

BRAINWARE UNIVERSITY

Term End Examination 2019 - 20

Programme - Master of Science in Biotechnology

Course Name - Analytical Technique

Course Code - MBT104

(Semester - 1)

Time allotted: 2 Hours 30 Minutes

Full Marks: 60

 $20 \times 1 = 20$

[The figure in the margin indicates full marks. Candidates are required to give their answers in their own words as far as practicable.]

Group -A

(Multiple Choice Type Question)

1. Answer any twenty from the following Which of the following is not the function of buffer? (i) a. Prevent cell lysis b. Chelating the metals d. Prevent lipase action c. Prevent protease action Which of the following techniques uses sound waves for cell disruption? (ii) b. Sonication a. Homogenization d. Mortar and Pestle c. Blender Which of the following is not the product of cell disruption? (iii) b. RNA a. DNA d. Water c. Protein Which of the following is a physical method of cell disruption? (iv) b. Using solvents a. Using detergents d. Using osmotic shock c. Using French press Which of the following options is not a stage of product recovery? (v) b. Isolation of organism a. Removal of solids d. Cell disruption c. Purification and concentration Who employed the term 'Chromatography'? (vi)

a. Tsvet

c. Richard

b. Archer

d. Erika

Whic	ch of the following options is used to pa	ack co	lumns in adsorption chromatography			
	~ 4		Silica gel			
c.	Potassium hydroxide	d.	Aluminium oxide			
Which of the following stationary phases is not used in gel filtration chromatography?						
	•		Sephacryl			
c.	Bio-Gel	d.	Resin beads			
Immunoaffinity chromatography is used for the purification of						
			Interferons			
c.	Antibodies	d.	Carbohydrates			
The HPLC uses the application of						
		b.	Low temperature			
c.	High pressure	d.	Low pressure			
Mobile phase can be						
a.	gas or liquid	b.	solid or liquid			
c.	only solid	d.	only gas			
Which of the following is used in electron microscope?						
a.	electron beams	b.	magnetic fields			
c.	light waves	d.	electron beams and magnetic fields			
In 500 × g, what does g represent in accordance to centrifugation						
a.	Gravitational force	b.	Centrifugal force is 500 times greater than earthly gravitational force			
c.	Centrifugal force is 500 times less than earthly gravitational force	d.	Centrifugal force is 500 times same as that of earthly gravitational force			
Resolving power of a microscope is a function of						
a.	Wavelength of light used	b.	Numerical aperture of lens system			
c.	Refractive index	d.	Wavelength of light used and numerical aperture of lens system			
At what speed do you centrifuge blood?						
a.	2200-2500 RPM	b.	3000-3200 RPM			
c.	1000-1500 RPM	d.	4000 RPM			
Gram staining technique was developed by						
a.	Alexander Fleming	b.	Hans Christian Gram			
c.	Joseph Christian Gram	d.	Robert Gram			
The compound microscope consists of two lenses known as						
o Objective & Eveniese						
c.	Eyepiece & Occular	d.	None.			
	a. c. Whice a. c. Immu a. c. The I a. c. Mobi a. c. Whice a. c. In 50 a. c. Resol a. c. C. Cram a. c. The cc a.	a. Carbon c. Potassium hydroxide Which of the following stationary phases is a. Sephadex c. Bio-Gel Immunoaffinity chromatography is used for a. Lipoproteins c. Antibodies The HPLC uses the application of a. High temperature c. High pressure Mobile phase can be a. gas or liquid c. only solid Which of the following is used in electron m a. electron beams c. light waves In 500 × g, what does g represent in accorda a. Gravitational force c. Centrifugal force is 500 times less than earthly gravitational force Resolving power of a microscope is a functional wavelength of light used c. Refractive index At what speed do you centrifuge blood? a. 2200-2500 RPM c. 1000-1500 RPM Gram staining technique was developed by a. Alexander Fleming c. Joseph Christian Gram The compound microscope consists of two least opening the compound microscope consi	c. Potassium hydroxide Which of the following stationary phases is not use a. Sephadex c. Bio-Gel d. Immunoaffinity chromatography is used for the properties a. Lipoproteins b. c. Antibodies d. The HPLC uses the application of			

(xviii)		th of the following cannot be used natography?	l as	adsorbent in Column adsorption		
	a.	Magnesium oxide	b.	Silica gel		
	c.	Activated alumina	d.	Potassium permanganate		
(xix)		olumn chromatography, the stationary phe is made of	ase i	is made of and the mobile		
	a.	Solid, liquid	b.	Liquid, liquid		
	c.	Liquid, gas	d.	Solid, gas		
(xx)	In Thin layer chromatography, the stationary phase is made of and the mobile phase is made of					
	a.	Solid, liquid	b.	Liquid, liquid		
	c.	Liquid, gas	d.	Solid, gas		
(xxi)	In which of the following type of paper, chromatography does the mobile phase move horizontally over a circular sheet of paper?					
	a.	Ascending paper chromatography	b.	Descending paper chromatography		
	c.	Radial paper chromatography	d.	Ascending – descending chromatography		
(xxii)	The charged molecules can be separated by					
	a.	Ascending paper chromatography	b.	Descending paper chromatography		
	c.	Radial paper chromatography	d.	Ascending-Descending chromatography		
(xxiii) Which of the following is used to separate molecules based on affinity?				les based on affinity?		
	a.	Column chromatography	b.	Ion exchange chromatography		
	c.	Thin layer chromatography	d.	Affinity chromatography		
(xxiv)	Liquid chromatography can be performed in which of the following ways?					
	a.	Only in columns	b.	Only on plane surfaces		
	c.	Either in columns or on plane surfaces	d.	Neither in columns nor on plane surfaces		
(xxv)	Patte	rn on paper in chromatography is called		. ∠ <i>n</i> ′′ • ₹		
	a.	chroming	b.	chroma		
	c.	chromatograph	d.	chromatogram		

		Group – B	
		(Short Answer Type Questions)	$4 \times 5 = 20$
An	swer a	any four from the following	
2.		Write short notes on buffer solution.	5
3.		Write the Basics principle Of Chromatographic Process?	5
4.		What are the main differences between High Performance Liquid Chromatography?	5
5.		Explain the terms Rf Value and their importance.	5
6.		Briefly explain about the different methods of cell disruption.	5
7.		Briefly explain the different types of non-mechanical enzymatic methods.	5
		Group – C	2 10 20
		(Long Answer Type Questions)	$2 \times 10 = 20$
Ans	wer a	ny two from the following	
8.	(a)	Write short notes on French press.	5
	(b)	What do you mean by acidic buffer and how can you prepared in laboratory	y. 5
9.	(a)	Define Buffer.	2
	(b)	Write the Basic principle Of Chromatographic Process?	4
	(c)	What is buffer solution? Write the characteristics of buffer solution.	4
10.	(a)	What is cell disruption? Enlists the importance of cell disruption.	2+5
	(b)	What do you mean By The Term Rf Value?	3
11.	(a)	Explain the concepts of cell disruption? Enlists the importance of c disruption.	ell 2.5+2.5
	(b)	Write the Basic principle Of Chromatographic Process?	