M.Sc. 3rd Semester Examination BIOCHEMISTRY

PAPER -BIC-303

Full Marks: 40

Time: 2 hours

Answer all questions

The figures in the right-hand margin in Candidates are required to give their ar own words as far as practica

Illustrate the answers wherever no

GROUP-A

- 1. Answer any five from the following
 - (a) An individual with 47 chroincluding an additional chromos said to be (i) triplet (ii) trisomic (iv) tricycle.

- (b) What are the main objective of the study of 'Metabolomics'?
- (c) What are the different types of satellite DNA? What are their utility?
- (d) What is linkage mapping? Mention the principle of this procedure with its utility.
- (e) What different levels the translational regulations may be controlled?
- (f) Pattern 'baldness is more frequent in male and is assumed to result from (i) Y-linkage of trait, (ii) X-linked recessive mode of inheritance, (iii) sex-influenced autosomal inheritance.
- (g) What could be utility p53 knock out mice in research?
- (h) What is an 'anti-seme' RNA? Mention its working principle?

GROUP-B

Answer any two from the followi

- 2. Define 'polymorphism'. What are the sof SNPs? Explain with suitable exa
- 3. Explain with suitable diagram the mination procedure in *Drosophila Me*
- 4. Discuss the similarities and different miRNAs and siRNAs in terms of
 - (i) location and structure of genes them,
 - (ii) their function in post transcrip silencing.
- 5. What are the importance of the so ca part of the DNA sequences? How do evolution? Define natural selection

GROUP-C

Answer any two from the follow

6. (a) What are the reporter and ma

used in gene targetting procedure? Give Example.

- (b) With suitable diagram briefly describe homologous gene targetting procedures. 2 + 8
- 7. Briefly describe different gene transfer procedure in plant, bacteria and eukaryotive cells?
- 8. (a) What are nuclear orphan receptors? How do they function? What are the use of stem cells in RDT?
 - (b) Briefly define functional genomics.
 - (c) Briefly define 'epistasis' and heterosis? $(4+2)+1\frac{1}{2}+1\frac{1}{2}+1$
- 9. Both fragile X-syndrome and Huntington disease are caused by trinucleotide repeat expansion. Individuals with fragile X syndrome have at least 200 CCG repeats at the 5' end of FRM-1 gene. In contrast, individuals with Huntington disease have 36 or more in frame CAG repeats within protein-coding region of the Huntington gene.
 - (a) Do you expect gene expression at the two

genes to be affected in the same repeat expansion? Explain you

(b) Based on your answer why many syndrome be recessive, wherea disease to dominant.