AMAZON/PARANA/KARNAFULI/SURAMA/SIYANA/SINHAGAD- I (CBCS) (2012 Course) WINTER - 2014

SUBJECT: RESEARCH METHODOLOGY & BIOSCREENING

Day : Date :		ednesday 7-01-2015	me : 10:00AM·TO 1: ax. Marks : 60
2) :) :)	Attempt any THREE questions from Section – I and S Answers to both the sections should be written in SEP Figures to the right indicate FULL marks.	Section – II each. ARATE answer books.
		SECTION - I	
Q.1		Explain meaning, purpose of research and types of research	earch. (10)
Q.2		Explain the important considerations for selecting a rom. M.Pharm.	esearch problem for (10)
Q.3		What do you understand by quality by design (QBD) principles.	? And explain the (10)
Q.4	a)	Write elaborate notes on any TWO of the following: Descriptive research	(10)
	b)	References	
	c)	Objectives of research	
		SECTION – II	
Q.5		Explain the bio-screening of antihypertensive drugs.	(10)
Q.6		Explain the three 'R' principles of animal usage for ex	sperimentation. (10)
Q.7		In primary health centre 30% of women were found If from the batch of 100 women 30 women are select is the probability of atleast 20 women being malnouring	led at random. What
Q.8		Write elaborate notes on any TWO of the following:	(10
	a		
	b		
	c) Randomization	

AMAZON/PARANA/KARNAFULI/SURAMA/SIYANA/SINHAGAD-I (CBCS) (2012 COUTSE) WINTER - 2014

SUBJECT: RESEARCH METHODOLOGY & BIOSCREENING

Day Date	Wax Marks O		o 1:0	
N.B.	1) 2) 3)	Attempt any THREE questions from Section – I and Sec Answers to both the sections should be written in SEPAl Figures to the right indicate FULL marks.	ction – II each. RATE answer book	s.
		SECTION - I		
Q.1		Explain meaning, purpose of research and types of research	ch.	(10)
Q.2		Explain the important considerations for selecting a rese M.Pharm.	earch problem for	(10)
Q.3		What do you understand by quality by design (QBD)? principles.	And explain the	(10)
Q.4		Write elaborate notes on any TWO of the following:		(10)
Ų.·	a)	Descriptive research		
	b)	References		
	c)	Objectives of research		
		SECTION – II		
Q.5		Explain the bio-screening of antihypertensive drugs.		(10)
Q.6		Explain the three 'R' principles of animal usage for expe	rimentation.	(10)
Q.7	,	In primary health centre 30% of women were found to If from the batch of 100 women 30 women are selected is the probability of atleast 20 women being malnourished		(10)
Q.8	3	Write elaborate notes on any TWO of the following:		(10)
-	a)	Chi-square test		
	b			
	c	Randomization		

M- Pharm - Semi-I / 2015 AMAZON/KARNAFULI/PARANA/SURAMA/SINHAGAD

SIYANA - I (CBCS) : SUMMER - 2015
SUBJECT : ELECTIVE - I : PHARMACEUTICAL ADMINISTRATION

: Wednesday Day Time : 10:00AM. TO 1:00 P.M. Date 08-07-2015 Max. Marks: 60 N.B. 1) Attempt any THREE questions from Section – I and Section – II each. 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 Discuss various types of plans and planning process. (10)Q.2 Discuss departmentalization in detail. (10)Q.3 Enumerate various steps involved in staffing. Discuss performance (10)appraisal and career strategy in detail. Q.4 Write elaborate notes on any TWO of the following: (10)Formal and informal organizations b) Manager development process and training Management by objective (MBO) SECTION - II (10)Discuss McClelland's needs theory in detail. Q.5 Discuss productivity problems and their measurement in detail. (10)Q.6 (10)Discuss feed back and feed forward controls in detail. Q.7 Write elaborate notes on any TWO of the following: (10)**Q.8** Human factors in managing a) Preventive control

Basic control process

PARANA-I (CBCS - 2012 COURSE) : WINTER - 2014 SUBJECT : ADVANCED PHARMACEUTICAL CHEMISTRY-I

Day : Friday Date : 09-01-2015 Time: 10:00 AM TO 1:00 P.M.

Max. Marks: 60

N.B.:

- Attempt any THREE questions from Section-I and any THREE questions from Section-II.
- 2) Figures to the RIGHT indicate full marks.
- 3) Answers to both sections should be written in SEPARATE answer books.

SECTION-I

- Q.1 Describe protection and deprotection strategies for -OH and carbonyl groups. (10)
- Q.2 Explain the principle, various methods and advantages of catalysis by enzymes. (10)
- Q.3 a) Discuss the mechanism of homogeneous reduction. (05)
 b) Give applications of metal-ammonia reductions in synthetic chemistry. (05)
- Q.4 Write short notes on any TWO of the following: (10)
 - a) Bayer Villiger reaction
 - b) Clemmenson's reaction
 - c) Curtius rearrangement.

