# STELLA MARIS COLLEGE (AUTONOMOUS) CHENNAI 600 086 (For candidates admitted from the academic year 2015 – 2016 & thereafter)

**SUBJECT CODE: 15BI/PC/GP34** 

## M. Sc. DEGREE EXAMINATION, NOVEMBER 2018 BIOINFORMATICS THIRD SEMESTER

**COURSE : CORE** 

PAPER : GENOMICS AND PROTEOMICS

TIME : 90 MINUTES MAX. MARKS: 50

#### SECTION - A

### ANSWER ALL THE QUESTIONS:

(20x1=20)

- 1. Which of the following is incorrect regarding the advantages of Molecular data for phylogenetics study?
  - a. They are more numerous than fossil records.
  - b. They are easier to obtain as compared to fossil recors.
  - c. Sampling bias is involved
  - d. More clear-cut & robust phylogenetic trees can be constructed with the molecular data
- 2. The molecular clock is based on the finding that.
  - a. all mutations are either advantageous of disadvantageous to the organism.
  - b. new species have evolved at a kuniform rate throughout evolutionary time
  - c. genetic differences accumulate at a fairly constant rate.
  - d. all mutations are neutral to selection
- 3. Divergence is necessary for all of the following EXCEPT
  - a. speciation
  - b. the accumulation of different random mutations in the genes of two species
  - c. the use of the molecular clock.
  - d. the accumulation of random mutations in the DNA.
- 4. To use molecular data to reconstruct evolutionary history requires making a number of reasonable assumptions. Which of the following is incorrect about it?
  - a. The molecular sequences used in phylogenetic construction are homologous
  - b. The molecular sequences used in phylogenetic construction share a common origin
  - c. Phylogenetic divergence cannot be bifurcationg
  - d. Parent branch splits into two daughter branches at any given point
- 5. International Human Genome project was initiated by
  - a. National Institute of Health (NIH)
  - b. Celera genomics
  - c. US Department of Energy(D0E)
  - d. NOH and US D0E
- 6. DNA sequencing followed by genome annotation are steps of
  - a. Comparative genomics b. Structural genomics
  - c. Functional genomics d. transcriptomics
- 7. All are genome sequencing strategies except
  - a. Edman degradation method b. short gun library
  - c. Whole genome short gun sequencing d. Directed gene sequencing

- 8. Small cDNA sequence that represents a unique segment of an active gene is called
  - a. SNPs b. SnRNAs c. ESTs d. contig
- 9. The technique of subtractive hybridization allows identification of genes that are selectively activated under a certain set of conditions.
  - a. True b. False
- 10. Which of these might be an advantage to genetic testing of individuals via microarrays?
  - a. Many different potential mutations in a single gene could be tested at once.
  - b. Expression patterns of many different genes can be analyzed simultaneously.
  - c. Microarray analysis can provide relative levels of expression of particular genes
  - d. All of these
- 11. Which of these would not be a valid reason that use of microarray technology to differentiate between closely related bacterial species and subspecies is important?
  - a. Certain strains of bacteria are more pathogenic than other related strains
  - b. Some strains of bacteria are more active in bioremediation than other related strains.
  - c. Infection by different strains of bacteria may require different therapeutic approaches
  - d. In many cases, critical information about characteristics of a bacterium causing an infection needs to be immediately available.
- 12. The two most common processes that lead to production of multiple functional proteins from the same DNA sequence are:
  - a. RNA editing and alternative splicing.
  - b. Protein folding and posttranslational covalent modifications
  - c. Alternative splicing and posttranslational covalent modifications
  - d. Posttranslational covalent modification and transcriptional regulation.
- 13. Which of these conclusions might be drawn from the results of a 2D gel electrophoresis experiment?
  - a. Levels of mRNA expression for two different genes are lower under one set of conditions than another
  - b. In a mutant cell, the lack of protein expression is due to production of unstable mRNA, which is rapidly degraded.
  - c. A mutation prevents proper posttranslational modification of a protein
  - d. None of All of these are reasonable conclusions
- 14. Which of these concerns would apply to functional protein microarrays but not antibody microarray?
- 15. How many potential open reading frames are present in a DNA sequence?
- 16. Why might you want to search a database for a protein motif?
- 17. What is a 'proteotypic' peptide?
- 18. Isobaric tags are:
- 19. If your quantitative proteomics experiment contains a large number of samples, which of these would be a good method to chose?
- 20. Which of these is the most important aspect of planning and designing a good proteomics experiment?

### SECTION - B

## ANSWER ANY THREE QUESTIONS (Draw Diagrams wherever necessary)(3x10=30)

- 21. Describe the salient features of Human genome and compare it with chimpanzee genome.
- 22. Give brief account on computer mediated analysis for gene function.
- 23. Explain in details about the pathway regulatory networks with its importance.
- 24. Elaborate the mapping of protein modification with suitable example.
- 25. Write brief account on protein-protein interactions with Y2B method.

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