### MANIKGAD – III : SUMMER - 2016 SUBJECT : PHARMACOLOGY - II

Time : 2.00 P.M. TO 5.00 P.M : Monday Day Max. Marks: 70 :04.04.2016 Date Q.1 and Q.5 are COMPULSORY. Out of the remaining attempt any TWO N.B. 1) questions from each Section. Figures to the right indicate FULL marks. Answers to both the sections should be written in SEPARATE answer books. 2) 3) SECTION - I (08)Answer any FOUR of the following: 0.1 Classify antibiotics based on their mechanism of action. What is paracrine and autocrine signaling? ii) Give the mechanism of action of  $\beta$  lactam antibiotics. iii) Explain co-trimoxazole. iv) Mention the toxic effects of aminoglycoside antibiotics. v) (03)Write the mechanism of protein synthesis. Classify antineoplastic agents. Discuss the mechanism of action and adverse (12)Q.2 effects of folate antagonists. Give the mechanism of action, pharmacokinetic properties and clinical uses (07)Q.3 of spirinolactone. Define and classify immunosuppressants. Discuss the pharmacological (05)profile of cyclosporine. (12)Write short notes on ANY THREE of the following: Q.4 Treatment for UTI a) Antifungal agents b) Gene mutation c) Anticoagulants d) **Biosensors SECTION - II** (80)Answer any FOUR of the following: **Q.5** Write mechanism of action of sulphonamides. i) Give mechanism of action and clinical use of chloramphenicol. ii) Write a short note on tRNA iii) Write a short note on chromosomes. iv) Give reasons for the development of resistance to antihelminthic v) drugs. (03)What is acute and chronic toxicity? Give pathogenesis of cancer, classify anticancer drugs. Write mechanism of (12)anticancer drugs, and explain the reasons for the development of resistance to Q.6 anticancer drugs. What are tetracyclines? Classify them, write mechanism of action and (07)**Q.7** a) adverse effects of tetracyclines. Write a note on antiviral drugs. b) (12)Write ANY THREE of the following: 0.8 Write the steps involved in viral replication a) Describe in detail the loop diuretics b) Describe in detail various steps of mitotic cycle. c) Write chemotherapy for amoebiasis.

#### MANIKGAD – III: SUMMER - 2016 SUBJECT: PHARMACEUTICAL ANALYSIS

Time 2.00 PM. To 5.00 P.M. Day: Wednesday Max. Marks: 70 Date: 06.04.2016 N.B.; Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining solve 1) Any TWO questions from Section - I and Any TWO questions from Section - II. Answers to the two sections should be written in SEPARATE answer books. 2) Figures to the RIGHT indicate full marks. 3) SECTION - I (08)A) Attempt ANY FOUR of the following: 0.1State types of adsorbents used in TLC. Define specific conductance and molar conductance. Compare between Dropping Mercury Electrode and Rotating Platinum Electrode. d) Classify electrodes used in potentiometry. State types of gels used in gel chromatography. (03)Write methods to develop a chromatogram. B) (04)State theories involved in chromatography. Q.2 Discuss in brief Gas Chromatography instrumentation with an exhaustive b) note on columns used in GC. Discuss in brief degassing techniques and types of pumps in HPLC. (07)Q.3 a)

Explain polarographic apparatus in detail.

Conductometric Applications.

Amperometric titrations.

Applications of HPTLC.

ISO Elements.

Q.4

a)

b)

c)

Write short notes on ANY THREE of the following:

P.T.O.

(05)

(12)

#### **SECTION - II**

Q.5	A)	Attempt ANY FOUR of the following:	(08)
	i)	Write types of ions produced in mass spectrophotometer.	
	ii)	Explain Bragg's law.	
	iii)	Write types of molecular vibrations in IR spectroscopy.	
	iv)	State components of flurimeter.	
	v)	Write events occurring in flame photometer.	
	<b>B</b> )	Write factors influencing chemical shifts in NMR.	(03)
Q.6	a)	Discuss different types of interference in AAS.	(04)
	b)	State derive Beer- Lambert's Law. Explain deviations leading from it.	(08)
<b>Q.</b> 7	a)	Classify thermal methods of analysis. Describe instrumentation and applications of DSC.	(07)
	b)	Discuss applications of IR spectroscopy with examples.	(05)
Q.8		Write short notes on ANY THREE of the following:	(12)
	a)	ESR spectrophotometer	
	b)	Radiation sources in AAS	
	c)	Shielding and Deshielding in NMR	
	d)	Applications of XRD	

