PURUS – V (SEMESTER PATTERN): APRIL / MAY 2010 SUBJECT: PHARMACOGNOSY-II

Time: 10.00 A.M. TO 1.00 P.M : Monday Day Max. Marks: 80. Date : 17-05-2010 N.B.: Q. No. 1 & 5 are COMPULSORY. Out of remaining questions solve ANY TWO questions from Section I and ANY TWO questions from Section -II 2) Figures to the right indicate FULL marks. Answer to the two sections should be written in SEPARATE answer books. 3) Draw neat and labeled diagrams WHEREVER necessary. 4) **SECTION-I** Q.1 Attempt ANY FIVE of the following: [10] a) How do you detect the purity of Honey? What is meant by Saponification value? Give biological source, active constituents and uses of Arjuna. Differentiate between Phlobatannins and pyrogallotannins. Give Chemical test for Tannins. e) List out adulterants of Isapgol? f) Describe the general and distinguishing characters of fixed oils, essential oils [08] Q.2 a) and waxes. Write short note on solubility of fats. What are Gums? How are they classified? How does Gum acacia differ from [07] b) Tragacanth? What are Tannins? Differentiate between Hydrolysable and condensed [08] Q.3 a) What are starches? Describe microscopical structure, uses and method of [07] preparation of any one starch. Q.4 Write short notes on ANY THREE of the following: [15] Kaolin IP a) . Sesame oil b) Differences between Pale and Black Catechu c) Alginate **SECTION-II** Q.5 Attempt ANY FIVE of the following: [10] Differentiate between Alexandrian Senna and Indian Senna. What are Aloinosides? b) Give biological source, active constituents, and uses of Psoaralia. c) d) What is prepared digitalis? What are Isothiocyanate glycosides? e) What are Bitter Glycosides? Q.6 a) Define Glycosides. Outline its classification. What are steroidal glycosides? [08] Write a note on Solanum khaisanum. Give outline of drugs obtained from Mineral sources. What is Talc IP? b) [07]Describe cultivation, collection and drying of Aloe. [08] Q.7 a) Give a detailed account on preparation, identification and applications of [07] b) Bentonite. Write short notes on ANY THREE of the following: [15] T.S of Liquorice

PURUS -VI (SEMESTER PATTERN): APRIL / MAY 2010 SUBJECT: PHARMACOLOGY – III

Time: 10:00 AM. TO 1:00 P.M. : Thursday Max. Marks: 80. Date : 13-05-2010 N.B.: Q.No.1 and Q.No.5 are COMPULSORY. Out of remaining attempt ANY TWO 1) questions from each section. Answers to two sections should be written in SEPARATE answer books. 2) Figures to the right indicate FULL marks. 3) **SECTION-I** [10] Answer ANY FIVE of the following: Q.1 Enumerate anti-anxiety drugs. Give examples of analeptics. In which surgical conditions lignocine adrenaline combination is contraindicated. Enumerate stages of general anesthesia. Enlist uses of morphine. e) Give examples of atypical antidepressants. f) Which is the specific antagonist of benzodiazepine? Classify opioid analgesics. Describe the pharmacological actions, [08] Q.2 a) contraindications and precautions of morphine use. Classify neuroleptics. Describe the pharmacological actions and adverse [07] b) effects of chlorpromazine. Classify local anesthetics. Describe the various routes of administration of [08] Q.3 a) local anesthetics. Describe mechanism of action of barbiturates. What are their disadvantages as [07] b) sedative hypnotic drugs? [15] Write short notes on ANY THREE of the following: 0.4 Carbamazepine a) b) Benserazide c) New antiepileptics Intravenous general anesthetics **SECTION - II** Answer ANY FIVE of the following: [10] 0.5 Define antidote give examples. b) Enumerate main symptoms of acute lead poisoning. Give example of oral prostaglandin. In which conditions aspirin is contraindicated in children. Enlist symptoms of acute mercury poisoning. What are the side effects of prostaglandins? Discuss the mechanism of action, adverse effects, interactions, precautions and [08] Q.6 a) contraindications of allopurinol. [07] Describe the mechanism of actions and uses of aspirin. Explain the general principles of treatment of acute toxicity and acute [08] Q.7 a) Describe signs, symptoms, and treatment of acute poisoning with arsenic. [07][15] Q.8 Write short notes on ANY THREE of the following: Thromboxane

