

Roll No.

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Total No. of Questions : 10]

[Total No. of Pages : 02

B. Pharmacy (Sem. - 7th)
PHARMACEUTICS - VIII
(Pharmaceutical Technology - II)
SUBJECT CODE : PHM-4.7.2

Paper ID : [D0133]

[Note : Please fill subject code and paper ID on OMR]

Time : 03 Hours

Maximum Marks : 80

Instruction to Candidates:

- 1) Section - A is **Compulsory**.
- 2) Attempt any **Four** questions from Section - B.
- 3) Attempt any **Three** questions from Section - C.

Section - A

Q1)

(15 x 2 = 30)

- a) Microencapsulation.
- b) Lyophilization.
- c) Hemostatics.
- d) Aseptic area.
- e) Zero order release.
- f) Capsule.
- g) Gelatin.
- h) Absorbent cotton.
- i) Isotonicity.
- j) Quality control.
- k) Catguts.
- l) Preformulation.
- m) Coacervation.
- n) What is the need of granulation while preparing tablets?
- o) List any two criteria of drug(s) essential for microencapsulation.

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P.T.O.

Section - B

(4 x 5 = 20)

- Q2)** Highlight physics of making tablets.
- Q3)** Explain evaluation of capsules.
- Q4)** Explain techniques for the microencapsulation of drugs.
- Q5)** Enumerate sterility testing of injection containing antibiotics.
- Q6)** Highlight different types of transdermal controlled released drug delivery systems.

Section - C

(3 x 10 = 30)

- Q7)** (a) Explain packaging equipments for the packaging of oral solid dosage forms.
(b) How packaging testing could be correlated with stability of dosage forms.
- Q8)** (a) How aseptic area could be designed and evaluated?
(b) Enumerate IP method for the testing of pyrogen in parenterals.
- Q9)** Highlight formulation, packaging and evaluation of paracetamol tablet IP.
- Q10)** Write note on :
- (a) Stability testing.
(b) Wound dressing.
(c) Organ replacement materials.

