T.Y.B.PHARM. SEMESTER-V (2011 COURSE): SUMMER - 2018 SUBJECT: COSMETICOLOGY

Day Date	:	 Wednesday 02/05/2018 Time: 10.00 AM Max. Marks: 80 		O 01.00 1 W1	
N. B	.: 1) 2) 3)	TWO questions from each section. Answers to both the sections should be written in SEPARATE answer book.			
			SECTION – I		
Q.1	a) b) c) d) e) f)	Define cosmetics. Enlist the characteric Give the ideal characteric Explain depilatories	of the following: roperties for cleansing cream. stics for anti-dandruff shampooneteristics for hair colourants. based on thioglycollates. motor agents in shaving propert		
Q.2	a) b)	•	n and manufacturing of lather sh rmanent hair colorants.	naving cream. (08) (07)	
Q.3	a) b)		rfactants used in shampoos. cosmetics and drug formulation	(08) ns. (07)	
Q.4	a) b) c) d)	Write short notes or Cold cream Depilatories Aerosol shaving lath Tooth paste	n any THREE of the following:	(15)	
			SECTION – II		
Q.5	a) b) c) d) e) f)	Mention different ty Enlist various plasti What are cake eyeli Mention role of hen	requirements for baby cosmetic pes of waxes used in lipstick. cizers used for nail lacquers.		
Q.6	a) b)	<u>=</u>	e and psychometric evaluation. out hair waving of cosmetics.	(08) (07)	
Q.7	a) b)	Discuss in detail he Discuss formulation		(08) (07)	
Q.8	a) b) c) d)	Write short notes or Liposomes Baby powder Eye brow pencil Liquid lipsticks	any THREE of the following:	(15)	

T.Y.B.PHARM. SEMESTER-V (2011 COURSE): SUMMER - 2018

SUBJECT: DOSAGE FORM DESIGN - II

Day		Saturday		Time: 10.00 AM TO 01.00 PM	
Date	:	28/04/2018	S-2018-3965	Max. Marks: 80	
N.B.:					
	1)	Q.No.1 and	Q.No.5 are COMPULSORY	Y. Out of the remaining questions	
		attempt AN	Y TWO questions from each	section.	
	2)	Answers to	both the sections should be w	ritten in the SEPARATE answer boo)ks.
	3)	Figures to t	he right indicate FULL marks	s.	
			SECTION	- I	
0.1	A \	Angreen ANV	EIVE of the following.		[10]
Q.1	A) i)		FIVE of the following: ssify dermatological formulati		[10]
	ii)		a major component of capsulo		
	iii)		pathways of skin penetration of	-	
	iv)	What are adva	ntages and disadvantages of so	oft gelatin capsules?	
	v)		e of buffers in semisolid dosa	_	
	vi)	Enumerate va permeation.	rious physiochemical prope	rties of drugs that govern skin	
Q.2	a)	Discuss the de	rmatological factors affecting	selection of bases	[08]
Q.2	b)		_		[00] [07]
					. ,
Q.3	a)	Write down the	e different mechanisms of gel	formation.	[80]
	b)	Discuss variou	s capsule filling methods.	I	[07]
Q.4		Write short no	tes on ANY THREE of the fo	llowing:	[15]
Q.4	a)		nard gelatin capsule	nowing.	[15]
	b)	Penetration enl			
	c)	Defects in caps			
	ď)	Evaluation of s			
			SECTION -	- 11	
0.5	4.5	A A BITT	ENEXTED C41 - C 11 - '		r4 A1
Q.5	A)		FIVE of the following: Ivantages and disadvantages o	•	[10]
	i) ii)		bjectives of granulation?	i acrosors.	
	iii)		ating? Why it is carried out?		
	iv)		nfluencing pulmonary deposit	ion of drugs.	
	v)	What are dilue	nts? Give the ideal properties	of a tablet diluents.	
	vi)	Why methacry	lic acid co-polymers are used	for enteric coating?	
Q.6	a)	Discuss in deta	il dry powder inhalers.	ı	[08]
Q.U	b)		pating of tablets.	_	[07]
			C	·	
Q.7	a)	Discuss in deta	ils IPQC tests for tablets.	[[80]
	b)	Discuss proble	ms in tablet coating and the re	medies thereof.	[07]
0.0		Write short not	tes on ANY THREE of the fo	llowing:	[15]
Q.8	a)	Mouth dissolvi		nowing.	[13]
	a) b)		tests for aerosols		
	c)	Evaluation of c			
	ď)	Powders	·		

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T.Y.B.PHARM. SEMESTER-V (2011 COURSE): SUMMER - 2018 SUBJECT: MEDICINAL CHEMISTRY - I

Day Date	: :	Saturday S-2018-3963 Time: 10.00 AM TO 01.00 Max. Marks: 80		PM	
N. B.	:				
	1)		d Q.No.5 are COMPULSO ach from Section – I and So	DRY . Out of remaining solve any TV	VO
	2)	•		written in the SEPARATE answer l	books.
	3)	Figures to the	he right indicate FULL man	rks.	
			SECTION -	- I	
Q.1		Solve Any FI	VE of the following:		(10)
	a)	Define agonis	t and antagonist		
	b)	Draw the struc	cture of first neurotransmitt	er.	
	c)	Write scheme	of synthesis of carbachol o	r buthenechol.	
	d)	Write about th	e effect of partition coeffic	ient on drug action.	
	e)	Enlist importa	nt uses of neuromuscular b	lockers.	
	f)	List out drawb	backs of Ach as a therapeuti	c agent.	
	g)	Explain impor	tance of conformational isc	omerism in drug action.	
Q.2	a)		lionic agents with a note on of antimuscarinics with its n	neuromuscular nerve transmission.	(10)
	b)	Discuss SAR	of Cholinergic agents.		(05)
Q.3	a)		1 7	parameters that must be considered porate on the stereochemical aspects	(10)
	b)	Discuss biosyı	nthesis, storage, metabolisn	n of Ach.	(05)
Q.4		Write short no	tes on Any THREE of the	following:	(15)
	a)	Stereochemist	ry and biological action		
	b)	Classification	of antimuscarinics with stru	ucture	
	c)	Irresversible a	nticholinesterases		
	d)	Outline synthe	esis of Gallamine and Meca	mylamine	
	e)	Papaverine alk	caloids and their synthetic a	nalogue	

