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SEAT No.

[70]

No of Printed Pages: 02

# SARDAR PATEL UNIVERSITY

# M.Sc. Genetics, Third Semester Examination 1<sup>st</sup> November 2017, Wednesday 2.00 to 5.00 pm

	1 November 2017, wednesday					
	2.00 to 5.00 pm PS03CGEN01: Immunogenetics					
	Total Marks- 70					
No	te: i) Attempt all questions.					
	ii) Marks are indicated on the right hand side.	08				
Q.1						
1.	The antigens present in blood are presented to B cells in					
a. Lymph nodes b. spleen c. thymus d. MALT						
2.	The following cell surface marker is used to identify Tc population					
	a. CD4 b. CD6 c. CD8 d. C9					
3.	The major antibody class secreted in Peyer's patches is					
	a. IgA b. IgG c. IgE d. IgD					
4.	C1 inhibitor blocks following pathway of complement activation					
	a. Classical b. Alternative c. lectin d. all of these					
5.	The following interaction is not present between and antigen and antibody molecule					
	a. Covalent bond b. hydrophobic interactions c. hydrogen bonds					
	d. ionic bond					
6.	Luminex assay is done for					
	a. MHC matching b. minor antigen matching c. pre-existing antibodies					
	d. Blood group matching					
7.	The nervous system is affected in the following autoimmune disease					
	a. Diabetes mellitus b. Grave's disease c. Multiple Sclerosis d. Rheumatoid					
	arthritis					
8.	In bare lymphocyte syndrome, specific defect is observed in					
٠.	a. T cells b. B cells c. Neutrophils d. NK cells					
	d. Feelis S. Beelis C. Reddiophilis d. R. Celis					
Q.2	Attempt Any Seven of the following:	14				
1.	Enlist important differences between humoral and adaptive immunity.					
2.	What are germinal centers?					
2. 3.	-					
4.						
5.	What is class switching?					
5. 6.	Explain types of ELISA.					
7.	Mention characteristics of Myasthenia gravis.					
7. 8.	What are recombinant vaccines?					
9.	How is attenuated vaccine produced?					
Q.3 A	Give a detailed account of cells of myeloid lineage.	06				
Q.3 B	Write a note on: lymph node as a secondary lymphoid organ.	06				

### OR

Q.3 B	Explain the mechanism of killing by phagocytic cells.				
Q.4A	Explain the sequence of events occurring during gene rearrangement in the variable region of heavy chain.	06			
Q.4B	(i) Comment on polygenic and polymorphic nature of MHC genes.  (ii) Explain the structure of class I and class II MHC molecules.  OR	06			
Q.4B	Describe the production of human monoclonal antibodies by phage display.				
Q.5 A	Discuss the formation of C5 convertase during complement activation.	06			
Q.5 B	Explain methods for determination of antibody affinity.  OR	06			
Q.5 B	Explain " Memory and Specificity of graft rejection".				
Q.6A	Discuss the establishment of tolerance for self antigens in body.				
Q.6 B	Write a note on (i) Immunity to viral infections OR (ii) Primary Immunodeficiency	06			

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M. Sc. (Genetics) - Third Semester Examination (CBCS) Friday, 3rd November, 2017 2:00 p.m. to 5:00 p.m.

PS03CGEN02: Human Molecular Genetics Total Marks: 70 Note: (1) Figures to the right indicate marks. (2) Draw a neat and labeled diagram, wherever necessary. Q. 1 Choose the most appropriate answer from the four alternatives given: [08]i. Which one of this describes a contig? (a) Library of overlapping clones (b) A complete mRNA library (c) An ordered genomic library (d) None of these Linkage mapping can determine the distance between which of the following pairs of DNA sequences? (a) Two known genes (b) AFLPs and RFLPs (c) Two AFLPs (d) All of the above iii. Poly Q tract is a characteristic feature found in\_\_\_\_ (a) Cystic fibrosis (b) Huntington's chorea (c) Phenylketonuria (d) Hemophilia iv. Which one of the following is a temporary diabetes? (a) IDDM (b) Monogenic diabetes (c) Diabetes incipidus (d) Gestational diabetes v. Accumulation of branched chain amino acids in children causes (a) Cystic fibrosis (b) DMD (c) Phenylketonuria (d) MSUD vi. Which one of the following MPS has 4 subtypes? (a) Hurlers syndrome (b) Sanfilippo syndrome :(c) Morquio syndrome (d) Sly syndrome vii. Drug induced hemolytic anemia in African males are due to deficiency of\_ (a) Glutathione Reductase (b) NADPH (c) G6PDH (d) Glucose 6. Phosphatase viii. Which of the following is not a part of typical phase II conjugation reactions? (a) Acetylation (b) Hydroxylation (c) Methylation (d) Glucuronidation (iii. .. i. . n., 11.4. Satismilli Lille ich Q.2Answer any SEVEN from the following: [14] i. Differentiate between genetic mapping and physical mapping. ii. Enlist various positional independent strategies. iii. White functions of Htt protein. iv. Write examples of various epigenetic mechanisms. . Enlist X linked lipidosis and mucoploysachharidosis. P.T.O. 17:1. or Mary Land

