1/H-77 (i) (Syllabus-2015)

(2)

2022

(February)

BIOTECHNOLOGY

(Honours)

(Cell Biology and Genetics)

Marks : 56

Time: 3 hours

The figures in the margin indicate full marks for the questions

Answer Question No. **1** which is compulsory and *any* **four** from the rest

- **1.** Differentiate between the following: $3\times4=12$
 - (a) Primary and secondary lysosomes
 - (b) Membrane-bound ribosomes and free ribosomes
 - (c) Heteroplasmy and homoplasmy
 - (d) Maternal effect and maternal inheritance

| 2. | (a) | Give a comparison of the characteristics | | | |
|----|-----|------------------------------------------|-------------|-----|------------|
| | | between | prokaryotes | and | eukaryotes |
| | | (with diagrams). | | | |

6

(b) What is cell theory? There are exceptions to the cell theory.' Explain.

2+3=5

6

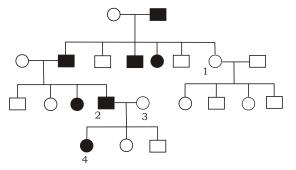
- **3.** (a) Explain genic-balance theory of sex determination.
 - (b) Write a note on cell cycle checkpoints. 5
- **4.** (a) What are centrioles? Explain their roles in a cell. 1+3=4
 - (b) Explain with diagrams the 'cis' face and 'trans' face of the Golgi apparatus. 4
 - (c) Differentiate between mitochondrial DNA and nuclear DNA.
- **5.** What is the role of the ClB balancer chromosome in the detection of recessive mutations on the X-chromosome in *Drosophila*? Compare and contrast between Muller's ClB method and Muller-5 method.

3+8=11

(4)

- **6.** (a) What is pseudodominance? How is it produced? 1+1=2
 - (b) Differentiate between the following: 3×3=9
 - (i) Aneuploidy and euploidy
 - (ii) Paracentric inversion and pericentric inversion
 - (iii) Silent mutation and neutral mutation
- **7.** Write short notes on the following : $5\frac{1}{2}+5\frac{1}{2}=11$
 - (a) Polytene chromosome
 - (b) Dosage compensation
- **8.** (a) If the recombinant frequencies between the gene loci mn is 38%, mo is 6% and no is 28%, find the order of the genes.
 - (b) Write on multifactorial inheritance giving a suitable example.

(c) In the following human pedigree, the filled symbols represent the affected individuals. You may assume that the disease allele is rare and therefore individuals marrying into the family are unlikely to have defective allele:



- (i) What is the most likely mode of inheritance for this pedigree?
- (ii) State the genotypes of individuals numbered 1–4 using the letter *A*. Use the uppercase letter to represent the dominant allele and lowercase letter to represent the recessive allele.

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