Time: 3 Hours]

CS/M.Pharm (Pharmaceutics)/SEM-2/MPT-206(II)/09 PHYSICAL PHARMACEUTICS (SEMESTER - 2)

1.	Signature of Invigilator								di di		O Lie Ĉ	o ah		.	
2.	Signature of the Officer-in-Charge	No.												$\underline{\mathbb{I}}$	
	Roll No. of the Candidate														
	CS/M.Pharm (Pha ENGINEERING & MAN PHYSICAL PHAR	AGE	ME	NT	EX	AM	IINA	ATIC	ONS	, Jt	JLY	- 2	009		

INSTRUCTIONS TO THE CANDIDATES:

- 1. This Booklet is a Question-cum-Answer Booklet. The Booklet consists of **32 pages**. The questions of this concerned subject commence from Page No. 3.
- 2. a) In **Group A**, Questions are of Objective type. You have to write the answer in the space provided **marked 'Answer Sheet'**.
 - b) For **Groups B** & **C** you have to answer the questions in the space provided marked 'Answer Sheet'. Questions of **Group B** are Short answer type. Questions of **Group C** are Long answer type. Write on both sides of the paper.

[Full Marks: 70

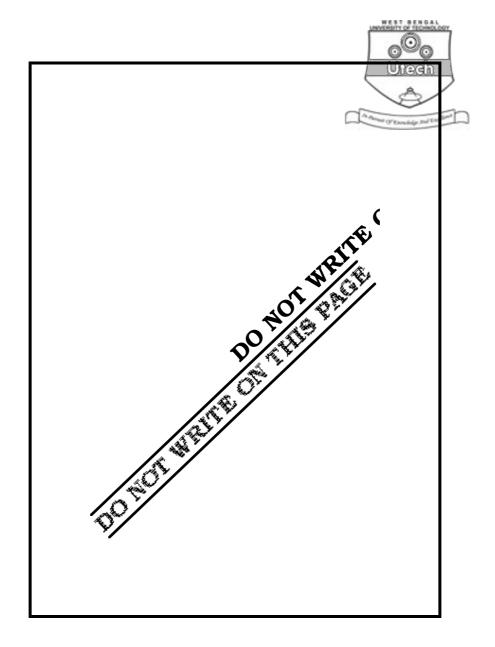
- 3. **Fill in your Roll No. in the box** provided as in your Admit Card before answering the questions.
- 4. Read the instructions given inside carefully before answering.
- 5. You should not forget to write the corresponding question numbers while answering.
- 6. Do not write your name or put any special mark in the booklet that may disclose your identity, which will render you liable to disqualification. Any candidate found copying will be subject to Disciplinary Action under the relevant rules.
- 7. Use of Mobile Phone and Programmable Calculator is totally prohibited in the examination hall.
- 8. You should return the booklet to the invigilator at the end of the examination and should not take any page of this booklet with you outside the examination hall, **which will lead to disqualification**.
- 9. Rough work, if necessary is to be done in this booklet only and cross it through.

No additional sheets are to be used and no loose paper will be provided

Head-Examiner/Co-Ordinator/Scrutineer

49004 (08/07)







CS/M.Pharm (Pharmaceutics)/SEM-2/MPT-206(II)/09 PHYSICAL PHARMACEUTICS SEMESTER - 2

Time: 3 Hours [Full Marks: 70

GROUP - A

(Objective Type Questions)

l.	Ansv	Answer any <i>ten</i> of the following :							
	i)	Which is the major mechanism for drug absorption ?							
		a)	Active transport	b)	Pore transport				
		c)	Passive diffusion	d)	Facilitated diffusion.				
	ii)	Which factor influences the solubility of salt form of drug?							
		a)	Size of particle	b)	Size of counter ion				
		c)	Amount of salt	d)	pH value.				
	iii)	Directly compressible tablet diluent is							
		a)	CaCO ₃						
		b)	Microcrystalline Cellulose						
		c)	Mannitol						
		d)	Lactose.						
	iv)	A hardness of kg is minimum for uncoated tablet.							
		a)	5	b)	10				
		c)	20	d)	2.				
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v)	The dissolution assembly is suspended in a liquid medium in a suitable vesse containing									
	a)	800 ml beaker	b)	900 ml beaker						
	c)	1000 ml beaker	d)	1100 ml beaker.						
vi)		ne triple point, <i>O</i> , in case of iclom is		r-vapour phase system, the degree of blank)						
vii)	If a g	gel is highly hydrated, diffusion	occurs	through the						
viii)	The rheological properties of solid are determined by measuring									
	a)	angle of repose	b)	carr index						
	c)	disintegration time	d)	none of these.						
ix)	(x) Heckel plot illustrates the relationship between the following :									
	a)	Density and moisture content								
	b)	ing agent								
	lubricating agent									
	d)	d) None of these.								
x) The equation, $J = D_f \cdot \frac{dc}{dx}$ is commonly known as										
	a) Fick's First Law of Diffusion									
b) Voyes and Whitney equation (law)										
	c)	Peppus equation								
	d)	Higuchi's law.								
xi)		temperature at which the s		y of the surfactant equals CMC is						
xii)	The particle diameter of individual particles can be measured by									



GROUP - B

(Short Answer Type Questions)

Answer any three of the following.



- 2. Briefly discuss the factors affecting drug dissolution.
- 3. Write about the Physics of Tablet compression.
- 4. Distinguish 'micellization' from 'solubilization'.
- 5. Critical micellization concentration (CMC) is influenced by certain factors. Elaborate, in short with example.
- 6. Write the method of determination of particle size (any *one*) and also explain the significance of particle size on powder flow property.

GROUP - C

(Long Answer Type Questions)

Answer any three of the following.

 $3 \times 15 = 45$

- 7. What are Hydrogels? How do you prepare synthetic hydrogels?
- 2 + 13
- 8. Discuss, in detail, about different stages involved in tablet compression.
- 9. a) Write an account on kinetic aspects of Swelling deriving relevant equations.
 - b) The diffusion coefficient, D, of a drug in a swollen hydrogel were measured at several pH values corresponding to different hydration values, H:

pH's 1 3 5 7
H (dimensionless, hydration value)
$$0.352$$
 0.337 0.630 0.880 $D \times 10^{-8}$ cm $^{-2}$ /sec. 2.50 3.58 44.60 139.00

Compute the diffusion coefficient, $D_{\rm o}$ in water and the constant, K_f , of the system using appropriate equation. $7 \, {1 \over 2} \, + \, 7 \, {1 \over 2}$



- 10. Write notes on any three of the following:
 - a) Uses of surfactants in increasing solubility of drugs
- Utech

- b) Kraft temperature and cloud point
- c) Association colloids
- d) Oral gels.
- 11. Discuss the role of surfactants in the contents of
 - a) percutaneous absorption
 - b) parenteral administration.



END