# J: Biotechnology

# Q. 1 - Q. 6 carry one mark each.

Q.1	A thermostable DNA polymerase that can carry out both reverse transcription reaction and polymerization has been isolated from				
	(A) Thermococcus	s litoralis	(B) Thermus aquai	ticus	
	(C) Thermotoga m	aritima	(D) Thermus thern	nophilus	
Q.2	When present in tissue culture medium, gibberellin				
	<ul> <li>(A) helps to break dormancy of buds and bulbs</li> <li>(B) promotes dormancy development in buds and bulbs</li> <li>(C) is regarded as plant growth inhibitor</li> <li>(D) prevents normal recognition of auxin molecule</li> </ul>				
Q.3	To promote attachment and spreading of anchorage-dependent animal cells, the surface of the culture vessel needs to be coated with				
	(A) trypsin	(B) collagen	(C) pronase	(D) polyglycol	
Q.4	For amplification of GC rich sequences by polymerase chain reaction, identify the reagent that binds and stabilizes AT sequences and destabilizes GC regions.				
	(A) Tetramethyl ammonium chloride     (B) Betaine     (C) 7-deaza-2'-deoxyguanosine triphosphate				
	(D) Sodium dodecyl sulphate				
Q.5	Which of the following statements is INCORRECT about immobilized plant cell cultures?				
	(A) It is possible to use high cell densities				
	(B) Cells remain active for long periods				
	(C) Cell products or inhibitors can be removed easily				
	(D) It provides low shear resistance to cells				
Q.6	All the cells that participate in immune responses originate from a population of				
	(A) neutrophils	(B) stem cells	(C) macrophages	(D) lymphocytes	

# Q. 7 - Q. 24 carry two marks each.

growing in a medium containing  (A) thiamine, hypoxanthine, aminopterin (B) thymidine, histidine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, histidine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (C) uridine, hypoxanthine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (D) the forum 2.  Group 2 (B) P-2, Q-3, R-1, S-4 (D) P-3, Q-1, R-2, S-1 (D) P-3, Q-1, R-1, S-4 (D) P-3, Q-1, R-2, S-1 (D) P-3, Q-1, R-2, S-4 (D) P-	Q.7	Identify the natural plant growth regulators from the following list.  (P) Zeatin  (Q) Benzylaminopurine (BAP)  (R) Indole-3-acetic acid (IAA)  (S) 2,4-Dichlorophenoxyacetic acid					
secreting B-lymphocyte (HPRT *) can be selected to produce monoclonal antibody by growing in a medium containing  (A) thiamine, hypoxanthine, aminopterin (B) thymidine, histidine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine (A) P-2, Q-3, R-1, S-4 (B) P-1, Q-2, R-4, S-3		(A) P, Q (B	) Q, S	(C) P, R	(D) R, S		
(B) thymidine, histidine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (C) uridine, hypoxanthine, aminopterin (D) thymidine, hypoxanthine, aminopterin (C) thymidine, hypoxanthine, aminopterin (C) thymidine, hypoxanthine, aminopterin (C) thymidine, hypoxanthine, aminopterin (E) the forum 1. Group 2. (B) P-2, Q-3, R-1, S-4 (C) P-3, Q-1, R-2, S-1 (D) P-3, Q-1, R-2, S-4 (D) P-3	Q.8	secreting B-lymphocyte (HPRT +) can be selected to produce monoclonal antibody by					
Group 1 P. VNTR sequence Q. Leader sequence Q. Leader sequence R. SD sequence S. cis-acting sequence Q. Leader sequence S. cis-acting sequence Q. DNA finger printing S. cis-acting sequence S. cis-acting sequence Q. DNA finger printing S. cis-acting sequence S. cis-acting site S		(B) thymidine, histidine, aminopterin (C) uridine, hypoxanthine, aminopterin					
P. VNTR sequence Q. Leader sequence Q. Leader sequence R. SD sequence S. cis-acting sequence Q. Leader sequence R. SD sequence R. SD sequence S. cis-acting sequence Q. Leader sequence R. SD sequence Q. Ribosome binding site R. SD sequence Q. DNA finger printing S. cis-acting sequence Q. Page of the following sequence A. Functions in attenuation  (A) P-3, Q-1, R-4, S-2 (B) P-2, Q-3, R-1, S-4 (D) P-3, Q-1, R-2, S-4  (D) P-3, Q-1, R-2, S-4  Q.10 Q.10 Q.10 Q.10 Q.10 Q.10 Q.10 Q.1	Q.9	Match items in group 1	Match items in group 1 with correct options from those given in group 2.				
controlled by appropriate sensor (probe). Match each probe in group 1 with the appropriate response mechanism in group 2.  Group 1 (Probe) P. Thermistor Q. Oxygen electrode R. Metal rod S. pH electrode Group 2 (Response) 1. Activation of acid / alkali pump 2. Activation of vegetable oil pump 3. Activation of hot / cold water pump 4. Increase / decrease in stirrer motor speed  (A) P-2, Q-3, R-1, S-4  (B) P-1, Q-2, R-4, S-3		P. VNTR sequence Q. Leader sequence R. SD sequence S. cis-acting sequence (A) P-3, Q-1, R-4, S-2	<ol> <li>Gene regula</li> <li>Ribosome t</li> <li>DNA finger</li> </ol>	pinding site printing a attenuation (B) P-2, Q-3, R-	-1, S-4		
P. Thermistor Q. Oxygen electrode R. Metal rod S. pH electrode  (A) P-2, Q-3, R-1, S-4  1. Activation of acid / alkali pump 2. Activation of vegetable oil pump 3. Activation of hot / cold water pump 4. Increase / decrease in stirrer motor speed	Q.10	controlled by appropriate sensor (probe). Match each probe in group 1 with the					
(C) P-3, Q-2, R-4, 3-1 (D) P-3, Q-4, R-2, 3-1		P. Thermistor Q. Oxygen electrode R. Metal rod S. pH electrode	<ol> <li>Activation</li> <li>Activation</li> <li>Activation</li> </ol>	of acid / alkali p of vegetable oil of hot / cold wat decrease in stirre	pump er pump r motor speed -4, S-3		
Q.11 Which of these mice fail to develop a thymus?  (A) Mountain mice (B) Beige mice (C) Knock out mice (D) Nude mice	Q.11	(A) Mountain mice	to develop a thyn	(B) Beige mice			

