MARKING SCHEME

CLASS-XII

BIOTECHNOLOGY (2025-2026)

Q.no	Questions	Marks
1.	a) Variable number of Tandem Repeats	1
2.	a) DNA ligase	1
3.	c) Single-Stranded	1
4.	b) Restriction Fragment Length Polymorphism	1
5.	b) Transfection	1
6.	d) PAGE	1
7.	Both A and R are true and R explains A	2
8.	Both A and R are true but R does not explain A	2

9.	D 41 A 1D 4 1D 1 ' A	
9.	Both A and R are true and R explains A	1
10.	Tissue Culture	1
11.	Plasmid	1
12.	Cryopreservation	1
13.	Virus-free plant production	1
14.	Micropropagation	1
15.	Sickle Cell Anemia	1
16.	What are Restriction Enzymes? Restriction enzymes, also known as restriction endonucleases, are specialized enzymes that can cut DNA at specific sequences, usually palindromic sequences (same forward and backward).	
	 Naturally found in bacteria to defend against viral DNA. They recognize specific nucleotide sequences (called recognition sites) and cut the DNA at or near those sites. 	
	★ Example:	

	EcoRI is a restriction enzyme that recognizes the sequence: GAATTC and cuts between G and A .
	1. Cutting DNA Precisely: Allows scientists to cut DNA at exact locations.
	 Creation of Recombinant DNA: Used to insert foreign genes into plasmids (vectors).
	3. Gene Cloning: Essential in inserting genes into bacteria or other organisms to replicate.
	 Genetic Engineering: Enables modification of DNA to create genetically modified organisms (GMOs).
	 DNA Fingerprinting & Diagnosis: Helps in identifying individuals or detecting mutations in DNA.
17.	What are Cry Proteins?
	Cry proteins (short for crystalline proteins) are toxic proteins produced by the bacterium <i>Bacillus</i> thuringiensis (Bt).
	These proteins have insecticidal properties and are used in genetically modified (GM) crops to protect them from insect pests.

How Do They Work? When an insect larva eats the Cry protein, it gets activated in the alkaline gut of the insect. The protein **creates pores** in the gut lining, causing the insect to **die** from dehydration or starvation. **Example of Cry Protein:** Cry1Ac and Cry2Ab: Used in Bt cotton to kill bollworms. Cry1Ab: Used in Bt corn to protect against corn borers. Or**Bioinformatics** is the field that combines **biology**, computer science, and information technology to analyze and interpret biological data. It plays a vital role in **modern biotechnology**. **▲ Major Uses of Bioinformatics in Biotechnology:** Helps in reading and interpreting DNA sequences of humans, plants, and microbes. 2. Prug Discovery & Development Identifies potential drug targets using molecular modeling and simulations. 3. Genetically Modified Organisms (GMOs) Assists in designing genetically

engineered crops by analyzing gene functions.

4. Gene Prediction & Annotation

 Identifies coding regions and predicts gene functions using DNA data.

5. **Comparative Genomics**

 Compares genomes of different species to find evolutionary relationships.

6. Protein Structure Prediction

 Helps model 3D structures of proteins for understanding their function.

7. Personalized Medicine

 Tailors treatments based on a person's genetic profile.

8. **Database Management**

Stores and organizes large volumes of biological data like NCBI, PDB, GenBank.

18.	Feature	Stirred Type Bioreactor	Sprayed Type Bioreactor	
	Working Principle	Uses mechanical stirrers or impellers to mix nutrients and cells	Uses spray nozzles to spray culture media over surfaces	
	Oxygen Supply	Supplied through spargers and agitation	Supplied through air or gas spray	
	Mixing	Efficient due to mechanical agitation	Limited or uneven mixing	
	Used For	Suspension cultures (microbial/animal cells)	Immobilized cell cultures or surface growing organisms	2
	Design	Typically a cylindrical tank with baffles and stirrer	Tall chamber or column with trays or surfaces	
	Example	Used for making antibiotics, vaccines, enzymes	Used in wastewater treatment or plant tissue cultures	
19.	_	vectors are DNA molecunetic material into a host ion.	•	
	• Two (Common Types of Clon	ing Vectors:	
	1. Plas	smids		
	C	Small, circular, double found in bacteria	e-stranded DNA	5
	C	Example: pBR322, pl	UC19	
	2. Bac	teriophages (Phage Vec	etors)	
	C	Viruses that infect bac	eteria and can carry	

