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UNIVERSITY OF RUHUNA - FACULTY OF ALLIED HEALTH SCIENCES

DEPARTMENT OF PHARMACY

SECOND BPHARM PART I EXAMINATION – NOVEMBER 2019 PH 2123 PHARMACEUTICS II B (SEQ)

TIME: TWO HOURS

INSTRUCTIONS

- There are four (04) questions in the SEQ paper.
- Part Λ and part B should be answered separately in the booklets provided.
- No paper should be removed from the examination hall.
- Do not use any correction fluid.
- Use illustrations where necessary.

Part A

1.		
	1.1. Define the term "pharmaceutical incompatibility".	(10 marks)
)	1.2. Mention three major types of pharmaceutical incompatibilities.	(15 marks)
	1.3. Define the term "parenteral administration".	(10 marks)
	1.4. Write a short note on "large volume parenteral fluids".	(25 marks)
	1.5. Discuss three major routes of parenteral administration.	(40 marks)
2.	2.1. State three major stages in formulation of tablet dosage form.	(15 marks)
		(25 marks)
	2.2. List five key attributes of tablet dosage form.	
	2.3. State two examples for each of the following excipients.	(30 marks)
	2.3.1. Binders	
	2.3.2. Diluents	
	2.3.3. Lubricants	
	2.3.4. Disintegrants	
	2.3.5. Flavoring agents	

2.4. Tabulate the steps in formulation of tablet dosage form using three different techniques.

(30 marks)

Part B

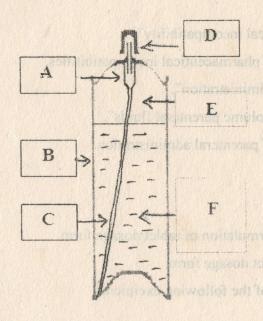
3.

- 3.1. What are the main considerations in the dosage form design? (15 marks)
 3.2. Define the term "microencapsulation". (15 marks)
 3.3. Give five reasons for microencapsulation. (20 marks)
- 3.4. Draw and identify all the parts of a hard gelatin capsule. (20 marks)
- 3.5. Briefly explain the main steps involved in making the production of empty hard gelatin capsules.

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4.

- 4.1. Define the term "pharmaceutical aerosol". (10 marks)
- 4.2. Name A-F components of the below schematic diagram. (12 marks)



- 4.3. Provide two labeling information that should be on the primary container of a "flammable" or "extremely flammable" acrosol container. (08 marks)
- 4.4. Briefly explain the advantages of sub dermal implant contraceptives. (20 marks)
- 4.5. List five parameters of a good toothpaste formulation. (20 marks)
- 4.6. Compare and contrast sustained release dosage form and controlled release dosage form.

(30 marks)

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