

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

**CLASS: BE
BRANCH: BIOTECHNOLOGY**

**SEMESTER : V
SESSION : MO/19**

SUBJECT: BT5021 BIOPROCESS ENGINEERING

TIME: 3 HOURS

FULL MARKS: 60

INSTRUCTIONS:

1. The question paper contains 7 questions each of 12 marks and total 84 marks.
 2. Candidates may attempt any 5 questions maximum of 60 marks.
 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 is balanced growth? [2]
- Q.1(b) Discuss the effect of substrate concentration on specific growth rate of a bacterial culture Growing in a chemostat. [4]
- Q.1(c) Discuss the model used to describe heat generation by bacteria in a batch culture. [6]
- Q.2(a) Differentiate between yield and productivity. [2]
- Q.2(b) Show that mixed growth associated product formation follows Leudeking-piret equation. [4]
- Q.2(c) Production of single cell protein from hexadecane in presence of ammonia was done as follow. If, RQ=0.43, determine stoichiometric coefficients. [6]
- $$C_{16}H_{34} + aO_2 + bNH_3 \rightleftharpoons cCH_{1.66}O_{0.27}N_{0.20} + dCO_2 + eH_2O$$
- Q.3(a) What do you mean by enzymes? Write the role of enzymes in food sector. [2]
- Q.3(b) Discuss the immobilization of enzyme by membrane entrapment methods. [4]
- Q.3(c) Derive an expression for rate of reaction (V_o) for single substrate enzyme catalysed reaction. [6]
- Q.4(a) What is decimal reduction time with respect to medium sterilization? [2]
- Q.4(b) Design a continuous sterilization system for medium using heat exchangers. [4]
- Q.4(c) What do you mean by 'Del' factor? Calculate the holding time at 121 °C in an batch autoclave (Initial and final concentration of bacteria were $1 \times 10^{14} L^{-1}$ and $1 \times 10^{-3} L^{-1}$, respectively). The value of sp. death rate, $Del_{heating}$ and $Del_{cooling}$ were recorded as $2.54 min^{-1}$, 9.8 and 10.1, respectively. [6]
- Q.5(a) Describe the features of Rushton turbine. [2]
- Q.5(b) Discuss the power requirements for a CSTR in ungasged condition. [4]
- Q.5(c) Calculate the $K_L a$ value from the given data of bacterial culture (2.5 g/l) in a CSTR. (Given that $C^*=7.1 mg/l$) [6]

DO (mg/l)	4.3	4.3	4.3	2.8	2.5	1.8	2.4	3.2	4.0	4.1	4.2
Time (min)	0	1	2	3	4	5	6	7	8	9	10
Air (on/off)	on	on	off	off	off	on	on	on	on	on	on

- Q.6(a) Differentiate between batch and continuous mode of CSTR operations. [2]
- Q.6(b) How will you change a batch CSTR into continuous CSTR ? Write your strategy. [4]
- Q.6(c) 'Most of the commercial fermenters are batch fermenter', justify the statement. [6]
- Q.7(a) Give a labelled diagram of bubble column reactor. How it works? [2]
- Q.7(b) Design a system for operation of a CSTR as chemostat. [4]
- Q.7(c) Describe the various methods for setup of fed batch bioreactor. Mention method of monitoring of bioreactor during its use. [6]

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