

J : BIOTECHNOLOGY

Q. 1 – Q. 7 carry one mark each.

- Q.1 The method used for prediction of three dimensional structure of a protein from known structure(s) of one or more related proteins is
- (A) Multiple sequence alignment
 - (B) Homology modeling
 - (C) Phylogeny
 - (D) Docking
- Q.2 To produce plants that are homozygous for all traits, the best choice is
- (A) Protoplast culture
 - (B) Cell suspension culture
 - (C) Anther and pollen culture
 - (D) Apical meristem culture
- Q.3 Restriction endonucleases from two different organisms that recognize the same DNA sequence for cleavage are called
- (A) Isoschizomers
 - (B) Isozymes
 - (C) Concatamers
 - (D) Palindromes
- Q.4 Caspases are involved in the process of
- (A) DNA replication
 - (B) Mutation and recombination
 - (C) Antibody synthesis
 - (D) Apoptosis
- Q.5 Baculovirus expression system is used to express heterologous genes in
- (A) Mammals
 - (B) Plants
 - (C) Insects
 - (D) Yeasts
- Q.6 A culture vessel in which physical, physicochemical and physiological conditions, as well as cell concentration, are kept constant is known as
- (A) Cell concentrator
 - (B) Biostat
 - (C) Batch bioreactor
 - (D) Incubator
- Q.7 Virus resistant transgenic plants can be developed by the expression of
- (A) Cowpea trypsin inhibitor
 - (B) Crystalline toxin protein
 - (C) Defective movement protein
 - (D) Snowdrop lectin

Q. 8 to Q.21 carry two marks each.

Q.8 Which of the following are commonly used as reporter genes ?

- P. NPT gene
- Q. Luciferase gene
- R. CFTR gene
- S. GFP gene

- (A) Q, S (B) R, S
(C) P, R (D) P, Q

Q.9 Which of the following statements are true about glyphosate tolerant transgenic plants ?

- P. Transgenic plants detoxify glyphosate.
- Q. Transgenic plants produce an altered enzyme that is not affected by glyphosate.
- R. Transgenic plants sequester glyphosate in vacuoles.
- S. Transgenic plants overcome the inhibition of aromatic amino acid biosynthesis.

- (A) P, Q (B) R, S
(C) Q, S (D) P, R

Q.10 Match the items in Group 1 with an appropriate description in Group 2 :

Group 1

- P. UPGMA
- Q. CLUSTAL
- R. SWISS-PROT
- S. RasMol

Group 2

1. Protein sequence database
2. Phylogenetic analysis
3. 3-D structure visualization
4. Multiple sequence alignment

- (A) P-4, Q-1, R-2, S-3 (B) P-2, Q-4, R-1, S-3
(C) P-2, Q-3, R-1, S-4 (D) P-2, Q-1, R-4, S-3

Q.11 Match the properties in Group 1 with the downstream operations in Group 2 :

Group 1

- P. Size
- Q. Density
- R. Volatility
- S. Solubility

Group 2

1. Extraction
2. Distillation
3. Filtration
4. Sedimentation

- (A) P-3, Q-4, R-2, S-1 (B) P-4, Q-1, R-2, S-3
(C) P-4, Q-3, R-1, S-2 (D) P-3, Q-2, R-4, S-1

Q.12 Match the items in Group 1 with their functions in Group 2 :

Group 1

- P. *rol* genes
- Q. Opines
- R. Virulence genes
- S. *Aux* and *cyt* genes

Group 2

1. Food and energy source
2. Tumor formation
3. Hairy root induction
4. T-DNA transfer and integration

- (A) P-4, Q-3, R-2, S-1 (B) P-3, Q-2, R-4, S-1
(C) P-1, Q-3, R-4, S-2 (D) P-3, Q-1, R-4, S-2

Q.13 Which of the following statements hold true for pluripotent stem cells (PSCs) under *in vitro* conditions ?

- P. PSCs can be maintained in an undifferentiated state.
- Q. PSCs exhibit abnormal and unstable karyotypes.
- R. PSCs can differentiate into a wide variety of cell types.
- S. PSCs cannot be passaged continuously.

- (A) P, Q
- (B) P, R
- (C) Q, R
- (D) Q, S

Q.14 Determine the correctness or otherwise of the following **Assertion (a)** and **Reason (r)** :

Assertion (a) : IPTG (Isopropylthiogalactoside) is a gratuitous inducer of *lac* operon.
Reason (r) : IPTG is an efficient inducer, but not a substrate of *lac* operon.

