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2022

BINA CHOWDHURY CENTRAL LIBRARY (GIMT & GIPS) Azára, Hatkhowapara,

B.Pharm. 7th Semester End-Term (Regular) Examination

NOVEL DRUG DELIVERY SYSTEMS (THEORY)

(For new regulation (w.e.f. 2017-18))

(New regulation)

Full Marks - 75

Time - Three hours

The figures in the margin indicate full marks for the questions.

1.	Answer	the	follow	ing	(MCQ)
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 $(20 \times 1 = 20)$

- (i) Liposomes consist of a bilayer of
 - (a) Hydrophilic molecules
- (b) Hydrophilic molecules
- (c) Both (a) and (b)
- (d) None
- (ii) Polymers formed from two or more different monomers are called
 - (a) Homopolymers
- (b) Copolymers

(c) Oligomers

- (d) None of the above
- (iii) Albumin is an example of
 - (a) Non biodegradable polymer (b)
- (b) Synthetic Polymer
 - (c) Natural Polymer
- (d) None of the above
- (iv) Which step is involved in the preparation of microcapsules by coacervation phase seperation?
 - (a) Formation of 3 immiscible chemical phases
 - (b) Deposition of liquid polymer coating upon core material
 - (c) Rigidizing the coating
 - (d) All the above

(1	v)	Wha	at is/are the size of Microspher	re?	the state of the state of the
		(a)	More than 200 micrometre	(b)	Less than 200 micrometre
		(c)	Both (a) and (b)	(d)	None of these
(vi)	Nan	noscience can be studied with	the he	lp of
		(a)	Quantum mechanics	(b)	Newtonian mechanics
		(c)	Macro dynamics	(d)	Geophysics
(vii)	Dru	g loading capacity of MDDS i	s	to mind the second second
	7.	(a)	Low	(b)	High
		(c)	Very low	(d)	Medium
(viii) Solv	vent evaporation is which type	e of mi	cro encapsulation techniques?
		(a)	Chemical	(b)	Physical
		(c)	Physico-chemical	(d)	All of the above
((ix)	Chi	tosan istype of p	olyme	
		(a)	Anionic	(b)	Cationic .
		(c)	Neutral	(d)	None of the above
	(x)	Tak	olets that are placed under the	e skin	are
		(a)	Enteric coated tablets	(b)	Film coated tablets
		(c)	Implants	(d)	Sublingual tablets
	(xi)	Fol	lowing is/are the physical metho	ds of n	nicroencapsulation techniques
		(a)	Polymerisation	(b)	Microencapsulation
		(c)	Air suspension	(d)	All of the above
	(xii) Fol	lowing are the theories of "my	ucoadh	esion" except
		(a)	Wetting agent	(b)	Absorption theory
		(c)	Fracture theory	(d)	Diffusion theory
	(xii	i) Sh	ellac is the coating materials	for mic	crocapsules of following categorie
		(a)		(b)	take of the same and
		(c)	Enteric coated resins	(d)	Wax resins
	(xiv	v) Wi	nich of the following is comme	ercially	used osmotic agents?
		(a)	Sorbitol	(b)	Osmogene
		(c)	Imogene	(d)	Aluminum chloride

(xv) W	nich of the following is implantable osmotic pump?
(a	Single chamber osmotic pump
(b	Push pull osmotic pump
(c)	Multiple chamber osmotic pump
(d	Higuchi leeper pump
	e time for which the floating dosage form floats on dissolution medium is led
(a	Floating time (b) Buoyancy lag time
(c)	Lead time (d) Transit time
(xvii) V	hich of the following is wrong statement for GERD?
(a	Useful to provide both local and systemic effects of the drug
(b	Useful to increase gastric retention of the drug
(c	Useful to reduce the first pass effect of the thugs
(d	Useful to target site specific release of drugs
	Thich of the following formulations would not be applicable to ocular ministration?
(a	Solution (b) Liniment
(c	Suspension (d) Ointment
	hich of the following oral liquid formulations would be considered as an opharyngeal formulation?
(a	Syrup (b) Elixir
(c	Mouthwash (d) Linctus
(xx) W	ich of the following GRDDS is also known as "Plug type system"?
(a	Floating system (b) Swelling system
(c	Bioadhesive system (d) Pulsatile system
Answe	any seven questions $(7 \times 5 = 35)$
(a) C	assify the polymers used to modify the drug release. (5)
(b) D	scuss the pharmaceutical applications of microspheres. (5)
(c) D	escribe osmotically regulated implants as a new drug. (5)
	rplain the biological factors affecting controlled release drug delivery stems. (5)
22	rite a note on pulmonary route as a promising route of drug ministration. (5)

- (f) Write the advantages and disadvantages of implants.
- (g) Discuss strategies and the components of targeted drug delivery systems. (5)
- (h) What are the advantages and disadvantages of intrauterine drug delivery system? (5)
- (i) Describe the monoclonal antibodies with its applications. (5)
- 3. Answer any two questions

 $(2 \times 10 = 20)$

(5)

(a) Write the concept of controlled drug delivery systems. Explain the approaches for the Controlled release formulations based on dissolution.

(3+7=10)

- (b) Describe various theories of mucoadhesion with their significance in designing mucoadhesive products. (10)
- (c) Define nanoparticles. State various methods to prepare nanoparticles. What are the advantages of nanoparticles in the drug delivery system?

 (2+5+3=10)