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		SEM: V - THEORY EXAMINA	ΤΙΟΝ (20	24 - 20	25)					
		Subject: Bioprocess En	ngineerin	g						
		Hours					Max	. M	arks	: 100
		structions:	م ماه ماه م	4				1	ala	a 4 a
		ly that you have received the question paper westion paper comprises of three Sections -A,								etc.
		(MCQ's) & Subjective type questions.	<i>D</i> , α c. <i>I</i>	i consis	is oj	WI U	шрте	Cn	oice	
_		m marks for each question are indicated on i	right -har	ıd side	of ea	ch q	juesti	ion.		
		e your answers with neat sketches wherever	_		Ü		•			
		suitable data if necessary.								
•		bly, write the answers in sequential order.					_			
		t should be left blank. Any written material a	fter a bla	nk shee	et wil	l no	t be			
evalud	itea/ci	checked.								
SECT	ION-	J-A								20
		all parts:-								
1-a.	_	What is true about aerobic bacteria? (CO1, K	1)		J'					1
	(a)	fluorish in the presence of free oxygen								
	(b)	consume organic matter as their food								
	(c)	oxidise organic matter in sewage								
	(d)	All of the above								
1-b.	` ′	The organic material of the solid waste will d	lecompos	e (CO1	K 1)				1
1 0.	(a)	By the flow of water	iccompos	c (CO1	, 111	,				1
	(a) (b)	By the soil particles								
	(c)	By the action of microorganisms								
	(d)	By oxidation								
1-c.	` ′	Which of the following is incorrect for a cata	lvst? (CC)2 K2)						1
1 0.	(a)	Bio-chemical reactions are mostly catalyz	•							1
	(b)	Catalyst does not start a reaction	ica by chi	Lymes						
	(c)	Catalyst changes the equilibrium constant	of a reac	tion						
	(d)	Co-enzymes increase the activity of an en		uon						
1 .1	` ′		· ·	1		1-19	(CO)) IZ	1)	1
1-d.		Which of the following is INCORRECT for t	ле юск-а	ша-кеу	шос	161 ?	(CU ₂	۷, K	1)	1
	(a)	It is used to describe the binding process		.1 1						
	(b)	The active site of the enzyme is complementation of the enzyme is complementation.	•	the sub	strat	e				
	(c)	It demonstrates enzyme-substrate complex	V							

	. 1 \		
	(d)	The binding of the substrate produces a conformational change in enzyme	
1-e.		rom the following volumes, which capacity the Pilot-scale bioreactor holds? CO3, K1)	1
	(a)	100-1000L	
	(b)	1000-10000L	
	(c)	1-100L	
	(d)	less than 1L	
1-f.	W	which of the following physicochemical factor does not affect SSF? (CO3, K1)	1
	(a)	Pressure	
	(b)	Temperature	
	(c)	рН	
	(d)	Moisture content	
1-g.		he production of bio ethanol is by fermenting the and starch omponents. (CO4, K1)	1
	(a)	Acid	
	(b)	Milk	
	(c)	Sugar	
	(d)	Alcohol	
1-h.	T	o make transport fuel the bio ethanol is blended with(CO4, K2)	1
	(a)	Diesel	
	(b)	Petrol	
	(c)	oil	
	(d)	kerosene	
1-i.		o maintain aseptic conditions during fermentation which of the following is eeded? (CO5, K2)	1
	(a)	Sterilization of fermentor	
	(b)	Sterilization of air supply	
	(c)	Aeration and agitation	
	(d)	All of these	
1-j.	in	onionizing radiation and ionizing radiation are sterilization methods mainly used hospitals. Ultraviolet radiation is one example of nonionizing radiation, name le ionizing radiation? (CO5, K2)	1
	(a)	Infrared	
	(b)	X-rays and gamma rays	
	(c)	Halogens	
	(d)	Ethylene oxide	
2. Att	empt a	all parts:-	
2.a.	-	That is the major difference between nephelometry and turbidimetry? (CO1, K2)	2

2.b.	How enzyme catalyze the biochemical reaction? (CO2, K2)	2
2.c.	Write any four applications of solid-state fermentation? (CO3, K1)	2
2.d.	Write the name of all three scientist who won the nobel prize for penicillin discovery? (CO4, K1)	2
2.e.	Why thermocouples are most widely used as temperature sensors? (CO5, K2)	2
SECT	ION-B	30
3. Ansv	wer any <u>five</u> of the following:-	
3-a.	Discuss in detail about the indirect methods of determining cell number density? (CO1, K2)	6
3-b.	Dissolved oxygen is an important substrate in aerobic fermentation. Justify this statement? (CO1, K2)	6
3-c.	Explain briefly about the batch operation of a mixed reactor? (CO2, K2)	6
3-d.	Discuss in detail about the chemostat with immobilized cells in a bioreactor? (CO2, K2)	6
3.e.	Discuss about the scale-up process of a bioreactor? (CO3, K3)	6
3.f.	Write any seven applications of bioprocess engineering? (CO4, K1)	ϵ
3.g.	Discuss in detail about the common instruments used for process automation? (CO5, K3)	6
	ION-C	50
	wer any <u>one</u> of the following:-	
4-a.	Explain microbial growth curve with its different phases along with a suitable graph? (CO1, K3)	10
4-b.	Illustrate the working mechanism of particle counter in detail? (CO1, K2)	10
5. Ansv	wer any <u>one</u> of the following:-	
5-a.	Explain in detail about the different ways by which cells can be recycled in a fermentation process? (CO2, K3)	10
5-b.	Write the advantages of chemostat cascade and chemostat recycle with flow sheet? (CO2, K2)	10
6. Ansv	wer any <u>one</u> of the following:-	
6-a.	With the help of labelled diagram, explain the steps for oxygen transfer from gas bubble to individual cell? (CO3, K2)	10
6-b.	A fermentation broth with viscosity 10^{-3} Pa s and density 1000 kg m ⁻³ is agitated in a 100 m ³ baffled tank using a marine propeller 1.2 m in diameter. Calculate the power required for a stirrer speed of $5s^{-1}$. (Take k1 as 40 and Np ' as 0.35 for marine propeller) (CO3, K3)	10
7. Ansv	wer any <u>one</u> of the following:-	
7-a.	With the help of schematic diagram, discuss about the different stages of bioprocess development? (CO4, K1)	10
7-b.	Explain briefly about the downstream process of antibiotic production? (CO4, K2)	10

- 8. Answer any one of the following:-
- 8-a. What is sterilization? Why is there the need of sterilization? What happens if the fermentation medium gets contaminated? (CO5, K3)
- 8-b. Explain briefly about the factorial design and Plackett-Burman design for medium optimization? (CO5, K2)

