SAMPLE QUESTION PAPER

BIOTECHNOLOGY (045)

Class XII (2022-23)

Max.Marks:70 Time allowed: 3 hours

General Instructions:

- i) All questions are compulsory.
- ii) The question paper has five sections. All questions are compulsory.
- iii) Section—A contains 12 Multiple choice questions and 4 Assertion-Reasoning based questions of 1 mark each; Section—B has 5 short answer questions of 2 marks each; Section—C has 7 short answer questions of 3 marks each; Section-D has two casebased question of 4 marks; Section-E has three long answer questions of 5 marks each.
- iv) There is no overall choice. However, internal choices have been provided in some questions. A student has to attempt only one of the alternatives in such questions.

	SECTION A	
1.	Male sterility is widely used in crops such as maize, sunflower for hybrid production. Male sterile plants are created by introducing a gene encoding-	1
	(a) Barnase protein	
	(b) TA29	
	(c) Barstar protein	
	(d) Coat protein	
2.	Body builders prefer to drink buffalo milk to build muscle mass. Determine the reason for this?	1
	(a) Easier to digest	
	(b) Lower fat content	
	(c) Higher calcium and phosphorus content	
	(d) Balanced calorie source	

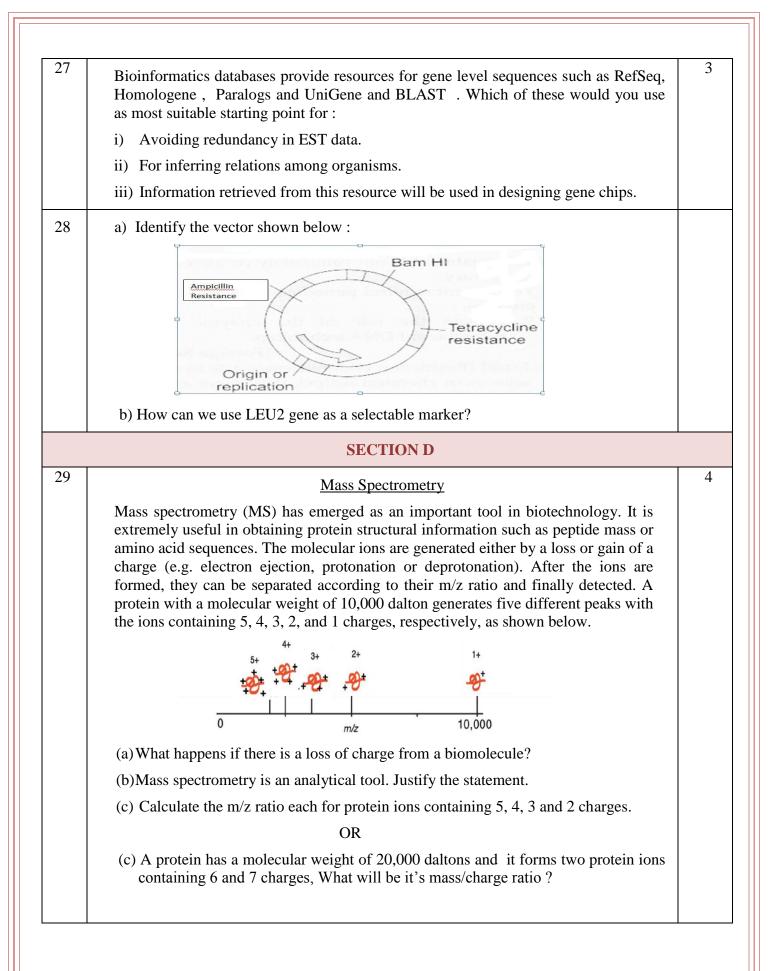
3.	An industrially important secondary metabolite which is used as a red pigment in lipstics and dye for silk is obtained from-	1
	(a) Datura stramonium	
	(b) Lithospermum erythrorhizon	
	(c) Digitalis lanata	
	(d) Coptis japonica	
4.	Proteome of a given cell is dynamic because :	1
	(a) In response to Internal and external changes the biochemical machinery of the cell could be changed.	
	(b) In response to Internal and external changes the biochemical machinery of the cell could not be changed.	
	(c) No direct relationship exists between Internal and external changes in the biochemical machinery of the cell.	
	(d) Indirect relationship exists between Internal and external in changes the biochemical machinery of the cell.	
5.	Artificial seeds are produced by-	1
	(a) Encapsulating somatic embryos in calcium alginate beads	
	(b) Desiccating the somatic embryos with or without coating	
	(c) Hydrating the somatic embryos	
	(d) Hydrating the zygotic embryos.	
6.	Being a researcher, you want to improve the deficiency of certain amino acids in cereals and legumes. Choose the technique out of the following which will be the best to achieve your goal:	1
	(a) Plant tissue culture	
	(b) Adding fertilizers to soil	
	(c) Protein engineering	
	(d) Vegetative Propagation	
7.	Foreign DNA is directly introduced into the recipient cell using a fine micro-syringe to transform it. The probable advantage this provides could be:	1
	a) No specialised equipment required	
	b) No damage to cells	
	c) Low transduction rate	
	d) Precision of delivery	