SECTION - II

Q.5		Outline synthesis of Any FIVE drugs of the following:	(10)
	a)	Guanethedine sulphate	
	b)	Salbutamol	
	c)	Prazocin	
	d)	Acetazolamide	
	e)	Terbutaline	
	f)	Methyldopa	
	g)	Isoproterenol	
Q.6	a)	Classify diuretics with examples and explain mode of action of carbonic anhydrase inhibitors.	(10)
	b)	Discuss Chemistry of cardiac glycosides.	(05)
Q.7	a)	Classify cardiotonics with examples and explain in detail about chemistry of cardenolides with mode of action of cardiotonics.	(10)
	b)	Outline biosynthesis of nor-adrenaline through chemical reactions.	(05)
Q.8		Write short notes on Any THREE of the following:	(15)
	a)	Purines and related heterocyclics	
	b)	Mercurial diuretics	
	c)	Direct sympathomimetics	
	d)	Antianginals	
	e)	Combination antihypertensive therapy	

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TY B. Phaym. Sem-I (CB.CS.) 2015 Course: SUMMER-2018 SUBJECT: MEDICINAL CHEMISTRY-I

: Saturday Time: 10:00AM-T01:00PM. : 21-04-2018 Date Max. Marks: 60 5-2018-3927 N. B.: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt any TWO questions 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** Attempt ANY FIVE of the following (10)Enlist various neurotransmitters with their structures. Write IUPAC names, structures and category of i) Physostigmine ii) b) Ticrynafen. Why acetylcholine is poor therapeutic agent. c) Write about structure specific drugs. d) Outline synthesis of for Bethenechol **OR** Hydrochlorthiazide e) f) Write significance of solubility which affect biological action. Discuss SAR of Cholinergic agents. 0.2 a) (07)Write an exhaustive note on potassium sparing diuretics. b) (03)Classify diuretics based on its chemistry. Explain chemistry of sulfonamides. **Q.3** a) (07)Explain reversible anticholine-esterases agents. (03)b) 0.4 Write short notes on ANY TWO of the following (10)Mercurials chemistry a) Outline synthetic schemes of Dicylcomine and Furosemide. b) Chemistry of phenoxy acetic acid derivatives. c) **SECTION II** Attempt ANY FIVE of the following (10)Q.5 Mention important uses of cardiac glycosides. a) Write IUPAC names, structures and category of i) Lobeline ii) Verapamil b) Write about solanaceous alkaloids. c) Outline synthesis for Mecamylamine OR Propranolol d) Write about chemistry of adrenergic nerotransmitters. **e**) Classify neuromuscular blocker on the basis of mechanism of action. f) (07)Classify adrenergic agonist. Discuss SAR of direct sympathomimetics. **Q.6** a) (03)Outline scheme to explain biosynthesis of nor adrenaline. b) Discuss in detail chemistry and mechanism of action of ACE inhibitors and (07)**Q.**7 a)

* * * * *

(03)

(10)

calcium channel blockers.

Anti-arrythmics

Cardiotonics.

Write short notes on ANY TWO of the following

b)

a)

b)

c)

Q.8

Write classification of antihypertensive with examples.

Outline synthetic schemes of Chlorzoxazone and Salbutamol.

T.Y.B.PHARM. SEMESTER-V (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACEUTICAL ANALYSIS - III

N.B.: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining questions solve any TWO questions from each section. 2) Figures to the right indicate FULL marks. 3) Answers to both the sections should be written in SEPARATE answer book. SECTION-I Q.1 Answer any FIVE of the following: (10) a) Enlist variables that lead to band broadening. b) Explain briefly column resolution and retention time. c) Enlist characteristics of an ideal Gas chromatographic detector. d) Discuss in brief Rota meter. e) Explain the term 'HETP'. f) Give the relation between Rf and Rx values. Q.2 a) Enlist all the detectors used in GC. Explain in detail any two. (08) b) Explain in detail Plate Theory and Rate Theory. (07) Q.3 a) Write a detail columns used in GC. b) Discuss the various elution techniques used in chromatography. Add a note on applications of GC. Q.4 Write short notes on any THREE of the following: (15) a) Classification of chromatographic methods b) Temperature programming in GC c) Types of carrier gas in GC d) Advantages of GC SECTION-II Q.5 Solve any FIVE of the following: (10) what are the various techniques used for detecting colourless spots in paper chromatography. b) Give types of solvents used in paper chromatography? c) What are the advantages offered by paper chromatography? d) Importance of Rf value in chromatography. e) Give preparation of gels in gel permeation chromatography. f) Give physical properties of Ion exchange resins. Q.6 a) Explain different development modes in Paper chromatography.	Day: Date:		uesday 1/04/2018	S-2018-3964	Time: 10.00 AM TO Max. Marks: 80	O 01.00 PM
solve any TWO questions from each section. 2) Figures to the right indicate FULL marks. 3) Answers to both the sections should be written in SEPARATE answer book. SECTION-I Q.1 Answer any FIVE of the following: a) Enlist variables that lead to band broadening. b) Explain briefly column resolution and retention time. c) Enlist characteristics of an ideal Gas chromatographic detector. d) Discuss in brief Rota meter. e) Explain the term 'HETP'. f) Give the relation between Rf and Rx values. Q.2 a) Enlist all the detectors used in GC. Explain in detail any two. b) Explain in detail Plate Theory and Rate Theory. (07) Q.3 a) Write a detail columns used in GC. b) Discuss the various elution techniques used in chromatography. Add a note (05) on applications of GC. Q.4 Write short notes on any THREE of the following: a) Classification of chromatographic methods b) Temperature programming in GC c) Types of carrier gas in GC d) Advantages of GC SECTION-II Q.5 Solve any FIVE of the following: a) What are the various techniques used for detecting colourless spots in paper chromatography. b) Give types of solvents used in paper chromatography? c) What are the advantages offered by paper chromatography? d) Importance of Rf value in chromatography. e) Give preparation of gels in gel permeation chromatography. f) Give physical properties of Ion exchange resins.	N.B.:	1)	O No 1 and O	N. 5 COMPUL CODY O	6.1	
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f) Give physical properties of Ion exchange resins.		d)	-			
		•		-	raphy.	
O.6 a) Explain different development modes in Paper chromatography. (08)		I)	Give physical prope	erties of Ion exchange resins.		
	Q.6	a)				(08)
b) Describe principle of separation in Ion exchange chromatography. Explain (07)		b)			romatography. Explain	(07)
with suitable example.			with suitable examp	ple.		
Q.7 a) Explain the mechanism of Gel permeation chromatography. Add a note on (08)	Q. 7	a)			graphy. Add a note on	(08)
applications of Gel Permeation chromatography.		_			' O.D. 1.'	(05)
b) What are the various types of separation by electrophoresis? Explain any one. (07)		b)	What are the variou	s types of separation by electropho	resis? Explain any one.	(07)
Q.8 Write short notes on any THREE of the following: (15)	Q.8					(15)
a) Sketch of schematic of capillary electrophoresis system and label major		a)		ic of capillary electrophoresis sy	stem and label major	
components		1.	-	ahuamata ayarke		
b) Advantages of Gel chromatographyc) Pharmaceutical Applications of Ion- Exchange chromatography		,			atography	
d) Explain the principle of separation by Paper chromatography		•				

