Reg. No		

B.Tech/M.Tech(Integrated) DEGREE EXAMINATION, NOVEMBER 2023

Third Semester

21BTC206T - GENETICS AND CYTOGENETICS

(For the candidates admitted during the academic year 2022-2023 onwards)

Note:

i. Part - A should be answered in OMR sheet within first 40 minutes and OMR sheet should be handed over to hall invigilator at the end of 40th minute.

ii Part - B and Part - C should be answered in answer booklet.

ii. Pa	rt - B and Part - C should be answered in an	swer booklet.			
Time	: 3 Hours		Max.	Mark	s: 75
	PART - A $(20 \times 1 = 2)$ Answer all Ques	•	Mark	as BL	CO
1.	When a disease is more predominant in a fais said to be (A) X-linked (C) autosomal dominant	amily irrespective of males and females it (B) Y-linked (D) autosomal recessive	1	2	, quan
2.	What are the possible offspring from blood (A) A, B and AB (C) O and AB	type A mother and AB father (B) B and AB (D) A and O	1	1	1
3.	If a daughter of a colour blind person marri-	es a normal man then their progenies will	1	4	1
	(A) Half of their daughters are colour blind(C) All the sons are colour blind	(B) Half of their sons are colour blind(D) All the daughters are colour blind			
4.	Test cross of a plant with yellow round seed yellow wrinkled: green round: green wringenotype of the yellow round parent? (A) YYRR (C) YYRr		1	2	1
5.	Genes a and b are 10 map units apart. An a^+b/a^+b individual. What gametes can the (A) $45a^+b$: $5ab$: $5a^+b^+$: $45ab^+$ (C) $40a^+b$: $10ab$: $10a^+b^+$: $40ab^+$	ab^+/ab^+ individual was mated with an e F ₁ produce and in what ratio? (B) $45a^+b^+$: $5a^+b$: $5ab^+$: $45ab$ (D) $40a^+b^+$: $10a^+b$: $10ab^+$: $40ab$	1	2	2
6.	A heterozygous gray-bodied (b^+) , normal with a recessive male gave the follow vestigial-10, black normal-10, and black vetthe genes b and vg ? (A) 10 map units (C) 5 map units	ving progenies: Gray normal-90, gray	1	2	2
7.	The genetic markers present in homologindividual is $X+Z/+Y+$. What is the arraindividual?	angement of the genes X and Y in the	1	2	2
	(A) cis (C) coupling	(B) trans (D) different chromosomes			
8.	In somatic cell hybridization, HAT selection (A) Parent cells (C) synkaryon	n procedure is used to eliminate growth of (B) hybrid cells (D) heterokaryon	1	1	2

9.	he arrangement of genes in a chromosome is $\underline{A} \underline{B} \underline{C} \underline{D} \underline{F} \underline{\bullet} \underline{F} \underline{G}$, where $\underline{\bullet}$ represents e centromere. What type of chromosome mutation is required to change the rangement in this chromosome to $\underline{A} \underline{E} \underline{D} \underline{C} \underline{B} \underline{\bullet} \underline{F} \underline{G}$?			2	3
	(A) Deletion (C) Translocation	(B) Inversion (D) Duplication			
10.	Some people with Turner syndrome are 45, (A) Duplication of chromosomes (C) Translocation		1	2	3
11.	During FISH analysis if a person exhibit specific probe, then the person is said to be (A) monosomic (C) trisomic	its two signals on hybridization with a (B) normal (D) tetrasomic	1	1	3
12.	Exchange of chromatid segments between as (A) Deletion (C) inversion	non-homologous chromosomes is called (B) duplication (D) reciprocal translocation	1	1	3
13.	A random segment of the donar DNA is tra: (A) Generalized transducing phage (C) Specialized transducing phage	•	1	2	4
14.	Integrated state of the F plasmid that increase (A) prophage (C) lysogen	ses the frequency of recombination is (B) Hfr (D) extrachromosomal DNA	1	1	4
15.	During transformation, mapping of bacter frequency of (A) Co-transformation (C) double transformation	(B) co-transduction(D) lysogen	1	2	4
16.	Recombination of the donor DNA with a during (A) Single cross over (C) Three cross over	the recipient genome in bacteria occurs (B) Double cross over (D) Five cross over	1	2	4
17.	If the frequency of the L_M allele is 0.3 in what is the frequency of the heterozygous c. (A) 0.42 (C) 0.7	a population, assuming random mating arriers? (B) 0.5 (D) 0.23	1	2	5
18.	The incidence of recessive albinism is 0.0 frequency of the recessive allele? (A) 0.2 (C) 0.4	(B) 0.02 (D) 0.04	1	2	5
19.	The frequency of M, N and MN phenotypes What is the frequency of L_M allele? (A) 0.78 (C) 0.39	(B) 0.22 (D) 0.61	1	4	5
20.	A certain gene has three alleles A_1 , A_2 and 0.2 respectively. What is the frequency of A (A) 0.16 (C) 0.04	A ₃ . The frequency of A ₁ , and A ₂ is 0.4 and A ₂ A ₃ hetrozygotes? (B) 0.32 (D) 0.14	1	2	5
	PART - B ($5 \times 8 = 40$) Answer all Ques		Marks	s BL	CO