	larger DNA fragments	
	 Example: Lambda (λ) phage 	
	✓ Option 2: Define the Terms	
	a) Callus	
	A callus is an unorganized, mass of undifferentiated plant cells that forms when plant tissues are cultured on a nutrient medium.	
	• It can later differentiate into shoots or roots under specific conditions.	
	b) Explant	
	An explant is a small piece of plant tissue (like leaf, root, or stem) that is taken from a plant and used to start a tissue culture.	
20.	Gene cloning is the process of making identical copies of a gene using biological tools and host organisms (usually bacteria).	
	⋄ Steps in Gene Cloning:	
	1. Isolation of Gene of Interest	
	 Extract DNA and isolate the gene you want to clone. 	
	2. Cutting DNA with Restriction Enzymes	
	 Use enzymes to cut the gene and plasmid at specific sites. 	
	3. Insertion into Vector (Recombinant DNA)	

 Insert the gene into a plasmid/vector using DNA ligase.
4. Transformation
 Introduce recombinant DNA into a host cell (like <i>E. coli</i>).
5. Selection of Transformed Cells
 Use markers (e.g., antibiotic resistance) to identify successful clones.
6. Cloning and Expression
 Allow transformed cells to multiply and express the cloned gene.
Role of Enzymes in PCR (Polymerase Chain Reaction)
PCR is a technique to amplify DNA. Enzymes play a vital role at each step.
Key Enzyme Used:
1. Taq DNA Polymerase
 Heat-stable enzyme from <i>Thermus</i> aquaticus.
 Synthesizes new DNA strands during extension step.
Role in PCR Steps:
1. Denaturation (94–95°C)
 DNA strands separate – no enzyme involved.

	2. Annealing (50–65°C)	
	o Primers bind to template DNA.	
	3. Extension (72°C)	
	 Taq polymerase adds nucleotides to build the new DNA strand. 	
21.	Genetic Engineering (also called Recombinant DNA technology) plays a key role in developing crops with better yield, disease resistance, and improved nutrition by modifying their genetic material.	
	1. Improved Yield	
	Genes responsible for faster growth or larger produce are inserted.	
	Crops can be made to withstand drought, extreme temperatures, or salinity, ensuring consistent yield.	
	• Example: High-yield Bt cotton and Golden Rice.	
	* 2. Disease Resistance	
	Crops are engineered to produce proteins that kill pests or viruses.	
	Reduces the need for chemical pesticides.	
	• Example:	
	 Bt crops (cotton, corn) produce Cry proteins that kill insect larvae. 	

 Virus-resistant papaya (resistant to Papaya ringspot virus). 	
3. Nutritional Enhancement (Biofortification)	
• Crops are modified to produce more vitamins, minerals, and essential nutrients.	
Helps fight malnutrition and deficiency diseases.	
• Example:	
 Golden Rice is enriched with Vitamin A (β-carotene). 	
 Iron-enriched wheat and zinc-enriched rice. 	
 Advantages of Using Genetic Engineering in Crops: 	
Reduces use of chemical fertilizers and pesticides	
Increases crop shelf-life	
Supports sustainable agriculture	
Helps in feeding a growing population	
OR	3
Potential Applications of Plant and Animal Cell Culture	
∠ Applications of Plant Cell Culture	
Plant cell culture involves growing plant cells or tissues in a controlled, sterile environment. It has a wide range of agricultural, pharmaceutical, and	

industrial uses.

- 1. Micropropagation
- Rapid multiplication of disease-free plants
- Used in agriculture, horticulture, and forestry
 - 2. Production of Secondary Metabolites
- Produces valuable compounds like alkaloids, flavonoids, and essential oils
- Example: Shikonin, Taxol, Ajmalicine
 - 3. Germplasm Conservation
- Storage of rare or endangered plant species in vitro
- Useful in biodiversity conservation
 - 4. Genetic Engineering
- Used for gene transfer and creation of transgenic plants
- Improves traits like drought resistance or pest tolerance
 - 5. Somatic Hybridization
- Fusion of two different plant cells to create hybrid plants
- Combines traits from different species

🦬 Applications of Animal Cell Culture

Animal cell culture involves growing animal cells in a nutrient medium under sterile conditions. It plays a vital role in biotechnology,

	medicine, and research.	
	1. Vaccine Production	
	Cultured animal cells are used to produce vaccines like Hepatitis B, polio, and rabies.	
	2. Monoclonal Antibody Production	
	Used in diagnostics and cancer therapy	
	• 3. Tissue Engineering & Regenerative Medicine	
	Helps in creating artificial organs and tissues for transplantation	
	Example: Skin grafts, artificial cartilage	
	 4. Drug Testing & Toxicology 	
	Used to test new drugs and chemicals on cultured cells before animal or human trials	
	• 5. Genetic Studies & Cancer Research	
	Helps in understanding cell behavior, cancer development, and gene functions	
22.	Biosafety regulations are a set of guidelines and laws that ensure the safe handling, use, transport, and release of Genetically Modified Organisms (GMOs) to protect human health and the environment.	
	These regulations are essential for maintaining ethical and scientific standards in biotechnology.	
	Why Biosafety Regulations Are Important:	
	1. Prevents Health Hazards	

