PURUS - V (2011 COURSE): WINTER - 2014 SUBJECT: COSMETICOLOGY

(For the Student of 2012-13 Batch and Onwards) Day: Monday Time: 10,00 A.M. To 1,00 PM Date: 17/11/14 Max. Marks: 80 N.B.: 1) Q. No. 1 and 5 are COMPULSORY. 2) Figures to the RIGHT indicate full marks. 3) Solve any TWO questions from each section. Answers to both the section should be written in SEPARATE answer books. SECTION-I 0.1 Solve any FIVE of the following: (10)Define cosmetics. a) b) What are viscosity modifiers? Give two examples. c) Define emollients. d) Enlist factors to be considered for design of successful cosmetic formula. e) Name agents banned for use in cosmetics applications. Give a example of regulatory agency of cosmetics. f) What are normal functions of skin? Classify oral hygiene products. Give formulation information for these. O.2 a) (10)Discuss about skin cleansing creams with examples. (05)Q.3 a) Explain importance of physiological considerations in cosmetic product (08) development. Describe vanishing cream and its modifications to get other cosmetics products. (07)0.4 Write short notes on any THREE of the following: (15)Vehicles in cosmetics Permanent hair dyes Viscosity modifiers in skin care products d) Humectants **SECTION-II** Q.5 Solve any FIVE of the following: (10)What are different herbal materials used in cosmetics? a) What is the principle of nail lacquer? b) Why stearic acid based creams require addition of humectants? d) What are the differences between lip gloss and lip slave? Classify and define lipstick products. What are ideal requirements of manicure products? f) How lotions are different from creams. Discuss in detail psychometric evaluation of any cosmetics product. Q.6 a) (10)Describe different baby products along with examples. (05)What are different mechanisms of herbal ingredients in cosmetics? (08)Discuss about lipcare product along with examples. (07)0.8 Write short notes on any THREE of the following: (15)

a) Lip Balm b) Nail paints

c) Polymers in cosmetics

WINTER - 2014

PURUS - V (2011 COURSE): ... WINTER - SUBJECT : PHARMACOGNOSY - I

		SUBJECT : PHARMACOGNOSY - I						
	Day Date	: 1	Friday Time: 10:00 A: M Max. Marks: 80	TO1.00PM				
			-11.2017					
	N.B.:	1)	Q.No.1 and Q.No.5 are COMPULSORY. Out of remaining questions attempt ANY TWO questions from each section.					
		 Answers to both the sections should be written in the SEPARATE answer Draw neat and labeled diagrams WHEREVER necessary. Figures to the right indicate FULL marks. 						
			SECTION - I					
	Q.1		Attempt ANY FIVE of the following: Name any two drugs obtained from mineral origin. Draw structure of 2, 4D.	[10]				
			What is role of ethylene in plant growth? Write mechanism of action of auxins. Write different methods of drying of crude drugs.					
		f)	What is chemotaxonomic classification?					
	Q.2	a)	What is significance of evaluation of crude drugs? Explain chemical evaluation detail.					
		b)	Give chemical and pharmacological classification of crude drug with example.	suitable [07]				
	Q.3	a) b)	What is physical evaluation of crude drug? Give examples. Explain in detail collection and processing of crude drug.	[08] [07]				
	Q.4	a) b) c) d)	Biological evaluation of crude drug Natural sources of crude drugs	[15]				
-			SECTION - II					
	Q.5	b) c) d)	Write biological source of Shark liver oil. Write chemical test for Acacia.	[10]				
	Q.6	a) b)	Write in detail pharmacognostic account of Agar. What are carbohydrates? Give biosynthesis, classification chemist chemical tests for carbohydrates.	[08] try, and [07]				
	Q.7	a)	lipids with suitable example.					
		b)	Write in detail pharmacognostic account of Pectin and Cod liver oil.	[07]				
	Q.8	a) b)	Write short notes on ANY THREE of the following: Chinese system of medicine Mevalonic acid pathway	[15]				

