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Total No. of Pages: 02

Total No. of Questions: 08

## M.Sc.(Ph. Chem) (Sem.-3) DRUG DESIGN AND DEVELOPMENT

Subject Code: MSPC-205 M.Code: 20511

Time: 3 Hrs. Max. Marks: 80

## **INSTRUCTIONS TO CANDIDATES:**

- 1. Attempt any FIVE questions out of EIGHT questions.
- 2. Each question carries SIXTEEN marks.

1.	a) Discuss the historical development of drug design approaches.	6
	b) Give five postulates of quantum mechanics.	5
	c) Give Schrodinger equation and explain the significance of wave function ( $\psi$	<sup>2</sup> ). 5
2.	a) Explain the drug action on the basis of concept of receptor.	5
	b) Describe the extrathermodynamic model of traditional QSAR.	6
	c) Explain Topliss operational scheme for the designing of test series in QSAR	. 5
3.	a) Describe the role of data retrieval techniques in drug design with example.	8
	b) Discuss the principle of molecular mechanics and its applications in drug de	sign. 8
4.	a) Describe the single compartment pharmacokinetic model.	8
	b) Describe Lipinski rule and its applications.	8
5.	a) Write short note on environmental pharmacokinetics.	8
	b) Discuss the role of biopharmaceutics in drug design.	8
6.	a) What are peptidomimetics? Describe Farmer's rule to convert per peptidomimetic.	otide into 8
	b) Give brief note on oligonucleotide based antiviral chemotherapy.	8

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7.	a)	What are the basic steps of rational prodrug design?	6
	b)	With the help of suitable examples describe the concept of prodrugs for site speand prolonged action drug delivery system.	ecific 10
8.	Wı	rite short note on any two:	
	a)	Validation of QSAR model.	8
	b)	Perturbation theory of drug action.	8
	(c)	Limitation of CADD	8

NOTE: Disclosure of Identity by writing Mobile No. or Making of passing request on any page of Answer Sheet will lead to UMC against the Student.

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