PURUS - VI (2011 COURSE): SUMMER - 2017

SUBJECT: MEDICINAL CHEMISTRY - II Day : Saturday Time 10.00 AM TO 01.00 PM Date : 22/04/2017 Max. Marks: 80 N.B. 1) Q.1 and Q.5 are COMPULSORY. Out of the remaining, attempt any TWO 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 Q.1 Attempt any FIVE questions of the following: (10)a) Draw structures of any two triazolo-benzodiazepines. b) Outline synthesis and two uses of trimethadione. c) Explain mode of action of anticonvulsants. d) Explain physiology of sleep in brief. e) Explain aromatic hydroxylation. f) Draw structures of any two ultra-short acting barbiturates. g) Explain the term enzyme induction with examples. Q.2 a) What is metabolism? Discuss characteristic features of Phase - I and Phase -(07)II metabolism and add a schematic representation of their summary. b) Write in details about glucuronic acid and sulfate conjugation. (08)Q.3 a) Classify general anesthetics with one representative structure from each class (05)and explain stages of general anesthesia. Discuss chemistry of anticonvulsants. (05)c) With a short note on nonbarbiturates. (05)Write short notes on any THREE: **Q.4** (15)a) Barbiturates b) Classification of epilepsy and anticonvulsants c) Reduction and Hydrolytic pathways d) Mode of action of general anesthetic e) Analeptics **SECTION - II** Q.5 Answer the following any FIVE: (10)a) Classify Neuroleptics with examples. b) Outline the synthesis of Haloperidol with its category. c) Outline the synthesis of Imipramine with its uses. d) Draw any two structures of Tricyclic antidepressants. e) Draw any two structures of local anesthetics belonging to benzoic acid ester class. Outline the synthesis of Diazepam. g) Explain the terms epidural and caudal anesthesia. Q.6 a) Define and classify anxiolytics. (05)b) Describe SAR of Benzodiazepine class of anti-anxiety agents. (05)c) Comment on mode of action and therapeutic uses of Benzodiazepines. (05)Q.7 a)Classify antidepressants in details with one representative structure from each (05)

b) Classification of prodrugs c) MAO inhibitors d) Pharmacokinetic applications of prodrugs e) Outline the synthesis of Benzocaine and Lignocaine

c) Discuss chemistry and SAR of tricyclic antidepressants.

b) Comment on SAR of phenothiazines.

Write short note on any THREE:

a) SAR of local anaesthetics

Q.8

(05)

(05)

(15)

PURUS -VI (2011 COURSE): SUMMER - 2017 SUBJECT: PHARMACEUTICAL ANALYSIS - IV

| Da ₂ | - | Tuesday 25/04/2017 | | | : 10.00 A Marks : 80 | | 01.00 P | М |
|-----------------|--------------------|---------------------------|---|----------|--------------------------------|-----------|------------|------|
| N. | B.: 1 2 3 |) Answers to both | No.5 are COMPULSORY. WO from each section. In the sections should be writing the indicate FULL marks. | | | | | S. |
| | | | SECTION - I | | | | , | |
| Q.1 | | Solve Any FIVE o | f the following: | | | | | (10 |
| | a) | Why is thin layer c | chromatography superior to | other ty | pes of ch | romat | ographic | (10) |
| | b) | | tages of High performance I | | | | | |
| | c) | How solvents are se | elected in TLC? | orquia (| | rapny | / : | |
| | d) | | cteristics of detectors used in | HPLC | ? | | | • |
| | e) f) | Compare between a | dsorption TLC and partition armaceutical applications of | TLC | | | , | |
| Q.2 | a) | Discuss in detail var | rious types of pumps used in | HPLC | | , | . (| (07) |
| | b) | Describe the various TLC. | s techniques used for the pr | eparatio | on of chro | mato | plates in | (08) |
| 2.3 | a) | Explain the various | sample injection systems in | HPLC. | | · ; ·, | | (07) |
| | b) | | methods used for locating t | | | Ÿ | to the | (08) |
| .4 | | Write short notes on | Any THREE of the follow | ing : | | | • | (15) |
| | a) | Refractive index dete | ector | | | | | |
| | b) | Compare between HI | PLC and TLC | | | | | |
| | c) | Reverse phase partition | on TLC | | | | | |
| | d) | Applications of TLC | | | | | | |

P.T.O.

