



10. Insulin is used to treat diabetes; which of the statement regarding Insulin use is **CORRECT**?  
 compensate for inadequacies                       cure a condition  
 treat symptoms                                               alter the patient's emotional state
11. Which lipophilic substituent increases lipophilicity in aromatic group but reduces lipophilicity in aliphatic system \_\_\_\_\_.  
 -methyl                       -fluorine                       -chlorine                       -hydroxyl
12. Which of the statement regarding Allosteric drugs is **INCORRECT**?  
 binds to primary site on the target  
 alters the shape of the target  
 alters the vibrational motion of the target  
 donate or accept electrons or protons
13. How many degrees of freedom are allowed for rigid docking methods?  
 None                                               Three  
 Six                                               Depends upon the no. of atoms
14. The value of energy attributed to Stretching for force field calculation is \_\_\_\_\_.  
 1.017 KJ/mol                       9.406 KJ/mol                       -6.084 KJ/mol                       -0.121 KJ/mol
15. Drugs that are absorbed by placing them under the tongue is defined as \_\_\_\_\_.  
 Buccal Administration                       Sublingual Administration  
 Subcutaneous Administration                       Intravenous Administration

SECTION "B"

[5Q. × 0.5 = 2.5 marks]

**Define in one sentence.**

16. Multiple drug interactions:

17. Suicide inhibitor:

18. Biased Agonism:

19. Soft docking:

20. Drug bioavailability:

KATHMANDU UNIVERSITY  
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Year : IV  
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Course : BIOT 413  
Semester: I  
F.M. : 40

SECTION "C"

(Long answer questions)

[4Q. × 6 = 24 marks]

Attempt *ANY FOUR* questions.

1. Explain the process of allosteric modulation. How does binding to allosteric sites changes the concentration effect curve? [3+3]
2. Explain how protein structures can be determined by X-ray crystallography.
3. Explain how the de novo structure-based ligand design works?
4. Explain the methods for handling ligand flexibility during molecular docking.
5. Explain structure activity relationships and quantitative structure activity relationships. How adding new substituents alters the behavior of the molecule? [3+3]

SECTION "D"

(Short answer questions)

[8Q. × 2 = 16 marks]

6. Write short notes on (*ANY EIGHT*):
  - a. Caco-2 assay
  - b. DNA binding Zinc Finger Motif
  - c. Two-state receptor model
  - d. Homology Modeling
  - e. Biological Assemblies
  - f. Octanol/water partition coefficient
  - g. Pharmacophore
  - h. DNA binding drugs
  - i. Small Molecule Preparation during Docking
  - j. Adverse Drug Reaction