PURUS - VI (Semester Pattern) April/May - 2010 SUBJECT: PHARMACEUTICAL BIOTECHNOLOGY (Including Molecular Biology)

: Saturday Day Time: 10:00 AM-TO 1:00 RM. Date : 15-05-2010 Max. Marks: 80 N.B.: 1) Q.No.1 and 5 are COMPULSORY. Out of remaining questions attempt ANY TWO from each section. Answers to both the sections should be written in the SEPARATE answer books. 2) Figures to the right indicate FULL marks to the question. 3) SECTION-I 0.1 Attempt ANY FIVE: [10] a) Give complementary sequence of GATTCAAGCCTTATCAT. b) What is a primer? c) Draw and label double helix. d) What is a chromosome? e) Define plasmid. What is ligation? f) Define mutation. Q.2 What do you understand by rDNA technology? Give various uses of the same [15] in pharmacy. Q.3Elaborate DNA as master molecule of life and describe Griffith's experiment [15] that established DNA as the transforming principle. Q.4 Write short notes on ANY THREE: [15] a) Taq DNA polymerase b) Principle of gel electrophoresis c) DNA hybridization d) Plasmid as vector SECTION - II 0.5 Attempt ANY FIVE: [10] a) Give any two methods of immobilization of enzymes. b) What is continuous fermentation? c) Give applications of amylase. d) Differentiate encapsulation and adsorption. e) Enlist aims of protein engineering. What is site directed mutagenesis? f) g) Define enzyme. Discuss the salient achievements already made and future prospects in protein [15] Q.6 engineering. Describe enzymes as biocatalyst and discuss various factors that affects rate of [15] Q.7 enzymatic reaction. Write short notes on ANY THREE: Q.8 [15] a) Fluidized bed fermentation b) Single cell protein c) Limitation of enzyme immobilization

PURUS-VI (SEMESTER PATTERN): APRIL/MAY 2010 SUBJECT: MEDICINAL CHEMISTRY-II

Time: 10:00 AM. TO 1:00 P.M. : Tuesday Day Date 04-05-2010 Max. Marks: 80. 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) Answers to the two sections should be written in SEPARATE answer books. 3) Figures to the RIGHT indicate full marks. SECTION-I Q.1 Give structure and explain mode of action of any FIVE drugs of following: (10)Fluphenazine Tranylcypromine sulfate b) Doxepin c) Alprazolam d) Phenobarbital e) f) Spironolactone Phensuximide. Q.2 a) Discuss SAR and chemistry of benzodiazepins and list out their important (07) therapeutic applications. Classify antidepressants and give an exhaustive account of tricyclic (08) antidepressants. 0.3 Classify sulfonamides and add an exhaustive note on thiazides. (07)Write synthesis of (08)Ethacrynic acid ii) Acetazolamide iii) Meprobamate iv) Haloperidol. Q.4 Write short notes on any THREE of the following: (15)Anticonvulusants for grand mal epilepsy b) Barbiturate chemistry and sedative action High ceiling diuretics c) Phenothiazines d) Purines and heterocyclic diuretics. P.T.O.

SECTION-II

Q.5	Q.5 Attempt any FIVE questions of the following:		
	a) b) c) d) e) f)	Explain hydralytic reactions of metabolism with suitable example. Define and classify drug metabolic pathways. Draw any two structures from Phenylethylamine class of analeptics. Give broad mechanism of action of CNS stimulants. Give any two structures from aminoalkyl esters of p-aminobenzoic acid class of local anesthetics. Differentiate between the local anesthetics and general anesthetics. Give the various stages of general anesthesia.	
Q.6		Give an exhaustive account of oxidation-reduction metabolic pathways with examples.	(15)
Q.7	a)	Classify local anesthetics with structure. Discuss their mode of action, SAR and therapeutic uses.	(10)
	b)	Outline synthesis of procaine and benzocaine	(05)
Q.8	Q.8 Write short notes on any THREE of the following:		(15)
	a) b) c) d) e)	Chemistry of General anaesthetics Hallucinogens Factors affecting drug metabolism Analeptics Drugs of abuse	

PURUS-VII: (Semester Pattern) A pril-May. 2010 SUBJECT: PHARMACEUTICAL ANALYSIS-IV

Time: 2.00 P.M. To 5.00 P.M.

