

L 1034

B.E./B.Tech. DEGREE EXAMINATION, MAY/JUNE 2006.

Fourth Semester

Bio-Technology

BT 1251 — BASIC INDUSTRIAL BIOTECHNOLOGY

(Regulation 2004)

Time : Three hours

Maximum : 100 marks

Answer ALL questions.

PART A — (10 × 2 = 20 marks)

1. Why are secondary metabolites referred to as “Biochemical Appendices”?
2. Give the scheme of a plant for the production of absolute alcohol.
3. Differentiate between liqueurs and spirits.
4. With the help of a flow chart show the production of citric acid by the precipitation method?
5. Define mutasynthesis.
6. Give an example of an aminoglycoside antibiotic, its application and a producing strain.
7. Name two methods by which enzyme productivity can be increased many-fold.
8. What is nisin and its application?
9. Give two methods by which L-glutamate production in C. glutamicum be increased.
10. Differentiate between “Callus” cultures and “Suspension” cultures.

PART B — (5 × 16 = 80 marks)

11. (i) Write a note on the commercial use of amino acids. (3)
- (ii) Discuss strategies to improve production of amino acids. (3)
- (iii) Give a scheme for the material flow in L-glutamate production plant and discuss the production process along with a scheme of the main reactions in L-glutamate formation from glucose. (10)

12. (a) How is citric acid produced? Give a flow chart for the manufacture of citric acid.

What are the applications of citric acid?

Or

- (b) (i) Write a note on PHA, PHB and biopol and its importance in medicine and commerce. (4)
- (ii) How is PHB metabolism regulated? (2)
- (iii) Discuss production of PHA by plants microbial sources. (6)
- (iv) Describe the 2-stage bioprocess for the production of PHA. (4)

13. (a) (i) What are the different types of reactors used in the biotech industry? (10)

- (ii) On what basis is the selection of a reactor made? (6)

Or

- (b) (i) Give a flow chart for the production of cheese. (6)
- (ii) Write a detailed note on the manufacture of cheese. (10)

14. (a) (i) How can the production of secondary metabolites be enhanced? (6)

- (ii) What is the process for production of β -lactam antibiotics and the recovery of the product? (10)

Or

- (b) How can you recover and isolate α -amylase from *Bacillus amyloliquefaciens*?

15. (a) (i) Outline the process for the cultivation of mammalian cells. (6)
- (ii) How are mammalian cells genetically engineered? (6)
- (iii) How can perfusion cultures be used in biomass production of mammalian cells? (4)

Or

- (b) Discuss a strategy for the production of biopesticides from Bacillus sphaericus.