 Ensures GMOs do not cause allergies or toxic effects in humans or animals. 	
2.	
 Avoids unintended harm to non-target organisms, biodiversity, and ecosystems. 	
3. Regulates Field Trials	
 Controls where, how, and when GMOs can be tested or released. 	
4. Risk Assessment	
 Scientific evaluation is done before approving any GMO for commercial use. 	
5. Monitoring and Labeling	
 Helps in tracking GMO products and ensures proper labeling for consumer awareness. 	
6. Waste Disposal and Containment	
 Ensures safe disposal of GMO materials and prevents accidental spread. 	
Agencies Involved in India:	
GEAC (Genetic Engineering Appraisal Committee)	
RCGM (Review Committee on Genetic Manipulation)	
DBT (Department of Biotechnology)	
Or	3
What Are Transgenic Plants?	

Transgenic plants are those into which one or more foreign genes (transgenes) have been inserted using genetic engineering techniques to give them desirable traits, such as pest resistance, drought tolerance, or better nutrition.

Steps to Create Transgenic Plants:

1. Gene Identification

Select the desired gene (e.g., insect resistance gene from *Bacillus thuringiensis* – Bt gene).

2. Gene Cloning

 The selected gene is isolated and inserted into a vector (like a plasmid).

3. Gene Insertion into Plant Cells

- The gene is introduced into plant cells using methods like:
 - Agrobacterium-mediated transformation
 - Gene gun (biolistics)
 - Electroporation

4. Selection

 Use of marker genes (e.g., antibiotic resistance) to select successfully transformed cells.

5. Regeneration

 Transformed plant cells are grown on a nutrient medium to regenerate into a whole transgenic plant.

6.	6. Testing and Propagation o Transgenic plants are tested for the trait, grown in controlled fields, and multiplied for large-scale use.			
• E	Benefits of Trans	sgeni	ic Plants:	
✓ I	Benefit		T Example	
Pest	resistance		Bt cotton kills bollworm larvae	
Disea	ase resistance		Virus-resistant papaya	
Herb	oicide tolerance		Roundup Ready Soybeans	
Impr	oved nutrition		Golden rice with Vitamin A	
	Higher yield & stress tolerance		Drought-resistant maize or rice	
Redu	aced chemical u	se	Less pesticide and herbicide needed	
♪ P	otential Risks o	of Tra	ansgenic Plants:	
×	X Risk Co		Concern	
			n to non-target species, eversity loss	
Gene	Gene transfer Genes may transfer to wild plants (superweeds)			
Hum	ian health		gic reactions or unknown term effects	

	Ethical concerns Mixing genes across species boundaries Sand dependency on historia
	Economic issues Seed dependency on biotech companies
23.	PCR stands for Polymerase Chain Reaction. It is a laboratory technique used to amplify (make many copies of) a specific DNA segment — even from a very small amount.
	 Denaturation (94–95°C) The double-stranded DNA is heated to separate into single strands.
	 Annealing (50–65°C) Primers (short DNA sequences) bind to the target DNA strands at specific sites.
	 Extension/Elongation (72°C) The enzyme Taq DNA polymerase adds nucleotides to extend the primers and synthesize new DNA strands.

Cycle Repeats
These steps are repeated multiple times to exponentially increase the DNA quantity.
Or
DNA isolation (or extraction) is the process of separating DNA from cells or tissues in pure form for analysis or experiments.
Basic Steps of DNA Isolation:
1 Cell Lysis (Breaking the Cells)
The cell membrane and nuclear membrane are broken using a lysis buffer containing detergents like SDS (sodium dodecyl sulfate).
This releases DNA, proteins, and other cell contents.
2 Removal of Proteins and Cell Debris
Protease enzymes or chemicals (like chloroform or phenol) are added to break down proteins.
Centrifugation is used to separate the clear DNA- containing solution from the debris.
3 DNA Precipitation
Cold alcohol (usually ethanol or isopropanol) is added.
DNA is insoluble in alcohol, so it precipitates

	(becomes v	visible) as white threads or clumps.	
	remove im	pellet is washed with 70% ethanol to purities and salts.	
	•	ension NA is dissolved in TE buffer or sterile torage or further use.	
	★ Summary Tal	ble:	
	Step	Purpose	
	Cell lysis	Break open the cells	
	Removal of proteins	Remove unwanted proteins/debris	
	Precipitation	Make DNA visible and collectable	
	Washing	Purify the DNA	
	Resuspension	Store DNA in usable form	
24.	individuals based The principle is b (Variable Numbe Tandem Repeats)	ng is a technique used to identify on unique patterns in their DNA. eased on the presence of VNTRs or of Tandem Repeats) or STRs (Short — these are highly variable regions g part of the genome.	