8.	A piece of young hypocotyl was cultured in MS medium in a plant tissue culture lab. This is a type of-	1
	(a) Organ culture	
	(b) Callus culture	
	(c) Explant culture	
	(d) Mass cell culture	
9.	Molecular Biologists prefer to use artificial vectors with MCS. List a benefit for this choice.	1
	(a) Flexibility in choice of insert size	
	(b) Flexibility in choice of vector size	
	(c) Flexibility in choice of host organism	
	(d) Flexibility in choice of restriction enzyme	
10.	Native enzyme Subtilisin is inactivated by bleach, in detergents because of oxidation of methionine at position 222. Choose a strategy that will help overcome this problem:	1
	(a) Use Pepsin instead of Subtilisin	
	(b) Eliminate use of bleach	
	(c) Substitute another amino acid at position 222	
	(d) Use Amylase instead of Subtilisin	
11.	Culture based approaches for detecting pathogens, as compared to PCR based assays are	1
	(a) Faster, safer but less specific	
	(b) Slower but safer and more specific	
	(c) Slower, less safe and less specific	
	(d) Slower, less safe but more specific	
12.	A 100 Kb DNA fragment has to be cloned in a host cell. Which vector should be used for this experiment?	1
	a) Plasmid	
	b) Cosmid	
	c) BAC	
	d) Bacteriophage lambda	

Question No. 13 to 16 consist of two statements – Assertion (A) and Reason (R). Answer these questions selecting the appropriate option given below:	
A. Both Assertion and Reason are true and the reason is the correct explanation of the assertion	
B. Both Assertion and Reason are true but the reason is not the correct explanation of the assertion	
C. Assertion is true but Reason is false	
D. Both Assertion and Reason are false	
Assertion-The functional property of whey protein exploited in confectionery is browning.	1
Reason -Whey proteins undergo maillard reaction providing colour and aroma to food items	
Assertion - Foaming is a problem in most microbiological processes.	1
Reason - It is caused due to the presence of fatty acids and silicones in the culture medium.	
Assertion - Whey mixed with herbs and honey is administered to the sick to treat ailments like jaundice and infected skin lesions.	1
Reason - Whey proteins elevates the levels of glutathione which protects the cells from harmful oxygen intermediates.	
Assertion- It's difficult to count genes even if we know where the genes are in a given genome	1
Reason- There is no simple correlation between the intuitive complexity of an organism and the number of genes in its genome.	
SECTION B	
Depict the production and mode of action of tissue plasminogen activator through diagram or flowchart.	2
X is a valuable tool in plant breeding, wherein variation in tissue culture regenerated plants from somatic cells can be used for the development of crops with novel traits. Identify 'X'. State any one example where this tool can be used for crop improvement. OR	2
Leaf explants of brinjal are showing multiple shoot regeneration in a plant tissue culture laboratory. Which plant regeneration pathway is depicted here? In this process, what would happen if either auxins or cytokinins are high in the medium?	
	Answer these questions selecting the appropriate option given below: A. Both Assertion and Reason are true and the reason is the correct explanation of the assertion B. Both Assertion and Reason are true but the reason is not the correct explanation of the assertion C. Assertion is true but Reason is false D. Both Assertion and Reason are false Assertion-The functional property of whey protein exploited in confectionery is browning. Reason-Whey proteins undergo maillard reaction providing colour and aroma to food items Assertion- Foaming is a problem in most microbiological processes. Reason- It is caused due to the presence of fatty acids and silicones in the culture medium. Assertion- Whey mixed with herbs and honey is administered to the sick to treat ailments like jaundice and infected skin lesions. Reason - Whey proteins elevates the levels of glutathione which protects the cells from harmful oxygen intermediates. Assertion-It's difficult to count genes even if we know where the genes are in a given genome Reason- There is no simple correlation between the intuitive complexity of an organism and the number of genes in its genome. SECTION B Depict the production and mode of action of tissue plasminogen activator through diagram or flowchart. X is a valuable tool in plant breeding, wherein variation in tissue culture regenerated plants from somatic cells can be used for the development of crops with novel traits. Identify 'X'. State any one example where this tool can be used for crop improvement. OR Leaf explants of brinjal are showing multiple shoot regeneration in a plant tissue culture laboratory. Which plant regeneration pathway is depicted here? In this process, what

