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GUIARAT TECHNOLOGICAL UNIVERSITY

	•	BE - SEMESTER-IV (NEW) EXAMINATION – WINTER 2018 Code:2143601 Date:05/12	2/2018
Subject Name:Medicinal Chemistry & Physio-pharmacology Time: 02:30 PM TO 05:00 PM Total Mar Instructions:			
	2.	Attempt all questions. Make suitable assumptions wherever necessary. Figures to the right indicate full marks.	
			MARKS
Q.1	(a)	What is Partition Co-efficient ? How is it determined ?	03
	(b)	What are the different types of receptors ? Discuss how the Nuclear Receptors work.	04
	(c)	Write the synthesis of any two drugs describing the manufacturing process: (i). Tripellanamine HCl, (ii). Antazoline HCl, (iii). Chlorcyclizine HCl, (iv). Promithazine HCl	07
Q.2	(a)	Draw a neat & labeled structure of Nephron. Mention the functions of kidney.	03
	(b)	Discuss Hammett Constant. pKa value of benzoic acid is 4.11, pKa value of 4-Nitrobenzoic acid is 3.41. Find the Hammett Constant (σ) for the 4-Nitro substituent.	04
	(c)	What is Fragment Based Drug Design (FBDD)? Explain why FBDD leads to drugs with least side-effects. Four fragments A,B,C and D on screening, yielded a HIT molecule C-A. Show the steps involved.	07
	(c)	Define "solubility" and "rate of dissolution" of a substance. What are the factors affecting the rate of solubility of a drug ? What is "Maximum Absorbable Dose" (MAD) ?	07
Q.3	(a)	What are the characteristics of a "HIT" molecule ?	03
	(b)	Discuss Hansch first model and the improvement on the first model, for biological response.	04
	(c)	Discuss in brief the 04 types of Reversible Enzyme Inhibitors.	07
		OR	
Q.3	(a)	Explain the process of "Docking"	03
	(b)	Write a note on "Cell-based" assay in High Throughput Screening (HTS).	04
	(c)	Explain the SAR of Anilide class of Local anesthetics with suitable examples wherever necessary.	07

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(b)	What are Anticoagulants and what are their classifications? Give suitable examples. Write the MOA of Sodium citrate anticoagulant.	04
(c)	Draw the schematic flow diagram for Drug Discovery Process and mark relevant stages.	07
(\mathbf{a})		07
(a)	1 0	03
(b)	Write the synthesis of (i). Benzocaine, (ii). Lidocaine.	04
(c)	What are the different types of forces involved in Drug-Receptor interactions? Discuss the role of Hydrophobic Forces in the interaction	07
(a)	What is the general equation for biological response of a 3D QSAR model ?	03
(b)	Write a note on: (i). Anemia & Hematinics, (ii). Thrombolytics, (iii). Antiplatelet drugs. Give suitable examples	04
(c)	Describe "Loop & High Ceiling" Diuretics. What are its disadvantages?. Give two examples. Write the synthesis of any one drug.	07
	OR	
(a)	Discuss the filing of Abbreviated New Drug Application (ANDA).	03
(b)	Write the synthesis of anticoagulant drug-Warfarin. Write its MOA.	04
(c)	Write the SAR of H1 receptor Antihistamines. Give suitable examples wherever required.	07
	 (c) (a) (b) (c) (a) (c) (a) (b) (c) 	 (c) Draw the schematic flow diagram for Drug Discovery Process and mark relevant stages. OR (a) Describe "Acidotic Diuretics" with suitable examples and MOA of any one drug. (b) Write the synthesis of (i). Benzocaine, (ii). Lidocaine. (c) What are the different types of forces involved in Drug-Receptor interactions? Discuss the role of Hydrophobic Forces in the interaction (a) What is the general equation for biological response of a 3D QSAR model ? (b) Write a note on: (i). Anemia & Hematinics, (ii). Thrombolytics, (iii). Antiplatelet drugs. Give suitable examples (c) Describe "Loop & High Ceiling" Diuretics. What are its disadvantages?. Give two examples. Write the synthesis of any one drug. (a) Discuss the filing of Abbreviated New Drug Application (ANDA). (b) Write the synthesis of anticoagulant drug-Warfarin. Write its MOA.