	Answer any 1 Questions			
	 (b) How does random genetic drift contribute towards change in allele frequency in a population? Explain its effects. PART - C (1 × 15 = 15 Marks) 	Marks	BL	СО
25.	 (a) A locus has three alleles A₁, A₂, and A₃ with frequencies of 0.6 for A₁ and 0.3 for A₂. Assuming random mating, what is the expected frequency of all the heterozygotes and homozygotes in the population? (OR) 	8	3	5
	(b) How does a lytic phage help in gene transfer in bacteria? Explain its use in gene mapping.			
	(OR)			
	⁵ -H-4min-A-1min-L-3min-J->	**		
	4-A-1min-L-3min-J-9min-D->			
	3 -K-2min-D-4min-Y-8min-L->	,		
	1 -A-4min-H-2min-E-5min- R-> 2 -D-2min- K-4min-S-3min-R->	. P		
	the makers in the F+ strain.			
	interrupted mating experiments. From the following data, for several Hfr strains derived from the same F+, determine the order and distance between			
24.	(a) An F+ strain marked at 10 loci give rise spontaneously to Hfr progeny. For any Hfr strain the order of markers entering early can be determined by	8	4	4
	(b) How would you classify mutation that leads to alteration in the arrangement of genes in the chromosomes? Explain.			
	(OR)			
23.	(a) Explain the technique that can be used to detect aneuploids in humans in the early embryonic stage?	8	3	3
	(b) How would you map genes in humans to its specific linkage group using selection procedure?			
	(OR)			
22.	(a) Discuss the disadvantages of the two factor cross used in preparation of linkage maps.	8	3	2
	ii). Determine the genotypes of the family members. List the genotypes of each individual in the pedigree.			
	and they have twin daughter. i). Create a pedigree for the family.			
	(2 boys and a girl). The carrier daughter has one son with hemophilia who dies. One of the non-carrier daughters son marries a women who is a carrier			
	a carrier. Both daughters marry men without hemophilia and have 3 children			
	(b) A man and woman marry. They have five children, 2 girls and 3 boys. The mother is a carrier of hemophilia, an X-linked disorder. She passes the gene on to two of the boys who died in childhood and one of the daughters is also			
	(OR)			
	ii) What is the expected ratio of offspring from a recessive backcross with a heterozygous parent (CcPp)?			
	i) What is the expected ratio of the progenies resulting from a cross between cCpp X cCPP?			
	produce white flowers, with the purple color resulting from the presence of both factors.			
21.	(a) Given the genotypes of the parents, what will be the flower colour of the offspring of the following crosses? In sweet peas, genes C or P alone	8	4	1
21		0		

26. A cross was made between scute or loss of certain thoracic bristles (sc), echinu or roughened eye surface (ec), vestigial wing (vw) female (sc ec vw/sc ec vw) flies and wild type males. The F₁ females were test crossed. The progeny are listed as gametic genotypes derived from the heterozygous females

15 4 2

Gametic genotypes	Number
+++	169
sc + +	9
+ + vw	1
sc ec vw	172
+ ec vw	10
sc + vw	18
sc ec +	0
+ ec +	21

Are these genes linked? What is the correct order of the 3 genes on the chromosome? What is the genetic map distance between these three genes? Calculate the coefficient of interference?

27. How is a meroztgote used in mapping of closely related genetic markers in Bacteria? 15 4
