SEAT No.

Number of printed pages: 02

[65]

SARDAR PATEL UNIVERSITY

M. Sc Pharmaceutical Chemistry (Semester-III) Examination Thursday, 02/11/2017; Time-2:00 PM to 5:00 PM

SUBJECT CODE: PS03CPCH01

SUBJECT TITLE: Drug Design and Development

Maximum Marks: 70

Note: (1) All questions are compulsory.

(2) Figure to right indicates total marks of question.

1×8

Choose the correct option for the following: Q-1

Term measures the e-withdrawing or e-donating influence of substituents and can be measured experimentally is known as:

a. Taft's Steric Factor

b. Hammett Substituent Constant

c. Williamson's constant

d. None

Comperative Molecular field analysis is known as

a. CoMFA

b. CMFA

c. CMoFA

d. CMFiA

Maximize the interaction and minimize the side effect is:

a. Drug Optimization

b. Drug regulation

d. None

c. Drug targeting The important functional group of drug responsible for interaction is: b. Pharmacophore

a. Prodrug

c. Functionality

d. Can't say

Phenyl group interact with binding sites by means of

a. Hydrophobic region

b. Electrovalent bond

Covalent bond

d. Hydrogen bond

If drugs are readily metabolized and stay moderately in the body can cause

a. Good therapeutic effect

b. Diagnostic effect

c. Side effect

d. Can't say

When an antagonist is bound to its receptor for long period of time is:

a. Desensitization

b. Tolerance

c. Coherence

d. Sensitization

Measure of the maximum biological effect that a drug can produce is:

a. Tolerance

b. Coherence

c. Dependence

d. Eficacy

Answer the following (Any Seven). Q-2

Describe the importance of (log P) value. 1.

Define the QSAR. 2

Define Molar Refractivity (MR)? 3.

Describe the Lead compound. 4.

Highlight the antisense approach of drug design.

 2×7

	6	Discuss the Process development.	
	7		
	8	· ·	
	9		
Q-,	3 A		
	В	Explain Hansch Equation.	6
		OR	6
	B	note on QSAR.	
Q-4	~	rational approach of Drug design.	6
	В.	Explain the Binding role of aromatic rings, esters and amines	6
	_	OR	U
	В.	Write a note on Drug Development.	6
Q-5	A.	Elaborate the various strategies to improve the absorption	6
	В.	What is meant by targeting drugs? How the drug on GIT infection and Tumor cells can be targeted?	6
		OR	
	В.	Describe the strategies adopted to making days leave	
Q-6	A.	metabolism. What is receptor? How does the message get received?	6
	В.	How the antagonists can be designed which	6
		How the antagonists can be designed which can act at the binding site and out of the binding site.	6
r		OR	
•	В.	Elaborate Competitive (reversible) inhibitors with suitable examples.	6

SEAT No._ SARDAR PATEL UNIVERSITY M. Sc Pharmaceutical Chemistry (Semester-III) Examination Monday, 06/11/2017; Time-2:00 PM to 5:00 PM SUBJECT CODE: PS03CPCH02 SUBJECT TITLE: Medicinal Chemistry-II Maximum Marks: 70 Note: (1) All questions are compulsory. (2) Figure to right indicates total marks of question. 1×8 Choose the correct option for the following: Q-1 The drug had a calming effect on all animals is known as: 1. b. Analgesic a. Tranquilizer c. Symptomatic d. None Motor system disorder of the nervous system which is outlined as a progressive disorder that affects movement and results in the loss of dopamine-producing brain cells: b. Parkinson's disease a. Alzheimer d. Hypoglycemia c. Hyperthyroidism Medicines that boost brain activity, increase energy, attention, and alertness, and elevate blood pressure, heart rate and respiratory rate: **b.** Alzheimer a. Hyperthyroidism d. CNS stimulants c. Muscle relaxant A diverse group of medicines that have the ability to relax or reduce tension in muscle is known as: b. Hyperthyroidism a. Muscle relaxants d. CNS stimulants c. Alzheimer Compound typically are not stored within cells but rather synthesized as required: b. Analgesic a. Antipyretics d. None c. Eicosanoids Substances that reduce fever: b. antipyretic a. Analgesic d. None c. Both can be used Drugs which mimic the action of stimulation of cholinergic receptors and also called as: b. Cholinomimetic a. Parasympathomimetics d. All the above c. Cholinergic drugs A drug that opposes the downstream effects of postganglionic nerve firing

Answer the following (Any Seven). Q-2

a. Sympatholytic drug

c. Cholinergic drugs

d. none

b. Parasympatholytic drug

in effector organs innervated by the sympathetic nervous system:

Give the uses of Sedatives and hypnotics. 1.