		SU	JBJECT : PHARMACEUTICA	L ANALYSIS-III		
Day Date		Tuesday 24/04/2018	S-2018-3928	Time: Max. Marks: 60	10.00 AM TO	O 01.00 PM
N. B.	:					
	1) 2) 3)	Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining solve An TWO questions each from section-I and Section-II. Answers to the two sections should be written in SEPARATE answer books Figures to right indicates FULL marks.				
			SECTION-I			·····
Q. 1	a)b)c)d)e)f)	Attempt any FIVE of the following Define the terms Chromatography and Partition coefficient. Explain the relation between Theoretical plates (N) and HETP (H). Give the principle of Column chromatography. How will you separate mixture of Amino acid by Paper chromatography. Give advantages and disadvantages of Paper chromatography. Channeled or loosely packed columns provide poor or insufficient separation, Why?				(10)
Q. 2	a)	Describe the	Rate theory of chromatography ir	detail.		(07)
	b)	Give advanta	ages and disadvantages of Column	chromatography.		(03)
Q. 3	a)	Discuss in de	etail packing techniques in Colum	n chromatography.		(07)
	b)	Give the vari	ous stationary phases used in Pap	er chromatography.		(03)
Q. 4	a) b) c)	Column char Applications	notes on any TWO of the following racteristics of Paper chromatography of chromatography	ng		(10)
			SECTION-II			
Q. 5	a)b)c)d)e)f)	Define the te Give the assa Give the idea Write note of Give advanta	FIVE of the following rms Retention time and Retention ay procedure for Hyoscine hydrobal properties of carrier gas used in separation of cations by Ion excludes of Gel permeation chromatog in sample injection system in Gas of	romide injection. Gas chromatograph hange chromatograp raphy.		(10)
Q.6	a)	Discuss the p	principle of Gas chromatography. ography.	Add a note on colu	ımns used in	(07)
	b)	Discuss the a	applications of Gas chromatograph	ny.		(03)
Q.7	a)	What do you of Ion exchan	n mean by Ion exchange chromatenge resins.	ography. Explain in	detail types	(07)
	b)	Explain in br	rief Ion exchange capacity.			(03)
Q. 8	a) b) c)	Temperature Applications	notes on any TWO of the following programming system in Gas chrown of Ion exchange chromatography used in Gel permeation chromatography	mmatography		(10)

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SUBJECT: PHARMACEUTICAL JURISPRUDENCE

Day: Time: 10.00 AM TO 01.00 PM Monday Date: Max. Marks: 60 07/05/2018 S-2018-3932 N.B: Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt ANY 1) **TWO** questions from Section-I and Section -II. Answers to both the Sections should be written in **SEPARATE** answer sheets. 2) Figures to the right indicate FULL marks. 3) Draw a neat and labeled diagram WHEREVER necessary. 4) **SECTION - I** (10)**Q.1** Answer **ANY FIVE** of the following: What is "First register" & "Subsequent Register" under Pharmacy Act 1948? a) Define Schedules FF, X, S and M-I. What are the functions of Drugs Technical Advisory Board? c) d) Define Drug and New Drug as per Drugs and Cosmetics Act 1940. e) Describe in brief provisions pertaining to import of drugs for personal use. Define Coca derivative and Opium derivative. f) Describe general conditions for grant or renewal of license to manufacture of (07) Q.2 a) drugs other than those specified in Schedule C, C1 and X. Define Medicinal Preparations and Toilet Preparations and add a note on (03) b) objectives of Medicinal and Toilet Preparations Act 1955. (07)Write brief account on Government Analyst. Q.3 a) Discuss in detail the objectives and salient features of Drugs and Magic (03)Remedies Act 1976. (10)Write a short note on **ANY TWO** of the following: **Q.4** a) State and Joint State Pharmacy Council Poisons Act 1919 b) Provisions for classes of Prohibited Advertisement and Exempted Advertisement **SECTION - II** (10)Answer **ANY FIVE** of the following: Q.5 What types of leaves are entitled for the employee under Shops and a) Establishments Act? Discuss Offences and Penalties under Prevention of Cruelty to Animals Act. b) Define Copyrights and Geographical Indications. c) Define Insecticide and Misbranded Insecticide. d) Enlist different Consumer Dispute Redressal Agencies. e) Explain criteria for obtaining a Patent. f) Describe the professional ethics in pharmacy practice framed by Pharmacy (07)Council of India. Describe the constitution and functions of Animal Welfare Board of India. (03)b) Describe in detail different mechanisms for protection of Intellectual Property (07)Q.7 aRights. Write procedure for analysis of Food Adulteration as per Act 1954. (03)b) (10)Write a short note on **ANY TWO** of the following: Q.8 Constitution and Functions of Central Insecticides Board a) Food Inspector b) USFDA and MHLW c)