- (A) Both (a) and (r) are true and (r) is the correct reason for (a).
- (B) Both (a) and (r) are true but (r) is not the correct reason for (a).
- (C) (a) is true but (r) is false.
- (D) (a) is false but (r) is true.

Q.15 Which of the following statements are true about bioreactors ?

- P. Continuous bioreactors provide less degree of control and uniform product quality than batch bioreactors.
- Q. Batch bioreactors are ideally suited for reaction with substrate inhibition.
- R. Choice of a bioreactor is dictated by kinetic considerations.
- S. Fed batch bioreactors are also called semibatch bioreactors.

- (A) P, Q
- (B) Q, S
- (C) R, S
- (D) P, R

Q.16 Match the items in Group 1 with correct options in Group 2 :

Group 1

- P. DNA footprinting
- Q. Yeast two-hybrid system
- R. DNA fingerprinting
- S. SAGE

Group 2

1. Protein-protein interaction
2. VNTR
3. DNA binding protein
4. Transcriptome analysis

- (A) P-1, Q-2, R-4, S-3
- (B) P-3, Q-1, R-2, S-4
- (C) P-3, Q-4, R-1, S-2
- (D) P-4, Q-2, R-1, S-3

Q.17 Determine the correctness or otherwise of the following **Assertion (a)** and **Reason (r)** :

Assertion (a) : Bacterial growth is called synchronous when majority of the cells are in same stage of the bacterial cell cycle.

Reason (r) : Synchronous culture can be obtained by growing bacteria in an enriched medium

- (A) Both (a) and (r) are true and (r) is the correct reason for (a).
- (B) Both (a) and (r) are true but (r) is not the correct reason for (a).
- (C) (a) is true but (r) is false.
- (D) (a) is false but (r) is true.

Q.18 Match the products in Group 1 with their possible applications in Group 2 :

Group 1

- P. Erythropoietin
- Q. Anti-fibrin 99
- R. Collagenase
- S. Transferrin

Group 2

- 1. Blood clot
- 2. Binding and transport of iron
- 3. Anaemia
- 4. Animal cell separation

- (A) P-3, Q-1, R-4, S-2
- (C) P-2, Q-3, R-1, S-4

- (B) P-3, Q-4, R-1, S-2
- (D) P-2, Q-1, R-4, S-3

Q.19 Match the products in Group 1 with their producer organisms in Group 2 :

Group 1

- P. Ethanol from glucose
- Q. Probiotics
- R. Citric acid
- S. Sauerkraut

Group 2

- 1. *Aspergillus niger*
- 2. *Leuconostoc mesenteroides*
- 3. *Saccharomyces cerevisiae*
- 4. *Bifidobacterium*

- (A) P-1, Q-3, R-2, S-4
- (C) P-3, Q-4, R-2, S-1

- (B) P-3, Q-4, R-1, S-2
- (D) P-1, Q-4, R-3, S-2

Q.20 A RNA polymerization assay was performed using ^3H UTP as the labelled nucleotide with a specific activity of $500 \mu\text{Ci} / \mu\text{mol}$ ($1 \mu\text{Ci} = 2.2 \times 10^9$ counts per min). After 10 min incubation, the trichloroacetic acid – insoluble radioactivity was found to be 692521 counts per min as determined in a liquid scintillation counter working at 60% efficiency for ^3H . The amount of UTP incorporated into the RNA will be

- (A) 15 nmol
- (B) 105 nmol
- (C) 150 nmol
- (D) 50 nmol

Q.21 One unit of glucoamylase enzyme activity is defined as the amount of enzyme required to produce $1 \mu\text{mol}$ of glucose per min in a 4% solution of Lintner starch at pH 4.5 and 60°C . If in a reaction mixture with 1 ml of the crude enzyme preparation containing 8 mg protein and 9 ml of 4.44% starch, $0.6 \mu\text{mol}$ of glucose/ml-min is produced, what will be the specific activity of the crude enzyme preparation ?

- (A) 1 unit/mg protein
- (B) 1.5 units/mg protein
- (C) 0.25 units/mg protein
- (D) 0.75 units/mg protein

END OF SECTION - J