

## Lab 6 – Gel Electrophoresis Extra Credit

**10 points possible**

**Exercises:**

1. Use a standardized DNA marker of known DNA fragment sizes to design a standard curve.
2. Apply the standard curve to estimate fragment size of your PCR fragment product

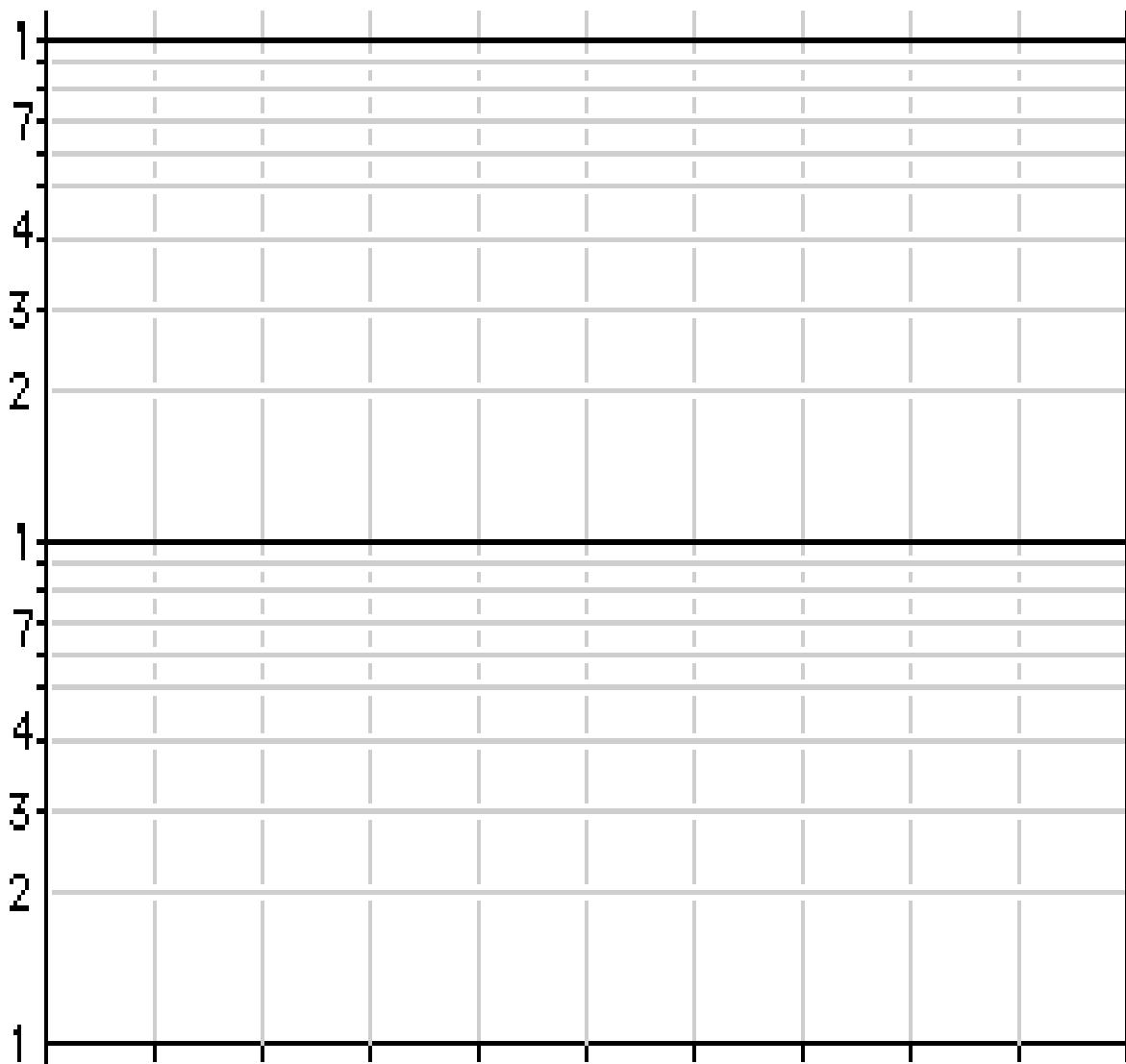
### **Exercise 1:**

**(6pts)**

1. Print out the picture of the gel from class with your PCR product results.
2. Using the lanes marked DNA Marker, measure from the wells the distance in **mm** each DNA marker band traveled and note the distance traveled in the tables below.
3. Draw a standard curve using the semi-log graph provided below. Must include proper axis labels and graph title.

<b>Standard DNA Marker</b>	
<b>Fragment size (bp)</b>	<b>Migration distance (mm)</b>
10,000	
8,000	
6,000	
5,000	
4,000	
3,000	
2,500	
2,000	
1,500	
1,200	

<b>Standard DNA Marker</b>	
<b>Fragment size (bp)</b>	<b>Migration distance (mm)</b>
1,000	
900	
800	
700	
600	
500	
400	
300	
200	
100	



**Exercise 2.**

**(4 pts)**

1. Using the correctly made standard curve above approximate the size of your PCR product. Draw on the graph how you obtained the size to receive full credit.

**Approximate size of PCR product** \_\_\_\_\_