Q.12							
	clone encoding a protei	n X that has b	been isolated and purif	ied?			
	(P) m-RNA isolation						
	(Q) Antibody preparation						
	(R) Cloning into an app						
	(S) Western blotting on	(S) Western blotting on transferred plaques					
	(A) P, S   (1	B) Q, S	(C) Q, R	(D) R, S			
Q.13	When electroporation is	s used for inte	roducing DNA into ma	ammalian cells			
	(P) a carrier for DNA is not required						
	(Q) the lipid bilayer (m	embrane) inte	eracts with an electric	pulse to generate			
	permeation sites	1945 74	70 (40)				
		<ul><li>(R) the viability of the cells becomes approximately zero percent</li><li>(S) the first step involves absorption of DNA on the cell membrane</li></ul>					
	(S) the first step involve	es absorption	of DNA on the cell m	embrane			
	(A) P, Q   (1	B) Q, R	(C) P, S	(D) Q, S			
Q.14				ires			
	(P) photosensitive poly		ol dimethacrylate				
	(Q) CNBr activation of		dennes et all the old the transfer and the Arthree than the area and				
	(R) polyfunctional reagent like hexamethylene diisocyanate						
	(S) radiation of polyvin						
	(A) P, Q	B) R, S	(C) P, S	(D) Q, S			
Q.15	Which one of the following monolayer culture systems have the highest						
	surface area: medium ratio?						
	(A) Roux bottle		(B) Spiracell roller bottle				
	(C) Hollow fibres		(D) Plastic bag/film				
Q.16	Q.16 Majority of the cereals are highly recalcitrant to Agrobacterium-mediated						
	transformation, and so direct transformation methods have been development						
	transform such plants. Which of the following direct transformation methods i						
	applicable to intact plan	nt tissues?					
	(A) Calcium chloride and PEG-mediated transformation						
	(B) Liposome-mediated	iated transformation					
	(C) Electroporation						
	(D) Transformation using microprojectiles						
Q.17							
	Group 1	The State of the second field of the State o	Group 2				
	P. Tissue plasminogen	activator	1. Immunomodulator				
	Q. Gamma interferon		2. Biodegradable pla	SUC			
	<ul><li>R. Podophyllotoxin</li><li>S. Polyhydroxyalkanoa</li></ul>	ate	<ol> <li>Clot dissolution</li> <li>Anti-tumor agent</li> </ol>				
	5. I orymydroxydikanor		rand-tunior agent				
	(A) P-4, Q-3, R-1, S-2		(B) P-1, Q-3,				
	(C) P-3, O-1, R-2, S-4		(D) P-3, Q-1,	K-4, S-2			