Given below is a list different organisms. conserved, and (b) are	Based on	this inform					
Position → Organism↓	1	2	3	4	5	6	
Human	D	I	P	G	Н	G	
Chicken	D	I	A	G	Н	G	
Alligator	K	L	P	Е	Н	G	
Turtle	D	L	S	A	Н	G	
Tuna	D	L	T	T	M	G	
Carp	D	F	E	G	T	G	
	2000	ith antigen	G G	9 0			2
	pleen Ce Hybridom	Ills Fu	Myel	loma Ce	lls		
Cultur	pleen Ce	Ills Fu	usion	loma Ce	noclona		

	S	ECTION C			
22	(a) Chymotrypsinogen is inactive form of enzyme chymotrypsin. Which molecular alteration converts it into active form?				
	(b) The catalytic triad in chymotrypsi	in leads to a charge relay system. Justify			
		OR			
	1	lividual has to be compared with that of a person hology laboratory. Represent the steps of the same, in the form of a flow chart.			
23		s approved by US Food and Drug Administration Name the genes A to F introduced for the	}		
	Crop Gene Imp	roved character			
	Canola A Hyb	orid production			
	Corn B Inse	ect resistance			
	Cotton C Inse	ect resistance			
	Papaya D Viro	as resistance			
	Potato E Inse	ect and virus control			
	Soyabean F Wee	ed control			
24		f the culture medium has significant role in cell ingredients decides osmolarity of the medium			
25	You have the gene sequence of a pr you establish through tools of bioinfo	otein which has a proteolytic activity. How will ormatics that this protein:	3		
	(a) Has homologues in other organ	nisms			
	(b) Belongs to the chymotrypsin f	amily			
	(c) Has a database that can we proteolytic protein	used to trace the evolutionary history of this			
26		cleases (RE)? Give an example of a type II RE sequence recognized by it. Mention two other experiment.	}		



	No. of cell division	0	1	2	3	n	
		_				2 ⁰	
	No. of cells	1	2	4	8	_	
	Mathematically	N_0	$N_0 \times 2^2$	$N_0 \times 2^2$	$N_0 \times 2^2$	$N_0 \times 2^2$	
	bling time which is ell division is inverse		-			ole through one i	ound
(a)	In a microbiology generation time 20 time 30 s. Which ba	s and	d other bac	terial cult	ure is marked		
(b)	Using the above tab undergone to increa					population must	have
(c)	Using the above tab population in which cells/ml during four	h the	number of	bacteria i	ncreases from	ng time) of a bac 10 ⁸ cells/ml to	terial 10 ¹⁴
OR							
(c)	Explain any two dif	ferent	ways to me	easure mi	crobial growth.		
			SEC	CTION E			
anii use	reral medically important and rell culture and rell for the production alar form.	ecom	binant DNA	A technolo	ogy. Represent	the animal cell	line
 (a) Erythropoietin (b) Factor VIII (c) Follicle Stimulating Hormone (FSH) (d) Interleukin 2 (IL 2) 							
(e)	Monoclonal antibod	ies (n	nAbs)				
			(OR			
(i)	Differentiate between Defined and Serum-Anchorage-depende	suppl nt and	d Anchorage	e-indepen		ion two advanta	