SECTION - II

| Q.5 | | Solve Any FIVE of the following: | (10) |
|-------------|----------|--|------|
| | a) | What is critical point in supercritical fluid chromatography? | (10 |
| | b) c) | Define the terms HETP and theoretical plate. What are the advantages of supercritical fluid chromatography over HPLC? | |
| | d) | Explain the term retardation factor and capacity factor. | |
| | e) | Give the properties of supercritical fluids. | |
| | f) | What do you mean by Edge effect in chromatography? | |
| Q.6 | a) | Give the various applications of HPTLC. | (07 |
| | b) | Discuss in detail the steps involved in HPTLC development. | (08) |
| Q. 7 | a) | Discuss in detail the methods for the determination of adulterants in dairy products. | (07) |
| | b) | Discuss in detail the instrumentation of supercritical fluid chromatography. | (08) |
| Q.8 | | Write short notes on Any THREE of the following: | (15) |
| | a) | Role of food inspector in analysis of food products | () |
| | b) | Applications of super critical fluid chromatography | |
| | c) | Quantitation techniques in HPTLC | |
| | d) | Determination of adulterants in turmeric products | |
| | | | |

PURUS-VI (2011 COURSE): SUMMER - 2017 SUBJECT: DOSAGE FORM DESIGN-III

Time: 10.00 AM TO 01.00 PM Day: Saturday Max. Marks: 80 Date: 29/04/2017 N.B: Q.No.1 and Q. No.5 are COMPULSORY. 1) Attempt ANY TWO questions from each section. 2) Use separate answer sheets for both the sections. 3) Figures to the right indicate FULL marks. 4) **SECTION-I** (10)Solve ANY FIVE of the following: Q.1 a) What is D value in sterilization? Show diagrammatically various environmental zones. b) State the significance of flow properties of solids in SVP. c) Explain clean in place and steam in place. d) How antioxidants are selected? Why ethylene diamine is necessary in aminophylline injection? Discuss leakage test of ampoules. Give a layout of a parenteral section and discuss the HVAC systems. (08)Comment on various additives in SVP. (07)Discuss various routes of parenteral drug administration. (08)Q.3 a) (07)Give an account on b) LAL testing ii) Sterility testing i) (15)Write notes on ANY THREE of the following: **Q.4** Formation of sterile suspension Freeze dried products Sterilization by radiation

Isotonicity testing

P.T.O.

SECTION-II

| Q.5 | | Solve ANY FIVE of the following: | (10) |
|-----|----|--|------|
| | a) | What is siliconization of glass containers? | |
| | b) | Give the formula for Ringer lactate solution. | |
| | c) | Calculate millimoles/ liter of 0.9% w/v NaCl solution. | |
| | d) | Why 15% mannitol solution is warmed before use? | |
| | e) | 5% dextrose solutions are stored in Type I glass containers-give reason. | |
| | f) | Give the composition of tear film in eye. | |
| | g) | What are ocular inserts? Give one example. | |
| Q.6 | a) | Discuss formulation principles of various large volume parenteral. | (08) |
| | b) | Write a note on IV admixtures. | (07) |
| Q.7 | a) | Draw a neat diagram of Extrusion screw and discuss flat sheet extrusion. | (08) |
| | b) | Discuss formulation of elastomeric closures. | (07) |
| Q.8 | | Write notes on ANY THREE of the following: | (15) |
| | a) | Mechanical and optical properties of plastic containers | |
| | b) | Sterility and pyrogen testing | |
| | c) | Dry serum | |
| | d) | Ophthalmic ointments | |
| | | | |