PURUS-V (2011 COURSE): WINTER - 2014 SUBJECT: PHARMACOLOGY-II

. Wednes day Time : 10.00A.M. To 1.00 P.M Date : 19-11-2014 Max.Marks: 80 N.B.: 1) Q.No. 1 and Q.No.5 are COMPULSORY. Out of remaining, attempt Any TWO questions from SECTION-I and Any TWO questions from SECTION-II. Answers to both the sections should be written in SEPARATE answer books. 2) 3) Figures to the right indicate FULL marks. SECTION-I 0.1 Answer Any FIVE of the following: (10)a) Classify hematinics with suitable examples. b) Describe the biosynthesis of nitric oxide. Classify blood coagulants. c) d) Explain the mechanism of heparin. Define thrombolytics and give suitable examples. e) Enlist the contraindications for heparin therapy. Q.2 Classify antiarrhythmics. Explain in detail the pharmacology of calcium (08) channel blockers. Classify antihyperlipidemic drugs. Explain in detail the pharmacology of (07) statins. Classify antianginal drugs with suitable examples. Explain in detail (08) Q.3 a) pharmacology of nitrates. Explain drug therapy of hypertension. (07)Write short notes on Any THREE of the following: Q.4 (15)Anticoagulants Potassium channel openers b) Fibrinolytics c) d) Nitric oxide **SECTION-II** Q.5 Answer Any FIVE of the following: (10)What are diuretics? Give 2 examples of loop diuretics. Classify antidiuretics. c) Define anti-tussives. Give examples. d) Enlist the uses of potassium sparring diuretics. Enlist the drugs used for GERD. e) Give the WHO recommended ORS formula. Classify diuretics. Explain the pharmacology, uses and adverse effects of high (08) O.6 a) ceiling diuretics. Classify anti-asthmatic drugs. Explain the significance of oral corticosteroids (07) in treatment of asthma. Classify anit-ulcer drugs. Explain in detail the pharmacology of proton pump (08) Q.7 Classify anti-emetics. Explain in detail the role of 5-HT antagonists as anti- (07) emetics.

Write short notes and A THEEDER O. .

PURUS-V (2011 COURSE): WINTER - 2014 SUBJECT: PHARMACOLOGY-II

. Wednes day Time: 10.00A.M. To 1.00 P.M Date : 19-11-2014 Max.Marks: 80 N.B.: 1) Q.No. 1 and Q.No.5 are COMPULSORY. Out of remaining, attempt Any TWO questions from SECTION-I and Any TWO questions from SECTION-II. 2) Answers to both the sections should be written in **SEPARATE** answer books. 3) Figures to the right indicate FULL marks. **SECTION-I** Q.1 Answer Any FIVE of the following: (10)a) Classify hematinics with suitable examples. Describe the biosynthesis of nitric oxide. b) Classify blood coagulants. c) d) Explain the mechanism of heparin. Define thrombolytics and give suitable examples. e) Enlist the contraindications for heparin therapy. Q.2 Classify antiarrhythmics. Explain in detail the pharmacology of calcium (08) channel blockers. Classify antihyperlipidemic drugs. Explain in detail the pharmacology of (07) statins. Classify antianginal drugs with suitable examples. Explain in detail (08) Q.3 pharmacology of nitrates. Explain drug therapy of hypertension. (07)Q.4 Write short notes on Any THREE of the following: (15)Anticoagulants Potassium channel openers b) Fibrinolytics c) Nitric oxide **SECTION-II** 0.5 Answer Any FIVE of the following: (10)What are diuretics? Give 2 examples of loop diuretics. a) Classify antidiuretics. b) Define anti-tussives. Give examples. c) Enlist the uses of potassium sparring diuretics. Enlist the drugs used for GERD. e) Give the WHO recommended ORS formula. Q.6 a) Classify diuretics. Explain the pharmacology, uses and adverse effects of high (08) ceiling diuretics. Classify anti-asthmatic drugs. Explain the significance of oral corticosteroids (07) in treatment of asthma. Classify anit-ulcer drugs. Explain in detail the pharmacology of proton pump (08) 0.7 inhibitors. Classify anti-emetics. Explain in detail the role of 5-HT antagonists as anti- (07) emetics.

Write chart notes on A TETETETE C.

08

PURUS -V WINTER - 2014 (old Course)

SUBJECT: MEDICINAL CHEMISTRY -I
Day: Mouday Time: 10.00 A'MITO1'00 Max marks: 80 Date: 10-11-2014 N.B: Question No. 1 & 5 are COMPULSORY. Out of remaining 1) solve any TWO from section I & any TWO from section -II. Figures to the right indicate FULL marks. 2) **SECTION-I** Write structures of any FIVE of following, mentioning their chemical class & (10) category. Captopril a) Isoproterenol b) Dibenamine c) Propranolol d) Atenolol e) Guanabenz f) Verapamil g) (05)Write in details about cardiac glycosides. a) Classify antianginals giving one structure for each class. (05)b) Discuss mode of action of calcium channel blockers & organic nitrates. (05)c) Classify prodrugs on the basis of their carriers giving one structure for each (05)Q.3 a) Discuss pharmacokinetic applications of prodrugs. (05)b) Write in details about brain targeting by prodrugs. (05)c) (15)Write short notes on any THREE Biosynthesis & metabolism of noradrenaline a) b) SAR of direct sympathomimetics Indirect acting sympathomimetics c) Synthesis of guanethidine and terbutaline d) α- antagonists e) **SECTION-II** (10)Attempt any FIVE questions: a) Draw structures of carbachol & bethanechol Out line the synthesis of dantroline sodium b) Write on importance of stereo-chemical purity of drugs. c) Give an account of ganglionic stimulants. d) State Ferguson principle. e) f) Write about muscurinic receptors. Outline the synthesis of gallamine. Describe in details chemistry, pharmacological actions, SAR and metabolism of (10) Q.6 a) acetyl choline & its analogs. Give the classification of neuro-muscular blocking agents on the basis of MOA. Enlist various physico- chemical parameters affecting drug action. Comment on (08) 0.7 a) role of solubility & partion coefficient in drug's action.

