Registration No.:					

Total number of printed pages - 2

B. Tech

**PCBT 4307** 

NTRAL LIB

## Sixth Semester Regular Examination – 2014 INDUSTRIAL MICROBIOLOGY AND ENZYME TECHNOLOGY

**BRANCH: BIOTECH** 

QUESTION CODE: F 267

Full Marks - 70

Time: 3 Hours

Answer Question No. 1 which is compulsory and any five from the rest.

The figures in the right-hand margin indicate marks.

1. Answer the following questions:

2×10

- (a) Name four different kinds of bioreactors used for industrial production of antibiotics and aminoacids.
- (b) Define solid state fermenttion. What is its significance?
- (c) How Leloir pathway is advantageous for synthesis of oligosaccharides?
- (d) Define the term "de-controlled mutant"? Why it is siginificant in commercial strain development?
- (e) Define the efficiency of microbial transformation. Deduce the mathematical expression for it.
- (f) How the stability of lysozyme is increased?
- (g) Differentiate between feed back inhibition and feed back repression using flow diagram.
- (h) What do you mean by diffusion coefficient of immobilized enzyme? Why it is significant?
- (i) Why a lactose deficit mutant strain of bacteria may grow in medium supplemented with glucose?
- (j) How the protein engineering of protease play significant role in detergent industry?

2.	Wh	at do you mean by enzyme stability? Briefly explain the approac	ches of					
	enz	zyme engineering increase the enzyme stability across temperature a	nd pH.					
3.	(2)	Describe the unstream process for elechal productions	3+7					
٥.	(a)	Describe the upstream process for alcohol production.	5					
	(b)	Define Enzyme engineering. Draw the flow sheet for the product recombinant Taq polymerase using r-DNA technology.	ction of 5					
4.	Differentiate between: 2.5 ×							
	(a)	Enzymation and Fermentation						
	(b)	Entrapment and Cross linking						
	(C)	Membrane reactor and Column reactor						
	(d)	Broad spectrum and Narrow spectrum antibiotics.						
5.		efly explain the application of biocatalysts in biotransformation involving biochemical reactions:	volving 5×2					
	(a)	Group transfer redox reaction						
	(b)	Elimination of C-C, C-O and C-N bond by cleavage.						
6.	Wha	at do you mean by strain of bacteria? Briefly explain the strategies us	sed for					
	dev	elopment of commercially important strain with suitable examples.	2+8					
7.	Writ	te notes on :	5×2					
	(a)	Site directed mutagenesis						
	(b)	Purification of heterologous proteins.						
8.	(a)	What is the significance of reaction environment for industrial production various biomolecules? Explain.	ction of 5					
	(b)	Write short notes on 'kinetics of continuous culture'?	5					