BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI (END SEMESTER EXAMINATION)

CLASS: IMSC SEMESTER: IX
BRANCH: CHEMISTRY SESSION: MO/19

SUBJECT: SAC3009 MEDICINAL CHEMISTRY

TIME: 3 HOURS FULL MARKS: 60

INSTRUCTIONS:

- 1. The question paper contains 7 questions each of 12 marks and total 84 marks.
- 2. Candidates may attempt any 5 questions maximum of 60 marks.
- 3. The missing data, if any, may be assumed suitably.
- 4. Before attempting the question paper, be sure that you have got the correct question paper.
- 5. Tables/Data hand book/Graph paper etc. to be supplied to the candidates in the examination hall.

Q.1(a)	How do hydrophilicity/hydrophobicity impact as bio physico-chemical properties on metabolism? What are log P and logD values?	[6]
Q.1(b)	Define any two biological activity parameters and discuss their significance?	[6]
Q.2(a) Q.2(b)	What is Lipinski's rule of five? What are its exceptions? What are ion channel modulators? Discuss their mechanism of action along with examples?	[6] [6]
Q.3(a) Q.3(b)	Explain with schematic Calcium & Phospho inositol system for breakdown of phospholipids? Briefly describe about bioisosterism and explain the role of nonclassical bioisosteres in the discovery of Losarton as cardiovascular drug.	[6] [6]
Q.4(a) Q.4(b)	Define Therapeutic index and corelate it with C_{max} with the help of graphical representation? Discuss the mechanism of enzymatic action with special emphasis on promoters and inhibitors?	[6] [6]
Q.5(a)	What do understand with the term "Antimetabolite", describe anti-metabolite mechanism of pyrimidine/purine antagonist with an example.	[6]
Q.5(b)	Discuss the anti-metabolite theory proposed by Woods and Fildes and explain the bacteriostatic action of sulfa drugs.	[6]
Q.6(a) Q.6(b)	What is "Anti-vitamin" effect. Discuss with examples. Discuss the major reason or therapeutic agents are associated with deficiency of Vitamin B and C.	[6] [6]
Q.7(a) Q.7(b)	Draw a Dengue virus life cycle and suggest the possible site of action for drug or drug candidate. What is binding mode of drug. What type of non-covalent bonds are responsible in drug-receptor interaction.	[6] [6]

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