(07)

(15)

0.1

Q.2

0.4

Q.5

Q.8

Write about SAR of Anti-muscarinics.

Write short notes on :(Any THREE)

Dissociation constant & hiological action

PURUS – V (2011 COURSE): WINTER - 2014 SUBJECT: PHARMACEUTICAL ANALYSIS – III

Time: 10.00 A. M. TO 1.00 Day: Wednesday Max. Marks: 40 Date: 12-11-2014 N.B.: Q. No. 1 and Q. No. 5 are COMPULSORY. Out of remaining solve ANY TWO from section - I and Section - II each. Both the sections should be written in the SEPARATE answer books. 2) Figures to the right indicate FULL marks. 3) SECTION-I (10)Answer ANY FIVE of the following: Q.1 Explain in short 'Retention Time'. Define 'Column Resolution'. Explain the term 'Eddy Diffusion'. What is the difference between concentration sensitive and mass sensitive What properties should a stationary phase for gas chromatographic column possess? Explain briefly 'Number of Plates'. (08)Enlist all the detectors used in G.C. Explain in detail any two. Q.2 Draw a diagram showing parts of G.C. Give applications of the same. (07)Explain in detail the theory behind band broadening. How can it be reduced to (15) 0.3 improve column efficiency? (15)Write short notes on ANY THREE of the following: Q.4 GC-MS a) Open tubular columns b) Selectivity factor c) Classification of chromatographic methods **SECTION - II** (10)Attempt ANY FIVE of the following: Q.5 Write factors affecting retention factor, Rf. "Ion exchange materials are classified in terms of acidic or basic strength of functional groups attached to the polymer matrix". Why? Types of solvents used in paper chromatography. Discuss ion exchange capacity. d) Write procedure for separation of Amino Acid by paper chromatography. Applications of Immunoelectrophoresis. Give the principle involved in electrophoresis and a note on different modes (07) 0.6 of electrophoretic techniques. Write applications of electrophoresis add a detail note on different modes of (08) electrophoretic techniques. Write classification of Paper chromatography? Explain different modes of (07) Q.7 Paper Chromatography. Explain Choice of filter papers in paper chromatography. (08)(15)Write short note on ANY THREE of the following. Q.8 Partition Paper Chromatography

Applications of Ion Eychange Chromatography

PURUS - V (2011 COURSE): . WINTER - 2014 SUBJECT: DOSAGE FROM DESIGN - II

Time: 10.00 A. MITO 1.00 P.M Day: Friday Max. Marks: 80 N.B.: Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt any 1) TWO questions from each section. 2) Figures to the right indicate FULL marks. 3) Answers to both the section should be written in SEPARATE answer book. 4) Draw neat diagrams WHEREVER necessary. SECTION-I 0.1 Answer any FIVE of the following: (10)What are the formulation factors affecting drug availability from capsules? Write various sizes and volumes (cm³) filled in capsules. Mention the advantages of transdermal drug delivery systems. Define gels. Explain types of gels. Give the flow chart sequence of two piece hard gelatin capsule shell manufacturing. Write advantages and disadvantages for soft gelatin capsules. f) Mention IPQC checks during gelatin shells manufacturing Q.2 a) Discuss in detail Evaluation of Hard gelatin capsules. (10)Discuss the formulation of soft gelatin capsules. (05)Classify the defects of capsule, explain the defects in detail. 0.3 (08)a) Discuss the concept of topical and transdermal formulations. (07)0.4 Write short notes on any THREE of the following: (15)Percutaneous absorption a) Machines/ Equipments for manufacturing of semisolids Industrial filling of hard gelatin capsule c) Manufacturing process for gelatin **SECTION-II** Q.5 Answer any FIVE of the following: (10)Mention the disadvantages of powders as a dosage form. What is rational for tablet coatings? b) What are causes and remedies of orange peel effects? c) d) Mention the factors affecting the choice of inhaler device. What is sealing? Why is it applied? What are propellants? Give two examples of it. f) Why HPMC is most widely used cellulosic polymer in Immediate release coating. Discuss the formulation and filling of aerosols. (10)0.6 a) Explain different methods of granulations. (05)Q.7 a) Discuss the formulation of chewable tablet. (08)b) Discuss in detail Sugar coating of tablet. (07)Write short notes on any THREE of the following: Q.8 (15)a) Super Disintegrants b) Dispersible tablet

Dry powder Inhalation