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	

Series

Questions Booklet

B

[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।
- 3. 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)

(Remaining instructions on the last page)

(Only for Rough Work)

1.	The sequence alignment tool provided by	5.	Incorrect 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
2.	The procedure of aligning many	6.	MEGA 7 is:
	sequences simultaneously:		(A) Molecular Evolutionary Genetics
	(A) Multiple Sequence Alignment		Analysis
	(B) Pairwise alignment		(B) Creates dendrogram
	<u>-</u>		(C) Useful for phylogenetic
	(C) Global alignment		relationship
	(D) Local alignment		(D) All of the above
3.	Nucleotide sequence databases include :	7.	WWW 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
4.	SwissProt is:	8.	CENSOR 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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9.	Transp	posons :	12.	Which of the following is a feature of database?		
	(A) (can transfer genetic information				
	f	form one position to another		(A)	To deposit and store data	
		•		(B)	To retrieve data	
	F	position in the genome.		(C)	To analyse and interpret data	
	(B) c	can operate through DNA or RNA		(D)	All of the above	
	S	sequences.	13.	Which	of the following is a nucleotide	
	(C) a	are important factor for evolution.		databa	ise ?	
	(D) A	All of the above		(A) l	EMBL	
				(B)	Pfam	
10.	Which	Which of the following tools is used to			Swiss-Prot	
	find repetitive sequences?			(D)	Uni-Prot	
	(A) I	Repbase	14.	Which	n of the following databases is	
	(B) (CENSOR		useful	for whole genome sequences?	
	(C) I	Dfam		(A) l	Ensembl	
	(D) A	All of the above		(B) l	Expasy	
	(D) 1	Thi of the above		(C)	SwillProt	
11.	GCG s	software tool is useful for:		(D)	UniProt	
	(A) A	Analysis of gene and protein	15.	How n	many different open reading frames	
	S	sequences		are pos	ssible in an mRNA strand?	
	(B) I	Performing GO annotation		(A) Z	2	
	(C) I	Identifying SNPs		(B) 3	3	
	(C) I	dendrying bivi s		(C) 4	4	
	(D) A	Analyzing repetitive sequences		(D) :	5	

	(A)	DNA Data Bank of Japan		prepa	aring virtual library :
	(B)	DNA Database of Japan		(A)	Google scholar
	(C)	DNA Domain Bank of Japan		(B)	PubMed
	(D)	Data Domain Bank of Japan		(C)	Science Direct
17.	Wha	t is the size limitation for email		(D)	All of the above
	attac	hment ?	21.	Swis	sProt is operated by:
	(A)	25 MB		(A)	University of Geneva
	(B)	250 MB		(B)	University of California
	(C)	500 MB		(C)	University of Stanford
	(D)	1000 MB		(D)	NCBI
18.	Profi	ile analysis is used for :	22.	Whic	ch of the following is an example for
	(A)	Multiple sequence alignment		analy	ysing structure of macromolecules?
	(B)	Protein modelling		(A)	DDBJ
	(C)	Structure prediction		(B)	PDB
	(D)	Sequence retrieval		(C)	NCBI
19.	Publ	Med is:		(D)	None of the above
	(A)	A search engine for journal	23.	Journ	nal search can be done using:
		references		(A)	Science direct
	(B)	Repository of protein sequences		(B)	Pubmed
	(C)	Database of nucleotide sequences		(C)	Web of science
	(D)	None of the above		(D)	All of the above

(5)

20.

The following tool(s) will be helpful for

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Expand DDBJ:

16.

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								AQ stands for :		
	inclu	ıde:				(A)	Isobaric Tags for Relative and			
	(A)	VNTR				(-)	Absolute Quantitation			
	(B)	RAPD					(B)	Isometric Tags for Relative and Absolute Quantitation		
	(C)	RFLP				(C)	Isotopic Tags for Relative and			
	(D)	D) All of the above						Absolute Quantitation		
							(D)	None of the above		
25.	Expa	and RAPD :				28.	Ram	achandran plot is used for:		
	(A)	Rapid	Amplification	on	of		(A)	Analysing the angles of amino		
		Polymorphi					acids in a polypeptide			
	(D)	Random Amplified Polymorphic	hio		(B)	Analysing protein-protein				
	(B)				ilic			interaction		
		DNA	DNA					Analysing quaternary structure of		
	(C)	Regular Amplified Polymorphic		hic			protein			
		DNA					(D)	Analysing protein-ligand interaction		
	(D)	Rare Amplification of Polymorphic		hic	29.	Acco	ording to IUPAC, purines are			
	` /				indicated by the symbol:					
		DNA				(A)	R			
26.	Meth	Methods of proteomic analysis include :					(B)	Y		
	(A)	iTR A O					(C) (D)	K M		
		iTRAQ				30.	FASTA sequences are preceded with:			
	(B)	2-DE			30.		(A)	<		
	(C)	Both (A) an	nd (B)			(B)	>			
	(D)	None of the	e above				(C)	=		
							(D)	:		

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31.	UPC	GMA protocol is used for generation	35.	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
32.	EMI	BL is :		(D) All of the above
	(A)	European Molecular Biology	26	The board former to
		Laboratory	36.	Text based format to represent a
	(B)	Located at Germany		nucleotide or amino acid is
	(C)	(A) and (B) are true		(A) BLAST
	(D)	None of the above		(B) FASTA
33.	Ac/I	Os elements are :		(C) Multiple sequence alignment
	(A)	Molecular markers		(D) PROSITE
	(B)	Primer sequences		(D) TROSITE
	(C)	Transposons	37.	Which of the following is incorrect about
	(D)	Peptide fragments		ENTREZ ?
34.	Whi	ch of these is incorrect for stop		(A) It provides a series of forms that
	code	on ?		can be filled out to retrieve a
	(A)	Amber		Medline reference related to the
	(B)	Umbel		
	(C)	Opal		molecular biology sequence
	(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.
- 38. 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

- 39. 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.
- 40. 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
- 41. The term 'in vitro' refers to:
 - (A) Within the lab
 - (B) Within the cell
 - (C) Within the glass
 - (D) Outside the glass

42.	The laboratory work using computers	46.	Bioin	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
43.	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	47.	Dege	neracy of genetic code explains:
	(D) Metabalomics		(A)	Each amino acid is coded by
44.	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
45.	Types of FTP include :	48.	Ident	ify 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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- 49. The extension used for file transfer protocol:
 - (A) .doc
 - (B) .xls
 - (C) .ftp
 - (D) .ptx
- 50. 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
- 51. Which of the following is useful for construction of phylogenetic tree ?
 - (A) Dendrogram
 - (B) Cladogram
 - (C) Phylogram
 - (D) All of the above

- 52. 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
- 53. 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

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

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

55. 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

56. 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) $RNA \rightarrow DNA \rightarrow Protein$

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

58. BLASTx is useful for:

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

59. 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
- 60. All are sequence alignment tools, except:
 - (A) Rasmol
 - (B) BLAST
 - (C) MultAlin
 - (D) CLUSTAL W

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।