Give the stages of anesthesia. 2

1

 2×7

	3.	Define tranquillizers.	
	4.	Give the limitation of opioid Analgesics.	
	5.	Define the Skeletal Muscle relaxants.	
	6.	Draw the structure of Paracetamol.	
	7.	Draw the structure of caffeine and also give its uses.	
	8.	Describe autonomic nervous system.	
	9.	Define Antipyretic analgesics with suitable example.	
Q-3	A.	Give the classification of General anesthetics. Give the metabolism and mode of action of Barbiturates.	(
	В.	Write a note on the Tranquillizers.	(
		OR	
	В.	Describe the drugs used in Alzheimer's disease	6
Q-4	A.		•
	В.	Give the general mode of action of Narcotic analgesics and SAR of Morphine.	4
-		OR	C
	В.	Give the classification skeletal muscle relaxants. Give the synthesis and mode of action of Mephensin.	6
Q-5	A.	Give the Classification of NSAID and also describe the SAR of Salicylates.	6
	В.	Give the general structure of PG, its mode of action and side effect.	6
		OR	
	В.	Give the synthesis and metabolism of:	6
		i. Celecoxib ii. Ibuprofen	
Q-6	Α.	Give the synthesis of Epinephrine and describe the structure activity relationship of sympathomimetic.	6
	B.	Give the synthesis and mode of action of atropine.	(
		OR	6
	B.	Elaborate the parasympathomimetic drugs? Give the synthesis of Acetylcholine and its mechanism of action.	6

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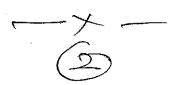
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M. Sc. Pharmaceutical Chemistry, Third Semester Examination

	Wednesday, 8 th November, 2017 02:00 a.m. – 05:00 p.m.				
		PS03CPCH03:	Spectroscopic Te	os:00 p.m. echniques for Quality	Combin
No	ote:			ques for Quality	Max Marks: 70
	1.	Figures to the right indic	ate marks.		INION INION VO. VO.
-		Draw neat and labeled d		r necessary.	
Q-	1 .	Attempt the followings			[08 X 01 =08]
		1. A molecule can change in its	absorb IR freq	uency only if its a	absorption causes a
		a) Electric dipole	b) Magnetic o	dipole c)Both a &	b d) None
		 Induced magnetic causing Shielding of pro c) Shielding of elect 	c field in NMR oton	reinforces the ap	plied magnetic field
				d) Deshielding of	electron
		presence of	are pattern o.	1 M : M+2 :: 100	: 98 indicates the
		a) 2 Br	b) 1 Cl	c) 2 Cl	d) 1 Br
		b) Ionisation, accel c) Acceleration, ion d) Acceleration, de	leration, deflect lisation, deflect lisation, ionisat	ion, detection ion, detection ion, detection	ectrometer is
	5	· identifies	the functional g	groups present in th	ie samnle
		aj NMR	b) Mass spectr	a c) FTIR	d) None of these
	- 6	a) Nucleic acid c) Ammonium sulpi	hate	b) Quinine sulpha d) None of these	te
	7.	In order to perform solvent in order to a) get higher signal c) increase reactivit	to noise ratio	the sample is preparations read the sample is protons read None of these	
	8.	Sigma bonded elect		-y 1.0210 of these	
0.0		a) Visible rays	b) UV rays	c) X – rays	d) γ – rays
Q-2	An 1. 2. 3. 4. 5. 6. 7. 8.	Compare the compo Classify different an What is mull technic Explain bathochrom Discuss the source of Why TMS is used as Differentiate between Define Beer's law.	nent of UV and alytical method que? ic and hypsoch of radiation used internal referen	Visible spectromets. romic shifts. d in NMR.	

Enlist the different types of analyzer used in mass spectroscopy.

