PURUS – VI (2011 COURSE) : SUMMER – 2016 SUBJECT : MEDICINAL CHEMISTRY – II

Day Date	: 5	SUBJECT: MEDICINAL CHEMISTRY - II aturday Time: 10.000 Max. Marks: 80	AMT
	2	73-04-2016 Max. Marks : 80)
N.B.	1)	Q.1 and Q.5 are COMPULSORY. Out of the remaining, attempt any TWO)
	2)	questions from Section – I and any TWO questions from Section – II.	
	3)	Figures to the right indicate FULL marks. Draw diagrams, structures and give reactions wherever necessary.	
		SECTION – I	
Q.1		Attempt any FIVE questions of the following:	(10)
	a)	Define the terms analeptics and hallucinogens with one example.	
	b)	Draw structures of any two long acting barbiturates.	
	c)	Explain the term status epilepticus and suggest the therapy for the same.	
	d)	Explain the terms REM and NREM giving their full forms.	
	e)	Outline synthesis of Ketamine hydrochloride giving its category.	
	f)	What is the role of Phase – I and Phase – II metabolic pathways?	
	g)	Draw the basic molecular frame work of benzodiazepines and succinimides.	
Q.2	a)	Elaborate in details oxidative pathways with examples.	(10)
	b)	Write an exhaustive note on Glutathione conjugation.	(05)
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Q.3	a)	Classify non-barbiturates giving one representative structure for each class.	(05)
	b)	Discuss Sandburg's hypothesis in details.	(05)
	c)	Outline synthesis and main uses of:	(05)
		i) Pentobarbital sodiumii) Phenytoin sodium	
		n) I henytoin soulum	
Q.4		Write short notes on any THREE:	(15)
	a)	Anticonvulsants	
	b)	Respiratory stimulants	
	c)	Chemistry of general anesthetics	
	d)	Preanesthetic medication	
	e)	Factors affecting drug metabolism	
0.5		SECTION - II	(10)
Q.5	a)	Answer the following any FIVE: Describe symptoms of depression.	(10)
	b)	Draw any two structures of phenothiazine class of antipsychotics.	
	c)	Draw two structures of local anesthetics belonging to amino alkyl esters of	
	~)	PABA.	
	d)	Outline the synthesis of Meprobamate.	
	e)	Outline the synthesis of Dibucaine.	
	f)	Draw the structures of any two local anaesthetics from anilide class.	
	g)	Enlist different neurotic disorders.	
Q.6	9)	Classify tricyclic antidepressants with structures.	(05)
Q.0	a) b)	Describe SAR of tricyclic antidepressants.	(05) (05)
	c)	Comment on MAO inhibitors.	(05)
	0)		. ,
Q.7	a)	Classify local anaesthetics in details.	(05)
	b)	Describe the SAR of ester class of local anaesthetics.	(05)
	c)	Give synthetic route for Lignocaine and Benzocaine.	(05)
Q.8		Write short notes on the following any THREE:	(15)
Q.O	a)	Site specific delivery through prodrugs	/
	b)	SAR of phenothiazines	
	c)	Pharmaceutical applications of prodrugs	

PURUS –VI (2011 COURSE) : SUMMER-2016 SUBJECT : PHARMACEUTICAL ANALYSIS - IV

Time : 10:00AM:T0 1:00 P.M. Day : Friday Date : 29-04-2016 Max. Marks: 80 N. B.: Q.No.1 and Q. No.5 are COMPULSORY. Out of the remaining questions 1) attempt Any TWO from each section. 2) Answers to both the sections should be written in SEPARATE answer books. 3) Figures to the right indicate FULL marks. SECTION - I Solve Any FIVE of the following: (10)0.1 What are the advantages of TLC? a) What is HPLC? How is it superior to other chromatographic techniques? Give the principle of TLC. c) Compare between normal phase chromatography and reverse phase chromatography in HPLC. Draw a well labeled diagram of high performance liquid chromatographic instrument. Give important applications of TLC in pharmacy. (07)Explain in detail columns used in HPLC. Q.2 Discuss the various types of development methods of separation in TLC. (08)Discuss in detail properties and working of electrochemical detector and (07) Q.3 fluorescence detector. Compare between TLC and paper chromatography. (08)Write short notes on Any THREE of the following: (15)Q.4 a) Types of pumps used in HPLC b) Methods for the preparation of thin layers on plates Sample injection systems in HPLC Reverse phase adsorption TLC