SUBJECT: PHARMACEUTICAL TECHNOLOGY-I

Time: 10.00 AM TO 01.00 PM Saturday Day S-2018-3929 28/04/2018 Date Max. Marks: 60 N. B.: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt any **TWO** questions from each section. Both the sections should be written in **SEPARATE** answer books. 2) Figures to the RIGHT indicate full marks. 3) **SECTION-I** Answer the following (ANY FIVE) 0.1 (10)Give the requirements of building and water supply according to GMP. Give flow chart for liquid manufacturing process. b) Define Intrinsic solubility. c) Write importance of GMP in pharmacy. d) What are different characteristics of well formulated suspension? e) Enlist the preformulation tests to be conducted on new drug substance. f) Q.2Discuss in detail formulation of emulsion. (07)a) Explain in brief the effect of temperature on stability of drug. (03)Q.3 a) What is zeta potential? How zeta potential affect on flocculation of dispersed (07)particles of suspension? b) Explain polymorphism and crystal properties. (03)Write short note on any **TWO** of the following: **Q.4** (10)a) Evaluation of monophasic liquids b) Theories of emulsification Good practices in quality control **SECTION-II** Answer the following (ANY FIVE) Q.5 (10)Write down the names of gelling agents and mention the type of gels. Mention the ideal properties of sunscreen agent. What are the ideal characteristics of penetration enhancer? c) What is epilation and depilation? d) Mention the various plasticizers used for nail lacquers. e) What are the liquid mascaras? f) Discuss the formulation and manufacturing of Lipsticks. (07)Q.6 a) Why rheological property of semisolid product is important? b) (03)Discuss the formulation of shampoos. (07)**O.**7 **b)** Explain the semi-permanent hair dyes. (03)Write short note on any **TWO** of the following: (10)0.8 a) Nail paint removers **b)** Moisturizing lotions c) Evaluation of semisolids.

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T.Y.B.PHARM. SEMESTER-V (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACOGNOSY-I

Day Date	:	Monday 07/05/2018	S-2018-3969	Time: 10.00 AM TO 0 Max. Marks: 80.	1.00 P
N.B.:	1) 2)	Q. No. 1 a questions for Section-I at Figures to the section of th	nd Q. No. 5 are COMPULSORY. Our section-I and any TWO questions and Section-II should be written in SEPA the right indicate FULL marks. labeled diagrams WHEREVER necess	from Section-II ARATE answer book.	rwo
			SECTION-I		
Q.1	Atto a) b) c) d) e) f) g)	Write any What is sto What is or Which is b What is tax Which are	VE of the following: four drugs obtained from mineral source matal index? ganoleptic drug evaluation? sest time for collection of bark? konomic classification? different naturally occurring auxins? ulteration?	e.	(10)
Q.2	a) b)		erapeutic classification of crude drug? E etail storage of crude drug.	Explain in detail.	(08) (07)
Q.3	a) b)		erent steps involved in collection of crucolant growth stimulants? Explain.	de drug.	(08) (07)
Q.4	Wria) b) c) d)	Ethylene of Scope of p Abscisic a	harmacognosy cid pharmacognosy		(15)
			SECTION-II		
Q.5	Atto a) b) c) d) e) f) g)	What is eff Which are Write any Write diffe What is pr What is 'tr	WE of the following: fect of aromatherapy on body? different compounds/ isotopes used in four drugs used in TCM. erence between Shark liver oil and Cod inciple of Homoeopathy? ridosh' theory? different secondary metabolites? Give	liver oil.	(10)
Q.6	a) b)		ipids? Give biosynthesis, chemistry, classetail Shikkimic acid pathway in plant bi		(08) (07)
Q. 7	a) b)	biosynthet	racer techniques? Explain role of tracer ic pathway. etail Mevalonic acid pathway.	techniques in elucidation of	(08) (07)
Q.8	Wr a) b) c) d)	ite short note Sun flowe Arachis oi Acacia Agar			(15)

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T.Y.B.PHARM. SEMESTER-V (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACOLOGY-II

Time: 10.00 AM TO 01.00 PM Day: Friday Date: 04/05/2018 S-2018-3968 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) Answer to the two sections should be written in **SEPARATE** answer books. Figures to the **RIGHT** indicate full marks. 3) **SECTION-I** Q.1 Answer ANY FIVE of the following: (10)Enlist adverse effect of iron preparations a) b) Enlist therapeutic uses of anticoagulants Classify hypolipoproteinemic drugs. c) d) Enlist the types and therapeutic uses of vasodilators e) What is the mechanism of action of cardiac glycosides Enlist the membrane stabilizing agents with suitable examples. f) **Q.2** Describe synthesis, pharmacology, therapeutic uses and adverse effects of a) (08)nitric oxide. b) Classify antiarrhythmic drugs with suitable examples. Explain mechanism of (07)action and pharmacology of class II antiarrhythmic drugs. Explain pharmacology, adverse drug reactions and contraindications of Q.3 (08)calcium channel blockers. b) Explain pharmacology and therapeutic uses of aldosterone antagonists. (07)Write short notes on ANY THREE of the following: **Q.4** (15)Angiotensin antagonists a) Erythropoietin b) Potassium channel openers c) d) HMG-CoA reductase inhibitors. **SECTION-II** Answer ANY FIVE of the following: Q.5 (10)Enlist therapeutic uses of diuretics a) Classify antidiarrhoeal drugs with suitable examples b) Explain mechanism of action of ulcer protective agents c) What are antacids? Classify with suitable examples d) Describe the composition and rationale of ORS e) Classify antitussive drugs with suitable examples f) Classify antiasthmatic agents. Explain the mechanism of action, (08)**Q.6** pharmacology, adverse drug reactions of Beta 2 sympathomimetics Discuss the mechanism of action, pharmacology and therapeutic uses of (07)b) thiazide diuretics. What are antiemetics? Classify antiemetics with suitable examples. Discuss **Q.7** (08)a) the pharmacology of 5-HT3 antagonists. Explain the principles of management of pharmacotherapy of diarrhoea. b) (07)Write short notes on **ANY THREE** of the following: **Q.8** (15)Leukotriene antagonists a) b) H₂ antagonists Antidiuretic hormone c) Mucolytics d)