# Third Year- Pharm D. - 2016.

## MANIKGAD – III : SUMMER - 2016 SUBJECT : PHARMACOTHERAPEUTICS - II

Day	y : Saturday Time: 2:00 P.M. To		00
ate	1 (a. Marka + 70)		
l.B:	<ol> <li>Q.No.1 and Q.No.5 COMPULSORY. Out of remaining questions attempt ATTWO questions from each section.</li> <li>Answers to both the sections should be written in SEPARATE answer books.</li> <li>Figures to the right indicate FULL marks.</li> </ol>		
		SECTION – I	
Q.1	A)	Answer ANY FOUR of the following:  i) What are the indications for antibiotics?  ii) What is Chemotherapy?  iii) Differentiate between Sinusitis and rhinitis.  iv) What are different types of urinary tract infections?  v) What is septicemia?  vi) Name the first line antitubercular drugs.  vii) What are different types of dermatitis?	[08]
	B)	What is the role of corticosteroids in different skin diseases?	[03]
Q.2		Discuss etiology, diagnosis and specific antibiotics therapies for infectious endocarditis.	[12]
Q.3	a) b)	Discuss pharmacotherapy of Pulmonary tuberculosis.  Discuss pathophysiology and choice of antibiotic regimeNs for bacterial gastroenteritis.	[07] [05]
Q.4	a) b) c) d)	Write short notes on ANY THREE of the following: Complications of malaria Antibiotic Regimen for sinusitis Antiretroviral drugs Malaria cycle	[12]
		SECTION – II	
Q.5	A)	Answer ANY FOUR of the following:  i) What is the significance of Creatinine clearance?  ii) What is the etiology for RA?  iii) What is hemodialysis?  iv) What is Cancer?  v) Define SLE.  vi) What is chemotherapy?  vii) What is spondylitis?	[08]
	B)	Differentiate RA and OA.	[03]
Q.6		Discuss etiology, Malaria cycle and Pharmacotherapy for malaria.	[12]
Q.7	<b>a</b> ) b)	Define RA and write in detail about pharmacotherapy of RA.  Discuss management of chemotherapy induced nausea and vomiting.	[07] [05]
Q.8	a) b) c) d)	Write short notes on ANY THREE of the following: Local treatment of Cancer Pharmacotherapy of Gout Management of Spondylitis Management of CRF	[12]

M.

## MANIKGAD – III : SUMMER - 2016 SUBJECT : PHARMACEUTICAL JURISPRUDENCE

Day Date	Tuesday : 12-04-2016		Time: 2.00 P.M. To 5.00 P.M. Max. Marks: 70	
N. B. :	1) 2) 3)	Q. No. 1 and Q. No. 5 are COMPULSORY. ANY TWO questions from Section – I and II Figures to the right indicate FULL marks.  Both the sections should be written in the SEF		
		SECTION - I		
Q. 1	A) i) ii) iii) iv) v) vi)	Answer any FOUR of the following: What are the objectives of Narcotic Drugs and P. Who is a Registered Pharmacist? How will you define an "Adulterated Drug"? What is Bonded Manufactory? What are the conditions for storage of Schedule What are the building requirements according to	X drugs?	
	B)	Classify medicinal and toilet preparations contain	(02)	
Q. 2	a) b)	Discuss the constitution of Pharmacy Council of What is the qualification required for a Drug Ins		
Q. 3		Discuss the provisions for offences and penaltic Preparations Act 1955.	es under Medicinal and Toilet (12)	
Q. 4	<b>a</b> )	Write short notes on ANY THREE of the follow Standards of Ophthalmic products	wing: (12)	
	b)			
	c)	Duties of Government Analyst		
	d)	Drug import provisions in India		

#### SECTION - 11

0.5	A)	Answer any <b>FOUR</b> of the following:	(08)
Q. 5	i)	Define Magic Remedy.	
	ii)	What is the object of Prevention of Cruelty to Animals Act 1960?	
	iii)	What do you mean by Bulk Drug?	
	iv)	Define an Invention.	
	v)	What are the penalties under Patents Act 1970?	
	vi)	What is Patent of addition?	
	B)	Differentiate between Generic and Branded Formulations.	(03)
Q. 6	a)	Discuss the process of grant of a Patent according to Patents Act 1970.	(06)
	b)	Discuss writing of prescription in detail.	(06)
Q. 7	a)	What are the classes of exempted advertisements under Drugs and Magic Remedies Act 1954?	(06)
	b)	How the non obviousness (inventive step) of an invention is determined?	(06)
Q. 8		Write short notes on ANY THREE of the following:	(12)
	a)	Animal Welfare Board of India	
	b)	Provisional specifications w.r.t. Patents Act	
	c)	Penalties under Essential Commodities Act	
	d)	OTC products	

## MANIKGAD – III: SUMMER - 2016 SUBJECT: MEDICINAL CHEMISTRY

Time : 2.00 P.M. TO 5.00 P. M : Saturday Max. Marks: 70 Date : 16-04 2016  $\overline{N.B.}$ Q.1 and Q.5 are COMPULSORY. Out of the remaining attempt any TWO 1) questions from each Section. Figures to the right indicate FULL marks. 2) Answers to both the sections should be written in SEPARATE answer books. 3) SECTION - I (08)Answer any FOUR of the following: 0.1 Classify anti-viral agents. Write two structures of antiamoebic agents. ii) Classify topical antiinfective agents. iii) Define QSAR. Write methods of QSAR. iv) Write structures of pyrazinamide and ethambutol. (03)Classify antifungal agents. b) Classify antimalarials. Write note on quinolines as a antimalarial agents. (07)Q.2 a) (05)SAR of sulphonamides. Define and classify antibiotics. Write a detail note on tetracyclines. (12)Q.3 (12)Write synthesis of any THREE of the following: Q.4 Sulphamethoxazole a) 17 β-oestradiol b) Furosemide c) Isoniazide **SECTION - II** (08)Answer any FOUR of the following: Q.5 Draw structures of any two anticoagulants. i) Classify antiarrhythmic agents. ii) MOA of HMG-CoA reductase inhibitors. iii) Drugs used in angina pectoris. iv) Draw structures of any two antihypertensive agents. (03)Write note on antihyperlipidemic agents. Classify diuretics and write a note on thiazide diuretics. (07)Q.6 a) Write a note on Glucocorticoids. (05)Write a detail note on antineoplastic agents. (12)Q.7 Write note on any THREE of the following: (12)Q.8 Antihypertensive agents a) Synthetic hypoglycemic agents b) Thyroid hormones and antithyroid drugs Diagnostic agents

