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

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

HUMAN GENETICS

Paper Code							
BCH	4	0	0	3	(B)		

Questions Booklet Series

D

[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।
- 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)

- Chromosome walking: 1. (A) is used in FISH is important component of genetic (B) mapping requires a genomic DNA library (C) (D) occurs in mitosis Holocentric condition of chromosomes is 2. commonly present in: (A) Bugs (B) Insects (C) **Plants** All vertebrates (D)
 - 3. Which of the following statements is not true?
 - (A) Homologous recombination can be used to disrupt genes
 - (B) Transposons can be directed to disrupt specific genes
 - (C) Transcriptomes can be characterized by Serial Analysis of Gene Expression (SAGE)
 - (D) Open reading frames are only found in protein-coding genes

- 4. Limnaea shell coiling is due to:
 - (A) Maternal inheritance
 - (B) Cytoplasmic inheritance
 - (C) Extranuclear inheritance
 - (D) All of the above
- 5. Microarrays are:
 - (A) used for analysis of transcriptomes
 - (B) made up of only glass and silica
 - (C) very much smaller than DNA chip
 - (D) can often be detected by histochemical assays
- 6. A codon bias:
 - (A) is not found in prokaryotes
 - (B) is found in genome mapping
 - (C) is found in functional RNA
 - (D) is used to identify genes
- 7. Mark the correct statement:
 - (I) Holocentric chromosomes have hole in the centre
 - (II) Histonic proteins initiate RNA transcription

- (III) Chiasmata are the results, not a cause of crossing over
- (IV) Females can act as carrier of sexlinked traits.

Codes:

- (A) I, IV
- (B) II, III
- (C) III, IV
- (D) II, IV
- 8. Fluorescent *in-situ* hybridization (FISH):
 - (A) requires a DNA polymerase
 - (B) requires a labelled probe
 - (C) can be used in physical mapping of the genome
 - (D) Both (B) and (C)
- 9. Which of the following is a co-dominant marker?
 - (A) RAPD
 - (B) RFLP
 - (C) AFLP
 - (D) RFLP and RAPD

- Mapping technique used to determine the position of restriction sites in a DNA molecule is
 - (A) DNA markers
 - (B) Biochemical markers mapping
 - (C) Restriction mapping
 - (D) Genetic map
- 11. Two linked genes a and b show 20% recombination. The individuals of a dihybrid cross between ++/++ × ab/ab show gametes:
 - (A) ++80: ab 20
 - (B) ++50: ab 50
 - (C) ++40 ab: 40: +a10: +b10
 - (D) ++30: ab 30: +a20: +b20
- 12. Genetic variation can be introduced into bacteria by all of the following methods except:
 - (A) mutation
 - (B) DNA amplification
 - (C) transformation
 - (D) transduction

13.	Somatic hybridization is achieved	16.	The term 'allele' was proposed by:				
	through		(A) Morgan				
			(B) Bateson				
	(A) Grafting		(C) Johannsen				
	(B) Protoplast fusion		(D) Mendel				
	(C) Conjugation	17.	A cross between F1 generation and one of				
	(D) Recombinant DNA technology		the recessive parents is:				
14.	The cloning that has provoked the most		(B) Reciprocal cross				
	public consternation and media		(C) Monohybrid cross				
			(D) Test cross				
	attention is:	18.	A haploid set of all the genes present in a				
	(A) Research cloning		gamete is called :				
	(B) Animal/plant cloning		(A) Genotype				
	(C) Intervention cloning		(B) Phenotype				
			(C) Genome				
	(D) Reproductive cloning		(D) Linkage				
15.	Which one of the following is a	19.	Which is the non-Mendelian cross when				
	mutagen ?		both phenotypic and genotypic ratio				
	(4)		obtained is same ?				
	(A) Oxygen		(A) Incomplete dominance				
	(B) Acetic acid		(B) Multiple alleles				
	(C) Mustard gas						
	(D) Pagtin		(C) Polygeny				
	(D) Pectin		(D) Pleotropism				

(5)

