## SINHAGAD – II (2007 COURSE): JULY - 2013 SUBJECT: PHARMACEUTICAL BIOTECHNOLOGY - III

Day: Date:	Fri 05/	day 07/2013 Time: 10:00 A.M. TO Max. Marks: 100			
N.B.:	1)	Question No. 1 and 5 are <b>COMPULSORY</b> . Out of the remaining attempt Any <b>TWO</b> questions from Section – I and <b>TWO</b> questions from Section – II.			
	<ol> <li>Figures to the right indicate full marks.</li> <li>Answer to the two sections should be written in SEPARATE answer books.</li> <li>Neat diagrams wherever necessary.</li> </ol>				
		SECTION-I			
Q.1		What are recombinant vaccines? Discuss their advantages.	(10)		
Q.2		Explain key steps in production of monoclonal antibodies. Write an account of its application as diagnostic tool.	(20)		
Q.3		What are viral vectors? Discuss applications in gene delivery with suitable examples.	(20)		
Q.4	a) b) c) d) e)	Write note on any <b>TWO</b> of the following: ISCOMS Saponins as adjuvant MHC complex Disadvantages of DNA vaccines synthetic peptides	(20)		
		SECTION-II			
Q.5		What is personalized medicine? Add a note on Cytochrome P450.	(10)		
Q.6	a)	Explain what you learned from Human genome project.	(20)		
	b)	How would it influence future of drug discovery.			
Q.7		Elaborate applications of Bioinformatics in pharmaceutical research.	(20)		
Q.8	a) b) c) d) e)	Small interfering RNA	(20)		

## SINHAGAD-II (CBCS – 2012 COURSE): JULY- 2013 SUBJECT: ADVANCE CORE SUBJECT-II: ADVANCED PHARMACEUTICAL BIOTECHNOLOGY-II

Day Date	: Wednesday : 03-07-2013	Time: 10:00 A.M.TO 1:0 Max. Marks: 60	00 P
N.B.	<ol> <li>Attempt any THREE questions from Section-II.</li> <li>Both the sections should be written in SEP.</li> <li>Figures to the RIGHT indicate full marks.</li> </ol>		from
	SECTION-I		
Q.1	Describe in details MHC complex and the propresented on cell surface.	cess with which antigens are	(10)
Q.2	Differentiate monoclonal and polyclonal antib monoclonal antibodies.	odies. Write applications of	(10)
Q.3	Describe different cell types involved in immune res	ponse.	(10)
Q.4	Write elaborate notes on:  a) ELISA b) Adjuvants		(10)
	SECTION-II		
Q.5	What is site directed mutagenesis? Describe oligo b site.	eased method to mutate a target	(10)
Q.6	How enzymes are immobilized? Discuss various me	thods giving their advantages.	(10)
Q.7	Outline a down-stream process for recovery of recovery involved.	mbinant protein describing key	(10)
Q.8	Write elaborate notes on: a) Bioreactor b) Enzyme inhibitor		(10)