10.00 AM TO 01.00 PM

SUBJECT: DOSAGE FORM DESIGN-III

Time: Day Friday S-2018-3972 27/04/2018 Max. Marks: 80 Date N.B.: Q.No.1 and Q.No.5 are COMPULSORY. Out of the remaining questions 1) attempt ANY TWO questions from each section. Answers to both the sections should be written in **SEPARATE** answer books. 2) 3) Draw neat and labeled diagram WHEREVER necessary. Figures to the right indicate FULL marks. 4) SECTION - I Q.1 Solve ANY FIVE of the following (10)Enlist various parenteral routes of administration. What is nitrogen storage system in industry? b) What is thermal death time? c) Comment on sodium equivalency test. d) How clarity test of injectibles is performed? Show schematically environmental zones in a parenteral section. State merits of Plackett- Burman factorial design. Give a detailed account of HVAC system in a parenteral section. (08)**Q.2** Discuss formulation of formulation of sterile solutions for injectibles. (07)Discuss sterile suspensions and dry powders for suspensions. (08)Q.3 Discuss sterility testing of parenterals. (07)Write notes on (ANY THREE) **Q.4** (15)Pyrogen elimination and testing. a) b) Batch Vs continuous operation. Dynamic storage of sterile water for Injection. c) d) Physicochemical properties in design of SVP's. Additives in Injectibles. **SECTION-II** (10)Answer ANY FIVE of the following: Q.5 Concentrated RBCs. a) b) Formulation of contact lens solution. Siliconization of glass containers. c) Coring of rubber closures. d) Use of acid citrate dextrose in blood collection units. e) What are sleeve stoppers? f) Explain optical property of glass containers. g) (08)Give a detailed account of types of glass. Q.6 a) Discuss composition of rubber closures. Discuss galvanization process. (07)b) Discuss formulation of multiple electrolytes LVP. (08)a) **Q.7** (07)Discuss GMP in a parenteral section. b) (15)Write notes on (ANY THREE) **Q.8** Plasma fractions. a) FFS technology. b) Evaluation of HEPA filters. c) Cost effectiveness of LVP. d) Mechanical properties of plastic containers.

2018

SUBJECT: MEDICINAL CHEMISTRY - II Time : 10.00 AM TO 01.00 PM Day Friday Date Max. Marks: 80 20/04/2018 S-2018-3970 N.B. Q.1 and Q.5 are COMPULSORY. Out of the remaining, attempt any TWO 1) questions from Section – I and any TWO questions from Section – II. 2) Figures to the right indicate FULL marks. 3) Draw diagrams, structures and give reactions wherever necessary. SECTION - I Attempt any FIVE questions of the following: 0.1 (10)a) Outline synthesis of Phenobarbital sodium. b) Draw structures of any two short acting barbiturates. c) Explain mode of action of benzodiazepines. d) Explain the terms Grand mal and Petit mal epilepsy. List of characteristic features of Phase – II metabolic reactions. Give any four major uses of Sedatives-Hypnotics. f) Draw structures of any two halogen containing hydrocarbons from general anesthetic category. **Q.2** What is the goal of metabolism? Explain two main types of metabolic (05)Describe oxidation pathways for Carbon-Heteroatom systems giving with (10)examples. Classify Sedatives and Hypnotics giving one representative structure for each Q.3 (05)a) Discuss SAR of barbiturates. (05)Add a note on nonbarbiturates. (05)Write short notes on any THREE: Q.4 (15)Theories of general anesthesia b) Classification and chemistry of anticonvulsants c) Acetylation and Methylation d) Factors affecting metabolism e) Hallucinogens SECTION - II (10)Q.5 Attempt any FIVE questions of the following: Give the structures of 2 natural compounds from which local anesthetics were a) designed. b) Write synthesis of Lignocaine. c) Draw any two structures of benzodiazepine class of anxiolytics. d) Explain what do you mean by EPS? e) Draw any two structures of Tricyclic antidepressants. f) Draw the structures of products of partial hydrolysis of cocaine. g) Outline the synthesis of Benzocaine. Classify psychotherapeutic drugs with example. Give mode of action and SAR (10) **Q.6** phenothiazines. Describe in short local anaesthetic agents from ester category. (05)b) (08)Classify prodrugs with examples explaining each term. **Q.7** a) Explain pharmacokinetic applications of prodrugs. (07)Write short notes on any THREE: (15)0.8

Tricyclic antidepressants a) Anxiolytics b)

c) Mode of action of local anesthetics

Outline synthesis of Imipramine and Diazepam d)

Outline synthesis of Chlorpromazine and Haloperidol

SUBJECT: MEDICINAL CHEMISTRY-II

Time: 10.00 AM TO 01.00 PM Day: Friday S-2018-3933 Date: Max. Marks: 60 20/04/2018 N.B: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt ANY TWO questions from each Section. 2) Answer to the both sections should be written in **SEPARATE** answer book. Figures to the right indicate FULL marks. 3) **SECTION-I Q.1** Attempt ANY FIVE of the following: (10)a) Mention similarities & difference between barbitures & benzodiazepines. Write therapeutic applications of non barbiturates. b) Write physiology of sleep. c) Outline scheme of synthesis of Methohexital Sodium. d) e) Draw Structure mention category of Amobarbital & Propofol f) Draw Structure mention category IUPAC of Triazolam &Oxazepam. Q.2 a) Discuss SAR of Sedatives & Hypnotics with Classification. (07)Define metabolism. Write a note on Phase II metabolic pathways with (03) examples. $\mathbf{Q.3}$ a) Write types of epilepsies with note on chemistry of oxazolidinediones & (07) hydantoins. State ideal properties of general anesthetics with its MOA. b) (03)0.4 Write short notes on ANY TWO of the following: (10)Antiparkinson's agents. a) **b)** Factors affecting metabolism. Outline scheme of synthesis of Thiamylal sodium & Ketamine Hydrochloride. **SECTION-II** Attempt ANY FIVE of the following: **Q.5** (10)Draw Structure mention category of Desipramine & Clozapine. a) b) State uses of diphenylbutyl piperidines. Write biogenic theory of depression. c) d) Draw Structure mention category IUPAC of Dibucaine & Chlorpromazine. Outline scheme of synthesis of lignocaine. e) Comment on anxiolytics. f) What are local anesthetics? Give chemical classification with example. (07) Q.6 a) Explain SAR with examples. Discuss about non barbiturates. (03)b) Classify antipsychotics. Write classification & SAR with examples. (07)**Q.7** a) Write a note on COMT inhibitors. (03)b) Write short notes on ANY TWO of the following: (10)Q.8 a) MAO inhibitors Chemistry & SAR. b) Classification of prodrugs. Outline scheme of synthesis Meprobamate & Doxepin.

