



Name :
Roll No. :
Invigilator's Signature :

CS/B.Sc.(H)/BT/Gen/Micro-Bio/Mol-Bio/SEM-4/CBB-402/2013

2013

COMPUTATIONAL BIOLOGY & BIO-INFORMATICS

Time Allotted : 3 Hours

Full Marks : 70

The figures in the margin indicate full marks.

*Candidates are required to give their answers in their own words
as far as practicable.*

GROUP - A

(Multiple Choice Type Questions)

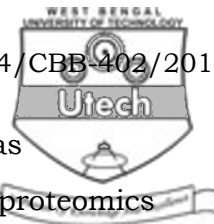
1. Choose the correct alternatives for any *ten* of the following :

$10 \times 1 = 10$

- i) Which of these is not a protein sequence database ?
 - a) PIR
 - b) GenBank
 - c) PDB
 - d) Swiss-Prot.
- ii) BCA stands for
 - a) Bacteria Artifice Chromosome
 - b) Bacterial Artifice Chromosome
 - c) Bacterial Artificial Chromosome
 - d) Bacterial Artificial Chromatid.
- iii) Which step is omitted in whole genome sequencing ?
 - a) Random shqedding of genome
 - b) Making phage library
 - c) Computational analysis
 - d) None of these.



- iv) Which of the following is not primary nucleotide sequence database ?
- a) DDBJ b) OWL
c) GEN BANK d) EMBL.
- v) URL for NCBI is
- a) www.ncbi.nlm.nih.gov b) www.ncbi.nih.nlm.gov
c) www.ncbi.gov d) www.ncbi.nlm.gov.
- vi) Clastal W
- a) multiple sequence alignment tool
b) protein secondary structure predicting tool
c) data retrieving tool
d) nucleic acid sequence analysis tool.
- vii) Which is data retiring tool ?
- a) KEGG b) EMBL
c) ENTERZ d) PHD.
- viii) BLAST & FASTA are used for
- a) global similarity
b) end free space alignment
c) local similarity
d) gap penalty.
- ix) Information of all known nucleotide & protein sequel are available on
- a) NCBI gene bank b) DDBJ
c) EMBL d) all of these.
- x) What is the primary goals of databases ?
- a) Minimizing data redundancy
b) Achieving data independence
c) Both (a) and (b)
d) None of these.
- xi) Phylogenetic relationship can be shown by
- a) Dendogram b) Data search tool
c) Data retrieving d) Gene bank.



- xii) Proteomics research can be categorized as
- structural proteomics & functional proteomics
 - structural functional & comparative functional
 - functional & comparative proteomics
 - none of these.

GROUP – B

(Short Answer Type Questions)

Answer any *three* of the following. $3 \times 5 = 15$

- Define WWW, HTTP, HTML, URL. $2 + 3$
- What is substitution matrix ? Write down the difference between PAM & BLOSUM matrix. $2 + 3$
- What is sequence alignment ? Write about *E* value & *Z* score. $2 + 3$
- Write down the difference between FASTA & BLAST. $2 + 3$
- Write about data retrieval tool. What is sequin & bankit ? $2 + 3$

GROUP – C

(Long Answer Type Questions)

Answer any *three* of the following. $3 \times 15 = 45$

- What is Secondary Database ? Name two Secondary Databases.
 - What is the difference between Accession Number and Gi Number ?
 - What are the basis of database searching in bioinformatics ?
 - What do you mean by BankIT ? $1 + 2 + 4 + 5 + 3$



8. a) Compare the features between Primary, Secondary and Tertiary database.
- b) What is PFAM and BLOCKS ?
- c) Write down the differences between Flat File and Hierarchical files.
- d) Why Biological Database are important ? 5 + 3 + 4 + 3
9. What is Dynamic Programming ? What are the general steps of it ? Name two database searching tools. Fill the matrix for DP and align the following sequences :
- GAATTCAGTTA (sequence#1) (Upto Matrix fill step only)
- GGATCGA (sequence#2) 2 + 1 + 2 + 7 + 3
10. What is substitution matrix ? In "PAM 250" the 250 stands for what ? Write down the difference of PAM 250 and BIOSM 62 matrix. What is humming and levenstein distance ? Write down one name of two small molecule database. What is Pubmed ? What is the name of the database where you get the inheritable disease of human ? What is word size ? Comment on sensitivity and selectivity ?
- 1 + 1 + 3 + 4 + 1 + 1 + 1 + 1 + 2
11. State the difference between homology and similarity ? Write down 6 emerging fields in bioinformatics. You have given a sequence of African elephant and Indian elephant how would you draw the phylogenetic relation between them ? What would you expect the result. What is relative mutability ?

3 + 3 + 4 + 2 + 3

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