SECTION - II

Q.5		Solve Any FIVE of the following:	(10)
	a)	Give the properties of CO ₂ as supercritical fluids.	
	b) c)	Explain the terms resolution and selectivity factor. How instrumentation for SFC does differ form HPLC?	
	d)	Why saturation of development tank is one of the essential steps in HPTLC development?	
	e)	Give the advantages of band application of sample over spot application.	
	f)	Give the chemical test for checking the adulterants in tea powder.	
Q.6	a)	Describe the duties of food safety officer.	(07)
	b)	Discuss the principle, technique and application of HPTLC.	(08)
Q.7	a)	Give the advantages and disadvantages of super critical fluid chromatography.	(07)
	b)	Give the chemical tests for checking adulterants present in various spices.	(08)
Q.8		Write short notes on Any THREE of the following:	(15)
	a)	Densitometric measurement in HPTLC	
	b)	Applications of super critical fluid chromatography	
	c)	Advantages of HPTLC	
	d)	Ideal properties of supercritical fluids	

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PURUS-VI (2011 COURSE): SUMMER 2016 SUBJECT: PHARMACEUTICAL BIOTECHNOLOGY (Including Molecular Biology)

Time: 10:00 AM-T01:00 P.M. Friday Day Max. Marks: 80. 06-05-2016 N.B.: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt any TWO questions from Section-I and any TWO questions from Section-II. 2) Both the sections should be written in SEPARATE answer books. Figures to the RIGHT indicate full marks. 3) Draw neat and labeled diagrams WHEREVER necessary. **SECTION-I** Q.1 Answer any FIVE of the following: (10)a) How would you define a plasmid? b) What are purines and pyrimidines? c) Give salient features of DNA double helix. Draw a neat labeled diagram of a bacterial cell. Write the importance of Thermophilus aquaticus. What are introns and exons? Differentiate between a probe and a primer. Q.2 Describe in details protein synthesis in a typical eukaryotic cell. (15)Q.3 Give details of Griffith's experiment and discuss its findings. (15)Q.4 Write short notes on any THREE of the following: (15)a) Recombinant DNA technology b) Differences in mRNA, rRNA and t-RNA c) Okazaki fragments d) Principle of Agarose gel electrophoresis e) Northern DNA hybridization. **SECTION-II** Q.5 Answer any FIVE of the following: a) What is induced mutation? b) What is shake flask culture? c) Draw a labeled diagram of batch fermentor. d) Differentiate a fermentor and bioreactors. What is single cell protein? Give key composition of fermentation media. Define strain improvement. Q.6 Define protein engineering and describe method used for site directed mutation. (15)Q.7 Describe enzymes as biocatalyst. Explain methods of enzyme immobilization. (15)Q.8 Write short notes on any THREE of the following: (15)a) Downstream process b) Applications of enzymes in pharmaceuticals Spray drying c) Whole cell immobilization Applications of proteases

PURUS- VI (2011COURSE): SUMMER - 2016 SUBJECT: PHARMACOGNOSY-II

Time: 10:00AM-TO 1:00P.M. Day: Thursday 12-05-2016 Max. Marks: 80 N.B: Q. No.1 and Q. No.5 are COMPULSORY. Out of remaining attempt ANY 1) TWO questions from each section. Answer to the TWO sections should be written in SEPARATE books. 2) SECTION-I Q.1 Answer the following question: (ANY FIVE) (10)a) What is infusion? b) What is dry extract? What is soft extract? c) State the methods of extraction of Fixed oil. Write down chemical constituents of Onion? e) Draw the diagram of Percolator. What is extraction? Explain Microwave assisted extraction. Q.2 (15)OR Explain the application of HPTLC in herbal drug evaluation. Q.3 a) Explain in detail about Super Critical Fluid Extraction. (08)What are the advantages of continuous hot extraction over conventional (07)methods of extraction? Q.4 Write notes on (ANY THREE): (15)Neutraceuticals a) b) Percolation c) Maceration d) HPLC **SECTION-II** Q.5 Answer the following question: (ANY FIVE) (10)What is Ash value? a) What is Foaming Index? b) c) Cosmetic functions of Aloe. Compare vein Islet number and Vein termination number. d) Explain functions of cytokinins. Phase of growth curve. Explain WHO guidelines for standardization of herbal drugs. Elaborate (15) Q.6 bitterness value and morphological characteristics crude drugs. OR Highlight WHO guidelines for standardization of herbal drugs. Explain pesticide residue and microbial load. Q.7 a) Explain the protocol to establish plant cells under in vitro conditions. (08)Highlight various nutritional requirements of plant tissue culture. b) Explain various biotechnological strategies to enhance secondary metabolites. (07)Q.8 Write notes on (ANY THREE): (15)a) Surface sterilization of explant b) Microscopical evaluation