SUBJECT: PHARMACEUTICAL ANALYSIS - IV

Day Date	:	Monday 23/04/2018	S-2018-3971	Time: 10.00 AM TO 01.0 Max. Marks: 80	00 PM
N. B.	 Q.No.1 and Q. No.5 are COMPULSORY. Out of the remaining questions attempt Any TWO from each section. Answers to both the sections should be written in SEPARATE answer books. Figures to the right indicate FULL marks. 				
			SECTION	- I	
Q.1	a) b) c) d)	Why TLC is su What are the di How adsorbent Compare between	and solvents are selected een HPLC and HPTLC.	olumn packing in HPLC technique. in TLC.	(10)
Q.2	e) f)	Give the basic	Ivantages and disadvantages requirements for TLC.		(07)
	b)	Describe the p HPLC.	roperties and working of	UV and refractive index detectors	in (08)
Q.3	a)b)		es factors affecting separate and page adsorption TLC and page		(07) (08)
Q.4	a)b)c)d)	Reciprocating J Visualizing age Guard column	ents used in TLC in HPLC	following: rse phase chromatography	(15)

SECTION - II

Q.5		Solve Any FIVE of the following:	(10)			
	a)	Enlist the supercritical fluids. Give their properties.				
	b)	Define the terms R_f value and R_x value.				
	c)	Give the adulterants found in turmeric powder.				
	d)	How instrumentation for SFC does differ form Gas chromatography.				
	e)	Give the advantages of HPTLC technique.				
	f)	Give the role of food inspector in food analysis.				
Q.6	a)	Discuss in detail the applications of HPTLC.	(07)			
	b)	Explain in detail the principle and instrumentation of SFC.	(08)			
Q.7	a)	Write in detail the advantages and limitations of supercritical fluid chromatography.	(07)			
	b)	Discuss in detail the development techniques of HPTLC in detail.	(08)			
Q.8		Write short notes on Any THREE of the following:	(15)			
	a)	Selection of adsorbent and solvent in HPTLC				
	b)	Determination of adulterants in milk and milk products				
	c)	Compare between band and spot applications of sample				
	d)	Responsibilities and duties of food safety officer.				
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SUBJECT: PHARMACEUTICAL ANALYSIS-IV

Day: Date:		Monday 23/04/2018	BJECT: PHARMACEUTICAL ANA S-2018-3934	Time: 10.00 AM TO 01 Max. Marks: 60
N.B:				
	1) 2) 3)	questions Figures to	and Q. No. 5 are COMPULSORY . Out each from Section-I and Section-II. the right indicate FULL marks. the both section should be written in S .	•
			SECTION-I	
Q.1		Attempt ANV	FIVE of the following:	(10)
χ	a)	_	Evalue for a spot on a TLC Plate calcu	
	b)	What is HPTL		
	c)	What could ha	ppen if you spot too much of compound	<u>*</u>
	d)		ical tests for checking adulterants in Te	
	e) f)		ntages of band applications of sample ov nappen if your solvent level is above	1 11
Q.2	a)		rious applications of TLC.	(07)
	b)	Give the advan	ntages and disadvantages of TLC.	(03)
Q.3	a)	Explain in deta	ail development techniques in TLC.	(07)
2. 0	b)	-	ical tests for checking adulterants in spi	• • • • • • • • • • • • • • • • • • • •
Q.4	a) b) c)	Role and respo	tes on ANY TWO of the following: onsibilities of Food Inspector d disadvantages of HPTLC ed in TLC	(10)
			SECTION-II	
Q.5		Attempt ANY	FIVE of the following:	(10)
	a)	Explain the ter	m Critical point in SFC.	
	b)		n Selectivity factor and Retention time.	whose LIDI C2
	c) d)		Phase HPLC is used more than Normal property and solute	
	e)		rties of Carbon dioxide as a supercritica	
	f)	Give the difference	ence between Isocratic and Gradient elu	ution techniques.
Q.6	a)	Discuss in deta	il pumps used in HPLC.	(07)
	b)	Explain in brie	f about sample injector system in HPLC	C. (03)
Q. 7	a)		trumentation of SFC.	(07)
	b)	Give in brief th	ne principle involved in SFC.	(03)
Q.8			tes on ANY TWO of the following:	(10)
	a)	Columns used		
	b) c)	Applications of UV- Visible De	t SFC etector and RI Detector in HPLC	
	~,	C, TISIOTO D	color and re Detector in the De	

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SUBJECT: PHARMACEUTICAL BIOTECHNOLOGY (Including Molecular Biology)

Day Date	:	Thursday 03/05/2018 S-2018-3	Time: 10.00 AM TO 01.00 PM Max. Marks: 80.
N.B.	2) 3) 4)	Q. No. 1 and Q. No. 5 are CO TWO questions from Section- Both the sections should be wr Figures to the RIGHT indicate	OMPULSORY. Out of the remaining attempt any I and any TWO questions from Section-II. itten in SEPARATE answer books. e full marks.
		SEC	CTION-I
Q.1	An a) b) c) d) e) f)	what is a palindrome? Define mutation in DNA. Give life cycle of T4 phage. Write the differences in prokaryote What are f [†] and f ⁻ bacteria? Enlist five rDNA products. What is reverse transcriptase?	and eukaryote.
Q.2	De	scribe transduction and write a note of	on significance of DNA transfer. (15)
Q.3	Dis	scuss the role of biotechnology in pha	armaceutical sciences. (15)
Q.4	Wr a) b) c) d) e)	ite short notes on any THREE of the Replication of lagging strand EtBr Watson and Crick's model of DNA Thermal cycler Southern blotting technique	
		SI	ECTION-II
Q.5	An: a) b) c) d) e) f)	what is strain improvement? Give applications of enzyme in text What is fermentation media? Draw a diagram of tubular flow rea What is disulfide bond in protein? Define enzyme. What is a single cell protein?	•
Q.6	Wh	at are objectives of protein engineer	ng? (15)
Q. 7	Wh	at are bioreactors? Describe a fed-ba	tch reactor. (15)
Q.8	Wr a) b) c) d) e)	ite short notes on any THREE of the Applications of amylase Lyophilization Site directed mutagenesis Enzyme immobilization Parameters affecting enzyme activi	

