

# RAMAKRISHNA MISSION VIDYAMANDIRA

(Residential Autonomous College under University of Calcutta)

B.A./B.SC. FIRST SEMESTER EXAMINATION, DECEMBER 2011

FIRST YEAR

MICROBIOLOGY (Honours)

Date : 16/12/2011

Time : 11 am – 2 pm

Paper : I

Full Marks : 75

[Use separate Answer Books for each group]

## Group – A

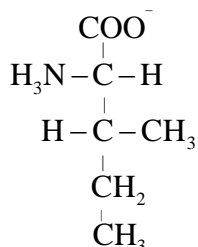
1. Answer **any five** questions : [2×5 = 10]
- What is pyrenoid?
  - Why is poly-β-hydroxybutyrate important for bacterial cell?
  - What do you mean by fruitbody?
  - How does rRNA homology help in modern bacterial systematics?
  - What is Tyndalization?
  - What is the limit of resolution of a compound light microscope?
  - What is the significance of calculating standard deviation in biological data?
2. Answer **any three** questions : [3×10 = 30]
- Describe with a ray diagram the working principle of a phase contrast microscope. [3]
    - What is the utility of oil immersion lens in microscopy? [2]
    - Differentiate between SEM and TEM [2]
    - “Basic dyes are often used in bacterial staining”— Justify. [2]
    - What is mycolic acid? [1]
  - Name one disease causing protozoa and mention the disease and also describe the life cycle of the mentioned protozoa. [2+4]
    - Write down the contribution of Robert Koch in the field of microbiology. [2]
    - Write the important differences between the Ascomycota and Basidiomycota. [2]
  - Name a bacterium devoid of cell wall. [1]
    - How does a capsule protect the bacterial cell? [2]
    - What is a spheroplast? [1]
    - Name an endospore forming bacterium. [1]
    - Describe the process of endospore formation. [4]
    - What is the role of pili in bacteria? [1]
  - What is hopanoid? Write down its two important functions. [2+2]
    - What is periplasm? [2]
    - Write four important functions of LPS in bacteria. [4]
  - What is histogram? [2]
    - What do you mean by coefficient of mean deviation? [2]
    - Find the mean and standard deviation of the following data. [6]

Number of bacteria	10	11	12	13	14	15	16
Number of colony	2	7	11	15	10	4	1

## Group-B

Answer **any five** questions from the following:

1. a) What are polyprotic acids? Explain with proper example. [2+1]  
b) How many milliliters of 5M H<sub>2</sub>SO<sub>4</sub> are required to make 1500ml. of a 0.002M H<sub>2</sub>SO<sub>4</sub> solution? [2]  
c) Calculate the pH of a buffer containing 0.1 mole acetic acid and 0.1 mole acetate ion.  
[K<sub>a</sub> = 1.8 × 10<sup>-5</sup>, -log 10<sup>-5</sup> = 5] [2]



2. a) How many chiral centres and optical isomers does it have? [2]  
b) Write short note on Butane Gauche interaction. [2]  
c) Mention the drawbacks of D, L nomenclature. [3]
3. a) Two amino acids are separated by thin layer chromatography and have *R<sub>f</sub>* values 0.6 and 0.5. A laboratory technician is asked to perform the same separation with different buffer system. Will he obtain the same *R<sub>f</sub>* values? Justify. [2]  
b) Super secondary structure is very important for protein stability — Justify. [2]  
c) Dansyl chloride treatment of a single polypeptide chain followed by its complete acid hydrolysis yields several dansylated amino acids. Explain. [2]  
d) Write down the structure of a biologically active tripeptide. [1]
4. a) Adenine – Thymine Watson – Crick base pairing. [2]  
b) Syn–Cytosine nucleoside. [2]  
c) What do you mean by C-value paradox? [2]  
d) Why is Melting Temperature (T<sub>m</sub>) of (G ≡ C) higher than (A = T)? [1]
5. a) Describe the preparations of 3 liters of 0.2M acetate buffer, pH 5.00, starting from solid sodium acetate trihydrate (MW 136) and a 1 M solution of acetic acid  
(Given, K<sub>a</sub> for acetic acid is 1.70 × 10<sup>-5</sup>) [3]  
b) Discuss the principle of "FORMOL" titration of glycine. [2]  
c) What happens when, peptide is treated with phenyl isothiocyanate? [2]
6. a) What do you mean by surface tension of water? Suggest a way to break it. [2+1]  
b) Define specific viscosity. Explain it's relation to relative viscosity. [1+2]  
c) What is the Helix Pitch for β-DNA? [1]
7. a) Point out the major differences between A, B and Z-DNAs. [3]  
b) Explain with suitable examples the role of H-bonding in stabilizing protein structure. Why are β-sheets pleated? [2+2]

