Roll No	 				Question Booklet Number
O. M. R. Serial No.					

M. Sc. (Biochemistry) (Fourth Semester) EXAMINATION, July, 2022

BIOINFORMATICS

Paper Code							
BCH	4	0	0	2			

Questions Booklet Series

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[Maximum Marks : 100

Time: 1:30 Hours]

Instructions to the Examinee:

- 1. Do not open the booklet unless you are asked to do so.
- 2. The booklet contains 60 questions. Examinee is required to answer any 50 questions in the OMR Answer-Sheet provided and not in the question booklet. If more than 50 questions are attempted by student, then the first attempted 50 questions will be considered for evaluation. All questions carry equal marks.
- 3. Examine the Booklet and the OMR Answer-Sheet very carefully before you proceed. Faulty question booklet due to missing or duplicate pages/questions or having any other discrepancy should be got immediately replaced.

परीक्षार्थियों के लिए निर्देश :

- प्रश्न-पुस्तिका को तब तक न खोलें जब तक आपसे कहा न जाए।
- 2. प्रश्न-पुस्तिका में 60 प्रश्न हैं। परीक्षार्थी को किन्हीं 50 प्रश्नों को केवल दी गई OMR आन्सर-शीट पर ही हल करना है, प्रश्न-पुस्तिका पर नहीं। यदि छात्र द्वारा 50 से अधिक प्रश्नों को हल किया जाता है तो प्रारम्भिक हल किये हुए 50 उत्तरों को ही मूल्यांकन हेतु सम्मिलित किया जाएगा। सभी प्रश्नों के अंक समान हैं।
- उ. प्रश्नों के उत्तर अंकित करने से पूर्व प्रश्न-पुस्तिका तथा OMR आन्सर-शीट को सावधानीपूर्वक देख लें। दोषपूर्ण प्रश्न-पुस्तिका जिसमें कुछ भाग छपने से छूट गए हों या प्रश्न एक से अधिक बार छप गए हों या उसमें किसी अन्य प्रकार की कमी हो, तो उसे तुरन्त बदल लें।

(शेष निर्देश अन्तिम पृष्ठ पर)

(Only for Rough Work)

1.	UPGMA protocol is used for generation	5.	The statistical packages among the					
	of:		following are :					
	(A) Phylogenetic tree		(A) SPSS					
	(B) Secondary structures		(A) 51 55					
	(C) Heat maps		(B) Sigma					
	(D) None of the above		(C) R					
2.	EMBL is:		(D) All of the above					
	(A) European Molecular Biology	6	Text based format to represent a					
	Laboratory	6. Laboratory						
	(B) Located at Germany		nucleotide or amino acid is					
	(C) (A) and (B) are true		(A) BLAST					
	(D) None of the above		(B) FASTA					
3.	Ac/Ds elements are :		(C) Multiple sequence alignment					
	(A) Molecular markers		(D) PROSITE					
	(B) Primer sequences		(2) 11102112					
	(C) Transposons	7.	Which of the following is incorrect about					
	(D) Peptide fragments		ENTREZ ?					
4.	Which of these is incorrect for stop		(A) It provides a series of forms that					
	codon?		can be filled out to retrieve a					
	(A) Amber		Medline reference related to the					
	(B) Umbel							
	(C) Opal		<i>.</i>					
	(D) Ochre		databases.					

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- (B) It provides a series of forms that can be filled out to retrieve a DNA or protein sequence.
- (C) It is a resource prepared only by the staff of the National Centre for Biotechnology Information.
- (D) One straightforward way to access the sequence databases is through ENTREZ.
- 8. Which of the following is not the objective to perform sequence comparison?
 - (A) To find the common motifs present in both sequences
 - (B) To study the physical properties of molecules
 - (C) To study evolutionary relationships
 - (D) To observe patterns of conservation

- 9. Which of the following is untrue about homology modelling?
 - (A) It doesn't involve the evolutionary distances anywhere.
 - (B) The principle behind it is that if two proteins share a high enough sequence similarity, they are likely to have very similar three-dimensional structures.
 - (C) Homology modelling predicts protein structures based on sequence homology with known structures.
 - (D) It is also known as comparative modelling.
- 10. The process of finding the relative location of genes on a chromosome is called:
 - (A) Gene tracking
 - (B) Genome walking
 - (C) Genome mapping
 - (D) Chromosome walking
- 11. The term 'in vitro' refers to:
 - (A) Within the lab
 - (B) Within the cell
 - (C) Within the glass
 - (D) Outside the glass

