### **SUBJECT: PATHOPHYSIOLOGY**

Day Date		Saturday 05/05/2018	S-2018-3920	Time : Max. Ma	<b>02.00 PM TO 05.00 PM</b> arks: 60
N.B.					
	1)		re <b>COMPULSORY.</b> Out of the remaining	g solve any	ГWО
	2)	questions from			
	2) 3)		right indicate FULL marks.	DATE	
		Answers to out	th the sections should be written in <b>SEPA</b>	RATE answ	er book.
			SECTION – I		
Q.1		Solve any FIVE q	mestions:		(10)
ν	a)		n by glycogen storage?		(10)
	b)		at are the symptoms of allergy?		
	c)	Define cell adapta			
	d)		s of hypersensitivity reactions?		
	e)	Define shock give	• •		
	f)	Define and classify	y of angina.		
Q.2		Define malignancy	y. Discuss pathogenesis of cancer.		(10)
Q.3	a)	Explain physical a	and chemical agents that can causes cell in	jury.	(07)
	<b>b</b> )	Enlist the various a	auto immune diseases. Write pathophysio	logy of any	one (03)
		auto immune disea	ases.		
Q.4		Write a note on an	v TWO:		(10)
ζ	a)	Glycogen infiltrati	<del></del>		,
	b)	Abnormalities in li			
	c)	Pathophysiology o			
			SECTION – II		
o =					(10)
Q.5	۵)	Solve any <b>FIVE</b> question Define bacillary dy			(10)
	a) b)	Explain wound hea	· ·		
	c)	What is Pneumonia			
	<b>d</b> )	What is swine flu?			
	e)	What is Ebola?			
	f)	What is peptic ulce	er?		
Q.6		Explain pathophys	iology of diabetes mellitus.		(10)
<b>Q.</b> 7	a)	Explain pathophys	iology of epilepsy.		(07)
	b)	Discuss pathophys	iology of bronchial asthma.		(03)
Q.8		Write a note on an	y <b>TWO</b> :		(10)
<b>~.</b> ··	a)	Acute renal failure			
	b)	Tuberculosis			
	c)	Parkinsonism			

### S.Y.B.PHARM. SEMESTER-III (2011 COURSE): SUMMER - 2018 SUBJECT: PATHOPHYSIOLOGY

Time: 02.00 PM TO 05.00 PM Day : Saturday S-2018-3956 Date : 05/05/2018 Max. Marks: 80 N. B. : 1) Q. No. 1 and Q. No.5 are COMPULORY. Out or remaining solve ANY TWO 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** Q. 1 Solve ANY FIVE of the following: (10)Explain risk factors of cancer. Explain about biological effects of radiation. b) Define cell adaptation. c) d) What do you mean by autoimmunity? e) Differentiate between benign and malignant tumors. Differentiate metaplasia from hyperplasia. Q. 2 Explain pathogenesis of cell injury. Give physical and chemical agents that causes cell injury. a) Discuss pathophysiology of malignancy. (08)Q. 3 b) Explain basic mechanism of inflammation and repair. (07)**Q.** 4 Write a note on **ANY THREE** of the following: (15)a) Allergy b) Glycogen infiltration c) Environmental carcinogenesis d) Glycogen storage disease **SECTION - II** (10)Solve **ANY FIVE** of the following: Q. 5 Define Bacillary dysentery. a) What is COPD? b) Define and classify angina. c) What do you mean by acute renal failure? d) What is shock? Give types of shock. e) Explain Schizophrenia. (15)Explain pathophysiology of HIV. Q. 6 (08)a) Explain pathophysiology of bronchial asthma. **Q**. 7 (07)b) Explain pathophysiology of Parkinsonism. Write a note on **ANY THREE** of the following: (15)Q. 8 a) Diabetes mellitus b) Paralysis c) Hepatitis d) Sleep disorders

## SUBJECT: PHARMACEUTICAL ANALYSIS-I

Day Date	:	Friday 27/04/2018	S-2018-3917	Time: <b>02.00 PM TO 05.00 PM</b> Max. Marks: 60	М
N.B.	: 1) 2) 3)	TWO question Both the section	Q. No. 5 are <b>COMPULSOR</b> ns from each section. ons should be written in <b>SEP RIGHT</b> indicate full marks.	Y. Out of the remaining attempt an ARATE answer books.	ny
			SECTION-I		
Q.1	Atte a) b) c) d) e) f)	0.1 N perchloric ac Write chemical re Sodium bicarbonat Why glycerine is a Write calibration o	pare and standardize 500 noid solution. eaction, principle involved e. dded in assay of boric acid.	al of 0.25 N HCl solution and in assay of Norfloxacin and on.	(10)
Q.2	a) b)	Derive an equation	for dissociation constant of ple involved in assay of aspir	a weak base. Give the chemical in.	(07) (03)
Q.3	a) b)	reaction principle a Give the equivalent NaHCO <sub>3</sub> , H <sub>2</sub> SO <sub>4</sub> , N	and assay procedure of sodium t weights of:	m acetate.	(07) (03)
Q.4	a)	te short notes on any Theories of Acid ba Hydrolysis of salt Non aqueous titrati			(10)
Q.5	Atte a) b) c) d) e) f)	Why masking and of How to prepare an solution. Why nitrobenzene	s redox indicator.  nd co-ordination number?  demasking agents are used.  d standardize 0.1 N KMnO  or dibutyl phthalate is added action principle and assay p	4 and 0.05 N Disodium EDTA	(10)
Q.6	a) b)	complex.	es and chelates? Explain sta Mohr's and Volhard's metho	·	(07) (03)
<b>Q.</b> 7	a) b)	Explain ceriometric	type of titrations.		(07) (03)
Q.8	a)	te short notes on any pM Indicators Fajan's method Permanganate titrat	TWO of the following:		(10)

