### 2012

### M.Sc.

### 1st Semester Examination

#### BIOTECHNOLOGY

PAPER-BIT-103

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.

# Group-A

- 1. Answer any five questions from the following:  $5\times2$ 
  - (a) What are porins? What role do they play?
  - (b) Mention the molecular mechanism of flagellar movement of bacteria.
  - (c) State the function of KDPG aldolase.

- (d) What is Plaque count?
- (e) Why Com EA is important during transformation in Bacillus subtilis?
- (f) State the role of sterols in mycoplasma membrane.
- (g) What is growth rate constant?

### Group-B

Answer any two questions from the following:  $5\times2$ 

- 2. (a) Is there any difference between a capsule and a clime layer? What increases the virulence of anthrox bacillus? State the role of UDP-glucose in capsule formation in Streptococcus pneumoniae. 2+1+2
  - (b) What is transcription antitermination in  $\alpha$  phage? How the molecular switch in the Or region works? Write a short note on "Prophage". 1+2+2
  - (c) How do sulfa drugs act as anti bacterial agents?

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(d) What is ED Pathway? Mention its benefit in terms of bioenergetics and anabolic pathways. 2+3

## Group—C

## Answer any two questions.

- 3. What are the different forms F factor takes in a E. Colicell? State the importance of tra gene products during conjugation. What are nonconjugative, mobilizable plasmids?
  4+5+1
- **4.** How 5-hydroxymethylcytosine help  $T_4$  bacteriophage? What are metastable structures?  $T_4$  DNA molecule is terminally redundant explain. Write short notes on
  - (i) Introns in T<sub>4</sub>:
  - (ii) Regulation of  $T_4$  gene expression. 3+1+2+4
- 5. How bacteria show resistance to oxidising biocides? How concatemers in viral genomes are formed? With a suitable diagram describe different major viral structural arrangements.
  2+2+6
- **6.** Mention different ecological types of biological  $N_2$  fixer with suitable examples. Write the regulatory process and the major components involved in  $N_2$  fixation. How do different bacteria protect their nitrogenase from  $O_2$  during  $N_2$  fixation?