VI.	write two major types of arbitish in humans.						
vii.	Why early diagnosis of IEM is crucial?						
viii.	Differentiate between pharmacogenetics and pharmacogenomics.						
ix.	Name genes affecting warfarin metabolism.						
Q.3(a)	Enlist various genetic markers. Discuss their importance in gene mapping.	[6					
(b)	Describe pulse field gel electrophoresis.	[6					
•	OR						
(b)	1. Explain synteny of genes.	[3]					
	2. Write a brief note on chromosome walking.	[3					
Q.4(a)	What are basis for the classification of trinucleotide repeat expansion disorders? Describe Friedreich's ataxia.	[6]					
·(p)	Justify that "obesity is a polygenic and multifactorial disease".	[6]					
	OR OR						
(b)	<ol> <li>Differentiate between hemophilia A, hemophilia B and hemophilia C.</li> <li>Write brief note on neurofibromatosis.</li> </ol>						
Q.5(a)	Describe molecular and biochemical aspects in PKU and alkaptonuria.	1.					
(b)	Describe human mitochondrial syndromes.	[6]					
(2)	OR	[6]					
(b)	Write short notes on the following:	: *′.					
(1)	I. Glycogen storage disease type II						
<i>1</i> 2	2. Gauchar disease	[3]					
(1.)	Re Say in	[3]					
Q.6(a)	Discuss practical implications of human genome projects.						
(b)	Discuss social and ethical issues in Medical genetics.	[6] [6]					
` ,	OR	וטן					
(b)	Explain SSCP and DGGE techniques for the scanning of genes for unknown mutation.						
	the sealing of genes for unknown mutation.	[6]					
# . 1,114							
	. *****						

1.4

1 15

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- (a) DNA sequencing based method
- (b) Restriction digestion based method

(c) PCR based method

(d) All of these

### (viii). Ideally distance between molecular marker and gene of interest or QTL is......

(a) < 1cM

(b) < 75 cM

(c) <25 cM

(d) <5 cM

### Answer any SEVEN from the following: Q.2

[14]

- (i). What do you mean by back cross method?
- (ii). Write a short note on polyploidy.
- (iii). Give a brief note on pure line selection in self pollinated crops.
- (iv). Differentiate between anther culture and ovule culture.
- (v). What do you mean by somatic hybridization?

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(	(vi).	Write any three factors influencing somaclonal variations.	
	vii).	Write a short note on pathogenesis related proteins.	
,	iii).	What do mean by flavr savr tomato?	
,	(ix).	Differentiate between AFLP and RAPD.	
Q.3	(a)	Write a detail note on back cross method in self pollinated crops.	[6]
	(b)	Discuss in detail about importance of mutational breeding in crop improvement	[6]
		programmes.	
		OR	
	(b)	What do you mean by male sterility in crops? Write various types of male sterility studied	[6]
_	4.5	by you.	[6]
Q.4	(a)	Write a detail note on methods for isolation of protoplast.	[6]
	(b)	Write a various applications of haploids and dihaploids in crop improvement.  OR	1~1
_			[6]
	(b)	Give a detail account on anther culture for haploids production.	[~]
Q.5	(a)	Define somaclonal variation? Discuss with in vitro scheme used for obtaining somaclonal	[6]
		variations in crop improvement programs.	r ća
	(b)	Discuss various applications and disadvantages of somaclonal variation.	[6]
		OR	
	(b)	Discuss the various approaches used for production of insect resistant transgenic plants.	[6]
Q.6	(a)	What do you mean by marker assisted selection? Write a detail note on nearly isogenic	[6]
		line (NL) strategy in crop improvement programs.	• . '
	(b)	Explain the following:	[6]
		(i) Improvement of starch and lipid quality in transgenic crops	:
		(ii) Role of herbicide resistance in crop improvement program with suitable examples	
		OR	. ,
	(b)	Write a detail note on drought resistance in transgenic crops.	[6]
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# M. Sc. GENETICS - Third Semester Examination

Thursday, 9<sup>th</sup> November, 2017.

2:00 p.m. to 5:00 p.m.