Ų-3	(A) (B)	Discuss in detail on automizers used in atomic absorption spectroscopy.	[06 [06]
	(B)	OR Draw neat and labeled schematic diagram of fluorimeter and explain.	[06]
Q-4	(A) (B)	Write a note on flame emission spectroscopy. Describe the different types of monochromator used in UV-Visible spectroscopy.	[06] [06]
	(B)	OR Discuss the various applications of UV-Visible spectrometry.	[06]
05	(0)		رددا
Q5	(A)	Discuss the types, characteristics and importance of vibrations observed in IR spectroscopy.	[06]
	(B)	Describe the spin-spin coupling of NMR in detail	[06]
	(B)	Giving reasons calculate the ¹ H NMR chemical shift of the following molecules	[06]
		$\begin{array}{cccccccccccccccccccccccccccccccccccc$	
Q6	(A) (B)	Derive the equation $m/z = B^2r^2/2V$ in Mass spectrometry Determine the molecular formula of compound by given molecular fragment data	[06] [06]
		M - M + 1	
		m/z 142 143	
		Abundance 100 1.14	
	(B)	What is mass spectroscopy? Discuss the EI method for obtaining the spectrum by mass spectroscopy with suitable example.	[06]



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SARDAR PATEL UNIVERSITY

M.Sc. Pharmaceutical Chemistry, Third Semester Examination Friday, $10^{\rm th}$ November 2017

2.00 p.m. to 5.00 p.m.

Advance Pharmaceutical Chemistry: PS03EPCH01

Total Marks: 70

Note:	(i) All questions are to be attempted. (ii) Figures to the right indicate marks.	
Q.1 (i)	Choose the correct option for the following: Which route of administration of a drug results in 100 % bioavailability? (a) oral (b) topical (c) parenteral (d) rectal	[08]
(ii)	Drug solubility can be enhance by	
(iii)	The Ozone disinfection	
(iv)	Gamma-rays are used to sterilize antibiotic, hormones, sutures, plastics and catheters etc, which is obtained from	
(v)	An increase in fluid intake over fluid output, and occurs when the increase in the volume of body fluids is due to water alone is called	
(vi)	Pilot plant studies include the close examination of the formula to determine (a) process modification (b) cost factor (c) both 'a' & 'b' (d) none	
(vii)	A dry powder blend that cannot be directly compressed because of poor flow or compression properties is known as	
(viii)	(a) slugging (b) capsuling (c) gelling (d) chipping Which type of plastics are commonly used for pharmaceutical products? (a) polyethylene (b) polypropylene (c) PVC (d) all	
Q.2 (i) (ii)	Answer the following: (Attempt any seven) Define systemic and relative bio availability. Explain the various parameter used in bioavailability determination by plasma level study.	[14]
(iii) (iv) (v)	What is the objective of Sewage treatment? Describe the parameters which are influencing D-Value. What is pilot plant?	
(vi)	What the processes must be included for transferring research to QA department?	
(vii)	Distinguish: Quality control and quality assurance.	i
(viii)	Write main function of QC department of Pharmaceutical Industry.	
(iv)	Write disadvantage of use of plastic containers	

Q.3 [A]	Answer the following: Define bioequivalence. Discuss in detail about bioequivalence experimental study	[06]
	design.	Ì
[B]	Discuss experimental study design for bioavailability measurement. OR	[06]
[B]	Discuss various methods for bioavailability enhancement via drug permeability factor.	[06]
Q.4 [A]	Answer the following: Discuss evaluation and in Process Monitoring of Sterilization procedures.	[06]
[B]	Describe the biological water treatment process.	[06]
-	OR	
[B]	Write a note on Reverse osmosis.	[06]
Q.5	Answer the following:	
[A]	Describe the Pilot Plant design for Tablets.	[06]
[B]	Give the methods for scale up for biotechnology derived products.	[06]
	OR	
[B]	How will you meet the requirement of Scaling-up of powder blending operations?	[06]
Q.6	Answer the following:	
[A]	Discuss in detail about main function of quality assurance department of industry.	[06]
[B]	Describe advantages and limitation of metal containers in pharmaceutical industry.	[06]
	OR	
[B]	Write a note on: Glass as packaging material in pharmaceutical industry.	[06]

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