12.	The laboratory work using computers	16.	Bioir	nformatics deals with:
	and associated with web-based analysis		(A)	Application of statistical tools for
	is referred to as:			analysis of biological data
	(A) In silico		(B)	Application of information
	(B) Dry lab			technology tools for analysis of
	(C) Wet lab			biological data
	(D) Pure lab		(C)	Application of biophysical
13.	Analysing or comparing entire genome			techniques for analysis of
	of organism:			biological data
	(A) Genomics		(D)	Entrepreneurial application of
	(B) Proteomics			biological research
	(C) Pharmacogenomics	17.	Dege	eneracy of genetic code explains:
	(D) Metabalomics		(A)	Each amino acid is coded by
14.	Which of the following is a mail client?			multiple codons
	(A) PINE		(B)	Each codon codes for single amino
	(B) Google			acid
	(C) Eudora		(C)	Triplet codon is without any gaps
	(D) All of the above		(D)	Genetic code is universal
15.	Types of FTP include :	18.	Ident	tify an operating system:
	(A) FTPES		(A)	Windows 10
	(B) FTPS		(B)	Linux
	(C) SFTP		(C)	Unix
	(D) All of the above		(D)	All of the above

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- 19. The extension used for file transfer protocol:
 - (A) .doc
 - (B) .xls
 - (C) .ftp
 - (D) .ptx
- 20. NCBI stands for:
 - (A) National Center for Bioinformatics
 - (B) National Center for Biotechnology

 Information
 - (C) National Center for Biology

 Information
 - (D) National Center for Biomedical

 Information
- 21. Which of the following is useful for construction of phylogenetic tree ?
 - (A) Dendrogram
 - (B) Cladogram
 - (C) Phylogram
 - (D) All of the above

- 22. Which of the following is true about Proteomics?
 - (A) Proteomics has enabled the identification of ever increasing numbers of protein.
 - (B) Proteomics generally refers to the large-scale experimental analysis of proteins and proteomes.
 - (C) Proteome is the entire set of proteins that is produced or modified by an organism or system.
 - (D) All of the above
- 23. Regarding structural proteomics which of the following is true?
 - (A) Structure domain is an element of proteins overall structure and often folds independent of rest of protein chain.
 - (B) Ribbon and Cartoon Diagram of protein structure gives information about various secondary structures that occurs in protein.
 - (C) Structure proteomics include the analysis of protein structure at large scale.
 - (D) All of the above

24.	The	database	useful	for	homology	
	mode	elling of pro	oteins :			

- (A) BLAST
- (B) EMBL
- (C) SwissMODEL
- (D) DDBJ

25. Which of the following statements is true?

- (A) Multiple Sequence Alignment(MSA) is useful to know the conserved regions of genes.
- (B) Alignment can be done for both genes and protein sequences.
- (C) Multalin is useful in performing the sequence alignment.
- (D) All of the above

26. Central dogma of molecular biology refers to:

- (A) DNA $\rightarrow c$ DNA \rightarrow Protein
- (B) DNA \rightarrow RNA \rightarrow Protein
- (C) Protein \rightarrow RNA \rightarrow DNA
- (D) $RNA \rightarrow DNA \rightarrow Protein$

- 27. The software tool used for sequence alignment:
 - (A) C++
 - (B) PRISM
 - (C) HTML
 - (D) CLUSTALW

28. BLASTx is useful for:

- (A) Translated nucleotide to protein
- (B) Protein to translated nucleotide
- (C) Protein to protein
- (D) Nucleotide to nucleotide

29. FASTA format is used to represent:

- (A) Nucleotide or amino acid sequences in standard format
- (B) Diagrammatic representation of protein structures
- (C) Phylogenetic tree
- (D) Sugar moieties in a glycoprotein
- 30. All are sequence alignment tools, except:
 - (A) Rasmol
 - (B) BLAST
 - (C) MultAlin
 - (D) CLUSTAL W

31.	The sequence alignment tool provided by	35.	Incor	rect statement for BLOSUM:
	NCBI is:		(A)	BLOSUM is substitution matrix
	(A) Chime		(B)	Rely on substitution sequences
	(B) BLAST		(C)	Rely on conserved sequences
	(C) Multalin		(D)	Do not measure the evolutionary
	(D) CLUSTALW			distance
32.	The procedure of aligning many	36.	MEG	GA 7 is:
	sequences simultaneously:		(A)	Molecular Evolutionary Genetics
	(A) Multiple Sequence Alignment			Analysis
	(B) Pairwise alignment		(B)	Creates dendrogram
			(C)	Useful for phylogenetic
	(C) Global alignment			relationship
	(D) Local alignment		(D)	All of the above
33.	Nucleotide sequence databases include :	37.	ww	W is:
	(A) PDB		(A)	World Wide Web
	(B) ExPASY		(B)	World Wired Web
	(C) SWISSPROT		(C)	World War Web
	(D) GenBank		(D)	World Wild Web
34.	SwissProt is:	38.	CEN	SOR program is used to find:
	(A) Protein database		(A)	Multiple sequence alignment
	(B) Nucleotide database		(B)	Repetitive elements
	(C) UniProt consortium		(C)	SNP
	(D) Both (A) and (C)		(D)	Protein modelling

