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# UNIVERSITY OF RUHUNA – FACULTY OF ALLIED HEALTH SCIENCES DEPARTMENT OF PHARMACY THIRD BPHARM PART II EXAMINATION - NOVEMBER 2022 PH 3233 PHARMACEUTICAL BIOTECHNOLOGY – SEQ PAPER

#### **TIME: TWO HOURS**

## INSTRUCTIONS

- There are four questions in part A and B in this SEQ paper.
- Answer all questions.
- No paper should be removed from the examination hall.
- Do not use any correction fluid.
- Use illustrations where necessary.

#### PART A

1. Same at the one privilents as been of and had minority munor to contractory of shall the

1.1.

	1.1.1. What is "fermentation" in biotechnology?	(10 marks)
	1.1.2. List five pharmaceutical products that can produce using fermentation.	(15 marks)
	1.1.3. Describe the "submerged fermentation".	(30 marks)
1.2.	uses of report binard DNA technology.	
	1.2.1. Write five advantages of immobilized enzyme.	(20 marks)

1.2.2. List five ideal properties of matrix used in immobilized enzyme system.(25 marks)

2.

2.1.	What is the difference between finite cell line and continuous cell line?	(10 marks)
2.2.	In cell culture, what is "subculturing"? Briefly describe.	(15 marks)
2.3.	List five limitations in cell culturing.	(20 marks)
2.4.	"Animal cloning allows to create a genetically identical individuals of an exist	ing animal".
	2.4.1. List five animal species that have been cloned successfully.	(10 marks)
	2.4.2. Write four applications of animal cloning in medicine.	(20 marks)
	2.4.3 Briefly describe the steps in animal cloning process	(25 marks)

### PART B

- 3. Downstream processing is used to purify protein of interest from a crude product from fermentation.
- v3.1. What are the two different types of cellular products available for downstream processing?

par ouesions in part A and B in this SEO paper.	(10 marks)
3.2. What are the two methods used for initial product recovery?	(10 marks)
3.3. Describe each step briefly mentioned in 3.2.	(20 marks)
3.4. State five different types of cell disruption methods.	(10 marks)
3.5. Describe the basis of ion exchange chromatography.	(20 marks)
3.6. State four practical limitations of affinity chromatography.	(20 marks)

3.7. State five properties of serum albumin that can be used as stabilizing excipient. (10 marks)

4. Recombinant DNA technology is used by several thousands of industr	ies for their
biotechnological productions.	
4.1. State five uses of recombinant DNA technology.	(10 marks)
4.2. State two types of enzymes used in recombinant DNA technology.	(10 marks)
4.3. Briefly describe the functions of each enzyme mentioned in 4.2.	(20 marks)
4.4. State the name of the source organism of following enzymes.	(10 marks)
4.4.1. <i>ECoR</i> 1	
4.4.2. Hind111	
4.4.3. Alu1	
4.4.4. <i>Pst</i> 1	
4.4.5. Not1	
4.5. What is the cloning vector?	(10 marks)
4.6. What are three essential components of a cloning vector?	(15 marks)
4.7. Briefly describe following.	(15 marks)
4.7.1. Genomic library	
4.7.2. cDNA library	
4.7.3. Blue script	
4.8. State five uses of transgenic animals.	(10 marks)

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