## LOYOLA COLLEGE (AUTONOMOUS), CHENNAI – 600 034



## M.Sc. DEGREE EXAMINATION - BIOTECHNOLOGY

## THIRD SEMESTER - NOVEMBER 2016

## **BT 3956 - FUNCTIONAL GENOMICS**

Date: 11-11-2016 Time: 09:00-12:00	Dept. No.		Max.: 100 Marks
		PART- A	
Answer all the questions	S		
I. Choose the correct an	swer		$(5 \times 1 = 5 \text{ Marks})$
1. Which among the follo	wing model organisms is	a vertebrate?	
a) D. melanogaster	b) Danio reiro	c) S.cerevisio	ae d) <i>E.coli</i>
2. The following are used	as substrates for microar	ray except	
a) Silica	b) Glass	c) Nylon	d) Quartz
3. Which of the following	g uses Bridge PCR for am	plification?	
a) Pyrosequencing	· <del>-</del>	-	equencing d) Em-PCR
4 is	the process of identifying	as many possible prot	eins in a test sample.
a) Mining		b) Profiling	
c) Protein network ma	11 0	d) Network expres	ssion
5. Which among the follo	=	= =	
a) RNAi b) Ge	ene knockout c) Ch	emical mutagenesis of	d) Insertional mutagenesis
II State whether the foll	owing statements are Ti	rue or False	$(5 \times 1 = 5 \text{ Marks})$
	C		,
6. Molecular and genetic	details of <i>C. elegans</i> is se	en in FlyBase.	
7. Oligonucleotide arrays	are more specific than cI	DNA arrays.	
8. Genome tiling arrays a	re used for genome wide	mapping of RNA.	
9. The term proteome was	· ·	_	
10. The primary miRNA	is converted into pre miR	NA by Dicer.	
III Complete the following			$(5 \times 1 = 5 \text{ Marks})$
11 is th	ne genome browser at NC	'RI	
2 destroys the unused dNTPs in pyrosequencing.			
is the study of the proteome, the protein complement of the genome.			
4 is a common drug metabolizing enzyme.			
15. The size of miRNA tr	_	•	
	•		
IV. Answer the following, each within 50 words			$(5 \times 1 = 5 \text{ Marks})$
16. What are NATs?			
17. Give an example for a	•		
18. Mention the tag based		A profiling.	
19. Write the principle of 20. Define metabolomics.			
20. Define metaudiumies.	•		

PART - B

 $(5 \times 8 = 40 \text{ Marks})$ 

Answer the following, each within 500 words. Draw diagram wherever necessary.

21. a) Explain the four levels of annotation in model organisms.

OR

- b) Draw the structure of an eucaryotic gene and mention any five properties that adds to the complexity of genome.
- 22. a) Write about facultative gene expression and constitutive gene expression.

OR

- b) Employ a suitable PCR method for real time quantification of gene expression.
- 23. a) Compare Northern blot and reverse northern blot.

OR

- b) Explain genome tiling microarray.
- 24. a) Elucidate with examples how online resources have enhanced the study of protein-protein interactions.

OR

- b) Comment on the different techniques to study protein- protein interactions.
- 25. a) Write about gene silencing using antisense RNA.

OR

b) Compare the response of people to the Codeines and Imatinib.

PART - C

 $(2 \times 20 = 40 \text{ Marks})$ 

Answer any TWO of the following, within 1500 words. Draw diagram wherever necessary.

- 26. Elaborate on pyrosequencing and Illumina sequencing.
- 27. Write in detail about microarray technology and add a note on its applications.
- 28. Post-translational modification in proteins increase the functional diversity of the proteome. Discuss.
- 29. Explain in detail about siRNA and miRNA mediated RNA interference.

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