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39. Transposons:

- (A) can transfer genetic information form one position to another position in the genome.
- (B) can operate through DNA or RNA sequences.
- (C) are important factor for evolution.
- (D) All of the above
- 40. Which of the following tools is used to find repetitive sequences ?
 - (A) Repbase
 - (B) CENSOR
 - (C) Dfam
 - (D) All of the above
- 41. GCG software tool is useful for:
 - (A) Analysis of gene and protein sequences
 - (B) Performing GO annotation
 - (C) Identifying SNPs
 - (D) Analyzing repetitive sequences

- 42. Which of the following is a feature of database?
 - (A) To deposit and store data
 - (B) To retrieve data
 - (C) To analyse and interpret data
 - (D) All of the above
- 43. Which of the following is a nucleotide database?
 - (A) EMBL
 - (B) Pfam
 - (C) Swiss-Prot
 - (D) Uni-Prot
- 44. Which of the following databases is useful for whole genome sequences?
 - (A) Ensembl
 - (B) Expasy
 - (C) SwillProt
 - (D) UniProt
- 45. How many different open reading frames are possible in an mRNA strand?
 - (A) 2
 - (B) 3
 - (C) 4
 - (D) 5

всн-	-4002	(10)			Set-D
	(D)	None of the above		(D)	All of the above
	(C)	Database of nucleotide sequences		(C)	Web of science
	(B)	Repository of protein sequences		(B)	Pubmed
		references		(A)	Science direct
	(A)	A search engine for journal	53.	Journ	nal search can be done using:
49.	PubN	Med is:		(D)	None of the above
	(D)	Sequence retrieval		(C)	NCBI
	(C)	Structure prediction		(B)	PDB
	(B)	Protein modelling		(A)	DDBJ
	(A)	Multiple sequence alignment		analy	ysing structure of macromolecules?
48.	Profi	le analysis is used for :	52.	Whic	ch of the following is an example for
	(D)	1000 MB		(D)	NCBI
	(C)	500 MB		(C)	University of Stanford
	(B)	250 MB		(B)	University of California
	(A)	25 MB		(A)	University of Geneva
	attacl	hment ?	51.	Swis	sProt is operated by:
47.	What	t is the size limitation for email		(D)	All of the above
	(D)	Data Domain Bank of Japan		(C)	Science Direct
	(C)	DNA Domain Bank of Japan		(B)	PubMed
	(B)	DNA Database of Japan		(A)	Google scholar
	(A)	DNA Data Bank of Japan		prepa	aring virtual library :
46.	Expa	nd DDBJ:	50.	The	following tool(s) will be helpful for

54.	Туре	es of	molecular	mar	kers	57.	iTRA	AQ stands for :
	inclu	ıde :					(A)	Isobaric Tags for Relative and
	(VAUTO						Absolute Quantitation
	(A)	VNTR					(B)	Isometric Tags for Relative and
	(B)	RAPD						Absolute Quantitation
	(C)	RFLP					(C)	Isotopic Tags for Relative and
	(D)	All of the	ahove					Absolute Quantitation
	(D)	All of the above					(D)	None of the above
55.	Expa	and RAPD:				58.	Ram	achandran plot is used for:
	(A)	Rapid	Amplifica	tion	of		(A)	Analysing the angles of amino
		Polymorphic DNA						acids in a polypeptide
	(B)	Torymorphic DTVT	((B)	Analysing protein-protein			
		Random Amplified Polymorphic					interaction	
		DNA				(C)	Analysing quaternary structure of	
	(C)	Regular Amplified Polymorphic						protein
		DNA					(D)	Analysing protein-ligand interaction
	(D)	Dara Amplification of Dalymamhia				59.	Acco	ording to IUPAC, purines are
	(D)	Rare Amplification of Polymorphic				indic	cated by the symbol:	
		DNA					(A)	R
56.	Meth	Methods of proteomic analysis include :					(B)	Y
	1,1001						(C)	K
	(A)	iTRAQ					(D)	M
	(B)	2-DE				60.	FAS	TA sequences are preceded with:
	(C)	Both (A) a				(A)	<	
	` '	None of the above			(B) (C)	>		
	(D)					` /	= :	
							` /	

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4. Four alternative answers are mentioned for each question as—A, B, C & D in the booklet. The candidate has to choose the most correct/appropriate answer and mark the same in the OMR Answer-Sheet as per the direction:

Example:

Question:

Q.1 (A) (C) (D)
Q.2 (A) (B) (C) (D)
Q.3 (A) (C) (D)

Illegible answers with cutting and over-writing or half filled circle will be cancelled.