Q.18	Match items in group 1 with correct options from those given in group 2				
	P. Amperometr Q. Evanescent R. Calorimetric S. Potentiometr	wave biosensor biosensor	Group 2 1. Light beam 2. Flux of redox electro 3. Field effect transistor 4. Exothermic reaction		
	(A) P-3, Q-4, R (C) P-3, Q-2, R		(B) P-2, Q-1, R-4 (D) P-2, Q-4, R-4		
Q.19	(P) homology r (Q) threading f (R) threading e	For prediction of three dimensional structure of protein (P) homology modeling tries many possible alignments (Q) threading first identifies homologues (R) threading evaluates many rough models (S) homology modeling optimizes one model			
	(A) Q, S	(B) P, Q	(C) R, S	(D) Q, R	
Q.20	Immobilization of enzymes  (P) increases the specificity of the enzyme for its reactants (Q) facilitates reuse of the enzyme in batch reactions (R) makes it unsuitable for its use in a continuous reactor system (S) decreases the operational cost of the industrial process				
	(A) Q, S	(B) Q, R	(C) R, S	(D) P, Q	
Q.21	<ul> <li>Which of the following would result in somaclonal variation in micropropagated plants?</li> <li>(P) Propagation by axillary branching in the absence of plant growth regulators</li> <li>(Q) Cell suspension maintained for five years before induction of somatic embryogenesis</li> <li>(R) Callus induction using 20μM 2,4-Dichlorophenoxyacetic acid followed by shoot organogenesis</li> <li>(S) Shoot organogenesis from an explant in the absence of an intermediate callus phase</li> </ul>				
	(A) P, Q	(B) Q, R	(C) P, S	(D) Q, S	
Q.22	The enzymes that can be used in 5'end labeling of DNA are (P) alkaline phosphatase (Q) DNA ligase (R) terminal transferase (S) polynucleotide kinase				
	(A) P, S	(B) R, Q	(C) P, R	(D) R, S	

#### Common Data Questions

#### Common Data for Questions 23, 24:

Lignocellulosic biomass was subjected to microbial composting. The microbial consortium produced an extracellular enzyme xylanase, which was a glycoprotein having a molecular weight of 68 kDa and a positive charge. An aqueous extract of the enzyme could be easily prepared from the compost.

Q.23	What techniques would you recommend for confirming the molecular weight of the			
	purified enzyme?			
	(P) Isoelectric focusing			
	Value and a second seco			

- (Q) SDS-PAGE
- (R) Native PAGE
- (S) Gel filtration
- (A) P, Q
- (B) Q, S
- (C) R, S
- (D) P, S
- Q.24 If Con A sepharose column was used for the purification of enzyme, the separation would be based on
  - (A) molecular exclusion
- (B) affinity binding

(C) ion exchange

(D) hydrophobic interaction

Linked Answer Question: Q. 25 to Q. 26 carry two marks each.

#### Statement for Linked Answer Question 25 & 26:

DNA content of Caenorhabditis elegans was analysed and found to contain 1.0 x 108 bp.

- Q.25 How many standard λ- phage vector carrying 20kb DNA fragments or YACs carrying 250kb DNA fragments are theoretically required to constitute a complete C. elegan genomic library?
  - (A) 500 λ- phage vectors or 40 yeast clones
  - (B) 400 λ- phage vectors or 5000 yeast clones
  - (C) 5000 λ- phage vectors or 400 yeast clones
  - (D) 5 x  $10^4$   $\lambda$  phage vectors or 4000 yeast clones
- Q. 26 How many λ- phage vectors / yeast clones should be prepared in order to ensure that every sequence is included in the library?
  - (A) 25 x 103 λ- phage vectors / 2000 yeast clones
  - (B) 20 x 10<sup>3</sup> λ- phage vectors / 1600 yeast clones
  - (C) 5 x 10<sup>4</sup> λ- phage vectors / 4000 yeast clones
  - (D)  $10 \times 10^4 \, \lambda$  phage vectors / 10000 yeast clones

### Statement for Linked Answer Questions 27 & 28:

A bioreactor of working volume 50 m<sup>3</sup> produces a metabolite (X) in batch culture under given operating conditions from a substrate (S). The final concentration of metabolite (X) at the end of each run was 1.1 kg m<sup>-3</sup>. The bioreactor was operated to complete 70 runs in each year.

Q.27 What will be the annual output of the system (production of metabolite (X)) in kg per year?

(A) 55

(B) 3850

(C) 45.5.

(D) 77

Q.28 What will be the overall productivity of the system in kg year 1 m-39

(A) 19250

(B) 38.50

(C) 3850

(D) 77

## END OF THE SECTION