### S.Y.B.PHARM. SEMESTER-III (2011 COURSE) : SUMMER - 2018 SUBJECT : PHARMACEUTICAL ANALYSIS – I

Day Date	:	Friday 27/04/2018	S-2018-3953	Time: <b>02.00 PM TO 05.00 P</b> M Max. Marks: 80	M
N.B.:	1)	ON 11	N. 5 COMPULGODY	0 + 64	
	1)	_	<del>-</del>	Out of the remaining questions	
	2)	-	TWO questions from each s	itten in <b>SEPARATE</b> answer bool	7.0
	3)		e right indicate FULL marks.		NS.
		Tigates to the			<u></u>
0.1		A A BIST TO	SECTION – I		r4.03
Q.1	(۵	Classify types of	IVE of the following:		[10]
	a) b)		standardize 0.25N HCl and 0.	25N NaOH solution?	
	c)		s added in assay of boric acid		
	d)		of non-aqueous titration.	•	
	e)		: i) Buffer ii) Buffer action	iii) Buffer index and pH.	
	f)	Enlist types of ir	idicators used in non-aqueous	S.	
Q.2	a)	Discuss in detail	about dissociation constant of	of strong acid and strong base.	[08]
	b)	Describe in detail	il salt hydrolysis.		[07]
Q.3	a)	Explain neutraliz	zation curve for weak acid an	d weak base.	[08]
_	<b>b</b> )	Discuss on types	s of solvents used in non-aqu	ueous. Write about differencing	[07]
		and leveling effe	ct.		
Q.4		Write short note:	s on ANY THREE of the fol	lowing:	[15]
	a)		eators used in Acid base titrat	ion	
	b)	Calibration of vo			
	c)		non-aqueous titration		
	d)	Buffering capaci	•		
			SECTION – II		
Q.5		*	IVE of the following:		[10]
	<b>a</b> )		during preparation of Iodine		
	b)	product of magn	ubility of magnesium hydrox esium hydroxide is 6.03 × 10 g-hydroxide is 58.33)	ide in mg/100ml, if the solubility -10.	
	c)	-		d in assay of H <sub>2</sub> O <sub>2</sub> and Sodium	
	d)	Why ammonium	buffer is added?		
	e)			Silver nitrate and 0.05N EDTA	
	υ,	solution.			
	f)	Write significan	ce of Ksp.		
Q.6	a)	How end point is	detected in complexometric t	citration? Explain types of EDTA	[08]
_	L)	titrations.	l to calculate equivalent weig	ht in raday titrations	[07]
	b)	Describe method	i to carculate equivalent weig	in in redox unations.	[0/]
<b>Q.</b> 7	a) b)		's method of precipitation in tric type of titrations.	detail.	[08] [07]
	U)	Explain ceriome	the type of thrations.		[0/]
<b>Q.8</b>			s on ANY THREE of the fol	lowing:	[15]
	a)	Fajan's Method			
	<b>b</b> )	_	rations		
	c) d)	Chelon effect Unit operations	in Gravimetry		
	u)	Omi operations	in Oraviniou y		

# 5.7. B. Phaym-Sem-III (CBCS-2015 COURSE): SUMMER-2018 SUBJECT: PHARMACEUTICAL BIOCHEMISTRY-II

Day: Monday Tim Date: 23-04-2018 5-2018-3916

Time: 2:00 PM TD 5:00 PM

Max. Marks: 60

N.B.:

- 1) Q. No. 1 and Q. No. 5 are **COMPULSORY**. Out of the remaining solve any **TWO** question 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 Attempt any FIVE of the following: (10)

- a) What is oxidative phosphorylation?
- **b)** What is oxidative deamination?
- c) What is favism?
- d) State transamination of aspartate.
- e) State biosynthesis of serotonin from tryptophan
- **f)** What is enzyme antibody conjugate?

Q.2 a) What is diagnostic PCR? Explain in detail. (07)

b) What is detoxication? Give examples. (03)

Q.3 a) What is Jaundice? Explain in detail and give different types. (07)

b) What is Ketosis? (03)

Q.4 Write notes on any TWO: (10)

- a) Catabolism of Arginine
- b) Hyperammononia
- c) Biochemical role of Vitamin –D

#### **SECTION-II**

Q.5 Attempt any FIVE of the following: (10)

- a) What is ATP cycle?
  - **b)** What is immunoprecipitation?
  - c) What is frame shift mutation?
  - d) How milk sugar is made available for glycolysis?
  - e) State the biochemical reaction where biotin in required as co-substrate.
  - f) What is osteoporosis?

Q.6 a) What is transcription? Explain in detail. (07)

b) What is chemical jaundice? (03)

Q.7 a) What is translation? Explain in detail. (07)

b) State biochemical role of vitamin ascorbic acid. (03)

Q.8 Write notes on any TWO: (10)

a) Glycolysis

**b)** Radio immunoassay (RIA)

c) ELISA

### SUBJECT: PHARMACEUTICAL CHEMISTRY-V (ORGANIC)

Day

Friday

S-2018-3915

Time: **02.00 PM TO 05.00 PM** 

Date : 20/04/2018

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) Both the sections should be written in **SEPARATE** answer books.
- 3) Figures to the **RIGHT** indicate full marks.

