King Abdul-Aziz university Faculty of science biological science department Women section Mark: 11



Introduction to proteomics (Bio 405) Lab final exam (written exam)

2<sup>nd</sup> term of 1434-1435 Date: 08/07/1435

Time: 1 hour

Name:	Computer Number				
<u>ivairie.</u>	First question  Please read the following carefully and mark it with True or False and correct the False:  ( /1 Mark)				
	1. The cationic form is green or red with absorbance at 470 nm. (T)				
	Correction:				
	2. Phosphoric acid is one of the Bradford reagents. (T)				
	Correction:				
	Second question				
	Choose only one correct answer on the following questions: ( /5 Marks)				
	<ol> <li>In SDS-PAGE experiment, before loading the samples into the gel</li></ol>				
	<ul> <li>In order to visualize the separated proteins on SDS-PAGE, proteins must be</li> <li>A. Stained with coomasiie brilliant blue R.</li> <li>B. Stained with bromophenol blue.</li> </ul>				

D. Placed in the destaining solution for 1-2 hours, and then placed in the

coomassie staining solution for 30 minutes to 2 hours.

C. Heated for 5 minutes at 100°C.

3)	SDS-Page is used in the molecular biology for A. Extracting protein. B. Lysing cells and extracting the total proteins.
	C. Measuring protein concentration
	D. Analyzing protein concentration.
4)	Bradford assay :
,	<ul><li>A. Depends on the sodium dodecyl sulfate that is present in the reagents.</li><li>B. Is a spectroscopic analytical procedure used to measure the concentration of protein in a solution.</li></ul>
	C. Depends on bovine serum albumin protein in the solution.
	D. Is a colorimetric protein assay that can analyze the number and the size of polypeptide subunits.
5)	One of the following is <b>not</b> one of the protein quantification methods:
	A. Bicinchoninic acid.
	B. Near UV absorbance.
	C. Sonication.
	D. 2-D Quant kit.
6)	Detergent which is used with protein to be separated:
٠,	A. Nonidet P-40.
	B. Triton x-100.
	C. Sarkosyl.
	D. All of the above.
7)	induce the formation of large pore gels.
.,	A. Polymers such as polyethylene glycol.
	B. Tris.
	C. Glycine.
	D. Glycerol.
8)	begins to breakdown almost immediately when dissolved in water, therefore, the accumulation of water in results in a rapid loss of reactivity. This is why solutions should be prepared fresh daily.  A. Acrylamide.

B. Ammonium persulfate.

D. Bis acrylamide.

C. Urea.

9)	factor that can affect the homogeneity of polyacrylamide gels.  A. Protein size.				
	B. Isoelectric point of polypeptides.				
	C. Gel thickness.				
	D. None of the above.				
10) 7	The BSA standard curve is called:				
	A. Bovine spectrophotometer analytical of	curve.			
	B. Protein purity curve.				
	C. Calibration curve Bradford macro assa	ay.			
	D. Melting curve assays.				
Third qu	uestion				
Fill in th	ne blanks with the correct words:	(	/ 5 Marks)		
1	low acrylamide concentrations are to separatehigh molecular weight proteins, whilehigh acrylamide concentrations are used to separate proteins oflow molecular weight.				
2	In SDS-PAGE , the polymerization is initiated byTEMED andAPS				
3	Gels less than3% acrylamide are almost fluid . So, the effective range of a polyacrylamide gel is between5-20% for a uniform gel concentration.				
4					
5	The effective resolving range of polyacryl pore size,buffers,pH	=			

- 6. Polymerization that is too fast ...< 10 minutes.... Or too slow .....> 60 minutes.... Leads to non uniform polymerization.
- 7. ....estimation of protein size... and..... estimation of protein purity.... are two of the reasons of SDS-PAGE usages.

مع تمنياتي لكن بالتوفيق والنجاح استاذه ريم الشريف