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Seat N	0.:	.: Enrolment No CULARAT TECHNOLOCICAL UNIVERSITY	
		BE - SEMESTER-VI(NEW) – EXAMINATION – SUMMER 2019	
Subject Code:2160508 Date:1			/2019
Subie	ect Na	ame:Biochemical Engineering	
Time	:10:3	0 AM TO 01:00 PM Total Mar	ks: 70
Instruc	ctions:		
	1. A 2. U 3. P 4. F	Attempt all questions. Inderstand the question carefully and write in proportion to marks it carri Provide explanation wherever required. Figures to the right indicate full marks.	ies.
Q.1	(a)	State three differences between a chemical reaction and its	03
	(b)	biochemical counterpart.	04
	(U)	phase is the longest?	04
	(c)	Discuss with examples the application of fermentation technology in	07
		food and beverage, industrial solvents, and antibiotics production.	
Q.2	(a)	Name three industrially important biochemical reactions and mention	03
		the name(s) of the microorganism involved therein.	0.4
	(D)	State and explain different types of solid and liquid media used for microbial growth	04
	(c)	Discuss various methods of cell disruption to get intracellular	07
	(•)	products. Give two examples of such intracellular products.	
		OR	
	(c)	State and briefly explain various steps, either sequential or	07
		concurrent, of an integrated bioprocess in general.	
03	(9)	State three methods for the measurement of microbial growth	03
Q.J	(b)	Write down Monod equation of microbial growth kinetics. Explain	03
		various terms in the same equation. How do you determine the kinetic	-
		parameters of the above equation graphically?	
	(c)	Using substrate and biomass balance derive the design equation for a	07
		continuous stirred tank system for blochemical process. State the	
		OR	
Q.3	(a)	What are the temperature, pressure and time conditions for steam	03
-		sterilization? Name a few methods for sterilization of medium other	
		than using steam.	
	(b)	Calculate the steady state substrate and biomass concentration in a	04
		continuous fermenter which has an operating volume of 25 L. When the starile feed stream contains limiting substrate at 2000 mg/L and	
		enters the vessel at 81 /h. The values of $K_{\rm a}$ and $\mu_{\rm m}$ are 10.5 mg/L and	
		0.45 h^{-1} respectively and the yield coefficient may be taken as 0.48.	
	(c)	Illustrate various probable configurations of fed-batch bioreactors	07
		with schematic diagrams.	

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- 0.4 (a) Write down the differences between prokaryotic and eukaryotic cells 03 with examples.
 - (b) How does an enzyme work? Classify enzymes with examples. 04
 - (c) Draw a schematic diagram of a fermenter, label its different parts and 07 state the functions of major parts.

OR

- (a) What are different types of controlling system in fermenter? **O.4**. 03 04
 - (b) Write a note on biosensors for fermentation control.
 - (c) Discuss various monitoring and control parameters of an ideal 07 fermenter.
- (a) Explain why oxygen needs to be supplied at a sufficient rate during Q.5 03 aerobic fermentation.
 - (b) Explain various physico-chemical parameters affecting volumetric 04 mass transfer coefficient K_La.
 - (c) State various methods of determination of volumetric mass transfer 07 coefficient K_La in a fermenter and explain any one.

OR

- (a) Name various unit operations used in upstream and down stream Q.5 03 processing in fermentation.
 - (b) Discuss briefly the rheological behaviour of fermentation broth with 04 appropriate examples.
 - n methe (c) Explain differential centrifugation method for separation of cells 07 from fermentation broth.