SECTION-II

- Q.5 Describe Fischer indole and Modelung indole synthesis. (10)
- Q.6 Discuss stereo-selective and stereo-specific formation of enolate anions citing (10) examples
- Q.7 Complete the following reactions giving reaction mechanism and major products. (10)

a)
$$\downarrow$$
 Li/NH3
 $+-BuoH$
 $-78^{\circ}C$
A
TMS-CI
B

- Q.8 Write short notes on any TWO of the following:
 - a) HUMO conservation of orbital symmetry.
 - b) Preparation of α -methylene lactone.
 - c) Role of Li in regioselective formation of enolate anions.

SIYANA - I (CBCS): WINTER - 2014 (2012 COUTSE) SUBJECT: ADVANCED QUALITY ASSURANCE TECHNIQUES - I

Time : 10:00AM.TO1:00P.M. : Friday Day Max. Marks: 60 : 09-01-2015 Date N.B. Attempt any THREE questions from Section - I and Section - II each. 1) Answers to both the sections should be written in SEPARATE answer books. 2) Figures to the right indicate FULL marks. 3) **SECTION - I** Discuss personnel hygiene and clothing requirements with reference to Q.1 cGMP. Discuss various requirements of pharmaceutical equipment. (10)Q.2 Discuss principles involved in purchasing material for pharmaceutical (10)Q.3 manufacturing. (10)Write elaborate notes on any TWO of the following: Q.4 Reference and working standards Outsourcing of analytical services b) Washing and toilet facilities **SECTION - II** (10)What are the basic concepts of QA and QC? Q.5 (10)Discuss control of mix-ups and cross contamination. Q.6 Discuss WHO requirements of documentation in detail. (10)Q.7 Write elaborate notes on any TWO of the following: (10)**Q.8** Pharmaceutical product distribution a) Recalled, returned and rejected goods b) Expiration dating

KARNAFULI - I (CBCS): WINTER - 2014 (2012 COURSE) SUBJECT: ADVANCED PHARMACOLOGY - I

Day: Friday
Date: 09-01-2015

Time: 10:00 AM TO 1:00 P.M.

Max. Marks: 60

N.B.:

1) Attempt ANY THREE questions from Section – I and Section – II each.

2) Each section should be written in SEPARATE answer books.

3) Figures to the right indicate FULL marks.

SECTION - I

Q.1	Discuss the regulations for laboratory animal care. Explain the role of IAEC	(10)
	in the animal experiments.	, , ,

- Q.2 Describe in detail the pre-clinical evaluation methods for anti-convulsants. (10)
- Q.3 Describe in detail the pre-clinical evaluation methods for anti hypertensive (10) agents.
- Q.4 Write short notes on ANY TWO of the following: (10)
 - a) High throughput screening
 - b) Organization of safety assessment tests
 - c) Screening of analgesics

SECTION - II

- Q.5 Explain in detail the preclinical evaluation methods for bronchodilators. (10)
- Q.6 Explain in detail the preclinical evaluation methods for anti-diabetic agents. (10)
- Q.7 Explain in detail the preclinical evaluation methods for diuretics. (10)
- Q.8 Write short notes on ANY TWO of the following: (10)
 - a) In vitro testing of drugs
 - b) Applications and Limitations of transgenic animals
 - c) Preclinical screening of androgens

AMAZON-I (2012 COURSE – CBCS) WINTER - 2014 SUBJECT ADVANCE CORE SUBJECT-I – ADVANCED PHARMACEUTICS-I

Time: 10:00 AM. TO 1:00P. M. Max. Marks: 60. Date N.B.: Answer any THREE questions from Section-I and any THREE questions from 1) 2) Both the sections should be written in SEPARATE answer books. 3) Figures to the RIGHT indicate full marks. **SECTION-I** Q.1 Explain the principle and instrumentation of DSC. Add a note on its application to (10) determine T_g of polymer. Q.2 Explain the Box Behenken approach for optimization of pharmaceutical (10) formulation. Q.3 Give an account of different mathematical models to describe dissolution kinetics. (10)Q.4 Write notes on: (10)a) Stability testing of biotechnological formulations as per ICH guidelines b) Polymorphism and its significance. **SECTION-II** Q.5 Describe the mathematical models for compact evaluation. (10)**0.6** What are dendrimers? Elaborate on their pharmaceutical applications. (10)0.7 Explain the thermodynamic aspects of polymer solutions. (10)

Q.8 Write notes on:

a) Mechanism of biodegradation

b) Liquid crystalline phase and methods to identify the same.

SINHAGAD-I :(CBCS): WINTER - 2014 (2012 COURSE)

SUBJECT: ADVANCED PHARMACEUTICAL BIOTECHNOLOGY-I Time: 10:00AM . TO 1:00 P.M. Day: Friday Max Marks: 60 Date: 09-01.2015 N.B: Attempt any THREE questions from Section-I and Section-II each. 1) Both the sections should be written in SEPARATE answer book. 2) Figures to the right indicate FULL marks. 3) **SECTION-I** Discuss central Dogma of molecular biology and describe how DNA controls (10) Q.1 the cellular metabolism. Describe how DNA is packaged into chromosome; elaborate the role of (10) Q.2 Histone proteins in process of gene expression.