 Core Principle: Every individual (except identical twins) has a unique DNA sequence. Specific regions of DNA have repeating sequences that vary in number between individuals. These regions are extracted, amplified (via PCR), and compared to create a DNA profile. 	
Use in Paternal Disputes (Paternity Testing): In cases where the identity of a child's biological father is disputed: 1. DNA is extracted from the child, mother, and alleged father. 2. The child's DNA profile is compared with both parents. 3. Since a child inherits half DNA from each parent, the father's DNA should match with the child's non-maternal bands. 4. If there is no match, the alleged person is not the biological father.	3
 ★ Applications in Paternity Cases: Settling legal disputes about biological fatherhood. Used in custody cases, inheritance claims, and adoption confirmation. 	

	Admissible as legal evidence in court.	
25.	GEAC is the Genetic Engineering Appraisal Committee, which functions under the Ministry of Environment, Forest and Climate Change (MoEFCC), Government of India.	
	 Main Role: GEAC is the apex body in India responsible for: Approving research and release of genetically modified organisms (GMOs) and products. Ensuring biosafety in the use of GM crops and biotechnology products. 	
	 Key Functions: Evaluate GM Research Projects Approves lab and field trials of transgenic plants. Assess Environmental Impact Analyzes risk to humans, animals, and biodiversity. Authorize Commercial Release Gives final clearance for GM crops like Bt cotton. 	3

	A Promo Dion CA D. 14' C. 1'	
	 4. ✓ Ensure Biosafety Regulations Compliance 	
	 ★ Important Point: GEAC works according to the rules under the Environment (Protection) Act, 1986. It plays a crucial role in balancing biotechnology progress with public safety and ethics. 	
26.	Animal cell culture refers to the in vitro (outside the body) growth of animal cells under controlled conditions, and it plays a key role in biopharmaceutical production, especially for therapeutic proteins.	
	What are Therapeutic Proteins? These are proteins used to treat diseases by replacing a deficient or abnormal protein in the body. Examples: Insulin, Interferons, Monoclonal antibodies, Human Growth Hormone	
	Applications of Animal Cell Culture in Therapeutic Protein Production:	_
	 1. Production of Recombinant Proteins Animal cells like CHO (Chinese Hamster Ovary) cells are genetically modified to produce: 	

 Insulin for diabetes Erythropoietin (EPO) for anemia Interferons for viral infections and cancer therapy 	
 2. Monoclonal Antibody Production Used in treatments for cancer, autoimmune disorders, and infectious diseases. Example: Trastuzumab (Herceptin) for breast cancer. 	
 Vaccine Production Cultured animal cells are used to produce safe and effective vaccines. Example: Polio, Hepatitis B, and COVID-19 vaccines. 	
 4. Gene Therapy Products Animal cells are used to grow viral vectors that carry therapeutic genes to treat genetic disorders. 	
 5. Tissue Engineering and Regenerative Medicine Culturing cells to create artificial skin, cartilage, or even organs, often with proteins that aid healing. 	5
Stem cells are undifferentiated cells that have the	

 unique ability to: Self-renew (divide and make more stem cells) Differentiate into various specialized cell types (like muscle, nerve, or blood cells) 	
What is Regenerative Medicine? Regenerative medicine is a field of medicine that focuses on repairing, replacing, or regenerating damaged tissues and organs using cells, genes, or biologically engineered materials.	
Role of Stem Cell Technology in Regenerative Medicine:	
 Tissue Repair and Regeneration Stem cells can replace damaged cells in tissues like skin, liver, heart, or nerves. Example: Treating burn victims using skin stem cells. 	
 Organ Regeneration Research is ongoing to grow entire organs (like liver, kidney) in the lab using stem cells — a future solution to organ donor shortage. 	
Treatment of Degenerative DiseasesHelps in treating diseases like:	

