P P SAVANI UNIVERSITY

Fifth Semester of B.Sc. Examination Dec2020-Jan 2021

SSBT3130-Recombinant DNA Technology I Time: 10:00a.m. to 12:30 p.m.

06.01.2021, Wednesday

Section-A (Total Marks - 20) (20) Objectives (20 MCQ Compulsory-20 mark each) If a double stranded DNA is treated with ______ enzyme, single stranded DNA product can be obtained after the treatment Bal31 A B Exonuclease-III C Endonuclease V D All the options are correct 2 Which of the following can be cleaved by S1 nuclease $\,$ Only Single stranded DNA B Only double stranded DNA with a nick Both double stranded DNA and single stranded DNA Both double stranded DNA with a nick and single stranded DNA 3 Recognition sequence for enzyme Hinfl is/are: GAATC В GAGTC C GATTC All of the options are correct BsrFI recognizes RCCGGY, where _ R= A or G and Y= C or T R= C or T and Y= A or G C R= A, G, C or T and Y= A, G, C or T All the options are correct 5 If the first 323 amino acids of DNA pol I polypeptide is removed. Polymerase function will be lost Nuclease activity will be lost Both polymerase and nuclease activity will be lost None of the activity will be lost 6 enzyme adds one or more deoxyribonucleotides onto the 3' terminus of a DNA molecule Alkaline phosphatase Polynucleotide kinase Terminal deoxynucleotidyl transferase None of the options are correct The vaccinia topoisomerase cuts DNA at the sequence 雄蕊 CCTTT A B CCCTT C CCCCT D CTTTT Which of the following plasmids do not have any apparent effect on the phenotype of host 8 R plasmid A B Cryptic C Col Cryptic Which of the following is an expression plasmid pBR322 В pUC18 pUC19 C

Maximum Marks: 60

	D	pET14-b	1
10		controlled by a set of <i>tra</i> genes	
	A	Conjugation and plasmid transfer	
	В	Only conjugation	
	C	, P	-
	D	All the options are correct	
11	In λ gt10 vector insert can be inserted into a unique EcoRI site located in the cI gene.:		
	A	clear rather than turbid plaques.	
	В	Insertional inactivation of this gene means that recombinants are distinguished as turbid rather than clear plaques.	
	C	and the state of this gene means that recombinants are distinguished as clear	
	D	rather than turbid plaques.	48.60
		Note of the options are correct	790 307
12	M13, after entering to E. coli host, converts into		
	A	single stranded DNA molecule called non-replicative form	
	В	double stranded DNA molecule called non-replicative form	
	C	double stranded DNA molecule called replicative form	
	D	Single stranded DNA molecule called replicative form	
13	In which of the following area(s) M13 vectors are useful		29_42
	A	DNA sequencing	-000
	В	Mutagenesis study	
	C		
	D	probe generation	
	Ъ	All of the options are correct	
14	In pEMBL8 vector the cloning sites are present within the lacZ' gene, so the visual screening can		
	DC	performed by.	
	A	Identifying recombinant plasmids on agar containing X-gal	40.0
	В	Identifying recombinant plaques on agar containing X-gal	
	C	Identifying non-recombinant plaques on agar containing X-gal	
	D	Identifying non-recombinant plasmids on agar containing X-gal	
15	In YAC, the presence of the insert DNA in the vector can be checked by testing for insertional		
	IIId	cuvation of:	
	A	SUP4 gene	
	В	URA3 gene	
	C	TRP1 gene	-
	D	LEU2 gene	
16	In YEp13 vector region codes for a protein that can convert A form to the B form		
	A	REP	
	В	FLP	
	C	REP	
	D	LPF	
17	C)	on a Section of the Long and Montale in the Control of the Control	
17		ning in Drosophila can be done by:	
	A	The N element	
	В	The P element	
	C	The C element	
	D	The T element	
8	For	using URA3 as a selectable marker in a cloning experiment	
		- a coming experiment	100

The yeast vector should have functional URA3 gene, whereas the host cell must have nonfunctional URA3 gene The yeast vector should have non-functional URA3 gene, whereas the host cell must have functional URA3 gene The yest vector and host both should have non-functional URA3 gene The yeast vector and host both should have functional URA3 gene In order to use LEU2 as a selectable marker, the host must be an _ auxotrophic mutant that has a non-functional gene for b-isopropyl-malate dehydrogenase В auxotrophic mutant that has afunctional LEU2 gene C auxotrophic mutant that has a non-functional gene for orotidine 5'-phosphate decarboxylase D None of the above 20 To clone the rabbit $\beta\text{-globin}$ gene, in simian virus 40 (SV40) ${\bf A} \quad \text{ The HindIII to BamHI restriction fragment of Early gene region was replaced} \\$ The HindIII to BamHI restriction fragment of Late gene region was replaced ${f C}$ The EcoRI to BamHI restriction fragment of Late gene region was replaced $\label{eq:Def} \textbf{D} \quad \text{ The EcoRI to BamHI restriction fragment of Early gene region was replaced}$ Section-B (Total Marks - 40) Short Notes (attempt allfour compulsory- 3 marks each) (12)Adaptors Vs linkers A В Replacement vector Yeast Episomal Plasmid (YEp13) C D Degenerate recognition sequence Explain in detail (attempt anyfour compulsory-7 marks each) Q.2 (28)A What are the basic steps of gene cloning? Explain each stepin detail? Discuss some of the strategy to introduce sticky end onto a blunt-ended DNA molecule? В What are Cosmids? Explain how concatemer is formed in EMBL4 vector? C 25 Explain YAC in detail. Why three different selection markers are required for cloning in YAC? D Describe two cloning plasmid vectors based on E. coli?