- 5. Each question carries equal marks. Marks will be awarded according to the number of correct answers you have.
- 6. All answers are to be given on OMR Answer sheet only. Answers given anywhere other than the place specified in the answer sheet will not be considered valid.
- 7. Before writing anything on the OMR Answer Sheet, all the instructions given in it should be read carefully.
- 8. After the completion of the examination candidates should leave the examination hall only after providing their OMR Answer Sheet to the invigilator. Candidate can carry their Question Booklet.
- 9. There will be no negative marking.
- 10. Rough work, if any, should be done on the blank pages provided for the purpose in the booklet.
- 11. To bring and use of log-book, calculator, pager and cellular phone in examination hall is prohibited.
- 12. In case of any difference found in English and Hindi version of the question, the English version of the question will be held authentic.
- Impt.: On opening the question booklet, first check that all the pages of the question booklet are printed properly. If there is ny discrepancy in the question Booklet, then after showing it to the invigilator, get another question Booklet of the same series.

4. प्रश्न-पुस्तिका में प्रत्येक प्रश्न के चार सम्भावित उत्तर—
A, B, C एवं D हैं। परीक्षार्थी को उन चारों विकल्पों में से
एक सबसे सही अथवा सबसे उपयुक्त उत्तर छाँटना है।
उत्तर को OMR आन्सर-शीट में सम्बन्धित प्रश्न संख्या में
निम्न प्रकार भरना है:

उदाहरण :

प्रश्न :

प्रश्न 1 (A) (C) (D) प्रश्न 2 (A) (B) (D) प्रश्न 3 (A) (C) (D)

अपठनीय उत्तर या ऐसे उत्तर जिन्हें काटा या बदला गया है, या गोले में आधा भरकर दिया गया, उन्हें निरस्त कर दिया जाएगा।

- 5. प्रत्येक प्रश्न के अंक समान हैं। आपके जितने उत्तर सही होंगे, उन्हीं के अनुसार अंक प्रदान किये जायेंगे।
- 6. सभी उत्तर केवल ओ. एम. आर. उत्तर-पत्रक (OMR Answer Sheet) पर ही दिये जाने हैं। उत्तर-पत्रक में निर्धारित स्थान के अलावा अन्यत्र कहीं पर दिया गया उत्तर मान्य नहीं होगा।
- 7. ओ. एम. आर. उत्तर-पत्रक (OMR Answer Sheet) पर कुछ भी लिखने से पूर्व उसमें दिये गये सभी अनुदेशों को सावधानीपूर्वक पढ लिया जाये।
- 8. परीक्षा समाप्ति के उपरान्त परीक्षार्थी कक्ष निरीक्षक को अपनी OMR Answer Sheet उपलब्ध कराने के बाद ही परीक्षा कक्ष से प्रस्थान करें। परीक्षार्थी अपने साथ प्रश्न-पुस्तिका ले जा सकते हैं।
- 9. निगेटिव मार्किंग नहीं है।
- 10. कोई भी रफ कार्य, प्रश्न-पुस्तिका के अन्त में, रफ-कार्य के लिए दिए खाली पेज पर ही किया जाना चाहिए।
- 11. परीक्षा-कक्ष में लॉग-बुक, कैलकुलेटर, पेजर तथा सेल्युलर फोन ले जाना तथा उसका उपयोग करना वर्जित है।
- 12. प्रश्न के हिन्दी एवं अंग्रेजी रूपान्तरण में भिन्नता होने की दशा में प्रश्न का अंग्रेजी रूपान्तरण ही मान्य होगा।

महत्वपूर्ण : प्रश्नपुस्तिका खोलने पर प्रथमतः जाँच कर देख लें कि प्रश्न-पुस्तिका के सभी पृष्ठ भलीभाँति छपे हुए हैं। यदि प्रश्नपुस्तिका में कोई कमी हो, तो कक्षनिरीक्षक को दिखाकर उसी सिरीज की दूसरी प्रश्न-पुस्तिका प्राप्त कर लें।