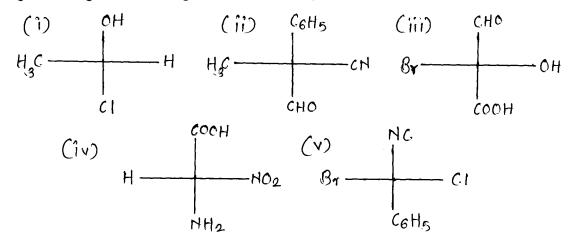
#### **SECTION-I**

Q.1 Attempt any FIVE questions of the following:

(10)

- a) Why racemic mixtures are optically neutral?
- b) Enlist the conditions of optical activity.
- c) Draw schematic diagram of polarimeter.
- **d**) What are *Z* and *E* isomers?
- e) Define and explain with example of enantiomer.
- f) Enlist the methods of racemic mixture preparation.

Q.2 Assign R & S configuration to following structures with reasons:



- Q.3 a) Draw the structure and show possible conformers using various projection formulae.
  - (i) 2-Bromobutane
- (ii) iso-Propylalcohol
- (iii)Ethylalcohol
- b) Comment on stability of free radicals.
- **Q.4** Write note on any **TWO** of the following:
  - a) Schmidt's reaction
  - b) Mannich reaction
  - c) Gabriel synthesis.

#### **SECTION-II**

Q.5 Attempt any FIVE questions of the following:

- (10)
- a) Explain how amines can be generated by Lossen rearrangement.
- b) What is role of  $H_2O_2$  in Baeyer-Villiger oxidation?
- c) What is limitation of Dakin oxidation?
- d) Explain how diazoketones converted in carboxylic acid in Wolff rearrangement.
- e) Name the rearrangement reactions which follows through isocyanate intermediate.
- f) What is role of acids in Pinacol-Pincolone rearrangement?
- **Q.6** Explain in detail mechanism orientation and stereochemistry for Hofmann and Schmidt rearrangement.
- Q.7 a) Complete the reaction and explain the mechanism.

$$(1) \qquad \frac{C_2H_5COCI}{AICI_3} \qquad ?$$

(ii) 
$$\frac{0}{11}$$
  $\frac{8\pi 2/HqOH}{6}$ 

- b) How  $\alpha$ -hydroxy carboxylic acids generated from 1,2-diketones.
- Q.8 Write note on any TWO of the following:
  - a) Fovorskii rearrangement
  - b) Beckmann rearrangement
  - c) Wittig rearrangement

\* \* \*

#### SUBJECT: PHARMACEUTICAL MICROBIOLOGY - I

Time: 02.00 PM TO 05.00 PM Day : Thursday Date Max. Marks: 60 : 03/05/2018 S-2018-3919 N.B.: 1) Q.No.1 and Q.No.5 are COMPULSORY. Out of the remaining questions 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 **Q.1** Answer ANY FIVE of the following: [10] State Koch Postulates. a) b) Draw a neat labeled diagram of Transmission Electron Microscope (TEM). c) Give pharmaceutical significance of micro-organisms. d) List morphological features of Escherichia. e) Explain principle of Acid Fast Staining. How bacterial cultures are preserved? f) **Q.2** a) How will you identify an unknown bacterial culture? [06] **b)** Draw and discuss Fluorescence Microscopes. [04]Q.3 a) Discuss scope and various applications of pharmaceutical microbiology. [06]How to measure bacterial growth? [04]Write short notes on ANY TWO of the following: **Q.4** [10] a) Discovery of Antibiotics b) Bacteriological Culture Media c) Bacterial Cell Wall **SECTION - II** Answer **ANY FIVE** of the following: [10] Q.5 Highlight characteristics of Rickettsia. a) **b)** Define and explain Z-value. Draw a well labeled diagram of Aspergillus niger. c) How halogens disinfect? d) Write general properties of viruses. e) Which biological indicators are used in moist heat sterilization? f) Discuss factors affecting disinfectant activity. [06] Q.6 a) b) Write in detail about sterility tests as per I.P. [04] Q.7 a) Draw and explain Lytic Cycle of viruses. [06]**b)** Explain isolation of Actinomycetes. [04] Write short notes on **ANY TWO** of the following: [10] Q.8 a) Phenol Coefficient Test b) Moist Heat Sterilization Cultivation of Viruses

### S.Y.B.PHARM. SEMESTER-III (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACEUTICAL MICROBIOLOGY - I

Time: 02.00 PM TO 05.00 PM Day: Thursday Date: 03/05/2018 Max. Marks: 80 S-2018-3955 **N.B.**: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of remaining questions, 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. 4) Draw neat, labeled diagrams wherever necessary. **SECTION - I** Q.1 Answer the following: (ANY FIVE) (10)Write function of flagella. a) b) How will you calculate microbial generation time? Differentiate between Gram + ve and Gram - ve bacterial cell. c) Write importance of *Penicillium* species. d) Enlist Koch postulates. e) Define Numerical Aperture and Resolving Power. f) **Q.2** Answer the following. (15)Write in detail Whittakar's five kingdom concept. b) Draw a ray diagram of compound microscope and explain the function of different parts of it. Write in detail different methods used for preservation of bacterial culture. 0.3 a) Explain the different methods used for isolation of pure cultures. (08)Describe different method for measurement of bacterial growth. b) (07)

Write short note **ANY THREE** of the following:

Reproduction of bacteria by asexual method

**Q.4** 

a)

b)

c)

Candida albicans

Importance of Actinomycetes

Phase contrast microscope

P.T.O.

