## 2013

# M.Sc.

## 3rd Semester Examination

## BIOMEDICAL LABORATORY SCIENCE AND MANAGEMENT

PAPER-BLM-301 (Unit-18)

Full Marks: 40

Time: 2 Hours

The figures in the right-hand margin indicate full marks.

Candidates are required to give their answers in their own words as far as practicable.

Illustrate the answers wherever necessary.

#### Module—I

# (CYTOTECHNOLOGY & CYTOGENETICS)

1. Answer any five of the following:

- 1×5
- (a) Write the full form of ABC technique,
- (b) What do you mean by primary antibody/monoclonal antibody in immuno cytotechnology?
- (c) Write the names of any two automated devices used in cytotechnology.
- (d) Write any one application of in-situ hybridization in cytotechnology.

- (e) Write the names of any two enzymes used in antibody labelling in cytotechnology.
- (f) Write the nature of image formed in compound microscope.
- (g) What do you mean by karyotyping?
- (h) Write the full form of Caspase.
- 2. (a) Write the fundamental principle of immuno enzymatic cytotechnology.
  - (b) State the fundamental steps adopted for the detection of specific marker by PAP technique in cytotechnology.
  - (c) Why immunofluorescence cyto technological method is preferred over immuno enzymatic cytotechnological method? 2+4+2

Or

- (a) Write the fundamental principle of real time PCR.
- (b) State the fundamental steps adopted in real time PCR for diagnosis of genetical disease in the field of cytotechnology.
- (c) Write the application of flow cytometry in cytotechnology. 2+4+2
- 3. (a) What is museum technology in cytology?
  - (b) Write the importances of museum technology in the field of cytotechnology.
  - (c) Write the fundamental principle adopted in in-situ hybridization. 2+3+2

- (a) State the basic steps followed for chromosome isolation in cytology.
- (b) Write the application of karyotyping in disease diagnosis.
- (c) Write the application of marker study in disease diagnosis. 2+3+2

#### Module-II

## (CYTOTECHNOLOGY)

- **4.** Answer any five questions of the following:  $1\times 5$ 
  - (a) Write the names of any two secondary fixatives.
  - (b) Write the objective of stropping.
  - (c) What do you mean by cryocut section?
  - (d) Write the names of any to dehydrating agents.
  - (e) What do you mean by controlled oxidizing agent?
  - (f) Write the application of 'Alcian-Blue' staining.
  - (g) Write forget molecules of PAS-stain.
  - (h) What do you mean by microwave technology in histology.

- 5. (a) Write the principle of PAS staining.
  - (b) State the basic steps followed for detection of biomolecule by PAS staining.
  - (c) How do you prepare Schiff reagent? 2+4+2

Or

- (a) Write the principle of Ninhydrine-Schiff method for detection of aminogroup in biomolecule.
- (b) State the routine steps followed in Ninhydrine-Schiff method in histology.
- (c) How is micro-organism in tissue section detected in histology? 2+4+2
- **6.** (a) When frozen section is preferred for diagnosis in histotechnology?
  - (b) Write the process of decalcification of bone tissue in histology.
  - (e) Write the objectives of clearing in tissue processing. 2+3+2

Or

- (a) Write the advantages of automated tissue processor over manual technique in histotechnology.
- (d) Write the procedure of Giemsa staining in histotechnology.
- (c) Why Millor reaction is applied in histotechnology. 2+3+2