SUBJECT: PHARMACEUTICAL BIOTECHNOLOGY

Day Date	:	Thursday 03/05/2018 S-2018-39	Time : 10.00 AM TO 01.00 PM Max. Marks : 60
N.B.			
 Q.No. 1 and Q.No.5 are COMPULSORY. Out of attempt ANY TWO questions from each section. Answers to both the sections should be written in Section at the right indicate FULL marks. 			m each section. Id be written in SEPARATE answer book.
		SECTION	ON – I
Q.1	a) b) c) d) e) f)	Answer ANY FIVE of the following: Define transduction. What are purins and pyrimidins? Differentiate between coding and non-What is action of a polymerase? What is a promoter? Draw a neat diagram of tRNA. What is reverse transcriptase?	coding DNA.
Q.2		Write in detail about DNA replication	[10]
Q.3		What is central dogma of molecular translation.	biology? Describe transcription and [10]
Q.4	a) b) c)	Write short notes on ANY TWO of the F+ plasmid and conjugation Mutation in sickle cell Applications of PCR	e following: [10]
		SECTIO	DN – II
Q.5	a) b) c) d) e) f)	Answer ANY FIVE of the following: What is shake flask culture? Define strain improvement. What is an inoculum? Draw labeled diagram of immunoglob What is a single cell protein? Enlist factors affecting enzyme activity. Differentiate between fermentor and b	7.
Q.6		Write an essay on innate immune syste	em. [10]
Q. 7	a) b)	What are monoclonal antibodies? How Applications of monoclonal antibodies	• • •
Q.8	a) b) c)	Write note on ANY TWO of the follow Sandwich ELISA Western blotting Fluidized bed reactor	wing: [10]

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SUBJECT: PHARMACEUTICAL TECHNOLOGY-II

Time: 10.00 AM TO 01.00 PM Day: Friday Date: 27/04/2018 Max. Marks: 60 S-2018-3935 N.B: Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt ANY 1) TWO questions from each Section. 2) Answer to the both sections should be written in **SEPARATE** answer book. Figures to the right indicate FULL marks. 3) SECTION -I **Q.1** Attempt ANY FIVE of the following: (10)Write productivity of tablet manufacturing per shift using 16 station rotary a) tablet machine Give the classification of powders according to particle size analysis b) Explain the significance of angle of repose and Carr's index test for granules c) Mention the essential characteristics of good face powder d) What are superdisintegrants? Give examples e) Water sensitive, unpleasant taste drug is to be formulated for patient of 3 years f) age, suggest suitable formulation. Discuss the construction and working of high speed granulator. 0.2 a) (07)Granules containing 10% moisture observes compression problems. Explain (03) b) various related problems and remedies. Tablet containing 800 mg drug is to be compressed, comment on possible Q.3 a) (07)difficulties in compression, ejection, patient compliance and ways to overcome these problems. Suggest face powder formulation to be used in humid and tropical weather (03)conditions **Q.4** Write short note on **ANY TWO** of the following: (10)Oral antibiotic dry syrup a) Tooth powder b) Effervescent tablets c) **SECTION-II Q.5** Attempt ANY FIVE of the following: (10)Enlist problems encountered in tablet coating a) Why gelatin is major component of capsule shell manufacturing? b) What are the advantages and disadvantages of soft gelatin capsule? c) What are the physiological considerations behind suppository formulation? d) Explain disintegration test of tablets e) Why do we measure the hardness of tablet? f) What are suppositories? Explain the selection criteria for suppository base. (07)Q.6 a) Write possible defects in hard gelatin capsules filled with hygroscopic drugs (03)b) Discuss the different equipments used in tablet coating (07)Q.7 a) Drug content test for 20 Paracetamol 500 mg/tablet is 455 mg. Does it passes (03)b) the test? **Q.8** Write short note on **ANY TWO** of the following: (10)Film coating of tablet a) Evaluation of hard gelatin capsule Method of preparation of suppositories

T.Y.B.PHARM. SEMESTER-VI (2011 COURSE) : SUMMER - 2018 SUBJECT: PHARMACOGNOSY-II

Day Date	:	Saturday S-2018-3975 05/05/2018	Time: 10.00 AM TO 01.00 I Max. Marks: 80	PM	
N.B.	:				
	1)	Q.No.1 and Q.No.5 are COMPULSO	RV Out of remaining quartiers and		
	_ `	And I wo questions from each section)n		
	2)	Answers to both the sections should be	written in SEPARATE answer boo	ks.	
	3)	Figures to the right indicate FULL ma	rks.		
		SECTION -	- I		
Q.1		Attempt ANY FIVE of the following		(10)	
	a) Write two main chemical constituents present in Onion and draw structure				
		the same.			
	b)	Write biological source and health benefits	e and health benefits of Arnica.		
	c)	What are neutraceuticals? Give examples.			
	d) e)	Write preparation of decoction. Write principle of TLC.			
	f)	Which are different advantages of HPTLC	in Herbal drug analysis?		
Q.2	a)	Which are different types of extracts? How	v will you standardize extracte?	(00)	
۷	b)	Applications of column chromatography in	herbal drug analysis.	(08) (07)	
	ŕ			(01)	
Q.3	a)	Write note on continuous hot extraction.		(08)	
	b)	Explain Successive Solvent Extraction and	supercritical fluid extraction.	(07)	
Q.4		Write note on ANY THREE of the followi	na	(15)	
ų. 4	a)	HPLC in herbal drug analysis	ng	(13)	
	b)	Cucumber			
	c)	Health benefits of Fenugreek			
	d)	Garlie			
	e)	pH gradient technique for alkaloid extraction	'n		
		SECTION -	П		
Q.5		Attempt ANY FIVE of the following		(10)	
	a)	Explain principle of totipotency.			
	b)	Give salient features of protoplast fusion.			
	c)	What is callus?	are for herbal shamnon?		
	d)	Which are different quality control paramet Write composition of herbal skin moisturize	er		
	e) f)	Explain history of tissue culture techniques			
	1)	•		(08)	
Q.6	a)	Explain the significance of quality control of	of herbal cosmetics. Give quality	(00)	
		control of any skin care product. Write different strategies for enhancement	of secondary metabolite production	(07)	
	b)	from plant cells.	of secondary means the f	, ,	
Q .7	a)	Explain WHO guidelines for standardization	n of herbal drugs.	(08)	
~~	b)	Give applications of different hair care prep	parations with examples.	(07)	
		ANN PITTER of the follows	no:	(15)	
Q.8		Write note on ANY THREE of the following Composition of tissue culture media	116.	` '	
	a)	Requirements for set up of tissue culture la	boratory		
	b) c)	Production of Hair care products			
	d)	Role of growth hormones in PTC			
	e)	Various hair dyes in herbal cosmetics	*		
		* * ^			

TYB. Pharm-sem-VI (CBCS) 2015 Course: SUMMER-2018

SUBJECT: PHARMACOGNOSY – III

5-2018-3936

Day: Monday

Time: 10:00 AM TO 1:00 PM

Date: 30-04-2018

Max. Marks: 60

N. B.:

- 1) **Q. No. 1** and **Q. No. 5** are **COMPULSORY**. Out of the remaining attempt any **TWO** questions from each section.
- 2) Figures to the right indicate **FULL** marks.
- 3) Answers to both the sections should be written in **SEPARATE** answer book.

