

[A-33]

SARDAR PATEL UNIVERSITY

M.Sc. (Integrated) Biotechnology (IGMBT), 10th Semester ExaminationFriday, 13th April 2018

10:00 A.M to 1:00 P.M

PS10CIGMB3: Drug design and development

Total Marks: 70

- Note: (1) Figures to the right indicate marks.
(2) Draw a neat and labeled diagram, wherever necessary.

Q. 1 Choose the most appropriate answer from the four alternatives given:

[8]

- (1) Curacin A is isolated from which of the following source?
(a) Brazilian viper (b) puffer fish (c) Ecuadorian poison frog (d) none of the above
- (2) Artemisinin drug used to cure:
(a) arthritis (b) Rheumatic Fever (c) Malaria (d) inflammation
- (3) Which of the following amino acids present in active-site of Enzymes?
(a) Serine (b) Histidine (c) Cysteine (d) All of the above.
- (4) Serotonin receptor agonists are used in treatment of which of the following diseases?
(a) Migraine (b) Parkinson's Disease (c) Cholera (d) None of the above
- (5) Palmitate ester of chloramphenicol is used to
(a) increase polarity (b) decrease polarity- (c) target drug (d) none of them
- (6) Carbidopa is a example of :
(a) prodrug (b) Sentry drug (c) none of them (d) all of them
- (7) Which of the following drug withdrawn from the market during phase IV clinical trial? : (a) Atenalol (b) Practolol (c) stematolol (d) A and B both
- (8) During Phase IV clinical trial Rofecoxib was associated with :
(a) Increase Risk of heart pain (b) increase in B.P (c) increase rate of heart attack
(d) all of them

[14]

Q-2 Answer any **SEVEN** from the following:

- (1) Pitfalls
- (2) What are Orphan receptors?
- (3) High-throughput screening
- (4) How antifreeze poisoning occurs?
- (5) Define Transition state analogues.
- (6) Differentiate: Sentry drugs and Prodrugs
- (7) Enlist the various methods used for drug targeting.
- (8) Define steric shields?
- (9) What is EPO and PCT?

PTO

Q.3 (a) How do bioassay helps in identifying the lead molecule. Write detail note on Screening by NMR. [6]

(b) Define lead molecule. How can you obtain it from natural or synthetic scaffold. [6]

OR

(b) What is an ideal drug target? Describe six properties of an ideal drug target. [6]

Q.4 (a) Explain drug action at carrier protein with suitable examples. [6]

(b) Discuss the various parameters used to design agonist. [6]

OR

(b) Discuss mechanism of action of 6 mercaptopurine and Aspirin in detail [6]

Q.5 (a) How drugs can be made less resistant to drug metabolism? [6]

(b) How drugs can be made to increase solubility and membrane permeability. [6]

OR

(b) What are Prodrugs? Write in brief purposes for which prodrugs are designed. [6]

Q.6 (a) Write detail account drug metabolism study for development of new drug. [6]

(b) Give detail account on phase III and IV clinical trial. [6]

OR

(b) Explain chemical development process for development of new drug in brief. [6]