32	a) Dr. Sharma discovered first restriction enzyme ever from a bacteria called <i>Thermus aquaticus</i> , strain DR 15. Name the enzyme.	5
	b) Design two primers (5 nucleotide long each) for the given sequence:	
	5'GATTCATTGCGCGCATTACTCGCATT3'	
	c) Recognition sites are generally palindromic in nature. Does it point towards the structure of restriction enzymes being that of a homodimer or heterodimer? Give reason for your answer.	
	d) A bacteriophage is known to infect <i>E.coli</i> with pili. How can it be modified to serve as a suitable vector?	
	(1+1+1+2)	
	OR	
	a) Schematically explain the formation of recombinant plasmid. (2)	
	b) Selection is an important step in genetic engineering. You are given ampicillin and tetracycline antibiotics. Using these antibiotics, which selection technique could be used to differentiate between recombinant and non-recombinant cells? (3)	
33	(a) A group of students are trying to isolate recombinant insulin .After processing the fermentation broth, they observed no yield .What could be the most possible reason for this?	5
	(b) A recently discovered microbial strain gives us the desired metabolite in nanomolar concentration. Suggest two ways of improving the production of the desired metabolite.	
	(c) <i>Pichia pastoris</i> has many advantages as a eukaryotic expression host. Justify giving two reasons.	
	OR	
	a) A professor told her students to ready a bacterial culture in 12 hours sharp. Suggest her students two ways to enhance the growth of bacterial cells in the lab so that they are able to fulfill the requirement.	
	b) Write any two commercial significance of microbial cell culture.	
	c) There are many ways of measuring microbial growth. Which technique is	

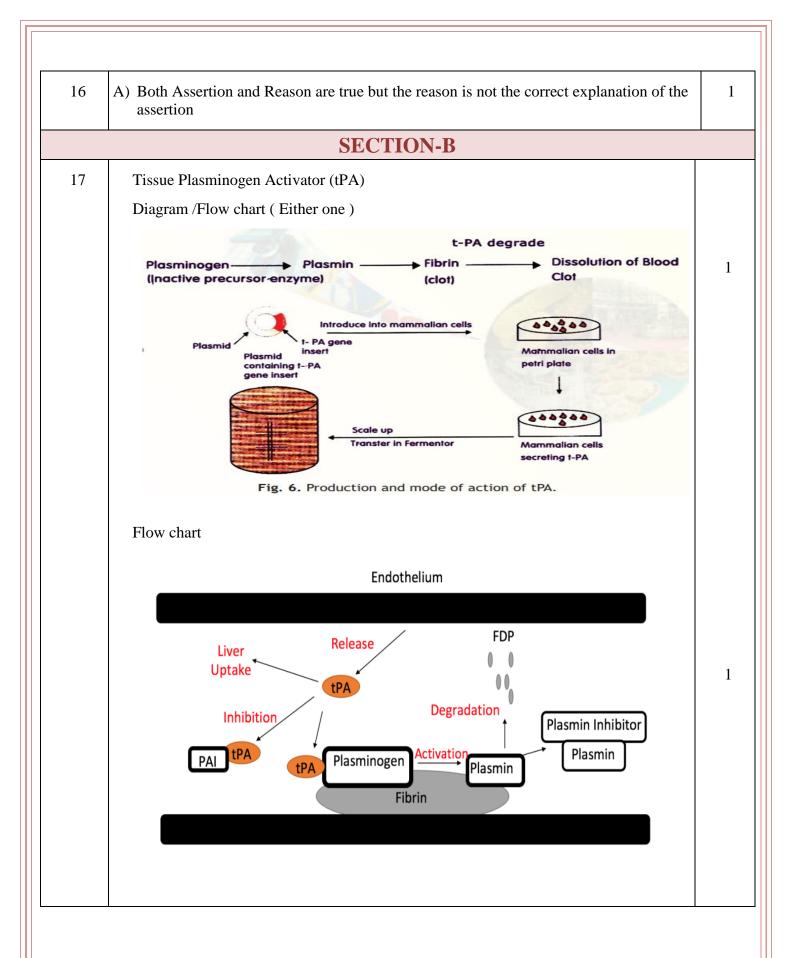
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Marking Scheme

