bands on the Bench 1 gel.)

Gel image (Label which samples are in each lane):

Calculations:

Bio Lab Section 2genetics Answers

Daniela Niemeyer



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