(15)

### **SECTION - II**

Q.S		Answer Art Five of the following.	(10)				
	a)	Draw a neat labeled diagram of T-even bacteriophages.					
	b)	What are biohazards? Give its symbol.					
	c)	How HEPA filters are validated?					
	d)	Compare between sterilization and disinfection.					
	e)	Define and explain interferons.					
	f)	How will you sterilize zinc oxide powder and fixed oil?					
Q.6	a)	Classify disinfectants. Explain mechanisms and applications of various disinfectants.	(08)				
	b)	Discuss different tests used in testing of Aseptic Areas.	(07)				
<b>Q.</b> 7	a)	Explain different sterilization indicators.	(08)				
	b)	Classify viruses. Describe the morphology of viruses in detail.	(07)				
Q.8		Write short notes on <b>ANY THREE</b> of the following:	(15)				
	a)	Non-thermal Sterilization Techniques					
	b)	Phenol Coefficient Tests					
	c)	Multiplication of Human Viruses					
	d)	Design of Aseptic Area					

\* \* \* \* \*

SUBJECT: PHYSICAL PHARMACY - I

02.00 PM TO 05.00 PM Time: Day : Monday Date : 30/04/2018 S-2018-3918 Max. Marks: 60 **N.B.:** Q.No.1 and Q.No.5 are COMPULSORY. Out of the remaining questions 1) attempt ANY TWO questions from each section. 2) Answers to both the sections should be written in the **SEPARATE** answer books. Figures to the right indicate FULL marks. 3) SECTION - I 0.1 Answer **ANY FIVE** of the following: [10] a) What is Joule Thompson effect? b) Derive ideal gas equation. c) Explain the term 'Phase'. d) Define Molarity. e) Differentiate between ideal and real solution. What is effect of dilution on equivalent and specific conductance? Q.2 a) Explain in detail binding forces between molecules. [06]Give an account of kinetic molecular theory of gases. [04]Q.3 a) Define Raoult's Law. Explain deviations from Raoult's Law. [06] Prove that lowering of the vapor pressure is a colligative property. [04] **Q.4** Write notes on ANY TWO of the following: [10] Arrhenius theory a) **b)** Two component system c) Determination of critical constants SECTION - II **Q.5** Answer **ANY FIVE** of the following: [10] a) What is effect of temperature and pressure on solubility of gases in liquid? b) Give significance of partition co-efficient. c) Define order of reaction. d) What is the unit of rate constant of first order reaction? Classify energy of thermodynamic system. Define half-life of a reaction. f) Derive and expression for rate constant of second order reaction. [06]0.6 a) Explain transition state theory. [04]b) What is Nernst distribution law? Explain effect of molecular association and  $\mathbf{Q.7}$  a) [06]dissociation on partitioning of molecules. **b)** Describe in detail solute – solvent interaction. [04]Write notes on ANY TWO of the following: Q.8 [10] a) Methods to determine order of reaction **b)** Solubility of slightly soluble electrolytes c) Effect of temperature on rate of reaction

SUBJECT: PHYSICAL PHARMACY- I

Time: 02.00 PM TO 05.00 PM Day: Monday S-2018-3954 Date: 30/04/2018 Max Marks: 80 N.B: Q. No 1 and 5 are COMPULSORY. Out of remaining attempt ANY TWO 1) Ouestions from each section. 2) Answers to both the sections should be written in **SEPARATE** answer book. Figures to the right indicate FULL marks. 3) **SECTION-I Q.1** Answer **ANY FIVE** of the following: (10)Explain the term 'Component'. a) Differentiate between ideal & real solutions. b) What is Joule Thompson effect? c) State rules for drawing ternary phase diagram. d) Give wrong assumptions of ideal gas law. e) Explain in detail methods used for liquefaction of gases. (08)Q.2 a) Discuss kinetic molecular theory. b) (07)What do you mean by critical constants? Explain different methods used for (08) Q.3 a) determination critical constants. b) Explain in detail one component three phase system. (07)Write short notes on **ANY THREE** of the following: **Q.4** (15)Compressibility factor a) b) Raoult's law & its deviation Phenol – water system c) Colligative properties d) **SECTION-II** Answer **ANY FIVE** of the following: **Q.5** (10)What is half life of a reaction? a) Define molality & mole fraction. b) What are advantages of conductometric titrations? c) Enlist methods of decomposition of medicinal agents with examples. d) Define equivalent & specific conductance. e) What is common ion effect? f) Explain in detail accelerated stability studies. (08)Q.6 a) Give detailed account of solute- solvent interactions. b) (07)Derive an expression for rate constant of second order reaction. (08)**Q.**7 a) Write a note on reaction theories. b) (07)Write short notes on ANY THREE of the following: **Q.8** (15)

\* \* \* \* \*

Effect of pH and solvents on solubility of weak electrolyte

Determination of energy of activation

Nernst distribution law

Debye Huckel theory

a)

b)

c)

d)