BIOTECHNOLOGY (045)

Class-XII (2022-23)

	SECTION-A	
1	(a) Barnase protein	1
2	(c) Higher calcium and phosphorus content	1
3	(b) Lithospermum erythrorhizon	1
4	(a) In response to Internal and external changes the biochemical machinery of the cell could be changed.	1
5	(a) Encapsulating somatic embryos in calcium alginate beads	1
6	(c) Protein engineering	1
7	(d) Precision of delivery	1
8	(c) Explant culture	1
9	(d) Flexibility in choice of restriction enzyme	1
10	(c) Substitute another amino acid at position 222	1
11	(c) Slower, less safer and less specific	1
12	(c) BAC	1
13	A) Both Assertion and Reason are true and the reason is the correct explanation of the assertion	1
14	(C) Assertion is true but Reason is false	1
15	(A) Both Assertion and reason are true and reason is the correct answer for the assertion.	1



18	Somaclonal variations	1+
	It helps in production of mutants e.g. disease resistance in Potato	
	OR	
	Organogenesis	
	If auxins are high in the medium, it promotes rooting while if cytokinins are high, shoot formation is promoted.	
19	G amino acid is most conserved	1
	A amino acid is most variable.	1
20	Essential amino acids and BCAA profile: Essential amino acids are those amino acids which have to be obtained from food and cannot be made in our cells.	1
	The branched chain amino acids (BCAA) are essential for the biosynthesis of muscle proteins. They help in increasing the bio-availability of high complex carbohydrates intake and are absorbed by muscle cells for anabolic muscle building activity.	1
	Biological value (BV) measures the amount of protein nitrogen that is retained by the body from a given amount of protein nitrogen that has been consumed. It has been observed that the BV of whey proteins is the highest compared to rice, wheat, soya and egg proteins.	
	Protein efficiency ratio (PER)- PER is used as a measure of growth expressed in terms of weight gain of an adult by consuming 1g of food protein. The PER value of the following proteins are arranged in decreasing order- whey,milk, casein, soya, rice, wheat. (Any two)	
21		2
21	a) Production of MoAb (0.5 mark)	2
	b) This technology has revolutionized the area of diagnostics and antibody-based therapies.	
	1) The availability of monoclonal antibodies has helped in the early detection of many infectious diseases like hepatitis and AIDS.	
	2) Therapeutic mAb –	
	OKT3 Therapeutic mAb - Herceptin OKT-3 is monab-CD3, an immunosuppressant drug given intravenously to reverse the acute rejection of transplanted organs such as the heart, kidney and liver.	
	Herceptin (trastuzumab) is a monoclonal antibody approved for therapy of early-stage breast cancer that is Human Epidermal growth factor Receptor 2-positive (HER2+). (1.5 marks)	

		CECETON		
	Т	SECTION-	C	
22	is inactichymotry substrate (b) Asp 102, The negatively His, leaving	ve. In-situ activation of trypsic psinogen which results in a color binding pocket. His 57 and Ser 195 lie in this order y charged aspartate carboxylate resilit with a negative charge Ser195	site is blocked and hence the enzyme in involves a proteolytic cut in informational change, exposing the forming a charge relay; due pulls the Ser –OH proton through becomes acidic due to the unique cause the protein has folded uniquely	1+2
		OR		
		Purify H	aemoglobin Sickle cell RBC	
		Trypsin Hemoglobin Hemoglobin is cleaved by protease trypsin. Try bonds adjacent to a lysi	psin breaks peptide	
		-	etrophoresis omatography	½ x 6
		Result : All peptides we samples except one (market)		
		Peptide se	equencing	
		Fig. 6. Protein f	ingerprinting	
		Protein fingerprinting	/ peptide mapping	
23	Crop	Gene	Improved Character	½ x 6
	Canola	(A) Barnase Barstear	Hybrid production	
	Corn	(B) BtCrylA(c)	Insect Resistance	
	Cotton	(C) BtCrylA(c)	Insect Resistance	
	Papaya	(D) Coat protein	Virus Resistance	
	Potato	(E) BtCrylllA & Coat protein	Insect & virus control	
	Soyabean	EPSP synthase	Weed control	

