

**2016**

**M.Sc. 1st Semester Examination**

**ZOOLOGY**

**PAPER—ZOO-104**

**Full Marks : 40**

**Time : 2 Hours**

*The figures in the margin indicate full marks.*

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

*Illustrate the answers wherever necessary.*

**Use separate Answer-scripts for Group-A & Group-B**

**Group—A**

*(Cytogenetics)*

**1. Answer any two questions of the following : 2×2**

**(a) In E. coli, four Hfr strains donate the markers shown in the order given :**

Strain 1	M	Z	X	W	C
Strain 2	L	A	N	C	W
Strain 3	A	L	B	R	U
Strain 4	Z	M	U	R	B

*(Turn Over)*

All of these Hfr strains are derived from the same F<sup>+</sup> strain. What is the order of these markers on the circular chromosome of the original F<sup>+</sup>.

- (b) About 70% of all white North American can taste the chemical phenylthiocarbamide and the remainder can not. The ability to taste is determined by the dominant allele T and the inability to taste is determined by the recessive allele t. If the population is assumed to be in Hardy-Weinberg Equilibrium, what are the genotypic and allelic frequencies in this population ?
- (c) In what sense is pRB a negative regulator of E2F transcription factor ?
- (d) During the cell cycle, the p16 protein is an inhibitor of cyclin/CDK activity. Predict the phenotype of cells homozygous for a loss of function mutation in the gene that encode p16. Would this gene be classified as a protooncogene or a tumor suppressor gene ?

2. Answer any *two* of the following :

4×2

- (a) In a HW equilibrium population, out of 100 people 17 have A type blood group, 17 have B type, 2 have AB type and 64 have O type. Calculate the allelic frequencies.
- (b) The data in the following table obtained from a 3 point transduction test. A gene encode tryptophan synthetase, anth is linked unselected marker. What is the linear order of anth and three mutant alleles of A gene in the table ?

4

Cross	Donor markers	Recipient markers	anth allele in Recombinant	% anth <sup>+</sup>
1.	anth <sup>+</sup> - A34	anth <sup>-</sup> - A223	72anth <sup>+</sup> : 332 anth <sup>-</sup>	18%
2.	anth <sup>+</sup> - A46	anth <sup>-</sup> - A223	196anth <sup>+</sup> : 180 anth <sup>-</sup>	52%
3.	anth <sup>+</sup> - A223	anth <sup>-</sup> - A34	380anth <sup>+</sup> : 379 anth <sup>-</sup>	50%
4.	anth <sup>+</sup> - A223	anth <sup>-</sup> - A46	60anth <sup>+</sup> : 280 anth <sup>-</sup>	20%

- (c) Explain the role of p53 in the cellular response to DNA damage.
- (d) All pur alleles result in defective enzyme P and map at one genetic locus. A complementation test among six mutant pur strains produce the following results where + indicate complementation and - indicate no complementation.

	1	2	3	4	5	6
1	-	-	-	-	+	-
2	-	-	-	-	+	+
3	-	-	-	-	-	-
4	-	-	-	-	-	+
5	+	+	-	-	-	+
6	-	+	-	+	+	-

Draw a complementation map and comment what kind of mutant might mutant 3?

3. Answer any *one* of the following :

8×1

- (a) You cross a number of *rII* deletion mutations in all possible combinations in *E. coli* B and plate them on *E. coli* k 12( $\lambda$ ) to determine whether  $r^+$  recombinants are formed. The formation of  $r^+$  recombinants indicates that the mutations can recombine and so, if they are deletions, they must be non-overlapping. The results are given in the accompanying table, in which a through f indicate an *rII* mutation and + indicates the formation of  $r^+$  recombinant progeny in the cross.

Assemble a deletion map for these mutations using a line to indicate the DNA segment that is deleted in each mutant.

	a	b	c	d	e	f
a	-	-	-	-	-	-
b		-	-	+	+	-
c			-	+	-	+
d				-	-	+
e					-	+
f						-

- (b) A cross is made between  $Hfr\ arg^+ bio^+ leu^+ X F^- art^- bio^- leu^-$ . Interrupted mating studies show that  $arg^+$  enters the recipient last, so that  $arg^+$  recombinants are selected

on a medium containing bio and leu only. These recombinants are tested for the presence of bio<sup>+</sup> and leu<sup>+</sup>. The following number of individuals are found for each genotype :

arg <sup>+</sup> bio <sup>+</sup> leu <sup>+</sup>	320
arg <sup>+</sup> bio <sup>+</sup> leu <sup>-</sup>	8
arg <sup>+</sup> bio <sup>-</sup> leu <sup>+</sup>	0
arg <sup>+</sup> bio <sup>-</sup> leu <sup>-</sup>	48

- (a) What is the gene order ?  
 (b) What are the map distances in recombination units ?
- 5+3

---

**Group—B**

( Immunology )

4. Answer any *two* of the following : 2×2
- (a) Write the principle and application of Immunohistochemistry
- (b) Mention the functional significance of the following :
- (i) Psoriasin,
  - (ii) Thymosin,
  - (iii) Secondary lymphoid organ,
  - (iv) IgA.
- (c) Define Agretope with diagram.
- (d) Distinguish between Central tolerance and Peripheral tolerance.

5. Answer any *two* questions of the following : 4×2
- (a) Distinguish between Necrosis and Apoptosis with proper diagram.
  - (b) Write the properties of T-cell epitope.
  - (c) What is Adjuvant ? Give exdample. Add a note on ADCC. 1+3
  - (d) State the characteristics of peptides that are presented through MHC class I molecule.
6. Answer *one* question of the following : 8×1
- (a) What is Hybridization Probe ? Write the principle of Southern Blotting Hybridization. How it differs from Western Blotting Hybridization ? 1+1+6
  - (b) (i) Describe the structure of class II MHC molecule with suitable diagram.
  - (ii) Illustrate the cytosolic pathway for processing of antigen. 3+5
-