PURUS - VI: (2011 COURSE): SUMMER - 2017 SUBJECT: PHARMACOLOGY - III

Time: 10.00 AM TO 01.00 PM Day: Wednesday Max. Marks: 80 Date: 03/05/2017 N.B.; Q. No. 1 and Q. No. 5 are COMPULSORY. Out of remaining questions 1) attempt Any TWO 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) |
| V. 2 | a) | Classify anti-anxiety drugs. Give examples. | |
| | b) | Classify opioid analgesics. | |
| | c) | Enlist the techniques of local anaesthesia. Define depression. Give examples of anti-depressants. | |
| | d) e) | D. Considerations | |
| | f) | Define status epitepticus. Define bipolar disorders. Give examples of drugs for treatment of bipolar disorder. | |
| | g) | Classify general anaesthetics. | |
| Q.2 | a) | Classify anti-Parkinson's drugs. Explain the role of anti-cholinergics in | (08) |
| | b) | treatment of Parkinsonism. What are cognition enhancers? Explain the pharmacology of cognition enhancers. | (07) |
| Q.3 | a) | Compare and contrast the mechanism of action, pharmacology, adverse effects | (08) |
| Q.D | b) | and toxicity of benzodiazepines and barbiturates. Classify local anesthetics. Explain in detail the pharmacology of amides used as local anesthetics. | |
| Q.4 | | Write Short notes on Any THREE of the following | (15) |
| | a) | Preanaesthetic medication | |
| | b) | | |
| | c) | Conversion and exthesia | |
| | d) | Phenothiazines | |
| | | SECTION - II | |
| Q.5 | | Answer Any FIVE of the following: | (10) |
| Ų.S | | will the signal of Give suitable examples. | |
| | a) b) | D. C analysis and antipyretics. | |
| | c) | Define toxicity. Give the composition of universal antidote. | |
| | ď | Court of American and for the treatment of goul. | |
| | e f | Figure 5 Figure 2 Figure 3 Fig | |
| | | Describe the mechanism of lead poisoning. | |

P.T.O.

| Q.6 | a) b) | Classify NSAIDs. Describe in detail pharmacology and adverse effects of non-selective COX inhibitors. Describe in detail the pharmacotherapy of rheumatoid arthritis. | (08) (07) |
|-----|----------|--|--------------|
| Q.7 | a) b) | Explain the biosynthesis of prostaglandins and leukotrienes. Describe the pathophysiological role of prostaglandins. Explain the general principles of treatment of acute poisoning. | (08) (07) |
| Q.8 | | Write Short notes on Any THREE of the following | (15) |
| | a) | Mercury poisoning | |
| | b) | Salicylates | |
| | c) | Treatment of barbiturate poisoning | |
| | d) | Pharmacology of leukotrienes | |

PURUS- VI (2011 COURSE): SUMMER - 2017 SUBJECT: PHARMACEUTICAL BIOTECHNOLOGY (INCLUDING MOLECULAR BIOLOGY)