### S.Y.B.PHARM. SEMESTER-IV (2011 COURSE): SUMMER - 2018 SUBJECT: DOSAGE FORM DESIGN – I

Time: 02.00 PM TO 05.00 PM Day : Wednesday Date : 02/05/2018 Max. Marks: 80 S-2018-3960 N.B.: Q.No.1 and Q.No.5 are COMPULSORY. Out of the remaining questions 1) attempt ANY TWO questions from each section. 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 Answer **ANY FIVE** of the following: [10]Define solubility. Explain any one method of solubility improvement. **b)** Explain the factors which affects the solubility and dissolution rate. Give comment on flocculating agents. Write in short about QA and QC. What is importance of partition coefficient in liquid formulation? e) Explain the role of colouring agents and dyes in preparation. f) Define preformulation. Explain the physico-chemical properties of [08] **Q.2** a) preformulation. Comment on common ion effect and polymorphism. b) [07]Q.3 Define solution. Explain in detail methods of preparation of solutions. [80]Enlist the types of emulsion. Explain test for identification of emulsions. b) [07]**Q.4** Write short note on **ANY THREE** of the following: [15] Stability of suspension a) Phase inversion b) HLB scale c) d) **CMC** SECTION - II **Q.5** Answer **ANY FIVE** of the following: [10]What are different protein structure? a) Differentiate between multiple and micro-emulsion. b) Explain dry suspension formulation. Give one example. c) Write different polymorphic forms of coca butter. d) Explain creaming of an emulsion. Enlist various preservative used in emulsion. f) Define suppository. Explain in detail formulation and evaluation of [08]  $\mathbf{Q.6}$ a) suppository. b) Explain role of emulsifying agent. [07] Explain in detail theories of emulsification. [08]**Q.7** a) Discuss factors affecting preservation of emulsion. Give two commercial [07]examples of emulsions. Write short note on **ANY THREE** of the following: Q.8 [15] Hydrocarbon base a) QC of emulsions b) Displacement value c) Factors affecting emulsion stability

### SUBJECT :PHARMACEUTICAL ANALYSIS - II

Day Date	:	Saturday 28/04/2018	S-2018-3923	Time: Max. M		<b>00 PM TO 05.0</b> : 60
N. B.	1)	any Two fi	o – 1 and 5 are COMPULSORY. Or com section - I and any Two question	ns from section - II		-
	2) 3)		both the sections should be written in the right indicate <b>FULL</b> marks.	n SEPARATE ansv	wer b	ooks.
			SECTION - I			-
<b>).1</b>		Attempt ANY	FIVE of the following:			(10)
	<ul><li>a)</li><li>b)</li><li>c)</li><li>d)</li><li>e)</li></ul>	Classify variou What is acid er What is 'maxin	ntiating pulse polarography s electro analytical techniques ror and alkaline error in potentiometr na' in polarography? How is it reduce alibration of pH electrode done?	•		
	f)		sary to remove oxygen from polarogu	caphic equipment?		
Q.2	a) b)		account of Amperometric titrations aciple involved in biamperometry			(7) (3)
Q.3	a)	Draw and expla	nin construction and working of DME			<b>(7)</b>
	b)	Give IIkovic ed	uation and its significance			(3)
<b>).4</b>		Write short not	es on ANY TWO of the following:			(10)
	a) b) c)		Coulometry rodes in potentiometry y various modes			
			SECTION - II			
<b>).</b> 5		Attempt ANY	FIVE of the following:			(10)
	a) b) c) d) e) f)	Explain the terr How and with w Explain the effective comparison	s factors affecting angle of refraction in ORD and CD what is calibration of conductometer pect of dilution on molar and equivaler on between co-precipitation and post performent iteration of Weak Acid Vs. Section 2015	nt conductance precipitation		
<b>Q.6</b>	a)	Give principle,	instrumentation and application of A	bbe's Refractomete	r	(10)
2.7	a)	Explain the prin	nciple and instrumentation of polarim	eter		(7)
	b)	Give applicatio	ns of Polarimetry			(3)
	a) b) c)	Write short not Gasometric ass Conductivity co Steps in gravim	ell			(10)

### S.Y.B.PHARM. SEMESTER-IV (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACEUTICAL ANALYSIS - II

Day Date	:	Tuesday 24/04/2018 S-2018-3958	Time: <b>02.00 PM TO 05.00 PM</b> Max. Marks: 80	I				
$\frac{Batte}{N.B.:}$								
	1)	Q.No.1 and Q.No.5 are COMPULSORY. Out of the remaining questions						
		attempt ANY TWO questions from each section.						
	2)	Answers to both the sections should be writt	en in <b>SEPARATE</b> answer book	S.				
	3)	Figures to the right indicate FULL marks.						
		SECTION – I						
Q.1		Attempt ANY FIVE of the following:						
	<b>a</b> )	State Ilkovic equation.						
	<b>b</b> )	<u> </u>	How pH meters are calibrated, write composition of buffers?					
	c) d)	State merits and demerits of instrumental analy Define Migration current and diffusion current.						
	e)	Classify indicator electrodes used in potention						
	f)	Why nitrogen gas is bubbled in polarographic a	•					
<b>Q.2</b>	a)	What is half wave potential? How it is calcula diffusion current.	ited? Explain factors affecting	[08]				
	b)	Describe Polarographic apparatus.		[07]				
	υ,	Desertoe Folarograpine apparatus.		[0/]				
Q.3	a)	Explain theory, principle involved in potention	netry. Discuss instrumentation	[08]				
		in detail.						
	b)	Discuss about ion selective electrodes used in p	otentiometry.	[07]				
Q.4		Write short notes on <b>ANY THREE</b> of the follo	wing:	[15]				
ν	a)	Dropping mercury electrode	······································					
	b)	Rotating platinum electrode						
	c)	Potentiometric titrations						
	d)	Applications of Amperometry						
		SECTION – II						
Q.5		Attempt <b>ANY FIVE</b> of the following:		[10]				
	a)	Give significance of inorganic precipitants in gr	ravimetry.					
	b)	Write about cell constant and its significance.						
	c)	, 1	arly polarized light					
	<b>d</b> )	Explain the terms specific refraction and molar	refraction.					
	e) f)	Explain factors affecting refractive index. Write about cotton effect.						
	1)	write about cotton effect.						
Q.6	a)	State principle involved in measurement ang	gle of refraction and give its	[08]				
		application.						
	b)	Explain the theory of optical activity. Discuss	about polarimeters.	[07]				
<b>Q.</b> 7	a)	Explain principle involved in gravimetric and	alysis. Discuss various steps	[08]				
~.	,	involved in gravimetric analysis.		[]				
	b)	Discuss in detail conductometric titration curve	S.	[07]				
$\mathbf{v}_{\mathbf{o}}$		Write short notes on ANY TUDEE of the follo	avina.	[15]				
Q.8	a)	Write short notes on <b>ANY THREE</b> of the follo High frequency titration	wing.	[15]				
	b)	Dipping refractometer and Pulfrich refractomet	er					
	c)	Circular Dichroism						
	ď)	Occlusion and Mixed crystal formation						
			*					

