LOYOLA COLLEGE (AUTONOMOUS), CHENNAI – 600 034



M.Sc. DEGREE EXAMINATION - BIOTECHNOLOGY

THIRD SEMESTER – NOVEMBER 2015

BT 3956 - FUNCTIONAL GENOMICS

Date: 13/11/2015 Time: 09:00-12:00	Dept. No.			Max. : 100 Ma	arks	
		Part- A				
Answer all the questions:			(20 Marks)			
I. Choose the correct answer				(5 ×	$(5\times1=5)$	
1. What is the reason behind	_		1\ M4	-4'11J		
a) C value paradox	b) Variation	c) Inbreeding	a) Mut	ational load		
2. Which of the following su a) Nylon	bstrate is suitable for b) Silica	or fluorescent probe c) Glas		d) Quartz		
3. Among the following which technology can be used to study proteins EXCEPT? a) SPR b) MS/MS c) NMR d) SAGE						
4. Which of the following usa) SPRc) Immunoprecipitation	es antibodies to stud b) FRET d) Pull dov					
5. Which among the following technique was used to study globin gene?a) RNAib) Gene knockout c) Ectopic expressiond) Insertional mutagenesis						
II State whether the following are True or False; if false give reason (5×1)					$(5\times1=5)$	
6. Luciferase is used to degrade the excess nucleotide bases.7. Operons are not found in prokaryotes.8. Genes are upregulated in cancers.9. Nitrosylation is very important for cell cycle progression.10. Metabolomics and metagenomics are same.						
III Complete the following					$(5\times1=5)$	
11. The longer arm of chrom12. Cy 3 gives13. The technique that can be	_ fluorescence.		ssion lev	vels is		
14 and 15. Most drug metabolizing	are o	common transient p	protein t	ransient binding dor	nains.	
IV Answer the following, ea	ach in about 50 wo	rds only:			$(5\times1=5)$	
16. What is alternate splicing 17. List one application of m 18. Name the enzymes used 19. Mention any two biologi 20. What is metabolic profili	icroarray technolog in SAGE. cal significance of p					



Answer the following, each in about 500 words;

 $(5 \times 8 = 40)$

Draw diagrams wherever necessary

21. a) Describe the four levels of annotation.

OR

- b) Illustrate the principle behind pyrosequencing.
- 22. a) Comment on the various substrates and dyes used in DNA microarrays.

OR

- b) Discuss about gene expression in eucaryotes.
- 23. a) Explain nuclease protection assay for RNA analysis.

OR

- b) Compare SAGE and MPSS.
- 24. a) Mention any four online sources to study protein interactions.

OR

- b) Review about the different types of protein-protein interactions.
- 25. a) Briefly explain gene knockout through homologous recombination.

OR

b) Discuss the pharmacokinetic and pharmacodynamic properties of drugs.

Part C

Answer any two of the following, each in about 1500 words; Draw diagrams wherever necessary

 $(2\times20=40)$

- 26. Elaborate RNA analysis using realtime qPCR.
- 27. Write in detail about any four techniques to study protein-protein interaction.
- 28. Explain about any two methods of genome sequencing and add a note on *de novo* genome assembly.
- 29. Describe about antisense RNA technology and RNAi.