(INCLUDING MOLECULAR BIOLOGY) Day: Friday Time: 10.00 AM TO 01.00 PM Date: 05/05/2017 Max Marks: 80 N.B: 1) Q.1 and Q.5 are COMPULSORY. Out of remaining answer any TWO Questions from each section. Figures to the right indicate FULL marks. 2) Answer to each section should be written in SEPARATE answer 3) **SECTION-I** Answer any FIVE of the following: Q.1 (10)Draw a neat labeled diagram of a prokaryotic cell. b) In a DNA analysis 60% Adenin was found, calculate cytosine content. Enlist rDNA products available in the market. c) d) What is primase? Differentiate between DNA and RNA. e) f) What is RNA dependent DNA polymerase? g) What are histone proteins? Q.2 a) Describe molecular events during PCR amplification. (08)Write a short note on PCR applications. (07)Q.3 Describe eukaryotic protein synthesis. (15)Q.4 Write note on ANY THREE (15)a) Frame-shift mutation Mutation in sickle cell anemia b) Griffith's experiment c) d) RNA splicing **SECTION-II** Q.5 Answer any FIVE of the following: (10)Define enzymes a) What is a single cell protein? b) Give applications of streptokinase c) Enlist factors affecting enzyme reaction. d) Differentiate encapsulation and adsorption. e) What is a fermentation media? f) Write different types of enzyme inhibitions. g) What is protein engineering? What are its objectives? Q.6 a) (07)Describe various methods involved in site-directed mutagenesis. b) (08)Q.7 Describe different types of bioreactors. Explain their advantages and (15) applications. **Q.8** Write note on ANY THREE (15)

Applications of enzymes in food industry

Whole-cell immobilization

Spray-drying

Microbes in extreme environments

a)

b)

c)

d)

PURUS – VI (2011 COURSE): SUMMER - 2017 SUBJECT: PHARMACOGNOSY-II

| Day Date | | Withitian | : 10.00 AM TO 01.00 PM Marks : 80 |
|-------------|----------|--|--------------------------------------|
| N.B.: | | | |
| | 1) | Q.No.1 and Q.No.5 are COMPULSORY. Out of | remaining questions attempt |
| | • | ANY TWO questions from each section. | SEDADATE anama basis |
| | 2) | Answers to both the sections should be written in S | SEPARATE answer books. |
| _ | 3) | Figures to the right indicate FULL marks. | |
| | | SECTION – I | |
| Q.1 | | Attempt ANY FIVE of the following | (10) |
| | a) | Write two main chemical constituents present in Garlio | c and draw structure of |
| | | the same. | |
| | b) | Explain percolation. | |
| | c) | What are neutraceuticals? Give examples. | |
| | , | What is maceration? | |
| | e) | Write principle of HPLC. | onby in Uarhal drug |
| | f) | Which are different advantages of column chromatogra analysis? | apny in Herbai drug |
| Ω2 | ۵) | Write history, current status and commerce of neutrace | uticals. (08) |
| Q.2 | a) b) | Role of HPTLC in herbal drug analysis. | (07) |
| | U) | Role of the Lee in herour drug undigoto. | , |
| Q.3 | a) | Write note on Microwave assisted extraction. | (08) |
| ~. | b) | What is SFE? Also explain continuous hot extraction. | (07) |
| | ~, | | |
| Q.4 | | Write note on ANY THREE of the following | (15) |
| _ | a) | Cucumber | |
| | b) | Principle and applications of TLC | |
| | c) | Onion | |
| | d) | Arnica | |
| | e) | Fenugreek | |
| | | SECTION – II | |
| Q.5 | | Attempt ANY FIVE of the following | (10) |
| Ų.S | a) | Define the term callus and totipotency. | |
| | b) | Write composition of nutrient media used in PTC. | |
| | c) | What is explant? | |
| | d) | Which are different quality control parameters for face | pack? |
| | e) | Write composition of herbal hair care product. | |
| | f) | Which are different advantages of herbal cosmetics? | |
| | -, | | (08) |
| Q.6 | a) | How will you control quality of herbal cosmetics? | (07) |
| - | b) | Role of growth hormone in PTC. | (07) |
| | | | oal drugs. (08) |
| Q.7 | a) | Write current status of regulatory affairs related to herb | , an anger |
| | b) | Write in detail composition of culture media along with | |
| | | Write note on ANY THREE of the following | (15) |
| Q.8 | | Protoplast culture | |
| | a) | Herbal skin care product | |
| | b) | Production of Hair care products | |
| | c) | History of tissue culture | |
| | d) | Organization of tissue culture laboratory | |
| | e) | Organization of tissue editors the * * * | |