### S.Y.B.PHARM. SEMESTER-IV (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACEUTICAL CHEMISTRY - VI (ORGANIC)

Time: 02.00 PM TO 05.00 PM Day Saturday S-2018-3957 Max. Marks: 80 Date 21/04/2018 N. B.; 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of remaining 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 books. **SECTION - I** Q. 1 Solve **ANY FIVE** of the following: (10)a) Annomeric forms of glucose, explain with structure. Hemiacetal formation of carbohydrates. Explain. b) Why sucrose not give tollen's test? c) Explain Fehling's solution test of carbohydrates? d) Write structure and properties for maltose. **e**) Write structure and properties for Lactose. f) Q. 2 Explain in detail with mechanism and example. Free radical chain reaction (15) and termination of reaction. How phenylosazone reaction with glucose mannose and fructose gives same (08) Q. 3 a) product? Explain in detail. Comment on Fluorination and Iodination of Alkens. (07)Q. 4 Write a note on **ANY THREE** of the following: (15)Reducing sugars a) Chlorine is more reactive and less selective than Bromine b) Hydrogen halides reaction with alkenes by radical mechanism c) Mutarotation d) **SECTION - II** Q. 5 Solve **ANY FIVE** of the following: (10)Draw structure and give numbering to following heterocyclic: (i) Imidazole (ii) Quinoline Name the heterocyclic and give the numbering: b) (i)

(i) Pyrimidine (ii) Indole d) Why Indole undergoes electrophilic substitution reaction at 2-position. Give general mechanism of electrophilic substitution reaction. e) How will you synthesize Glycine by phthalimide synthesis? Give any three methods of preparation and three chemical reactions of (15) **Q.** 6 pyridine and Indole. **Q.** 7 Explain any two methods of preparation of amino acids with examples. (08)Give structure, numbering, corresponding drugs and methods of preparation (07) b) of thiazole. Write a note on **ANY THREE** of the following: Q. 8 (15)Fischer Indole synthesis a) b) Secondary structure of protein Skraup synthesis c) Biologically important peptide d)

Name the corresponding drugs for following heterocycles:

c)

\* \* \* \* \* \*

#### SUBJECT: PHARMACEUTICAL CHEMISTRY - VI (ORGANIC)

Time: 02.00 PM TO 05.00 PM Day : Saturday S-2018-3921 Max. Marks: 60 : 21/04/2018 Date N. B.: Q. No. 1 and Q. No. 5 are COMPULSORY. Out of remaining solve 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** Q. 1 Solve **ANY FIVE** of the following: (10)Write structure and properties for maltose. a) Write structure and properties for cellobiose. b) Explain acid-base properties of amino acids. c) What is peptide linkage? d) Explain the fehling's test of glucose. e) What is annomerization of glucose? Write detailed classification of Amino acids with structures. Q. 2 (10)Q. 3 Write a detailed note on chemistry of Glucose. (07)a) Explain the iso-electric point of Amino acids (03)b) Write a note on **ANY TWO** of the following: Q. 4 (10)Separation of Amino acids by electrophoresis Ruff degradation of carbohydrates b) Peptide bond formation in Amino acids c) **SECTION - II** Solve **ANY FIVE** of the following: (10)Q. 5 Draw structure and give numbering to following structures a) i) ii) Imidazole Pyrrole b) Name the heterocycle and give the numbering: (i) (ii)

	c) d)	What are phospholipids? Define the term and give the example;				
	e) f) g)	<ul> <li>i) Synthon ii) Retrosynthesis</li> <li>What is Fischer indole synthesis?</li> <li>Give structures of two Sulphur containing heterocycles.</li> <li>Why pyrrole undergoes electrophilic substitution reaction at 2-position?</li> </ul>				
Q. 6		Give any three methods of preparation and two chemical reactions of furan and pyridine.	(10)			
Q. 7	a)	Explain rules of disconnections for retrosynthesis using synthesis of pyrimidine.				
	b)	Give the numbering and corresponding drugs for following structure:	(03)			
	·	i) Hydantoin ii) Quinoline				
Q. 8		Write a note on <b>ANY TWO</b> of the following:	(10)			
	a)	Methods of preparation of Isoquinoline				
	b)	Chemical properties of Imidazole				
	c)	Fat soluble vitamins				
	d)	Synthon approach in synthesis				