	 Parkinson's disease 	
	o Alzheimer's	
	 Spinal cord injuries 	
	o Type 1 diabetes	
	Bone Marrow Transplantation	
	The oldest and most common form of stem cell	
	therapy using hematopoietic stem cells to treat	
	blood cancers like leukemia.	
	5 Personalized Medicine	
	Patient's own stem cells can be used to reduce	
	the risk of rejection and create customized	
	treatments.	
	⚠ Challenges:	
	Ethical concerns (especially with embryonic	
	stem cells)	
	Risk of tumor formation	
	High cost and technical complexity	
27.	Both PCR (Polymerase Chain Reaction) and Gene	
	Cloning are used to amplify DNA, but they differ in	
	methods, tools, and applications.	
	Feature PCR (Polymerase Chain Gene Cloning	
	Reaction) Definition Definition Reaction In vitro method to amplify In vivo method to make	
	Definition DNA using enzymes and copies of a gene inside	

	temperature	a host cell	
Process Type	Artificial (test tube-based)	Biological (cell-based)	
Enzyme Used	Taq DNA Polymerase	DNA ligase, Restriction enzymes	
Time Required	Very fast (few hours)	Slower (may take days)	
Tool Used	Thermal cycler (PCR machine)	Vectors like plasmids, bacteria (E. coli)	
Accuracy	High, but may introduce errors with long sequences	Very accurate and maintains stable long inserts	
Purpose	Rapid amplification of DNA	Cloning, expression, or storage of desired genes	
Product	Only DNA copies	DNA integrated in living cells for further use	
	R is like photocopying D chine.		
	ne cloning is like putting letting the cell multiply		
	Or		5
enzyme th	nerase is a heat-stable Diat was originally isolate dic bacterium <i>Thermus a</i> ngs.	d from the	
(up	capable of withstanding to 95°C), making it idea lymerase Chain Reaction	al for use in PCR	
nuc	ynthesizes new DNA stra leotides to a DNA templ ension step of PCR.	•	
* Key P	roperties:		
	ermostable: Doesn't dena peratures	ature at high PCR	

	Optimal temperature: Works best at around 72°C For the Complete August 1 of the complete August 2 of the complete A	
	Fast: Can replicate thousands of base pairs in a few seconds	
	★ Important Application:	
	✓ Used in PCR (Polymerase Chain Reaction)	
	Taq polymerase is essential for DNA amplification in PCR because it can repeatedly synthesize DNA even after the high-temperature denaturation step.	
	Without Taq polymerase, PCR would not be possible, as regular enzymes would break down at high heat.	
28.	What is Biopiracy?	
	Biopiracy refers to the unauthorized use or patenting of	
	biological resources (like plants, animals, or traditional knowledge) by companies or researchers, often without	
	proper credit or compensation to the local communities	
	or countries from where they originated.	
	Famous Examples of Biopiracy:	
	1 Neem (Azadirachta indica) – India	
	A U.S. company tried to patent the antifungal properties of neem oil, a plant used in Indian traditional medicine for centuries.	
	The patent was later revoked after a legal	

challenge.	
 A U.Sbased company, RiceTec, tried to patent a variety of Basmati rice developed from Indian strains. This led to a major biopiracy dispute with India defending its traditional crop. 	
 Turmeric (Haldi) – India Two U.S. scientists were granted a patent on the wound-healing properties of turmeric, known in Indian Ayurvedic medicine. The patent was eventually canceled after evidence of prior traditional use was presented. 	5
 Hoodia – South Africa A plant used by San tribes to suppress hunger during long hunts. A pharmaceutical company patented it for weight-loss drugs without initially compensating the indigenous people. 	

29.	Gene transfer refers to the insertion of a foreign gene (transgene) into a plant's genome to modify or improve its traits (e.g., pest resistance, drought tolerance, nutrition).
	Two Major Methods of Gene Transfer in Plants:
	 L Agrobacterium-Mediated Gene Transfer (Biological Method) Uses Agrobacterium tumefaciens, a soil bacterium that naturally transfers genes into plant cells. Scientists modify the Ti plasmid of the bacterium to carry desirable genes (instead of tumorcausing genes). The bacterium infects plant cells, and the foreign gene is integrated into the plant genome.
	 Example: Used to develop Bt cotton (with insecticidal Cry gene) Used in virus-resistant papaya
	 2. Gene Gun or Biolistics (Physical Method) Tiny gold or tungsten particles coated with DNA are shot into plant cells using high pressure. The DNA enters the nucleus and integrates into the plant's genome. Example:

Used in rice, corn, and wheat, especially for monocots, which are hard to infect using Agrobacterium.
 → Other Gene Transfer Methods (Less Common): • Electroporation – Using electric pulses to open cell membranes. • Microinjection – Directly injecting DNA into plant cells (rare in plants).