Set-D

20.	The concept of phenotype and genotype	24.	Intragenic non-Mendelian inheritance		
	was given by:		includes:		
	(A) Bateson		(A) Dominance, codominance,		
	(B) Morgan		incomplete dominance		
	(C) Johannsen		(B) Codominance, multiple alleles,		
	(D) Punnet		incomplete dominance		
21.	Gamete normally contains:		(C) Epistasis, multiple alleles,		
	(A) Many alleles of a gene		dominance		
	(B) Two alleles of a gene		(D) Polygene, pleotropism,		
	(C) All alleles of a gene		codominance		
	(D) One alleles of a gene	25.	A pair of white sheep are mated and the		
22.	Inhibiting genes which suppress the other		offspring is black. What is the probability		
	genes are called:		for next generation being black? (White		
	(A) Epistatic genes		is dominant to black).		
	(B) Hypostatic genes		(A) 25%		
	(C) Recessive genes		(B) 50%		
	(D) Complementary genes		(C) 75%		
23.	Gene pool is the sum total of genes		(D) 0%		
	present in:	26.	The ultimate source of variation is:		
	(A) Cell		(A) Mitosis		
	(B) Organism		(B) Mutation		
	(C) Population		(C) Fertilization		
	(D) Ecosystem		(D) Meiosis		

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Set-D

27.	Marriages between close relatives are	31.	Haer	nophilia is more common in males			
	avoided because. It induces more:		beca	use it is:			
	(A) Blood group abnormalities		(A)	Recessive character carried by			
	(B) Multiple births			X-chromosome			
	(C) Mutations						
	(D) Recessive alleles to come together		(B)	Dominant character carried by			
28.	Human skin colour is controlled by three			Y-chromosome			
	separate genes. What will be the genotype		(C)	Dominant trait carried by			
	of intermediate (Mulatto) skin colour			X-chromosome			
	organism?		(D)	Recessive trait carried by			
	(A) AABbCc			Y-chromosome			
	(B) AaBbCc						
	(C) AABBCC	32.	Muta	ations are induced mostly by:			
	(D) aabbcc		(A)	UV radiations			
29.	Informosome is:		(B)	Beta rays			
	(A) mRNA + protein		(C)	Alpha rays			
	(B) rDNA + Histone		, ,				
	(C) DNA + Histone		(D)	Gamma rays			
	(D) RNA + DNA	33.	Meta	acentric chromosomes have :			
30.	Which is a sex-linked trait?		(A)	Unequal arms			
	(A) Colourblindness, haemophilia		(B)	Equal arms			
	(B) Nightblindness, albinism		` /	•			
	(C) Myxoedema, beri-beri		(C)	Only one arm			
	(D) Deafness tylosis		(D)	Elongated arm			
	•						

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Set-D

- 34. Balbiani rings are structural feature of:
 - (A) Lampbrush chromosomes
 - (B) Potytene chromosomes
 - (C) Allosomes
 - (D) Autosomes
- 35. The small part of chromosomes arm beyond secondary constriction is called:
 - (A) Centromere
 - (B) Satellite
 - (C) Chromonemata
 - (D) Kinetochore
- 36. The prevention or treatment of diseases through methods such as genetic testing, abortion of defective embryos, and germline therapy is known as:
 - (A) Positive eugenics
 - (B) Negative eugenics
 - (C) Reverse genetics
 - (D) Pro-eugenics

- 37. Which of the following women is the legal mother of a child?
 - (A) A woman who commissions a surrogate where the surrogate uses that woman's egg
 - (B) A woman who commissions a surrogate where the surrogate uses her own egg
 - (C) A surrogate mother
 - (D) None of the above
- 38. Research ethics committees are:
 - (A) Committees of scientists and researchers.
 - (B) Convened by organisations to monitor the ethical standards of research projects carried out under their auspices, under their name.
 - (C) Committees of researchers concerned with ethics.
 - (D) Concerned only with research conducted in the medical and paramedical sciences.

