(24)No. of Printed Pages: 2 SARDAR PATEL UNIVERSITY T.Y.B. Sc. (Genetics) - Fifth Semester Examination (CBCS) Monday, 25th November 2013 10:30 a.m. to 1:30 p.m. **US05CGEN06: Human Genetics** Total Marks: 70 Note: (1) Figures to the right indicate marks. (2) Draw a neat and labeled diagram, wherever necessary. Q1. Choose the most appropriate answer from the four alternatives given: [10] i) The current estimate for the number of human genes is_ A. 30 000 **C.** 1 000 000 **B.** 100 000 **D.** 30 0000 ii) Homologous repeats are involved C. Frameshifts **A.** Large deletions **B.** Missense mutations **D.** Splicing mutations iii) Unit for genetic map is_ A. cM C. Map unit **B.** Base-pairs D. Both A and C iv) Which of the following is not required in the whole-genome shotgun approach to genome sequencing? A. PCR reaction C. BAC library **B.** Physical map **D.** Plasmid library v) Which repetitive DNA includes an open reading frame for reverse transcriptase? A. LINE C. SINE B. DNA transposon D. VNTR vi) In human beings, multiple genes are involved in the inheritance of . C. Colour blindness A. Skin colour D. Phenylketonuria B. Sickle-cell anaemia vii) The most striking example of point mutation is found in a disease called A. Down's syndrome C. Thalassaemia B. Sickle cell anaemia **D.** All of these viii) Male XX and female XY sometime occur due to A. Transfer of segments in X and Y chromosome C. Hormonal imbalance D. None of these B. Deletion ix) Which of the following is not a hereditary disease? C. Cystic fibrosis A. Cretinism B. Haemophilia D. All of these x) Which of the following is a correct definition of genetics? C. The study of DNA A. The study of genes and traits defined by genes **B.** The study of variation between members of a species **D**. None of these

1 | Page

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01	A new on TEN from the following:	[20]
Q2.	Answer any TEN from the following: i. Define low-copy repeats.	[20]
	ii. What are coding sequences?	
	iii. Define physical map.	
	iv. What is quantitative inheritance?	
	v. Define chromosome walking.	
	vi. What is point mutation?	
	vii. Explain Tay Sach's Syndrome.	
	viii. What is haemophilia?	
	ix. Explain Ferric chloride test?	
	x. Define fetal chromosome.	
	xi. What is congenital disorder?	
	xii. Define overlapping DNA segments.	
Q3.	Discuss the potential benefits and problems of human genome project. OR	[10] 🛈
Q3.	Write notes on:	
•	a. Coding and non-coding sequences	[05]
	b. Gene families	[05]
Q4.	What is DNA construct? Discuss the applications of BACs library in genome projects. OR	[10]
Q4.	Write notes on:	
•	a. Physical mapping	[05]
	b. Positional cloning	[05]
Q5.	What are multifactorial disorders? Discuss the genetic basis of diabetes mellitus. OR	[10]
Q5.	Write notes on:	
	a. Monogenic diseases	[05]
	b. Obesity	[05]
Q6.	What is SAGE and how would you produces a snapshot of the mRNA population in the	[10]
~~~	sample of interest?	[10]
	OR	
Q6.	Describe genetic testing carried out at prenatal and neonatal stages of life with their significance.	[10]

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