Total Pages-3

## M.Sc.

# 2016

### 4th Semester Examination

#### **BIOMEDICAL LABORATORY SCIENCE AND MANAGEMENT**

#### PAPER-BLM-402

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.

Answer Q. No.1 and any three of the following.

1. Answer any ten questions of the following : 10×1

(a) What is selective media?

(b) Distinguish between solid media and liquid media.

(e) Write the principle of Dry heat mediated sterilization.

(d) Write the name of two basic stain.

(Turn Over)

- (e) What is chemotaxis?
- (f) How antibiotic solutions are sterilized?
- (g) What do you mean by mordant?
- (h) What is negative staining?
- (i) Define glycocalyx.
- (j) How bacterial endospore protect the genomic DNA?
- (k) Why bacterial population are constant in Lag phase?
- (l) Why petri plates should be inculated upside down in incubator?
- (m) Mention this role of Eosin methylene blue present in EMB agar?
- Describe in some detail the composition and structure of peptidoglycan, Gram positive cell wall and gram negative cell wall.
  5+2+3
- **3.** (a) Discuss the basic rules and regulations of Microbiology laboratory.
  - (b) Why mycobacterium is called as acid fast bacteria? Name one dye used to stain endospore. "Ampicillin (Amp) resistance gene in bacteria is extrachromosonal'— Justify. 5+(3+1+1)

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(Continued)

- 4. Write the different biochemical test for *E. Coli* and identification. 5+5
- 5. (a) Name the different generation of drugs used in urine culture.
  - (b) State the principle of a acidfast staining.
  - (c) Design an experiment for the enumeration of bacterial load in an infected urine sample along with their identification.
- 6. (a) You have given an unknown drug (DI). How will you check the efficacy of that particular dry on bacteria (both gram + ve and gram ve).
  - (b) How would you prepare a culture media for pathogenic bacteria. Write the identifying features of pathogenic bacteria. 5+3+2

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