24	Membrane integrity maintained	1x.
	Helps to maintain the shape and size of cells.	
	Salt, glucose and amino acids (any two) are the major ingredients that determine osmolality of the medium.	
25	(a) →BLAST search→ Find out→ homologous sequences in other organisms by looking for gene sequence of given proteolytic enzyme.	1
	(b) Look for conserved domain and find whether belongs to domain of Chymotrypsin or to other family of proteins	1
	(c) ALI database can be used for Phylogenetic (Evolutionary) analysis and alignment of proteins.	1
26	R.E. type II recognize a specific DNA sequence and cut within the sequence generating sticky/flush ends. In recombinant DNA technology, we use type II RE as they are highly specific in their action.	1
	Alu I with the restriction site (One strand) 5' AGCT'3 and Sma 1 with the restriction site 5 'CCC GGG' 3(flush ends) (One strand)	1
	The functions of a) Alkaline phosphatase b) DNA ligase.	
	*The role of alkaline phosphatase is to prevent self re-ligation of the vector	1/2
	*The role of DNA ligase is to make 3'-5' phosphodiester bond.	1/2
27	: i) UniGene database	1
	ii) Homologene database	1
	iii. RefSeq database	
28	a) p BR 322	1
	b) LEU2 gene codes for an enzyme required for the synthesis of amino acid leucine.	1
	Yeast cells having this plasmid can grow on a medium lacking leucine and hence	1/2
	can be selected e.g. Yep	1/2

	SECTION- D	
29	(a) The molecular ions are generated either by a loss or gain of a charge (e.g. electron ejection, protonation or deprotonation)	1
	(b) Mass spectrometry is used in-	1
	(i) Obtaining protein structural information such as peptide mass or amino acid sequence	
	(ii) Identifying the type and location of amino acid modification within proteins. (any one)	
	(c) $(c)m/z = (M+nH)^{n+}/n^{+}$	2
	For n=5, m/z= 10,000+ 5/5= 2001 For n=4,m/z= 10,000+ 4/4= 2501 For n=3, m/z= 10,000+ 3/3= 3334.3 For n=2, m/z= 10,000+ 2/2=5001	
	(c) $m/z = (M + nH)^{n+}/n^+$ For $n=6$, $m/z = 20,000 + 6/6 = 3334.33$ For $n=7$, $m/z = 20,000 + 7/7 = 2858.14$	
30	a) As generation time is inversely related to specific growth rate, hence bacterial culture marked "X" with generation time 20s will proliferate rapidly.	1
	b) $n = 3.3 \text{ (Log } 10^7 - \text{Log } 10^4)$ = 3.3 (7-4) = 10	1
	c) First calculate the number of divisions the population must have undergone to increase from 10 ⁸ to 10 ¹⁴ in 24 hours.	
	$n = 3.3 \text{ (Log } 10^{14} - \text{Log } 10^8)$ = 3.3 (6) = 19.8 $t_d = 240 \text{ minutes} / 20$ = 12 minutes	2
	OR c) (i) Measurement of Dry mass and Wet mass	
	(ii) Using spectrophotometer(iii) Using Slide counting Chamber(iv) Using Coulter chamber	
	(Any two)	

