

B.Sc. (Honours) Examination, 2019
Semester-VI
Botany
Course: BBC-62 (Core)
(Molecular Biology & Genetic Engineering)

Time: Three Hours

Full marks:40

Questions are of value as indicated in the margin

1. Answer **any eight** of the following: 1×8=8
- i) Name one enzyme used besides Taq DNA polymerase in PCR.
 - ii) What is meant by plasmid incompatibility?
 - iii) State the function of 5'-splice site in pre-mRNA splicing.
 - iv) State the function of SI nuclease.
 - v) What is SNP?
 - vi) What is a GenBank?
 - vii) Name two rep-initiator proteins required for plasmid replication.
 - viii) Which subunit of RNA polymerase contains catalytic activity?
 - ix) State the specific function performed by reverse transcriptase.
 - x) What is the recognition sequence for *Bam*HI?
 - xi) Point out the function of UvrB in nucleotide excision repair of DNA in *E. coli*
 - xii) What is wobble hypothesis?

Group-A (Molecular Biology)

Answer **any two** questions

- 2. What is a gene promoter? Why is sequence within the promoter highly conserved? What are the three major functions of sigma factor? 2+3+3=8
- 3. Why is it necessary to sequence the entire human genome? What is its size? What percentage of human genome is identical between any two individuals? 6+1+1=8
- 4. What is meant by inducible operon? Cite one example in which induction of promoter is cAMP dependent. What are the common strategies to achieve constitutive expression of β-galactosidase within the *lac* operon in *E. coli*? 2+1+5=8

Group-B (Genetic Engineering)

Answer **any two** questions

- 5. What is the difference between reporter and marker gene? Describe the significance of marker gene in cloning a gene. 4+4=8
 - 6. Distinguish between cDNA and genomic DNA library. Why do you clone cDNA to express a eukaryotic protein in bacteria? Mention two goals of constructing a cDNA library? 3+3+2=8
 - 7. Define cloning. Briefly explain four different strategies to confirm the presence of insert within the vector. 2+6=8
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