Name :	URAN
Roll No. :	Construction Provider and Conference
Invigilator's Signature :	

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2012-13

# 2012

## DNA TYPING, PROTEOMICS AND BEYOND

*Time Allotted* : 3 Hours

Full Marks : 70

The figures in the margin indicate full marks. Candidates are required to give their answers in their own words

as far as practicable.

## **GROUP – A**

### (Multiple Choice Type Questions)

1.	Choose	the	correct	alternatives	for	any	ten	of	the
	following	g :					10 :	× 1 =	= 10

i) The example of one metabolic pathway database is

- a) KEGG b) SWISS
- c) BLISS d) OMIMM.
- ii) Transcriptome are the
  - a) Whole genome b) Whole set of RNA
  - c) Total set of exons d) Whole set of protein.
- iii) Number of base pairs in hypervariable region 1 of *mt*DNA control region is
  - a) 268 b) 342
  - c) 400 d) 500.

5059

[ Turn over

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2012-13

- iv) DYS-390 is a
  - a) Single locus VNTR marker
  - b) Multilocus VNTR marker
  - c) STR marker
  - d) None of these.
- v) LINEs are more frequent in
  - a) Human b) Birds
  - c) Frogs d) none of these.
- vi) The size of the core sequence of microsatellie is
  - a) 2-4 bp b) 5-50 bp
  - c) 10-100 bp d) 100-1000 bp.
- vii) ESI is
  - a) Electro Spray Ionisation
  - b) Electron Spark Ionization
  - c) Electron Spark Initiation
  - d) None of these.
- viii) Cellular proteomics is the branch of proteomics whose goal is
  - a) about the analysis of whole genome
  - b) to map the location of proteins and protein-protein interactions in whole cells during key cell events
  - c) about the transcriptional regulation control
  - d) none of these.
- ix) One disadvantage of metabolomics is
  - a) high risk of false positive data
  - b) highly qualified expertise required
  - c) unavailability of biomarkers
  - d) problem of collecting body fluid.

5059

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## x) Protein array includes interactions of

- a) protein ligand b) protein drug
- c) protein antibody d) all of these.
- xi) To identify the location of labelled proteins, the technique used is
  - a) chromatography b) FRET
  - c) NMR d) X-ray tomography.
- xii) Analyser used in Tandem mass can be
  - a) TOF b) magnetic analyzer
  - c) quadrupole d) all of these.

#### **GROUP – B**

### (Short Answer Type Questions)

	Answer any <i>three</i> of the following.	$3 \times 5 = 15$
2.	What is the full form of RAPD ? Describe how it as genetic fingerprint.	it can be used 1 + 4
3.	Describe the process of protein array.	5
4.	Write down the use of SNP mapping.	5
505	9 3	[ Turn over

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- 5. Describe any PCR based DNA typing technique. 5
- 6. Write a short note on mitochondrial DNA (*mt*DNA) analysis. 5

#### $\mathbf{GROUP}-\mathbf{C}$

#### (Long Answer Type Questions)

Answer any *three* of the following.  $3 \times 15 = 45$ 

- 7. Write short notes on the following :
  - a) STR
  - b) VNTR
  - c) Satellite DNA.
- 8. What is DNA typing ? Discuss the characteristic features of DNA polymorphisms. What is SNP ? Write down the limitations of metabolomics. What are the key applications of metabolomics ? 2 + 3 + 2 + 4 + 4
- 9. What is repetitive DNA ? How does it differ from unique DNA ? Classify repetitive DNA and describe each class with suitable example.
  2 + 2 + 4 + 7
- 10. Write the procedure of Transcriptome analysis. Point out the application of proteomics. What is nuclear proteome ? Name one database used in forensic science laboratory.

6 + 5 + 2 + 2

 Write down the significance of colors in obligo-nucleotide microarray. Define synonymous, missence, frame-shift SNPs. What are the limitations of metabolomics. What is biochemical network. Explain briefly.

5059