## BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI (END SEMESTER EXAMINATION)

CLASS: BRANCH	(END SEMESTER EXAMINATION) : M. SC. & PRE-PHD SE CH: BIOENGINEERING & BIOTECHNOLOGY SE						MESTER : III SSION : MO/2022		
TIME:	3:00 Hours FL						LL MARKS: 50		
INSTRUC 1. The o 2. Atter 3. The r 4. Befor 5. Table	CTIONS: question paper conta npt all questions. nissing data, if any, re attempting the qu es/Data hand book/G	ins 5 questions may be assume estion paper, b raph paper etc.	s each of 10 ma od suitably. De sure that you to be supplied	rks and to 1 have got 1 to the ca	tal 50 marks. the correct q ndidates in th	uestion paper e examination	n hall.	-	
Q.1(a) (b)	<ul><li>a) Sketch the overview (outline) of any Bioseparation process.</li><li>b) Describe any one gradient centrifugation method.</li></ul>							CO 1 1	BL 2 2
(c)	(c) Explain various mechanical cell disruption methods.						[5]	1	2
Q.2(a) (b)	<ul><li>a) Write adsorption isotherm equations.</li><li>b) Justify the use of ammonium sulphate for precipitation of protein in salting out method.</li></ul>						[2] [3]	2 2	2 3
(c)	Calculate specific a data:	activity, purific	ation fold and	percentag	e recovery fr	om the given	[5]	2	5
	Steps	Total protein (mg)	Total activity (unit)	Specifi c activity	Purification fold	% recovery			
	Homogenate	2936	27028	activity					
	Sediments	1041	22846						
	Salt	32	18314						
		43	4185						
	Affinity	1.3	2137						
Q.3(a)	) What is reverse phase chromatography? How it is advantageous over normal phase chromatographic techniques.							3	1
(b)	Describe briefly the principle of anion exchange chromatography.							3	2
(c)	Two proteins of MW 2.5×10 <sup>5</sup> and 1×10 <sup>4</sup> were eluted out of a gel in gel filtration column at 220 mL and 300 mL respectively. Determine the molecular weight of a protein that elutes out at 270 mL under the same condition?						[5]	3	5
Q.4(a)	In a cross flow filtration, if inlet pressure is 8 atm., outlet pressure is 2 atm., calculate transmembrane pressure drop.						[2]	4	4
(b) (c)	Describe any one membrane module with its advantages and disadvantages. Explain the mechanism of transport of particles in a MF system.						[3] [5]	4 4	2 3
Q.5(a)	Write in detail abou	t the following					[5]	5	3

- (i) Tray dryer; (ii) Fluidized bed dryer
- (b) (i) Write about crystallization theory and explain in detail about the nucleation and [5] 5 3 crystal growth.
  - (ii) Write the working principle and operating procedure about oslo type crystallizer.

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