PS03EGEN01: Genetics of Mammalian Development

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		<del>-</del>			
(II) A	an que	estions are compulsory.		Total Marl	s: 70
Cho	ose th	e most appropriate alternative for th	e follow	ring:	(08)
1.	In o	ogenesis how many polar bodies are for	rmed at t	he end of meiotic division?	
	a)	One			
	c)	Three	d)	Four	
2.	In m	nammalian development, the embryo w	ill form f	from which population of	
	a)	the blastocoels	b)	the inner cell mass	
,	c)	the trophectoderm	d)	the extraembryonic membranes	
3.	Reti	noblastoma is associated with an abnor	mality of	f chromosome	•
	a)	14	b)	12	
	c)	13	d)	16	
4.	Serte	oli cells secrete hormone.			
	a)	Estrogen	b)	Testosterone	
	c)		d)	Ant-wolfian duct hormone	e
5.	Sing			<b>.</b>	
	a)		b)	syncitium	
	c)		d)	karyogamy	
6.	Peak	level of Bicoid regulatory protein	require	for structu	ire
	form	ation in Drosophila.			
	a)	Head	b)	Thorax	
	<b>c</b> )	Wing	d)	Segmented	
7.	Whi	ch one of the following is not a zygotic	gene?	•	
	a)	Gap gene	b)	Pair rule gene	
	c)	Segment polarity gene	d)	Selector gene	
8.		human conceptions do not succes	sfully de	velop to full term.	
	a)	1/2	•	1/3	
	c)	1/4			
	(ii) A Cho 1. 2. 3. 4. 5.	(ii) All que  Choose th  1. In of	<ol> <li>In oogenesis how many polar bodies are for a) One         <ul> <li>Three</li> </ul> </li> <li>In mammalian development, the embryo we cells?             <ul> <li>the blastocoels</li> <li>the trophectoderm</li> </ul> </li> <li>Retinoblastoma is associated with an abnormal at a polymorpherical secrets and a polymore.         <ul> <li>the trophectoderm</li> <li>Sertoli cells secrete hormone.</li> <ul></ul></ul></li></ol>	Choose the most appropriate alternative for the follow  1. In oogenesis how many polar bodies are formed at tanyone and tanyon	Choose the most appropriate alternative for the following:  1. In oogenesis how many polar bodies are formed at the end of meiotic division?  a) One b) Two c) Three d) Four  2. In mammalian development, the embryo will form from which population of cells? a) the blastocoels b) the inner cell mass c) the trophectoderm d) the extraembryonic membranes  3. Retinoblastoma is associated with an abnormality of chromosome a) 14 c) 13 d) 16  4. Sertoli cells secrete hormone a) Estrogen c) Anti-Mullerian Duct hormone d) Ant-wolfian duct hormone 5. Single cell with multiple nucei is called a) polymorphonucleus c) cleavage furrow d) karyogamy 6. Peak level of Bicoid regulatory protein require for structus formation in Drosophila. a) Head b) Thorax c) Wing d) Segmented  7. Which one of the following is not a zygotic gene? a) Gap gene c) Segment polarity gene b) Pair rule gene c) Segment polarity gene b) Pair rule gene c) Segment polarity gene c) Segment polarity gene d) Selector gene  8. human conceptions do not successfully develop to full term. a) 1/2 b) 1/3

1/2 - 2/3

d)

Q-2	Atter	npt ANY SEVEN from the following:	(14)
	1.	Draw and label the parts of a sperm cell, and list the functions of each.	
	2.	Write the basic steps of oogenesis.	
	3.	Classify the stem cells based on potency with example.	÷
	4.	Write about progeria.	
	5.	What is endocrine disruptor? give examples.	
	6.	Differentiate between cis and trans acting regulatory elements.	
	7.	Enlist major classes of developmental anomalies.	
	8.	Write names of any 2 transcription factors genes and their mutation phenotypes.	
	9.	Differentiate between primary and secondary infertility.	
Q-3	(a)	Discuss the types of cleavage patterns and the role played the yolk on cleavage.	(06)
	(b)	Describe the process of fertilization and add a note on prevention of polyspermy.  OR	(06)
	<i>(</i> )		
	(b)	Write short note on followings:	(03)
		<ol> <li>Development of human brain</li> <li>Cell movements during Gastrulation</li> </ol>	(03)
0.4	(-)	Explain the mechanisms for conversion of proto-oncogenes into oncogenes.	(06)
Q-4	(a)	How tumor suppressor genes lead to cancer? Explain with a suitable example.	(06)
	(b)	OR	( /
	(b)	Explain alcohol and retinoic acid as teratogens.	(06)
Q – 5	(a)	Describe cell to cell contact strategy for gene expression during development.	(06)
Q 3	(b)	Enlist techniques determining the functions of genes during development and Explain gene knock-out and anti-sense RNA techniques for the same.	(06)
		OR	
	(b)	Explain muscle differentiation in Sea squirt embryo by localized m-RNA.	(06)
Q – 6	(a)	Discuss various syndromes associated with human sex chromosomal aberrations.	(06)
	(b)	Give a detailed account on infertility.	(06)
	, ,	OR	
	(b)	<ol> <li>Explain positional cloning technique for identification of genes associated with human developmental anomalies.</li> </ol>	(03)
		2. Write a note on phenotypic heterogeneity.	(03)