## P. P. SAVANI UNIVERSITY

Fifth Semester of B.Sc. Examination December-2021

SSBT3090-Animal Biotechnology-II

Time: 12:30 p.m. to 03:00 p.m.

Maximum Marks: 60

Instruction	mer

10.12.2021, Friday

- 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 [10]
1.1 Objectives [05]

- 1.1a Stem cells can
  - A Renew
  - B Differentiate
  - C Multiply
  - D All of the above
- 1.1b Fertilized egg is
  - A Totipotent
  - B Multipotent
  - C Oligopotent
  - D Unipotent
- 1.1c To maintain pH of medium we add
  - A Salts
  - B Sodium
  - C Buffer
  - D Amino acids
- 1.1d To block mitosis we use
  - A Colchicine
  - B Colcemid
  - C Vinblastin
  - D Any one of the above
- 1.1e Animal Biotechnology includes
  - A Cell culture
  - B Tissue culture
  - C In vitro fertilization
  - D All of the above
- 1.1f DNA has high melting point when it has
  - A High AT content
  - B High GC content
  - C High AG content
  - D High TC content

		1				
1.1g	In animal tissue culture, the osmotic balance is maintained by	1				
,	A Sodium	1				
	B Potassium					
	C Calcium					
	D All of the above					
1.1h	•					
	A Hybridization					
	B Western Blotting					
	C Cell Separation					
	D Cell isolation					
1.1i	Conditioned media is					
	A Defined media					
	B Undefined media					
	C Synthetic media					
	D None of the above					
1.1j						
	A Cell separation					
	B Cell isolation					
	C DNA Hybridization					
	D Blotting					
1.2	Answer the Following:	[05]				
1.2a	SDS PAGE method is used					
1.2b	What is the Full form of VNTRs?					
1.2c	Define Karyotype.					
1.2d	What is animal cell characterization?					
1.2e	Transformed cells have limited life span (True/False).					
Q.2	Short Notes (Attempt any two)	[06]				
A	Feeder cells					
В	Conditioned media					
C	Northern Blotting					
Q.3	Explain in detail (Attempt any two)	[14]				
A	Explain how HAT media could be used as selective inhibitors.	[14]				
В	Explain different methods for animal cell separation.					
C	How do you prepare chromosome slide? & Write a detailed note on different					
	techniques of chromosome banding techniques.					
	Table of the office partning techniques.					

## Section-II (Total Marks - 30)

Q:1	Short Questions		[10]
1.1	Objectives		[05]
1.1a	We freeze the cells to avoid		
	A Genetic drift		
	B Transformation C Contamination		
	D All of the above		
111	Miliah af the faller single material water	t-2	
1.1b	Which of the following is not cryoprotectan A DMSO	L!	
	A DMSO B PEG		
	C PVP		
	D GMSO		
	D GM30		
1.1c	Inflammation causes due	NV is near the notice and world for the	
1.10	A Transformation		
	B Irritation		
	C Multiplication	· .	
	D None of the above		
	Tions of the above		
1.1d	Differentiated Fibrocytes produce		
	A Collagen	Menoral medical process of the control of the contr	
	B Albumin		
	C Hemoglobin	wagatersol antobar maketi in p	
	D Serum		
1.1e	Flow cytometry is used in		
	A Cell multiplication		
	B Cell culture		
	C Cell feeding		
	D Cell counting		
1.1f	is an example of terminal di	fferentiated cell.	
	A Zygote		
	B HSC		
	C RBC		
	D Myloid and Lyphoid		
1.1g		owing chemicals are used for	
	cellular communication		
	A Hormone interaction		
	B Chemical interaction		
	C Same type of cell interaction		
	D Different types of cell interaction	*	
111	DE2 is involved in		
1.1h	P53 is involved in		
	A Immortalization		
	B Malignancy	1.	
	C Anchorage independence D All of the above		
	D All of the above		

1.1i	Animal cells have following growth phases  A Lag  B Log  C Stationary  D All of the above	
1.1j	Marker for erythrocytes is  A Albumin  B Hemoglobin  C Transglutaminase  D invoclurin	
1.2 1.2a 1.2b 1.2c 1.2d 1.2e	Answer the Following: Define Differentiation Define Immortalization Thawing is done instantly (True/False) Full form of DMSO. Characterization is not required for Animal cells (True/False)	[05]
Q.2 A B C	Short Notes (Attempt any two) What are the rationales for freezing animal cells? What are the in vitro limitations of assays? What different growth phases are observed in animal cell culture?	[06]
Q.3 A B	Explain in detail (Attempt any two)  Define cryopreservation and write a detailed note about it.  Why do we count cells and what different methods are employed?  Write a detailed note on animal cell differentiation.	[14]