

2012**M.Sc.****3rd Semester Examination****MICROBIOLOGY****PAPER—MCB-304**

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 any two questions from each group.

Group — A

[Marks : 20]

1. (i) Classify fermentation process on the basis of cell mass production & product formation. 2
- (ii) Draw a schematic diagram of anaerobic batch STR. Mention the functions of its essential components. 3
- (iii) What is the basic difference between packed bed column reactor and fluidized bed reactor? 2
- (iv) Classify the fluids on the basis of Rheological Property. Define Renold's number and its significance. 3
2. (a) What are the various types of non-newtomian fluids? 2
- (b) How K_{1a} is measured by dyamic gassing out technique? 3
- (c) What is Leudeking-Piret model? 2

- (d) *Acetobacter aceti* are added to a vigorously aerated medium containing 10 g/l ethanol. After some time, ethanol concentration is 2 g/l and 7.5 g/l acetic acid is produced. How does the overall yield of acetic acid from ethanol compare with theoretical yield? 3
3. (i) What are the advantages and disadvantages of solid state fermentation process? What are the controlling parameters of this process? Write about the raw materials used in this process. 5
- (ii) What do you mean by scale up of a fermentor? What is the basis of scale up? Which parameters are involved in fermentas scale up? 3
- (iii) What do you mean by yield coefficient? Formulate different yield coefficient using cell mass (X), Product (P) and Substrete (S). 2

Group — B

[Marks : 20]

4. (a) Mention the steps of wine production through a flow diagram.
- (b) What do you mean by secondary wine fermentation? State its significance.
- (c) Why SO_2 is added in wine during Post fermentation treatment?
- (d) What is sparkling wine? How it is prepared? 4+3+1+2
5. (a) What is IPR? Discuss in details about the different forms of protection.
- (b) Write a note whole cell immobilization mentioning its advantages and disadvantages. (1+4)+5
6. (a) Mention the Bio-chemistry of citric acid production.
- (b) Name two centre where industrial strains are preserved.
- (c) Briefly describe the method commercial production of protease. 4+1+5