SECTION-I

Q.1 Answer any FIVE of the following:

(10)

- a) Give examples of auxins.
- **b)** What is adulteration?
- c) Define totipotency.d) Give the parameters consider in microscopical evaluation of crude drugs.
- e) What is Foaming Index?
- f) Enlist various leaf constants.
- Q.2 Explain the application of plant tissue culture. Give various strategies used for the enhancement of phytopharmaceuticals from plant tissue culture.
- Q.3 a) Describe physical and chemical methods of evaluation of crude drugs and give (06) their advantages and limitations.
 - b) Give the salient features of WHO guidelines for the standardization of crude (04) drugs.
- **Q.4** Attempt any **TWO** of the following:

(10)

- a) Determination of Pesticides residue
- b) Define and Differentiate between moisture content and loss on drying.
- c) Morphological evaluation of crude drugs
- d) Biological evaluation of crude drugs

SECTION-II

Q.5 Answer any FIVE of the following:

(10)

- a) What is Percolation?
- **b)** Define and differentiate between natural fibers and synthetic fibers with examples.
- c) Give biological source and chemical constituents of Ashoka.
- d) Give biological source and chemical constituents of Bramhi.
- e) Give biological source and chemical constituents of Ashwgandha.
- **f)** What is successive extraction?
- Q.6 Write an exhaustive note on extraction. Explain the principle, application and (10) advantages of ultra sound assisted extraction.
- Q.7 a) Write exhaustive note on microwave assisted extraction.

(06)

- b) Write the pharmacognostic details, toxicity and marketed formulations of (04) Amla.
- **Q.8** Attempt any **TWO** of the following:

(10)

- a) Give well labeled diagram of soxhlet apparatus
- **b)** Calamine
- c) Kaolin
- d) Shatavari

* * * *

SUBJECT: PHARMACOLOGY-III

10.00 AM TO 01.00 PM Time: Day Saturday S-2018-3938 05/05/2018 Max. Marks: 60 Date N.B.: Q.No. 1 and Q.No. 5 are COMPULSORY. Out of the remaining questions 1) attempt ANY TWO questions from each section. Answers to both the sections should be written in the **SEPARATE** answer books. 2) 3) Figures to the right indicate FULL marks. SECTION - I [10]0.1 Answer **ANY FIVE** of the following: What are CNS stimulants? a) b) Why levodopa is combined with carbidopa? c) Enlist the techniques of local anaesthetics. d) Give the adverse effects of phenytoin. e) Enlist the adverse effects of MAO inhibitors. f) Differentiate tonic clonic and absence seizures. g) Explain the mechanism of action of nitric oxide. Classify sedatives and hypnotics. Explain the pharmacology, therapeutic uses, [10] **Q.2** adverse effects and drug interactions of barbiturates. [06]a) Describe the pharmacology of levodopa. $\mathbf{Q.3}$ [04] b) Explain the stages of general anaesthesia. **Q.4** Write a short notes on **ANY TWO** of the following: [10] a) Lidocaine b) Selective serotonin reuptake inhibitors c) Cognition enhancers SECTION - II Q.5 Answer **ANY FIVE** of the following: [10] Enlist the symptoms of barbiturate poisoning. b) What are non sedative antihistaminics? Give examples. c) Brief the triple response of histamine. d) Give two examples of leukotrines antagonist. e) Enlist the symptoms of alcohol poisoning. f) Give two examples of opoid agonist and antagonist. **g)** Define antipyretics and analgesics. **Q.6** Classify NSAIDS with examples. Describe the Pharmacology of Aspirin. [10]a) Describe the pharmacotherapy of gout. $\mathbf{Q.7}$ [06] b) Describe the sign, symptoms and treatment of arsenic poisoning. [04]Q.8 Write short notes on **ANY TWO** of the following: [10] a) PAF b) Rheumatoid arthritis c) Prostaglandins

T.Y.B.PHARM. SEMESTER-VI (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACOLOGY - III

Day Date	:	Monday 30/04/2018	S-2018-3973	Time: 10.00 AM TO 01.00 F Max. Marks: 80	M					
N.B.:	1)	Q.No.1 and Q.No.5 are COMPULSORY. Out of the remaining questions attempt ANY TWO questions from each section. Answers to both the sections should be written in SEPARATE answer books.								
	3)	Figures to	Figures to the right indicate FULL marks.							
	SECTION – I									
Q.1	a) b) c) d) e) f)	Classify antia Define sedative Explain the management of the stage Enlist the stage Differentiate of Classify anti-	FIVE of the following: nxiety drugs. we and hypnotic with exame chanism and clinical uses of general anesthesia. grandmal and petitmal expanding parkinsonian drugs. chition enhancers?	es of amphetamine.	[10]					
Q.2	a)b)	actions, adver	se effects and contraindi	ain the mechanism of action, therapeutic cations of barbiturates. the pharmacology of phenytoin.	[08] [07]					
Q.3	a) b)	· ·	-	he pharmacology of liquid anesthetics. the pharmacology of phenothiazine.	[08] [07]					
Q.4	a) b) c) d)	Write short notes on ANY THREE of the following: Opioid analgesics Antidote of Benzodiazepine poisoning Preanesthetic medications Lidocaine								
	SECTION – II									
Q.5	a) b) c) d) e) f)	What are Eice Describe the s Define antipy Describe the r Classify the ty Enlist the sign	FIVE of the following: osanoids? Enlist their type synthesis of leukotrienes. retics. Give suitable examechanism of action and types of poisons. In and symptoms of insection and symptoms of insections are effects of NSAIDs.	mples. therapeutics uses of salicylates.	[10]					
Q.6	a)b)	and ibuprofen	•	used for the treatment of mercury	[08] [07]					
Q.7	a) b)	Enlist the drugs used for treatment of acute and chronic gout. Explain in detail pharmacology of colchincines and probenecid. Describe the signs and symptoms and treatment of arsenic poisoning.		[08] [07]						
Q.8	a) b) c) d)	Write short notes on ANY THREE of the following: Disease modifying anti-rheumatic drugs (DMARDs) Uric acid synthesis inhibitors Selective COX – 2 inhibitors Lead poisoning								