



NBG-003-006402 Seat No. _____

B. Sc. (Bioinformatics) (Sem. IV) (CBCS) Examination

April / May - 2017

BI - 402 : Structural Bioinformatics

(Old Course)

Faculty Code : 003

Subject Code : 006402

Time : 2½ Hours]

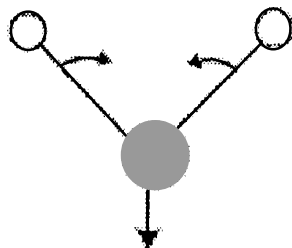
[Total Marks : 70

- Instructions :** (1) All questions are compulsory.
(2) The right side figure indicates total marks of the question.

SECTION – I

1 Answer the short answer questions : **20**

- (1) _____ first summarize a supposed relationship between chemical structures and biological activity in a data-set of chemicals
- (2) All amino acid residues in protein are remarkably in _____ configuration.
- (3) _____ amino acid is responsible for the formation of disulfide bridge.
- (4) Which bond is rigid and planar in protein?
- (5) An "oil drop with a polar coat" is a metaphor referring to the three-dimensional structure of _____,
- (6) X-rays provides the best resolution because Wavelength of X-rays is about the same length of _____ bond
- (7) Identify the vibrational mode in the following molecule



- (8) Globplot is a tool used for _____ prediction
- (9) Which method is used for protein-protein interaction study?
- (10) CAPRI is related to _____ interaction study
- (11) _____ represents a distinct protein sequence and its natural or artificial variants in SCOP
- (12) In distance matrix, each matrix position represents distance between _____
- (13) Which tool can be used for both structure visualization and superposition?
- (14) PSIPRED work based on _____ based method
- (15) What is the window length generally taken by secondary structure predictors?
- (16) What are the general assumptions in *ab initio* methods?
- (17) Which tool is considered as the most comprehensive tool for mitochondrial protein localization prediction?
- (18) Which method is used for protein domain analysis?
- (19) If the bond angles, bond lengths and torsion angles of the components are not modified at any stage of complex generation, it is known as _____
- (20) Gel Retardation assay is used for _____ interaction

SECTION – II

- | | | |
|----------|--|----------|
| 2 | <p>(a) Explain any Three :</p> <ol style="list-style-type: none"> (1) Protein folding (2) Turn or loops (3) Architecture of protein (4) SCOP (5) Double Dynamic Programming (6) Rational Drug,designing | 6 |
| | <p>(b) Explain any Three :</p> <ol style="list-style-type: none"> (1) Vibrational modes in IR Spectroscopy (2) SPDBV (3) CATH (4) Quaternary structure of Protein (5) Super secondary structures (6) Forces responsible for protein folding | 9 |

- (c) Attempt any **Two** : **10**
- (1) Distance matrix method for structure alignment
 - (2) Fold recognition
 - (3) X-Ray Crystallography
 - (4) Protein Secondary Structure Prediction
 - (5) How can you predict pockets and Post translational Modifications?
- 3** (a) Explain any **Three** : **6**
- (1) Protein domain
 - (2) Flexible docking
 - (3) Gel retardation assay
 - (4) Gene fusion
 - (5) Ab initio method
 - (6) Combinatorial Chemistry
- (b) Explain any **Three** : **9**
- (1) Post translational modification prediction
 - (2) Modification interference
 - (3) Protein-protein docking
 - (4) Protein localization prediction
 - (5) Phylogenetic profile method for protein-protein interaction
 - (6) What are the problems in experimental determination of protein structure
- (c) Attempt any **Two** : **10**
- (1) IR Spectroscopy
 - (2) Protein ligand docking
 - (3) Yeast two hybrid approach
 - (4) Homology modeling
 - (5) QSAR
-