



### IISEMESTER M.TECH (INDUSTRIAL BIOTECHNOLOGY) END SEMESTER EXAMINATIONS, JUNE 2022 (REGULAR)

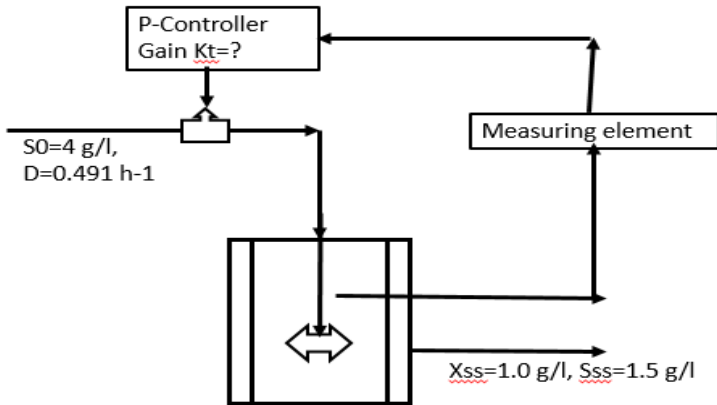
#### BIO5254: BIOREACTOR DESIGN AND ANALYSIS

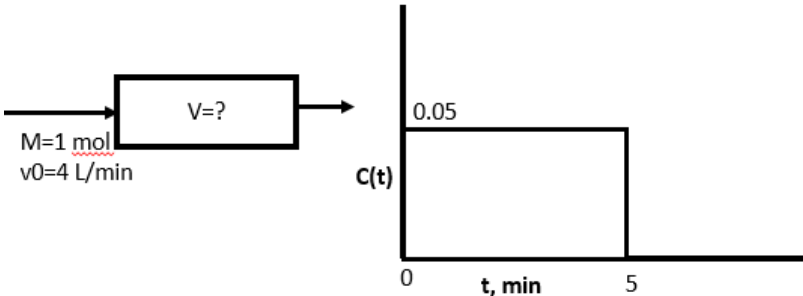
#### REVISED CREDIT SYSTEM ANSWER ALL QUESTIONS

Time: 3 Hours

MAX. MARKS: 50

SL NO	Question	CO	BTL	M
1A.	How do you define the reaction rate in heterogeneous reaction system? Explain various parameters affecting the reaction rate in heterogeneous reaction system.	1	3	4
1B.	Write on the various dimensionless parameters that are used in estimating (i) External Mass transfer effects (ii) Internal mass transfer effects	1	3	3
1C	Hydrolysis of palm oil is carried out in a CSTR with free lipase enzyme. The substrate sucrose ( $S_0=4M$ ) is pumped at 0.1 L/h. Find the volume of the reactor to achieve 45% conversion at steady state for (a) Michaelis-Menten Kinetics (b) Substrate Inhibition Kinetics. Kinetics data: $V_m=0.028$ M/min, $K_m=0.23$ M, $K_i=0.2$ M	2	3	3
2A.	What do you mean by constant feed rate policy in the operation of immobilized enzyme reactor system? Develop a suitable model for predicting the time course profiles of conversion due to enzyme deactivation for first order kinetics, no pore diffusion effects in PFR with immobilized enzyme.	2	3	4
2B.	Consider a 1000 L CSTR in which biomass is being produced with glucose as the substrate. The microbial system follows a Monod relationship with $\mu_m = 0.4$ h <sup>-1</sup> , $K_s = 1.5$ g/l and the yield factor $Y_{X/S} = 0.5$ g biomass/g substrate consumed. If the normal operation is with a sterile feed containing 10 g/l glucose at a rate of 100 L/h: i. What is the specific biomass production rate (g/L-h) at steady state? ii. If recycle is used with a recycle stream of 10 L/h and a recycle biomass concentration five times as large as that in the reactor exit, what would be the new specific biomass production rate?	2	3	4
2C	When do you prefer the Fed-batch reactor? Explain how quasi steady state is justified in Fed-batch reactor.	2	3	2
3A.	Differentiate between controllability and stabilizability of the bioprocess? Explain various steps involved in designing of controller for bioprocess.	4	3	4
3B	Consider that you have been working with Chemostat system where cell growth follows the substrate inhibition kinetics. Write the conditions for obtaining positive and negative real parts of Eigen values.	3	3	2