### MANIKGAD-III: SUMMER - 2016

SUBJECT: PHARMACEUTICAL FORMULATIONS
Day: Wednesday
Tim
Date: 20-04-2016
Max

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

Max Marks: 70

#### N.B:

- Q.1 and Q.5 are COMPULSORY. Out of remaining answer any TWO 1) Questions from each section.
- Figures to the right indicate FULL marks. 2)
- 3) Answer to each section should be written in SEPARATE answer book.

		SECTION-I	
Q.1	A) a) b) c) d) e) f) B)	Answer any FOUR questions Discuss term" Solubility" "Solublization" and "Dissolution". Write co- solvency technique used in monophasic liquid formulations. Discuss weight variation test for capsules. Explain physical stability of emulsions. Write concept of buccal tablet and their applications. Classify preservatives according its use into dosage forms. Discuss about colouring agents used in syrups.	(08)
Q.2	a) b)	Write about material of fabrication for hard capsules.  Discuss a method to manufacture soft gelatin capsules.	(06) (06)
Q.3	a) b)	Discuss methods to manufacture suspensions.  Write various physical and chemical stabilizers used in suspensions.	(06) (06)
Q.4	a) b) c) d)	Write note on ANY THREE Antioxidants Emulsifying agents Pre- formulation Sustained release tablets	(12)
		SECTION-II	
Q.5	A) a) b) c) d) e) f) B)	Answer any FOUR questions Enumerate advantages of dry granulation. Enlist advantages of sublingual tablets. Explain a method to sterilize thermolabile pharmaceuticals. Write benefits of enteric coated tablets. Define term: Isotonic, Sterile, Apyrogenic and Hypertonic. Discuss properties of water for injection and sterile water for injection. Describe techniques to determine flow property of granules.	(03)
Q.6	a) b)	Explain significance of tablet coating and enlist steps involved in sugar coating.  Write significance of glidants and lubricants in solid dosage forms.	(06) (06)
Q.7	a) b)	Explain steps involved in film coating of tablets.  Write detailed note on capsules formulation to be filled in hard capsules.	(06) (06)
Q.8	a) b) c) d)	Write note on ANY THREE Method for preparation of suppositories Factors affecting absorption of drug through skin Sterility test Jellies	(12)

#### SUPPLEMENTARY

MANIKGAD-III SUMMER-2016 SUBJECT: PHARMAECUTICAL FORMULATIONS

Time: 10:00 AM. TO 1:00 P.M. Max Marks: 70 07-07-2016 Date 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 the SEPARATE answer books. 2) Figures to the RIGHT indicate full marks. 3) **SECTION-I** (08)Attempt ANY FOUR of the following: Q.1 A) Write the advantages of tablet as dosage form. a) Write the classification of dosage form as per route of administration. b) Write the significance of suppository as dosage form. c) Differentiate between flocculated and deflocculated suspension. d) Explain the creaming and cracking of emulsion. (03)Write the criteria for selection of coloring agent in dosage forms. (12)Discuss in detail large scale manufacturing of emulsion and its evaluation. Q.2 (06)Write the factors affecting on percutaneous absorption. **Q.3** (06)Write the merits and demerits of sustained release formulations and write b) rational behind development of it. (12)Answer ANY THREE of following: **Q.4** Packaging of oral liquids. **a**) Implantable drug delivery system. b) Suspending agents. c) Solubility enhancement techniques. **SECTION-II** (08)Attempt ANY FOUR of the following: **Q.5** Define Large Volume Parenterals and Small Volume Parenterals. a) Enlist coating agent and write the significance of enteric coating. Write the significance of dissolution testing and how it is correlated with therapeutic effect. Write the role of tonicity modifiers and mention two examples of tonicity d) modifiers. How capsules are different dosage forms in comparison to tablets. e) Write the significance of disintegration time of different tablets. (03)Discuss the evaluation parameters of coated tablets. (12)Q.6 Discuss the need of formulating drug in the form of soft and hard gelatin (06)Q.7 a) capsules with their merits and demerits. Discuss the plastic containers used for parenterals. (06)(12)Answer ANY THREE of following: Q.8 a) Different "Class" of area as parenteral facility. b) Sugar coating of tablets. Filling of soft gelatin capsules. c) Approaches to minimize "Mottling" in tablets.