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Total Number of Pages : 02

B.Tech
PBT5D001

5th Semester Regular Examination 2018-19
GENOMICS, PROTEOMICS AND META-BOLOMICS

BRANCH : BIOTECH

Time : 3 Hours

Max Marks : 100

Q.CODE : E542

Answer Question No.1 (Part-1) which is compulsory, any EIGHT from Part-II and any TWO from Part-III.

The figures in the right hand margin indicate marks.

Part- I

Q1 Short Answer Type Questions (Answer All-10) (2 x 10)

- What is Pharmacogenetics and Pharmacogenomics?
- What are the factors affecting ionization of an analyte?
- What is molecular docking?
- What is reverse genetics? Write its utility.
- Name a database related to metabolomics and write its significance.
- What are the social implications of sequencing human genome?
- Write the principles of pyrosequencing.
- How many SNPs are exist in human protein coding genes approximately and about how many of these could affect protein structure?
- What do you mean by metagenomics?
- What is a single orphan gene? What is an orphan family?

Part- II

Q2 Focused-Short Answer Type Questions- (Answer Any Eight out of Twelve) (6 x 8)

- What is peptide mass finger-printing? Describe briefly.
- Describe Clone Contig assembly by chromosome walking.
- Briefly delineate the nomenclature used to describe the peptide ion fragmentation for protein analysis using Mass Spectrometry.
- What are the potential applications of metabolomics in biomedical research?
- Distinguish between nucleic acid programmable protein array (NAPPA) and DNA array to protein array (DAPA)
- What is genome mapping? Discuss the types of genome mapping techniques.
- What is Gene Ontology (GO)? Why GO is important in genome biology?
- What is functional genomics? What tools are used in this technology?
- What is posttranslational modification? How can you predict the disulfide bridge, signal peptide and subcellular localization of a protein by using Bioinformatics?
- Write down the principle and applications of NMR spectroscopy.
- What is Homology modelling? How can you build a 3D structure of a protein by using this technique?
- Discuss the various structure based drug design approaches.

Part-III

Long Answer Type Questions (Answer Any Two out of Four)

- Q3** Describe how proteomic analysis is accomplished using the MALDI-TOF Mass spectrometry. **(16)**
- Q4** What do you mean by High-throughput sequencing technology? Discuss the types, principles and advantages of Next generation sequencing technology. **(16)**
- Q5** How can you annotate the genes encoding “hypothetical proteins”? Discuss briefly. **(16)**
- Q6** How protein-protein interactions (ppi) help to understand the molecular interactions in living systems? Briefly describe different methods to predict protein network. **(16)**