BCH–4003(B) (8) Set-D

39.	Holai	ndric	genes	are	present	42.	The	functional unit of DNA undergoing
	on:						muta	ation is :
	(A)	Salivary	gland chro	omosome	es		(A)	Cistron
	(B)	X-chrom	nosomes				(B)	Muton
	(C)	Y-chrom	nosomes				(C)	Recon
	(D)	Lampbru	ısh chromo	osome			(D)	Genophore
						43.	A bo	by with normal brother and colour-
40.	Mito	chondria 1	myopathy	example	:		blind	I sister has his parents:
	(A)	Leigh sy	ndrome				(A)	Father normal, mother colourblind
	(B)	Mitocho	ndrial [NA o	lepletion		(B)	Both normal
		syndrom	e				(C)	Both colourblind
	(C)	Mitochondrial encephalon	ephalom	yopathy,	44.	(D)	Father colourblind, mother normal	
	` ,		1 7 1	roke-like		Dros	ophila having both male and female	
		episodes	,				trait	is:
		episodes					(A)	Homomorphic
	(D)	All of the	e above				(B)	Homozygous
41.	Diffe	rential e	xpression	of the	genetic		(C)	Gynandromorph
	mater	rial depe	nding on	its pare	ntage of		(D)	Hemizygous
	inher	itance giv	es			45.	Extra	anuclear chromosomes in eukaryotes
	(A)	Penetran	ce				are p	present in :
	(B)	Expressi	vity				(A)	All organelles
	` /	-	•				(B)	Membraneless organelles
	(C)	Imprintii	ng				(C)	Semiautonomous organelles
	(D)	Non-pen	etrance				(D)	Plasmids

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Set-D

46.	Enzy	mes used to regulate de novo	49.	The	first complete, gapless sequence of a		
	meth	ylation :		human genome came through:			
	(A)	DNMT1		(A)	T2F consortium		
	(A)	DINIVITI		(B)	T3T consortium		
	(B)	DNMT3a		(C)	T2T consortium		
	(C)	DNMT3b		(D)	T2Q consortium		
	(D)	Both (B) and (C)	50.	The	chemical composition of		
47.	A me	A man receives X-chromosome from his:			chromosome is:		
47.	AIII	un receives A-cinomosome from ms .		(A)	DNA, histonic protein, non-histonic		
	(A)	Mother			protein, ions		
	(B)	Father		(B)	RNA, histonic protein, ions		
(((C)	Partially from mother and partially		(C)	DNA, lipids, carbohydrates		
	(-)			(D)	DNA, RNA, pectin		
		from father Either from mother or from	51.	Whic	ch of these is dominant genetically?		
	(D)			(A)	Haemophilia		
		father		(B)	Colourblindness		
10	Chas			(C)	Albinism		
48.	Cnro	mosome cohesion:		(D)	Polydactyly		
	(A)	established during S phase	50	T 7 1			
	(B)	dissolved in metaphase	52.	Y-chromosomes which determine male			
	(C)	cohesion close to the centromere		sex c	sex of the individual are:		
	(C)			(A)	Androgen		
	facilitates bi-orientation of			(B)	Androsome		
		chromosomes		(C)	Gynosperm		
	(D)	All of the above		(D)	Androecium		

(10)

Set-D

53.	Which of th	e following is not a site on	57.	Exch	nange of segment between non-sister
	internet for a	lignment of sequence pairs ?		chro	matids of homologous chromosome
	(A) BLAS	ΓN		is:	
	(B) BCM S	Search Launcher		(A)	Non-disjunction
	(C) SIM			(B)	Crossing over
	(D) BLAS	ΓX		(C)	Translocation
54.	Expression o	f genes can be analyzed by:	~ 0	(D)	Transition
	(A) Souther	rn analysis	58.		extrachromosomal, circular DNA ng genes for sexuality are:
	(B) Northe	rn analysis		(A)	Plasmids
	(C) RNA in	nterference techniques		(B)	Plastids
	(D) Compa	rative genomics		(C)	Nucleoid
55.	Opponents of	f gene therapy insist that:		(D)	Mesosomes
	(A) Germ-l	ine therapy is permissible	59.	Prote	ein coding genes can be identified by:
	(B) Gene th	nerapy is harmless		(A)	ORF scanning
	(C) Reprod	uctive freedom has limits		(B)	Transposon scanning
	(D) Reprod	uctive freedom is a personal		(C)	Zoo-tan blotting
	right			(D)	Nuclease S22 mapping
56.	Heterochrom	atin is :	60.	All s	ex-linked traits always show:
	(A) Darkly	Stained band		(A)	Linkage
	(B) Genetic	cally inactive DNA		(B)	Crossing over
	(C) Replica	ating late in the S phase		(C)	Criss-cross inheritance
		he above		(D)	Dominance

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Set-D

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।
- ओ. एम. आर. उत्तर-पत्रक (OMR Answer Sheet) पर कुछ भी लिखने से पूर्व उसमें दिये गये सभी अनुदेशों को सावधानीपूर्वक पढ़ 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।