## P. P. SAVANI UNIVERSITY

Fifth Semester of B.Sc. Examination December-2021

SSBT3150-Recombinant DNA Technology-II

13.12.2021, Monday Time: 12:30 p.m. to 03:00 p.m. Maximum Marks: 60

## Instructions:

- 1. The question paper comprises of two sections.
- 2. Section I and II must be attempted in separate answer sheets.
- 3. Make suitable assumptions and draw neat figures wherever required.
- 4. Use of scientific calculator is allowed.

## Section-I (Total Marks - 30)

- Q.1 Short Questions
  1.1 Objectives
- 1.1a Which of the following cannot be studied using cDNA libraries
  - A Alternative splicing
  - B Regulatory regions of the genes
  - C Gene function
  - D All of the above
- 1.1b Which RNA is used for the construction of cDNA library
  - A rRNA
  - B tRNA
  - C mRNA
  - D sRNA
- **1.1c** Reverse transcriptase catalyzes the conversion of
  - A RNA to DNA
  - B RNA to Protein
  - C DNA to RNA
  - D Protein to RNA
- 1.1d Southern blotting is used for
  - A RNA hybridization
  - B Protein hybridization
  - C DNA hybridization
  - D All of the above
- 1.1e Which of the following is/are ideal vector for large inserts
  - A Cosmids
  - B λ phages
  - C BACs
  - D All of the above
- 1.1f Which of the enzyme is used for Nick Translation
  - A Klenow Fragment
  - B DNA Pol I
  - C Reverse Transcriptase
  - D Topoisomerase
- 1.1g In blue white screening, BLUE colonies reflect

[10]

/[05]

	A	Transformed and recombinant cells	
	В	Untransformed and recombinant	
	C	Transformed but non-recombinant cells	
	D	Untransformed and non-recombinant	
1.1h	Diet	hyl pyrocarbonate, is used during RNA extraction to:	
	A	extract protein	
	В	inhibit RNAase	
	C	degrade the cell wall	
	D	sepearte RNA from the mixture	
1.1i	Duri	ng Phenol-chloroform extraction, If pH is acidic (4.0)	
	A	RNA will stay in aqueous phase and DNA will be in organic phase	
	В	RNA will stay in organic phase and DNA will be in aqueous phase	
		RNA will stay in the interphase and DNA will be in aqueous phase	
		None of the options are correct	,
1.1j	Duri	ing RNA extraction,degrades proteins and inhibit RNAses	
		Guanidium thiocynate	
	В	Phenol- Chloroform	
		Isopropanol	
	D	All the options are correct	
1.2		wer the Following: (MCQ/Short Question/Fill in the Blanks)	[05]
1.2a		stands for	
1.2b		ng cDNA synthesis, enzyme is used to cut the loop of dsDNA at	
	one e		
1.2c	In Ri	NA extraction, 70 % ethanol is added for removing the salts (T/F)	
1.2d		genomic library construction and screening, a small genome will require	
1.0		r clones than a more complex one (T/F)	
1.2e	A gt1	0 has LacZ' for visual screening (T/F)	
Q.2	Char	t Notes (Attempt any true)	
A.Z		t Notes (Attempt any two) nutant	[06]
В	*	se and Carbon probability formula	
C		ening a cDNA library with a labeled oligonucleotide probe based on a	
C		vn peptide sequence	
	KIIOV	vii peptide sequence	
Q.3	Expl	ain in detail (Attempt any two)	[1.4]
A		uss different strategies to prepare Genomic library?	[14]
В		ribe different methods of radioactive and non-radioactive probe	
	label		
C		t is Colony Hybridization? Explain each sten-in detail	

	Section-II (Total Marks - 30)			
Q.1	Short Questions	[10]		
1.1	Objectives [0			
1.1a	In a given genome, the average spacing between HindIII sites (six-base-long			
	HindIII sequence) is approximately			
	A 0.25 Kb			
	B 4Kb			
	C 8 Kb			
	D 6 Kb			
1.1b	While cloning in EMBL4 vector, which of the following site in a recombinant			
	genome are cut to be packaged into phage heads			
	A Right arm			
	B Cos site			
	C Left arm	1		
	D All of the above			
1.1c				
	recombinant clones?			
	A Promoter			
	B Selection marker			
	C Origin of replication			
	D Multiple Cloning Site			
1.1d				
	with			
	A EcoRI			
	B Xbal			
	C BamHI			
	D HindIII			
1.1e				
	plating onto:			
	A agar containing only X-gal			
	B agar containing only IPTG			
	C agar containing both X-gal and IPTG D None of the above			
1.1f	In Nick translation, radioactive probes are labelled:			
1.11	A with P <sup>31</sup>			
	B with P <sup>35</sup>			
	C with P <sup>32</sup>			
	D with P <sup>33</sup>			
1.1g	P1-derived artificial chromosome, or PAC, is a DNA construct derived from the			
1.15	T derived artificial circumstance, or FAG, is a DIAA construct derived from the			

D DNA of P1 bacteriophages and Yeast artificial chromosome

1.1h During end labelling, \_\_\_\_\_ enzyme treatment of probe is done to

A DNA of P1 bacteriophages and Bacterial artificial chromosome
 B DNA of λ bacteriophages and Bacterial artificial chromosome
 C DNA of λ bacteriophages and Yeast artificial chromosome

	remove 5' phosphate	
	A Alkaline phosphatase	
	B Polynucleotide kinase	
	C DNase I	
	D Klenow fragment	
1.1i	Which of the following was the first widely adopted method for DNA	
	sequencing	
	A Sanger	
	B Maxam-Gilbert	
	C Ion-torent	
	D Solexa	
1.1j	Ion Torrent sequencing measures thefrom the incorporation of	
	individual bases by DNA polymerase	
	A Indirect release of H+	,
	B Indirect release of nitrogenous base	
	C direct release of nitrogenous base	
	D direct release of H+	
1.2	Answer the Following: (MCQ/Short Question/Fill in the Blanks)	[05]
1.2a	In Blue-White screening when insert is inserted within the lac Z region, it	[03]
	produces blue colonies (True/ False)	
1.2b	Insertional inactivation of cI gene produces turbid plaques (Ture/False)	
1.2c	Anti-digoxigenin antibodies with high affinities and specificity are used in a	
	variety of biological immuno-assays (Ture/False)	
1.2d	is a method to detect a polypeptide produced from a cloned gene	
1.2e	Define BACs	
Q.2	Short Notes (Attempt any two)	[06]
A	Nick translation	[00]
В	Maxam Gilbert sequencing	
С	Ion torrent sequencing	
Q.3	Explain in detail (Attempt any two)	[14]
A	What is NGS? Describe Solexa (Illumina) sequencing in detail.	1 10 1
В	Describe Immunoscreening technique in detail.	
C	What is Sanger sequencing? How is Sanger Sequencing different from Maxam	
	Gilbert sequencing?	