0.4	Write short notes on: (ANY TWO)	(10)

Explain in details semi-conservative model of DNA replication.

Write short notes on: (ANY TWO) Q.4

Promoter

Alternate splicing b) Transposable elements c)

Q.3

Q.8

a)

SECTION-II

- (10)What is a PCR? Explain how DNA is amplified during PCR. Q.5 Define a cDNA library. Write its significance in studying gene expression (10) Q.6
- Write an essay on a recombinant DNA technology. Outline a protocol to (10) Q.7
- produce recombinant insulin.
 - Hot-start PCR a)

pattern.

- Phage display b)
- Reverse transcriptase enzyme

Write short notes on: (ANY TWO)

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(10)

PARANA/SINHAGAD/AMOZON/KARNAEUL1/SURAMA/

SIYANA - I (CBCS): WINTER - 2014 (2012 COURSE)
SUBJECT: ELECTIVE - I: PHARMACEUTICAL ADMINISTRATION

Day Date	: Monday : 12-01-2015		Time : 10:00AM-T01:00P. Max. Marks : 60	
N.B.	1) 2) 3)	Attempt any THREE questions from Section – I and Section – II each. Answers to both the sections should be written in SEPARATE answer books. Figures to the right indicate FULL marks.		
-		SECTION - I		
Q.1		Discuss various function of management in detail.	(10)	
Q.2		Discuss structure and process of organizing.	(10)	
Q.3		Discuss process of decision making in detail.	(10)	
Q.4	a)	Write elaborate notes on any TWO of the following Performance appraisal of staff	g: (10)	
	b)	Management social responsibilities		
	c)	Human resource management		
		SECTION – II		
Q.5		Discuss human factors in managing in detail.	(10)	
Q.6		How will you control and improve productivity?	(10)	
Q.7		Discuss Abraham Maslow's theory of motivation.	(10)	
Q.8		Write elaborate notes on any TWO of the following	ng: (10)	
	a)	Control of overall performance		
	b)	Communication process		
	c)	Basic control process		

PARANA / SINHAGAD/AMAZON/KARNAFUL/SURAMA/SIYANA - I (2012 COURSE) (CBCS): WINTER - 2014 SUBJECT: ADVANCED PHARMACEUTICAL ANALYSIS

Time: 10:00 AM. TO 1:00 P.M. : Wednesday Max. Marks: 60 Date 14-01-2015 N. B.: Attempt ANY THREE questions from Section - I and Section - II each. 1) SECTION - I (10)Discuss the principle of NMR and explain chemical shift and spin - spin Q. 1 splitting in detail with suitable examples. (10)Q. 2 Discuss the fragmentation pattern involved in the mass spectrometry in detail. (10)Q. 3 Discuss the instrumentation involved in HPLC in detail with special attention to HPLC columns. (10)Q. 4 Write an elaborate note on: Applications of HPTLC b) Detectors in GLC **SECTION - II** Write a detailed note on applications of super critical fluid chromatography. (10)Q. 5 Write a brief note on immunoassays and explain radio immunoassay in detail. (10)Q. 6 (10)Q. 7 Write a note on: Applications of DTA b) Types of sources of X-rays

Discuss the instrumentation and applications of TGA.

Q. 8

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SIN HAGAD / AMAZON / PARANA/ KARNAFULI/ SURAMA/ SIYANA – I (CBCS): SUMMER - 2015

(CBCS): SUMMER - 2015 SUBJECT: ADVANCED PHARMACEUTICAL ANALYSIS

Day: Wednesday
Date: 01-07-2015

Time: 10:00AM TO1:00PM.

Max. Marks: 60

N.B.:

- Attempt any THREE questions from Section –I and section –II each.
- 2) Figures to the RIGHT indicate full marks.
- 3) Answer to both the section should be written in SEPARATE answer books.
- 4) Draw neat diagrams WHEREVER necessary.

SECTION-I

Q.1 a) Assign the chemical shifts and multiplicities for the following compounds. (10)

b) Calculate the λ max for the following compounds.

Q.2 Deduce the structure of a compound having following spectral data: (10)

Molecular Weight: 98

IR (cm⁻¹): 3429, 3385, 2900, 2210, 1660, 1620 ¹ HNMR (δ ppm): δ 2.14 (t, 9.8 squares)

3.40 (t, 9.6 squares)

4.03 (s, 10 squares)

Q.3 Discuss the sources of Band Broadening in view of van deemter equation. (10)

P. T. O.

Q.4		Write elaborate notes on:	(10)
	a)	HPLC coloumns	
	b)	GC- MS	
		SECTION-II	
Q.5		Write a detailed note on Radioimmuno assay.	(10)
Q.6		Write a detailed note on: i) Chiral chromatography ii) Ion pair chromatography	(10)
Q.7		Discuss the instrumentation and applications of TGA.	(10)
Q.8		Write a note on:	(10
	a)	Detectors used in XRD instruments	
	b)	Applications of DSC	