1STELLA MARIS COLLEGE (AUTONOMOUS) CHENNAI –600 086 (For candidates admitted from the academic year 2015 – 16)

SUBJECT CODE: 15BI/PC/BA44

M. Sc. DEGREE EXAMINATION, APRIL 2017 BIOINFORMATICS FOURTH SEMESTER

COURSE : CORE

PAPER : ADVANCES IN BIOINFORMATICS

TIME : 3 HOURS MAX. MARKS: 100

SECTION - A

ANSWER ALL QUESTIONS

 $(20 \times 1=20)$

- 1. Which of the following statements are not true The function of genes can be determined by
 - i. Gene inactivation ii. Homology search
- iii. Exon trapping iv. Zoo-blotting
- a. i and ii correct b. ii and ii
 - b. ii and iii correct c. i and iii correct
- d. ii and iv correct
- 2. Which of the following statements are not true Microarrays Are used for analysis of
 - i. transcriptomes
- ii. Contain RNA sequences
- iii. Contain DNA sequences iv. Are smaller than DNA chips
- a. i and ii correct b. ii and iii correct c. i and iii correct
- d. ii and iv correct
- 3. Which of the following statements are not true ORF scanning
 - i. Is used to find exons
- ii. Is used to find intergenic sequences
- iii. Is used to find gene homologies iv. Is used to find protein-coding genes
- a. i correct b. ii correct c. iii correct d. iv correct
- 4. Expand SMILES
- 5. Choose the correct chemical structure drawing packages
 - a. Rasmol
- b. chem Draw
- c. Pharma GKB
- d. Pubchem
- 6. Which of these projects would be best suited for Next Generation Sequencing?
 - a. To determine if a tumour sample contains a common missense mutation
 - b. To find the transcriptome of a tumour sample
 - c. To genotype ten genomic DNA samples for a known single nucleotide polymorphism
 - d. All of the above.
- 7. Automated DNA sequencing is an improvement of Sanger's method where
 - a) ddNTPS are used for chain termination
 - b) PCR is used for making sequencing templates
 - c) Fluorescently labelled dNTPs are used for chain termination
 - d) Fluorescently labelled ddNTPs are used for chain termination
- 8. MAML stands for_____
- 9. Comment on oligonucleotide.
- 10. The intensity of the pharmacological action of a drug mostly depends on the
 - a. Concentration of drug at the receptor site
 - b. Minimum toxic concentration of the drug
 - c. Minimum effective concentration of the drug
 - d. Elimination half life of the drug
- 11. Personalised medicine has the potential to yield plenty of health and economic benefits. Which of the following would not be a benefit of personalised medicine?
 - a. Increased number of medical jobs
- b. Improved medical decision making
- c. Delivery of most effective therapies
- d. Optimise disease prevention strategies

- 12. Mention a few database for toxicogenomics.
- 13. Which of the following is not true regarding pharmacogenomics?
 - a. The goal is to minimize drug toxicity
 - b. It focuses on individual candidate genes to identify markers that affect drug metabolism and drug effect
 - c. Drug-food interactions and drug-drug interactions are umimportant in pharmacogenomics
 - d. There is no variation of drug dose between patients
- 14. The most commonly occurring variant in the human genome is
 - a. tandem-repeat polymorphism.
- b. premature stop codon.
- c. nucleotide base insertion.
- d. single-nucleotide polymorphism.
- 15. CYP2D6 polymorphism can affect:
 - a. drug efficacy.

b. drug toxicity.

c. drug interaction potential.

- d. a, b, and c.
- 16. Genetic variations in drug targets may contribute to which drug property?
 - a. Bioavailability

- b. Half-life
- c. Racial differences in response
- d. Peak-dose area under the curve
- 17. Potential outcomes of pharmacogenetic research include all the following except
 - a. lower incidence of adverse drug effects. b. higher health care costs.
 - c. improved treatment outcomes.
 D. pretreatment screening for polymorphisms.
- 18. The R package which contains the bioinformatics modules is ____
 - a. NumR
- b. Bioconductor
- c. Bio R
- d. String R
- 19. What will be the output of following code snippet?

> paste("a", "b", sep = ":")

a. "a+b" b. "a=b'

b. "a=b" c. "a:b"

d. None of the mentioned

20. R Package was developed by

SECTION - B

ANSWER ANY FOUR QUESTIONS. EACH ANSWER SHOULD NOT EXCEED 500 WORDS. ALL QUESTIONS CARRY EQUAL MARKS. DRAW DIAGRAMS WHEREVER NECESSARY $(4 \times 10 = 40)$

- 21. What you mean safety metabolism?
- 22. Define the term preclinical toxicology
- 23. Briefly explain the tools used for construction of 2D structure
- 24. Explain the types of Next generation sequencing used of DNA
- 25. Briefly describe the History of DNA sequencing
- 26. Illustrate the term GEO
- 27. R is a Deluxe calculator justify

SECTION - C

ANSWER ANY TWO QUESTIONS. EACH ANSWER SHOULD NOT EXCEED 1200 WORDS. ALL QUESTIONS CARRY EQUAL MARKS. DRAW DIAGRAMS WHEREVER NECESSARY (2 X 20 = 40)

- 28. Describe the in details about the tools used microarray data analysis. What are the processes involved in visualizing microarray data?
- 29. Write the salient features involved in the Graphics in the R programming.
- 30. Explain the run types and analysis of data from the Next generation sequencing.
- 31. Find out the relationship between pharmacokinetics and metabolism with suitable example.