Reg	istra	tion No.:
Tota	al nur	mber of printed pages – 2  B. Tech
		PCBT 4307 (New)
		Sixth Semester (Back) Examination – 2013
	IND	USTRIAL MICROBIOLOGY AND ENZYME TECHNOLOGY
		BRANCH: BIOTECH
		QUESTION CODE: B268
		Full Marks – 70 Time: 3 Hours
/	Answ	er Question No. 1 which is compulsory and any five from the rest.
		The figures in the right-hand margin indicate marks.
1.	Ans	wer the following questions: 2×10
	(a)	Surface culture method.
	(b)	Secondary metabolites
	(c)	Auxotrophic mutant
	(d)	Enzyme encapsulation
	(e)	Batch fermentation
	(f)	Differentiate between Native PAGE and SDS-PAGE.
	(g)	β-lactam ring
	(h)	Enzyme immobilization
	(i)	Define catabolic repression.
	(j)	Broad spectrum antibiotics

Briefly describe industrial methods for citric acid production.

What is solid state fermentation? Discuss its significance in large scale

2.

(a)

(b)

operation.

5×2

3.	(a	Derive Michaelis and Menten kinetics for enzyme reaction.	5×2		
	(b	) Differentiate between broad spectrum and Narrow spectrum antibiot	ics.		
4.	(a	) Describe different methods of enzyme immobilization.	5×2		
	(b)	What are the applications of enzymes in medicinal therapy?			
5.	(a)	Differentiate between entrapment and cross linking.	5×2		
	(b)	What are the advantages and disadvantages of immobilized enzymes	s?		
6.	De	Describe the process of antibiotic production studied by you. Which type of bioreactor is used for antibiotic production.			
7.	Wri	ite down short notes on any <b>two</b> of the following :	5×2		
	(a)	Enzyme stabilization			
	(b)	Protein engineering and its significance			
	(c)	Site-directed mutagenesis in protein engineering.			
8.	Writ	Write down short notes on any two of the following:			
	(a)	Strain improvement			
	(b)	Overproduction of decontrolled mutant			
	(c)	Submerged culture.			