# S.Y.B.PHARM. SEMESTER-IV (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACEUTICAL MICROBIOLOGY (Including Immunology) – II

Time: 02.00 PM TO 05.00 PM Day : Friday Date : 04/05/2018 Max. Marks: 80 S-2018-3961 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) Figures to the right indicate FULL marks. SECTION - I Q.1 Answer ANY FIVE of the following: [10] Compare Cup-Plate and Turbidometric Assay methods. a) b) How to evaluate efficacy of a preservative? c) Enlist commercial probiotics products. d) What is TOC and COD? e) Draw a typical fermentation protocol. f) Mention microbial limits for Pure Water and Dried Aluminum Hydroxide. **Q.2** a) How to isolate and screen commercial micro-organisms? [08] b) Write various stages of downstream processing and discuss them. [07] Q.3 a) Discuss properties, mechanism and significance of probiotics. [80] How MIC of an antibiotics is determined? [07]**Q.4** Write short notes on **ANY THREE** of the following: [15] a) Trickling Filters Microbial Limit Tests b) Tray Fermenter c) d) Microbial Assay of Antibiotics **SECTION - II** Q.5 Answer **ANY FIVE** of the following: [10] Classify types of immunity. a) Give examples of hypersensitivity. b) Differentiate between Active and Passive Immunity. c) Write significance of Booster Dose. d) What is a Hapten? e) Define and explain vaccines. f) Explain various immunological preparations and products. [80]a) b) Describe Antigen-Antibody reactions and their significance. [07]Discuss Immediate Hypersensitivity in detail. [08]  $\mathbf{O}.7$ a) **b)** How monoclonal antibodies are produced? [07]Q.8 Write short notes on **ANY THREE** of the following: [15] a) BCG Vaccines b) Complement System c) Structure of Immunoglobulin d) Immunofluorescence

## SUBJECT: PHARMACEUTICAL MICROBIOLOGY – II

Day Date	:	: Tuesday : 24/04/2018 S-2018-3922 Time: 02.00 PM TO 05.00 Max. Marks: 60			M		
N.B.:	1) 2) 3)	attempt AN Answers to	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. Figures to the right indicate FULL marks.				
			SECTION – 1				
Q.1	a) b) c) d) e) f)	Which micro-o Enlist various s Define COD an Draw a well lab Give the micro	train improvement methods.		[10]		
Q.2	a) b)	•	and monitor fermentation proficance of MIC. How it is de		[06] [04]		
Q.3	a) b)		rmentation processing in deta preservative efficacy?	úl.	[06] [04]		
Q.4	a) b) c)	Microbial assay Fermentation M	Microbial Contamination		[10]		
0 -			SECTION – II	<b>I</b> .	F4.03		
Q.5	a) b) c) d) e)	Classify vaccin Define and exp What causes se Compare Activ What is Coomb	rum sickness? e and Passive Immunity.		[10]		
Q.6	a) b)	Discuss Hybrid What is Phagoc	oma Technology in detail. ytosis?		[06] [04]		
<b>Q.7</b>	a) b)		tion of bacterial vaccines in d ructure of Antibody.	letail.	[06] [04]		
Q.8	a) b) c)	Probiotics	es on <b>ANY TWO</b> of the follo pe Ag – Ab reactions sensitivity	owing:	[10]		

#### SUBJECT: PHARMACOGNOSY - I

Day : Friday Time : 02.00 PM TO 05.00 PM Date : 04/05/2018 Max. Marks: 60 S-2018-3925 N.B. 1) Q.1 and Q.5 are COMPULSORY. Out of the remaining THREE questions solve any TWO questions. 2) Figures to the right indicate FULL marks. Answers to both the sections should be written in **SEPARATE** answer book. 3) SECTION - I Q.1 Answer any FIVE questions: (10)Which are different types of soils? a) What is bhasma? b) What is main principle of Kampo system? c) Which are different quality control parameters for Asawa. d) Which are different natural auxins? f) Write advantages of chemical classification. What is drying? Write in detail advantage and disadvantage of different 0.2 a) (07)methods of drying of crude drug. Differentiate Primary and Secondary metabolites. (03)a) Explain Homoeopathic drug proving. Write in detail. (05)Q.3 Which are different factors influencing cultivation of medicinal plants. b) (05)**Q.4** Answer any **TWO** of the following: (10)Shikkimic acid pathway a) Mevalonic acid pathway b) Significance of Pharmacognostic parameter **SECTION - II** Q.5 Answer any FIVE questions: (10)Write biological source and main chemical constituent of Kokum butter. a) Write methods of preparation of Shark liver oil. What are uses of carotenoids? c) What is PUFA? d) Which are different uses of Guar gum? e) Write chemical constituents of Bees wax. What are carbohydrates? Write classification, biosynthesis, chemical tests (07)Q.6 a) for carbohydrates. (03)Write pharmacognostic account of Rice bran oil. b) (05)What are lipids? Write classification of lipids.  $\mathbf{Q.7}$ a) Write pharmacognostic account of Starch. (05)Answer any **TWO** of the following: (10)Q.8 a) Caster oil b) Neem oil c) Isapgol

