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#### **Enrolment No:**



## **UPES**

## **End Semester Examination, December 2024**

Course: Advanced Drug Delivery System
Program: Int. (B. Sc. + M. Sc. (Clinical Research)
Course Code: HSCR8008P
Semester: VII
Time: 03 Hours.
Max. Marks: 100

Instructions: All questions are compulsory.

#### Section A

# Short answer questions/ MCQ/T&F (20Qx1.5M= 30 Marks)

S. No.		30 Marks	co
Q1	Lacrisert is drug delivery system.  A. Erodible and ocular  B. Non-erodible and ocular  C. Erodible and mucosal  D. Non-erodible and mucosal	1.5	CO1
Q 2	is the toughest barrier for transdermal drug delivery.  A. Stratum corneum B. Hypodermis C. Dermis D. Mucosa	1.5	CO1
Q 3	Define gastroretentive drug delivery system.		CO1
Q 4	Implants should be sterile and contain no bio-load.  A. True  B. False	1.5	CO1
Q 5	Enlist three advantages of nasal drug delivery system.	1.5	CO1
Q 6	Enlist any three polymers used for transdermal drug delivery system.	1.5	CO1
<b>Q</b> 7	Define penetration enhancers.	1.5	CO1
Q 8	Select the theory that describes the mechanism of bioadhesion.  A. Molecular weight theory C. Vapor pressure theory D. Cohesion theory	1.5	CO2
Q 9	Give an example of a drug delivery system that has thread-like structure.	1.5	CO2
Q 10	Superoxide iron can be employed for designing drug delivery system.  A. high density B. low density C. magnetic D. floating	1.5	CO2
Q 11	Classify ocular drug delivery systems.	1.5	CO2
Q 12	Particles greater than 100 microns can easily be delivered to alveoli in lungs.  A. True  B. False	1.5	CO2
Q 13	is used to estimate <i>in-vitro</i> drug release from transdermal patches.  A. Paddle over disks apparatus  B. Disk over Paddle apparatus  C. Flow through apparatus  D. Basket type apparatus	1.5	CO2
Q 14	If the drug has good permeability through mucosal membrane but is hydrophobic, then suggest the suitable nanocarrier for drug delivery.	1.5	CO3
Q 15	Report any three types of delivery systems that are employed to avoid first pass metabolism.	1.5	CO3

Q 16	can be used for passive targeting of nanoparticles to tumor cells in cancer		
	treatment.	1.5	CO3
	A. Particle size B. Penetration enhancers	1.3	003
	C. Antibodies D. Pore size of capillary endothelium		
Q 17	Nebulizers are used to deliver volatile drugs to pulmonary system.	1.5	CO3
Q 18	A. True B. False  Identify and relate its proper use.		
Q 16	A. Mucosal drug delivery B. Nasal drug delivery C. Transdermal drug delivery D. Oral drug delivery	1.5	CO4
Q 19	Highly branched polymer can enhance bioadhesion.	1.5	CO4
Q 20	A. True  B. False  Draw a salamatic diagram of Consert shawing its components		
Q 20		1.5	CO4
	Section B (4Qx5M=20 Marks)		
Q	Draw a schematic diagram of Ocusert showing its components.  Section B (4Qx5M=20 Marks)  Short Answer Type Question  Describe any one polymer used for fabricating drug delivery systems.  Summarize the advantages of mucosal drug delivery systems.  Categorize and explain any two types of drug carriers for the targeted drug delivery.		CO
Q 1	Describe any one polymer used for fabricating drug delivery systems.	5	CO1
Q 2	Summarize the advantages of mucosal drug delivery systems.  5		CO2
Q 3	Short Answer Type Question  Describe any one polymer used for fabricating drug delivery systems.  Summarize the advantages of mucosal drug delivery systems.		CO4
Q 4	Microneedles can be used for transdermal drug delivery. Justify the statement.	5	CO3
	Section C (2Qx15M=30 Marks)		
Q	Two case studies 15 marks each subsection	30 Marks	CO
Q 1	Demonstrate the use of polymers in formulation of advanced drug delivery systems.		CO3
Q 2	<ul><li>a) Illustrate the anatomy of mucosal membrane and composition of mucus.</li><li>b) Discuss the significance of mucus in the human body.</li></ul>	7+8	CO4
	Section D		
0	(2Qx10M=20 Marks)	20	
Q	Long Answer type Questions	20 Marks	CO
Q 1	Explain various quality control parameters for transdermal drug delivery systems.	10	CO4
Q 2	Describe ideal characteristics of the drug for naso-pulmonary drug delivery system.	10	CO2