Total number of printed pages-7

3 (Sem-3/CBCS) BOT HC 3

2022

BOTANY

(Honours)

Paper : BOT-HC-3036

(Genetics)

'Full Marks : 60

Time : Three hours

The figures in the margin indicate full marks for the questions.

- 1. Answer **any seven** of the following questions: 1×7=7
 - (i) The transmission of characters or traits from one generation to another is called ______. (Fill in the blank)
 - (ii) The genotypic ratio of law of independent assortment is—
 - (a) 1:1:1:1
 - (b) 9:3:3:1
- (c) 9:3:3:3
 - (d) 3:9:1:3

(Choose the correct answer)

Contd.



(iii) When the phenotypic expression of a heterozygote is more extreme than that of either homozygous parent, then it is-

> Co-dominance (a)

Dominance (b)

- Overdominance (c)
- Incomplete dominance (d)
 - (Choose the correct answer)
- (iv) Genes which have little or no effect of their own but increase or decrease the expression of other major genes are known as-
 - (a) Pleotropic genes
 - (b) Modifying genes
 - (c) Over dominant genes
 - (d) Epistasis (Choose the correct answer)
- (v) Coupling and repulsion phases are two aspects of the same phenomenon called

(Fill in the blank)

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Femaleness (c) (d) Maleness (Choose the correct answer) (vii) Y-linked genes are called _ (Fill in the blank) _ is the change in frequency of (viii) _ an existing gene variant in the population due to random chance. (Fill in the blank)

(vi) Autosomes are concerned with-

(a) Sex determination

(b) Body characters

- (ix) The loss of a segment of genetic (material from a chromosome is termed as-
 - (a) Duplication
 - (b) Deficiency
 - Translocation (c)
 - (d) Inversion

(Choose the correct answer)

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(x) _____ is the smallest unit of DNA capable of recombination.

(Fill in the blank)

(xi) _____ refers to the number of processes by which a cell identifies corrects damage to the DNA molecules that encode its genome.

(Fill in the blank)

- (xii) Nullisomic is represented by-
 - (a) 2n-2
 - (b) 2n+1+1
 - (c) 2n+1
 - (d) 2n+2
 - (Choose the correct answer)
- 2. Answer **any four** out of the following questions : 2×4=8
 - (i) What are tetrasomics?
 - (ii) What are sex chromosomes?
 - (iii) What are exons?

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- (iv) What is the difference between complete and incomplete linkage?
- (v) What is pseudo-dominance?
- (vi) What is the purpose of a pedigree analysis?
- (vii) What is genetic variation?
- (viii) Differentiate between Mendelian and non-Mendelian inheritance.
- 3. Write short notes on **any three** of the following : 5×3=15
 - (i) Frameshift mutation
 - (ii) Epistasis
 - (iii) Mitochondrial DNA
 - (iv) Intercalating Agents
 - (v) Transposons
 - (vi) Speciation
 - (vii) Spontaneous mutation
 - (viii) Multiple Alleles

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4. Answer **any three** of the following questions: 10×3=30

(i) What are Mendel's law? Describe Mendel's second law with a suitable example. 2+8=10

- (ii) What is crossing over? Describe the cytological basis of crossing over with a suitable example. 2+8=10
- (iii) Explain with the help of diagram meiotic behaviour of paracentric and pericentric inversion. 5+5=10
- (iv) What do you mean by extra chromosomal inheritance? Describe with an example. 2+8=10
- (v) Define aneuploids. Discuss the causes of origin of aneuploids. 2+8=10
- (vi) With the help of suitable example discuss polygenic inheritance.

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(vii) What is induced mutation? Give a detailed account of physical mutagens. 2+8=10

(viii) Describe Hardy-Weinberg's law.

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