

**BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI
(END SEMESTER EXAMINATION)**

CLASS: M.Sc
BRANCH: BIOTECHNOLOGY

SEMESTER : II
SESSION : SP/2025

SUBJECT: BT416 ENZYME & BIOPROCESS TECHNOLOGY

TIME: 3 Hours

FULL MARKS: 50

INSTRUCTIONS:

1. The question paper contains 5 questions each of 10 marks and total 50 marks.
 2. Attempt all questions.
 3. The missing data, if any, may be assumed suitably.
 4. Before attempting the question paper, be sure that you have got the correct question paper.
 5. Tables/Data hand book/Graph paper etc. to be supplied to the candidates in the examination hall.
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- Q.1(a) What do you mean by growth of microbes? Design a method to estimate microbial biomass present in a 10000L bioreactor. [5] CO 1 BL 6

- Q.1(b) Calculate specific growth rate (μ) of *Escherichia coli* for its growth in batch bioreactor. Data from batch culture is provided below: [5] 1 4

Time of sample collection (h)	0	2	4	6	8	10	12
Produced Biomass (g/L)	0.006	0.018	0.126	0.542	1.148	2.868	3.152
Residual sugar (g/L)	10.000	9.650	8.310	7.743	5.267	2.434	0.032

- Q.2(a) Calculate V_{max} and K_m for an enzyme catalyzed reaction. Given $[E_0] = 0.025$ g/l. [5] 2 4

V_o (g/l-min)	0.18	0.48	0.66	0.78	0.93	0.98	1.32	1.56
S_o (g/l)	1.0	2.0	3.0	5.0	8.0	12	20	30

- Q.2(b) What do you mean by enzyme immobilization? Fungal derived α - amylase is purified from the culture broth. Elaborate encapsulation method in detail for its immobilization. Describe its merit, demerits and detailed methodology. [5] 2 3

- Q.3(a) What do mean by sterilization of medium in fermentation industry? Derive an expression to show the death kinetics in batch autoclave. [5] 3 3

- Q.3(b) Draw a schematic diagram for a system used for continuous sterilization by using two heat exchangers. Also, relate it with its methodology of sterilization. [5] 3 3

- Q.4(a) Design a system for aeration and agitation for a tank bioreactor, level the diagram and explain the process of aseptic aeration and agitation. [5] 3 3

- Q.4(b) What is volumetric oxygen transfer coefficient? Describe the dynamic method to calculate it. [5] 3 3

- Q.5(a) Draw the structure of a chemostat, level it and describe its working. [5] 3 2

- Q.5(b) Derive the expression to show that 'for production of primary metabolites, continuous culture is better'. [5] 3 3