75 MARKS 2 ½ HRS VCD/ SYBT SEM III BIOPROCESS TECHNOLOGY

NOTE:

- 1. Attempt all questions.
- 2. All questions carry equal marks.
- 3. Draw neat labeled diagrams wherever necessary.
- 4. For Q 2, Q 3 and Q 4 attempt A and B OR C and D.

Q 1	Do as directed (Any fifteen) State true or false. Enrichment culture technique is a secondary screening
	technique.
2.	Name any one Lactic Acid bacteria.
3.	Name any one of the strain improvement techniques.
4.	Media containing starch can be used for the screening ofenzyme
	producers.
5.	Name the medium used for the preservation of Clostridium spp. by serial
	subculture.
6.	oil is used for preservation by overlaying cultures.
7.	Name any one desirable property used for the screening of yeast for alcohol
	production.
8.	State the importance of sampling during fermentation.
9.	may be used for continuous sterilization.
10.	Adding feed and removal of spent broth at the same rate is a feature of
	fermentation.
11.	Give significance of aeration in a fermentor.
12.	Feed is added without removal of spent broth intype of
	fermentation.
13.	Solid state fermentation is preferably used for the cultivation of
14.	Name any one downstream production technique.
15.	Write any one disadvantage of bioassay methods.
16.	What is the full form of AUC?
17.	Write the formula used to calculate Rf value.
18.	Detection of amino acids separated by paper chromatography can be done by
ja Na	usingreagent.
19.	What are the different routes of drug administration?
20.	PCV is used for the determination ofin fermentation process.

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Q 2	A Discuss the role of fungi in different industrial production processes with suitable examples.	h 08
Q 2	B Write a note on primary screening of antibiotic producers from soil.	07
Q 2	C Describe lyophilisation as a preservation technique.	
Q 2	D Give significance of sugar and electrolity.	08
	Give significance of sugar and alcohol tolerance of yeast for ethanol production.	07
Q 3 A	and decount of collinions starilization	
Q 3 E	Give an account of fermentative production of all	08
	Give an account of fermentative production of ethanol using a flow-sheet. OR	07
Q3C	Discuss the different components of familiary	
Q 3 D	Describe pH and Temperature as fermentation process parameters.	08
	process parameters.	07
Q 4 A Q 4 B	time profile for a hypothetical drug administered extra-venously. What are physical-chemical and biological transfer to the serum concentration-	08
	op methods advantages and disadvantages.	07
Q4C Q4D	What is a biological assay? State its advantages and disadvantages and write	08
	Write a short note on turbidity analysis and cell yield determination.	07
Q 5	Write a short note on any three of the following	
a	Auxanography.	15
b	Turbidostat type of continuous fermentation.	
c	Maintenance of aseptic conditions in fermentation.	
d	Titrimetric and graving at	
	Titrimetric and gravimetric analysis.	
e	Half-life determination of a drug.	