		SECTION- E		
31				
	Proteins	Animal cell line used	Therapeutic use	½ x10
	Erythropoietin	CHO cells	Anemia	
	Factor VIII	CHO cells	Hemophilia A	
	Follicle Stimulating	CHO cells	Infertility	
	Hormone (FSH)			
	Interleukin 2 (IL 2)	CHO cells	Cancer therapy	
	Monoclonal antibodies (mAbs)	Hybridoma cells	Cancer therapy & Autoimmune diseases	
		OR		
	port growth of selection culture media and is (ii) Anchorage depende	has known chemicals, of fixed cted cells. Serum is an essentia a source of growth factors and tent cells grow as adherent cell as suspension cultures.	al component of animal cell l hormones.	2
		ring system used to maintain stem. Carbon dioxide from cells to drop in pH.		2
	$H_2O + CO_2$	$\langle \langle \rangle \rangle H_2CO_3 \langle \langle \rangle \rangle (H^+)$	+ (HCO3 ⁻)	
	Increase in Bicarbonate dioxide according to the	concentration neutralizes the e following equation:	effect of increased Carbon	
	J	NaHCo3> (Na+)+(HCO3-)		
	The increased HCO3- io reached at pH 7.4	ons derive the above equation t	o its left until equilibrium is	1
	Advantages:			
		o maintain in balance/ enzy s to cell surface receptors/Ion b		
32	(a) TaqDI			1
	(b) 5' AATGC 3' and 5' G	ATTC 3'		1
	(c) Palindromic means the Restriction enzyme is a	e DNA sequence reads same was homodimer.	when read from 5' to 3'. The	1/2
	As it cuts both the strand	ds of DNA simultaneously in 5	' to 3' direction.	1/2
		inserted into bacteriophage sirrupting any of the essential gen	-	
	_	nage which infects E. coli having cells containing a F plasmid (ca	~	2

	OR	2
	a)	
	Digention with Ecofil 3' Foreign DNA 5' Ori Plasmid vector with restriction enzyme (Ecofil) Ligate together with DNA ligase Treat with alkaline phospalite Insert Antibiotic resistance gene Sticky ends Treat with alkaline phospalite Insert Antibiotic resistance gene Sticky ends Antibiotic resistance gene Sticky ends Antibiotic resistance gene Sticky ends Antibiotic resistance gene Sticky ends	
	 b) Replica plating. Host cells are first plated (master plate) on solid media with the desired 	1/2
	 antibiotic overnight. Velvet paper is aligned, pressed on master plate. With the same alignment it is pressed onto the replica plate. 	1/2*:
	 Keep it overnight ,transformed colonies will not grow in replica plate The colonies having insert can easily be scored off from master plate by comparing the two plates. 	
33	(a) Recombinant insulin is an intracellular protein so we need to process the cell mass and not the fermentation broth.	1
	(b) Strain improvement is done in order to maximize metabolite production by:i) Mutant selection: There are two methods - Physical method & Chemical Method	1
	ii) Genetic engineering(c) i) It has strong inducible promotersii) It is capable of making post-translational modifications similar to those	1
	performed by human cells iii) Downstream processing is simpler as Pichia does not secrete its own proteins into the fermentation medium. (Any two)	1x2
	OR	

a) <u>Use</u>	of shake culture and Use of baffle flask
Baffle flas	sk: One of the simplest ways is to produce a V- shaped notch or indentation in the sides of the flask. Such flasks are called baffle flasks. This improves the growth of the microbes by improving the efficiency of oxygen transfer due to increased turbulence of the agitated culture medium.
Shakers:	Continuous agitation of the culture medium also greatly improves the efficiency of the oxygen transfer and this improves the growth of the microbes. In the laboratory, this is done by the use of shakers . Shakers may be end-to-end type or rotatory type. These may be designed for use at the ambient temperature or in a controlled temperature environment (incubator shaker).
b)	
1. Produ	ction of whole microbial cells (for food, vaccines)
2. Product	cion of primary metabolites (acids, alcohol)
3. Product	cion of secondary metabolites (antibiotics)
4. Biotran	sformation reactions (enzymatic, steroid)
5. Exploit	ation of metabolism (microbial leaching, biodegradable waste treatment)
-	sis of recombinant proteins (therapeutic proteins) Bioremediation/fermented ems/ recombinant proteins (Any two)
c) Viable	Plate Count is the best method since it does not count dead microbial cells.