Roll No. ..... Total Pages: 03

### BT-7/M-20

37062

# PHARMACEUTICAL BIOTECHNOLOGY BTT-403E

Time : Three Hours] [Maximum Marks : 100

**Note**: Attempt *Five* questions in all, selecting at least *one* question from each Unit.

#### Unit I

- 1. (a) What are organic therapeutic agents? Give their applications.
  - (b) Discuss in brief about the economics of drug development.
- 2. Explain the following with suitable applications:  $5\times4$ 
  - (a) Complex chemical conversion
  - (b) Condensation
  - (c) Fermentation
  - (d) Carboxylation.

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## Unit II

Write	brief notes on the following:	5×4	
(a)	Half-life of drugs		
(b)	Inborn errors of metabolism		
(c)	Single nucleotide polymorphism		
(d)	Drug metabolism.		
(a)	Explain various steps during action of drug	on	
	human body.	12	
(b)	Describe the applications of radioactive lab	eled	
	compounds in drug metabolism.	8	
Unit III			
Desci	ribe the methods of formulation of sustained ac	etion	
dosag	ge form. Write GMP requirements for cap	sule	
formu	ulation.	20	
Brief	ly explain the following:	5×4	
(a)	Packing techniques		
(b)	Hormones		
(c)	Tablet coating		
(d)	Oral liquids.		
_37062	2 2		
	(a) (b) (c) (d) (a)  Described of the control of th	<ul> <li>(b) Inborn errors of metabolism</li> <li>(c) Single nucleotide polymorphism</li> <li>(d) Drug metabolism.</li> <li>(a) Explain various steps during action of drug human body.</li> <li>(b) Describe the applications of radioactive lab compounds in drug metabolism.</li> <li>Unit III</li> <li>Describe the methods of formulation of sustained action dosage form. Write GMP requirements for cap formulation.</li> <li>Briefly explain the following: <ul> <li>(a) Packing techniques</li> <li>(b) Hormones</li> <li>(c) Tablet coating</li> <li>(d) Oral liquids.</li> </ul> </li> </ul>	

### **Unit IV**

7. (a) Give classifications of Vitamins. How can you estimate vitamins in a sample ? 12

(b) Write a note on nutritional status of vitamins. 8

**8.** What are therapeutic proteins? Explain their delivery and targeting. Explain their applications and regulatory aspects.

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