### S.Y.B.PHARM. SEMESTER-IV (2011 COURSE): SUMMER - 2018 SUBJECT: PHARMACOLOGY -I

Day: Time: 02.00 PM TO 05.00 PM Monday Date: Max. Marks: 80 07/05/2018 S-2018-3962 N.B.: 1) Q. No. 1 and Q. No. 5 are COMPULSORY. Out of the remaining attempt any **TWO** question from each section. 2) Figures to the right indicate FULL marks. Answers to both the sections should be written in **SEPARATE** answer books. 3) 4) Draw neat labelled diagrams WHEREVER necessary. **SECTION-I Q.1** Define with examples (any **FIVE**) (10)Therapeutic index a) b) Efficacy Noncompetitive antagonism c) Margin of safety d) Idiosyncrasy e) f) Cumulative effect Q.2 a) Discuss in detail the process of new drug development with special emphasis (08) on preclinical and clinical studies. Enlist the different routes of drug administration. Discuss the advantages and (07) disadvantages of transmucosal routes. Discuss in detail manifestation of adverse drug reaction. Q.3 a) (08)Define drug interaction. Discuss pharmacodynamics drug interactions with (07)examples. **Q.4** Write short notes on any **THREE** of the following: (15)Dose response relationship b) Excretion of drugs Drug tolerance c) Receptor regulation d) **SECTION-II** Q.5 Solve any **FIVE** of the following: (10)Explain the term cycloplegia. a) b) Classify cholinergic drugs. c) Write the tissue distribution of nicotinic receptors. d) Classify the drug acting on autonomic ganglia. e) Explain the term neurotransmitter and synaptic cleft g) Write the tissue distribution of  $\alpha$ -receptors. Q.6 a) Classify the anti-cholinergic. Discuss pharmacological action, adverse drug reactions and therapeutic uses of atropine. Discuss the pharmacological action adverse drug reactions and therapeutic uses (07)of Ephedrine. Write the classification, pharmacological action and adverse effects of (08)  $\mathbf{Q}.7$  a) adrenergic drugs. b) Write the clinical applications and adverse effects of beta-blockers. (07)(15)Write a note on any **THREE** of the following: **Q.8** a) Neuromuscular blockers b) Different parts and function of ANS Muscarinic receptors c)

d) Belladonna poisoning

### S.Y.B.PHARM. SEMESTER-IV (CBCS - 2015 COURSE): **SUMMER - 2018 SUBJECT: PHARMACOLOGY-I**

Time: 02.00 PM TO 05.00 PM

: Monday Date : 07/05/2018 Max. Marks: 60

S-2018-3926

Day

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

Q.1 Attempt ANY FIVE of the following: [10]

- Define Synergism. Give examples. a) b) Define pharmacodynamics.
- c) Define Therapeutic index.
- d) Define clinical trials. Name its phases.
- Define Bioavailability. e)
- f) Define drug dependence.
- **Q.2** Write in detail general mechanism of action of drugs. Give suitable examples. [10]
- Explain the role of drug distribution in drug action. Q.3 [07] a) Explain factors modifying drug effect. [03]
- Write short notes on **ANY TWO** of the following: **Q.4** [10]
  - a) Nature of drugs
  - Tolerance b)
  - Plasma protein binding c)

#### **SECTION - II**

- Attempt ANY FIVE of the following: [10] Q.5
  - Write the therapeutic classification of sympathomimetics. a)
  - Discuss the metabolic pathway of adrenaline. b)
  - Name the subtypes of nicotinic receptor and their tissue distribution. c)
  - Explain the terms neuroeffector junction and synaptic cleft. d)
  - Name the drugs used for the treatment of myasthenia gravis. e)
  - Write the co-transmitters of autonomic nervous system. f)
- Explain the pharmacological actions, adverse drug reactions and therapeutic [10] **Q.6** uses of atropine.
- Classify  $\alpha$  blockers. Write pharmacological account of ergot alkaloids. [07]**Q.**7 a)
  - Discuss the biosynthesis pathway of acetylcholine. [03] b)
- Write short notes on **ANY TWO** of the following: [10] **Q.8** 
  - Anti-cholinesterase poisoning a)
  - Mechanism of action of catecholamines b)
  - Neuromuscular junction blockers

SUBJECT: PHYSICAL PHARMACY - II

Time: 02.00 PM TO 05.00 PM Day : Wednesday Date Max. Marks: 60 : 02/05/2018 S-2018-3924 N.B.: 1) Q.No.1 and Q.No.5 are COMPULSORY. Out of remaining questions attempt ANY TWO questions from each section. 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 Answer ANY FIVE of the following: [10]What are different types of emulsions? Enlist methods to identify the same. a) What is HLB? Classify surfactants on the basis of HLB. b) Classify colloids with examples. Suspensions are thermodynamically unstable. Explain. d) Define syneresis and imbibition. Write an equation for Freundlich isotherm. Explain methods used to determine surface tension. 0.2 a) [06]Derive an expression for spreading co-efficient. b) [04] Give detailed account of solubilization. Q.3 a) [06]Add a note on controlled flocculation. b) [04] Write notes on **ANY TWO** of the following: **Q.4** [10] a) Theories of emulsification **b)** DLVO theory c) Preparation of colloids **SECTION - II** Q.5 Answer **ANY FIVE** of the following: [10] a) Classify crystals on the basis of bonds between molecules. Enlist different types of viscometers. b) What are bingham bodies? c) Give significance of Heckel plots. d) Explain dilatant flow behavior. e) Give applications of micromerities in pharmacy. a) Give an account of compaction of powders and methods to evaluate the same. Q.6 [06] Classify polymorphs with examples. b) [04]Explain in detail methods used to determine surface area of particles. [06]**b)** Give an account of viscoelasticity. [04]Write short notes on ANY TWO of the following: Q.8 [10] a) Derived properties of powder b) Thixotropy and methods to determine the same c) Measurement of diffraction angle