



UNIVERSITY OF RUHUNA – FACULTY OF ALLIED HEALTH SCIENCES DEPARTMENT OF PHARMACY

THIRD BPHARM PART II EXAMINATION – AUGUST/SEPTEMBER 2020 PH 3233 PHARMACEUTICAL BIOTECHNOLOGY (SEQ)

TIME: TWO HOURS

INSTRUCTIONS

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

PART A

01.

- 1.1 Bluescript is a cloning vector designed to simplify commonly used cloning and sequencing procedures.
 - 1.1.1 State different types of vectors and their maximum insert size.

(15 marks)

- 1.1.2 Briefly describe the properties of the bluescript vector? (20 marks)
- 1.1.3 What do you understand by multiple cloning sites? (10 marks)
- 1.1.4 Describe the basis of the visual test? (20 marks)
- 1.2 Transgenic animals are far useful as models of human diseases.
 - 1.2.1 What are transgenic animals? (10 marks)
 - 1.2.2 What are the advantages of transgenic animals? (10 marks)
 - 1.2.3 Briefly state the simplified procedure of creating a transgenic animal?

(15 marks)

- **02.** Downstream processing refers to the recovery and the purification of biosynthetic products from natural sources such as animal or plant tissue in a sterile culture medium.
 - 2.1 Briefly describe the steps of generation of water for injection? (20 marks)
 - 2.2 State five methods used in cell disruption? (10 marks)
 - 2.3 Briefly describe the method of diafiltration? (20 marks)
 - 2.4 Briefly describe the procedure of dye affinity chromatography and discuss its advantages and disadvantages? (30 marks)
 - 2.5 Write short note on "the stabilizing excipients"? (20 marks)

PART B

)3.	3.1 State the difference between monoclonal antibodies and polyclonal antibodies.			
	3.1 3	tate the difference between monocional anticodico and project	(10 marks)	
	3.2 Briefly explain the steps in the production of monoclonal antibodies.			
			(30 marks)	
	3.3 List five immunological techniques used in diagnostic immunology.			
			(15 marks)	
	3.4 I	Briefly explain one of the above techniques you mentioned.	(20 marks)	
	3.5 I	3.5 Enzyme Immobilization is a very important step in the production of enzymes.		
		3.5.1 What do you mean by Enzyme Immobilization?	(10 marks)	
		3.5.2 Briefly explain three methods used in Enzyme Immobilization	1.	
			(15 marks)	
04.				
	4.1	Briefly explain "cryopreservation" used in cell culture.	(15 marks)	
	4.2	Briefly explain three different types of artificial cloning.	(15 marks)	
	4.3	Discuss the contamination preventive measures that you can adapt	to prevent	
		contamination of a cell culture.	(25 marks)	
	4.4	State four types of plant tissue cultures.	(10 marks)	
	4.5	List three advantages of plant tissue culture.	(15 marks)	
		Briefly explain the three main steps in plant tissue culture process.	(20 marks)	

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