3C	<p>Certain bio-product is produced using <i>Bacillus</i> species in a cascade fermentation system of three chemostats in series with volume, <math>V_1=167</math> L, <math>V_2=40</math> L, <math>V_3=67</math> L. The feed flow rate is 100 L/h. Graphically determine the steady state biomass concentration in above all three chemostats. The growth kinetic data for <i>Bacillus</i> species is as follows:</p> <table><tr><td>X, g/l</td><td>0</td><td>0.2</td><td>0.3</td><td>0.4</td><td>0.55</td><td>0.60</td><td>0.70</td><td>0.80</td><td>0.90</td></tr><tr><td>dX/dt, g/l-h</td><td>0</td><td>0.1</td><td>0.2</td><td>0.3</td><td>0.38</td><td>0.36</td><td>0.3</td><td>0.2</td><td>0.05</td></tr></table>	X, g/l	0	0.2	0.3	0.4	0.55	0.60	0.70	0.80	0.90	dX/dt, g/l-h	0	0.1	0.2	0.3	0.38	0.36	0.3	0.2	0.05	2	3	4
X, g/l	0	0.2	0.3	0.4	0.55	0.60	0.70	0.80	0.90															
dX/dt, g/l-h	0	0.1	0.2	0.3	0.38	0.36	0.3	0.2	0.05															
4A.	<p>Lactic acid is produced using <i>Lactobacillus</i> in a Chemostat under sterile environment with glucose as the substrate, <math>S_0=4</math> g/l at dilution rate of <math>D=0.18</math> <math>h^{-1}</math>. Steady state substrate and biomass concentrations are 1.5 and 1.0 g/l respectively. Assume that growth follows the Substrate inhibition kinetics given by the following equation with, <math>\mu_m=0.53</math> <math>h^{-1}</math>, <math>K_s=0.12</math> g/l, <math>K_i=0.8</math> g/l and <math>Y=0.4</math></p> $\mu = \frac{\mu_m \cdot S}{(K_s + S + \frac{S^2}{KI})}$ <p>i. Find the elements of A matrix ii. Find the Eigen values and comment on the stability of the system</p>	3	4	4																				
4B.	<p>Proponoic acid is produced using <i>Bacillus licheniformis</i> in a Chemostat under sterile environment with glucose as the substrate, <math>S_0=4</math> g/l at dilution rate of <math>D=0.49</math> <math>h^{-1}</math>. Steady state substrate and biomass concentrations are 1.5 and 1.0 g/l respectively. Assume that growth follows the Monod's kinetics with, <math>\mu_m=0.53</math> <math>h^{-1}</math>, <math>K_s=0.12</math> g/l and <math>Y=0.4</math>. It is desired to control the biomass at steady state value by manipulating the dilution rate of feed.</p> <div><p style="text-align: center;"><u>Turbidostat operation of continuous fermenter</u></p><p>i. Find the <math>[A-BK_t]</math> matrix for the above process ii. Find the suitable <math>K_t</math> value that stabilizes the process in the case of disturbances</p></div>	4	4	4																				
4C	Name the Input, output and manipulated variable in Activated sludge process.	4	3	2																				
5A.	RTD results for the non-ideal bioreactor are shown in the following table for pulse input. Find the conversion for macro fluid with kinetics $(-r_A)=k$ , $k=0.5$ , $CA_0=10$ M	5	3	3																				

	<table><tr><td>Time, min</td><td>0</td><td>5</td><td>10</td><td>15</td><td>20</td><td>25</td><td>30</td><td>35</td></tr><tr><td>E (t)</td><td>0</td><td>0.03</td><td>0.05</td><td>0.05</td><td>0.04</td><td>0.02</td><td>0.01</td><td>0</td></tr></table>	Time, min	0	5	10	15	20	25	30	35	E (t)	0	0.03	0.05	0.05	0.04	0.02	0.01	0			
Time, min	0	5	10	15	20	25	30	35														
E (t)	0	0.03	0.05	0.05	0.04	0.02	0.01	0														
5B.	<p>A pulse input to a non-ideal bioreactor gives the results shown in the following fig.</p> <p>(a) Check the material balance with the tracer curve to see whether the results are consistent.</p> <p>(b) If the result is consistent, determine space time (<math>\tau</math>), Volume of bioreactor (V) and sketch the E curve.</p> <div><p>The diagram shows a rectangular bioreactor with an input arrow on the left and an output arrow on the right. The input is labeled with <math>M=1 \text{ mol}</math> and <math>v_0=4 \text{ L/min}</math>. Inside the reactor, it says <math>V=?</math>. To the right of the reactor is a graph of concentration <math>c(t)</math> versus time <math>t, \text{ min}</math>. The graph shows a rectangular pulse that starts at <math>t=0</math>, has a constant value of <math>0.05</math> until <math>t=5</math>, and then drops to zero.</p></div>	5	3	4																		
5C	Write on the adverse effects of Foaming in the operation of bioreactor. Explain how foaming is controlled in Industrial fermenters.	2	3	3																		