

- c) What is out-group sequence in phylogenetic tree and why it is so important in phylogenetic analysis ? Define the phylogenetic tree construction methods that directly based on sequences. 5+2+3

**M.Sc. (BIOTECHNOLOGY) PART - II EXAMINATION, 2019**

**COMPUTER AND BIOINFORMATICS**

**PAPER - 2/3**

Time : Four hours

Full Marks : 100

**PART - I**

Answer ***Q1*** and ***any two*** from rest

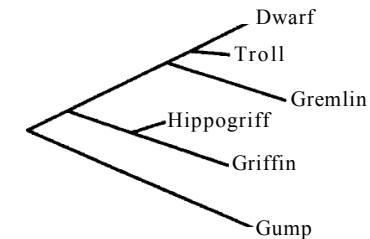
1. Answer ***any five*** questions : 5×4
  - a) What do you mean by algorithm ? What are its properties?
  - b) Differentiate between while and do while loop structure in C programming language.
  - c) Perform the following using BCD arithmetic :
    - i)  $27 + 79$                       ii)  $97 - 54$
  - d) Represent  $(-123.25)_{10}$  using IEEE 32-bit notation.
  - e) Differentiate between `i++` and `++i` in C programming language.
  - f) What are the differences between compiler and interpreter?
2. a) Write an algorithm to check a given integer is prime or not.

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- b) Draw a flowchart to find maximum between three numbers. 5+5+5
- c) What are the function of strlen(), strcat() and gets() in C language. 5+5+5
3. a) Write a C program to find maximum from a set of N numbers. N should be read from user.
- b) Write an algorithm to calculate  $1 + 1/X + 1/X^2 + \dots + 1/X^N$ .
- c) What are the different methods to represent a signed number in binary number system ? Represent  $(-45)_{10}$  using these methods with 16 bits. 6+5+4
4. a) Write a C program to convert a temperature from Celsius to Fahrenheit.
- b) Write a C program that accept a string and find number of vowels present in the string.
- c) Do you think that “1’s complement is better than 2’s complement”? Give reason to support your opinion. 5+6+4

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- c) What do you mean by similarity matrix ? Explain with example.
- d) Define the Total Score, Max Score, Query coverage and E-value, Identity in the context of sequence alignment.
- e) What’s the three major databases with their respective operative countries. Name the two structure databases. 3+2
- f) What are the differences between PAM and BLOSUM matrix. 5
8. Answer **any one** of the following : 10×1=10
- a) What are Databanks in the context of Bioinformatics ? How do you classify various types of databanks pertaining to bioinformatics. 5+5=10
- b) Answer the following questions about this tree :



- i) Which are the two most closely related species ?
- ii) Which of these is probably the outgroup ?
- iii) Which is most distantly related to the Gremlin other than the outgroup species ?
- iv) Write down the few basic character of outgroup.

[ Turn over

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- i) Submission to GenBank include
- i) Sequin and BankIt      ii) Sequin
  - iii) BankIt                  iv) BankIt and sequein
- j) Name two different methods used in pair wise sequence alignment.
6. Answer **any five** of the following :                  2×5=10
- a) Write the two most commonly used scoring matrix.
  - b) Why the alignment between two sequences is performed?
  - c) Write name of two web based tools used for restriction mapping.
  - d) What do you mean by Express Sequence Tags (EST) database ?
  - e) What is bioinformatics ?
  - f) What do you mean by local and global alignment ?
  - g) What do you mean by identity and similarity between two protein sequences ?
7. Answer **any four** of the following :                  5×4=20
- a) What is DNA sequencing chromatogram ? Explain how to interpret it ?    3+2
  - b) What is Homologs, Paralogs, Orthologs. Explains them with examples.    3+2

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## PART - II

Answer **all the** questions.

The figures in the margins indicate full marks.

5. Answer the following questions :                                  1×10=10
- a) What is primary and secondary database ?
  - b) What makes BLAST more useful than FASTA for database search ?
  - c) What are the basic structure of a GenBank and an EMBL entry ?
  - d) Write the name of scoring model used to build similarity between nucleotide sequences ?
  - e) For what purpose BLOSUM matrices are used ?
  - f) What is FASTA format ?
  - g) Which one is used for search all NCBI associated databases ?
    - i) BLAST                                  ii) CLUSTAL W
    - iii) ENTREZ                                  iv) PDB
  - h) If you want to BLAST the non-redundant database using a new protein sequences as query, which is the BEST search program to use ?
    - i) blastp    ii) blastn    iii) tblastx    iv) blastx

[ Turn over