: 07-05.2010 Max. Marks: 80. N.B.: 1) Q.NO.1 and Q.NO.5 are COMPULSORY. Out of remaining questions attempt ANY TWO questions from Section-I and Section-II. 2) Answers to both the sections should be written in the SEPARATE answer books. 3) Figures to the right indicate FULL marks. **SECTION-I** Q.1 Attempt ANY FIVE of the following: [10] Write the different regions of EMR. Explain the terms excitation and relaxation in spectroscopy. Name the different detectors for UV c) Write advantages and disadvantages of grating monochromators. d) Write advantages and disadvantages of a Barrier Layer Cell. What are the charge transfer transitions? f What are the different methods of quantitative analysis by UV? g) Write the construction and working of a PMT. [07] Q.2 a) What are the different transitions in UV spectroscopy? Explain transitions b) [08] involving pi and n electrons. State and explain Beer Lambert law. [08] 0.3 a) Write a note on deviations of Beer Lambert Law. [07] Write short notes on ANY TWO of the following: 0.4 [15] Different methods of spectroscopy. a) PDA detector in UV b) Spectrophotometric titartions. **SECTION-II** 0.5 Write short notes on ANY FIVE of the following: [10] Fluorimetry is more sensitive than UV Finger print region in IR b) Nephelometry is used for dilute solutions b) Temperature affects fluorescence intensity d) Fluorescence is instant while phosphorescence is delayed e) IR is not used for quantitative analysis. f) Role of attenuator in IR spectroscopy Explain Napheloturbidimetry giving principle instrumentation and [10] Q.6 a) applications. Write different molecular vibrations in IR. [05] b) What are the factors affecting fluorescence intensity? Explain. [08] Q.7 a) Give applications of fluorimetry. [07]b) Write short notes on ANY TWO of the following: [15] Q.8 ET.ID

: Friday

Day

PURUS -VIII (SEMESTER PATTERN): APRIL / MAY 2010 SUBJECT: DOSAGE FORM DESIGN - III

: Wednesday Time: 2.00 P.M. To 5.00 P.M. Max. Marks: 80. :12-05.2010 N.B.: Q.No.1 and Q.No. 5 are COMPULSORY. Out of remaining questions attempt 1) ANY TWO questions from each section. Answers to both the sections should be written in SEPARATE answer books. 2) Figures to the right indicate FULL marks. 3) SECTION-I [10] Answer ANY FOUR of the following: 0.1 a) Enumerate the merits of controlled release dosage form. b) Draw neat diagram of Alzet osmotic pump. c) What is pH activated DDS? d) Explain Dynamics of GI Tract. e) What is Nitro Dur system? What are magnetically activated DDS? Enumerate and discuss the principle, formulation and give biopharmaceutical [15] 0.2 consideration of Iontophoretic DDS. Discuss giving example the principle, formulation and kinetics of osmotic [15] Q.3 pressure controlled DDS. Write short notes on ANY TWO of the following: Q.4 [15] a) Feed back regulated DDS b) Hybrid Drug Delivery System c) Mucoadhesive DDS d) Multiunit Dosage forms. SECTION - II 0.5 Answer ANY FOUR of the following: [10] a) Give a schematic representation showing anatomy of skin. b) Give composition of nasal secretion. c) State the applications of Niosomes. d) Show with the help of diagram different components of aerosol packaging. What are different classes of antibodies? State the applications of metered dose inhalers. Q.6 Discuss the formulation, manufacturing and quality control of pharmaceutical [15] aerosols. 0.7 a) Discuss physico-chemical parameters influencing development of nasal DDS. [10] Comment in brief technologies used in development of transdermal DDS. [05] Write short notes on ANY TWO of the following: Q.8 [15] a) Aerodynamics of aerosols b) Formulations of Niosomes Formulations of liposomes

Rectal drug delivery formulations