UNIVERSITY OF JAFFNA, SRI LANKA BACHELOR OF PHARMACY

THIRD YEAR SECOND SEMESTER EXAMINATION – MARCH 2019 PHAMC 3214 MEDICINAL CHEMISTRY II – PAPER II

Date: 25.03.2019 Time: 02 Hours

Answer all six questions

1.			
	1.1	1.1 Draw the structure of morphine and discuss its structure activity relations hip (SAR).	(70 Marks)
	1.2	Give three examples of antagonist of morphine and draw the chemical	(10111111111111111111111111111111111111
		for each of them.	(30 Marks)
2.			
	2.1 2.2	Draw the structure of serotonin. 2.2.1 Structure 'X' is lead compound for 5- HT ₂ c receptor. Explain how structure 'X' is converted to structure 'Y' through drug	(20 Marks)
		optimization process.	
		CH ₂ CH ₃ CH ₂ CH ₃ H N N N N N CH ₃ CH ₃ CH ₃ CH ₃	
		Structure 'X' Structure 'Y'	
		Structure 'X' Structure 'Y'	(40 Marks)
		2.2.2 Discuss the ring variation strategy for activity of structure 'Y'.2.2.3 Draw the schematic diagram of molecular modelling studies for	(40 Marks) (20 Marks)
		2.2.2 Discuss the ring variation strategy for activity of structure 'Y'.	
2		2.2.2 Discuss the ring variation strategy for activity of structure 'Y'.2.2.3 Draw the schematic diagram of molecular modelling studies for	(20 Marks)
3.	3.1	2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y'	(20 Marks)
3.	3.1	2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y' Describe the structure activity relationship of aryl alkanoic acid	(20 Marks) (20 Marks)
3.	3.1 3.2	2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y'	(20 Marks) (20 Marks) (70 Marks)
		2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y' Describe the structure activity relationship of aryl alkanoic acid containing anti inflammatory drugs.	(20 Marks) (20 Marks)
3.	3.2	2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y' Describe the structure activity relationship of aryl alkanoic acid containing anti inflammatory drugs. Draw synthetic route of carbimazole.	(20 Marks) (20 Marks) (70 Marks) (30 Marks)
		2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y' Describe the structure activity relationship of aryl alkanoic acid containing anti inflammatory drugs. Draw synthetic route of carbimazole. Describe the different mechanisms of action of antiepileptic drugs. Classify antiepileptic drugs based on their chemical structure with	(20 Marks) (20 Marks) (70 Marks)
	3.2 4.1	2.2.2 Discuss the ring variation strategy for activity of structure 'Y'. 2.2.3 Draw the schematic diagram of molecular modelling studies for receptor binding of structure 'Y' Describe the structure activity relationship of aryl alkanoic acid containing anti inflammatory drugs. Draw synthetic route of carbimazole. Describe the different mechanisms of action of antiepileptic drugs.	(20 Marks) (20 Marks) (70 Marks) (30 Marks)

5.1	Explain why N^{α} - Guanyl histamine acts as a partial agonist?	(40 Marks)
5.2	Illustrate the mechanism of inhibition of proton pump by Omeprazole with the help of curly arrows.	(60 Marks)
6.1	Give two examples with chemical structure of	
		(20 Marks)
		(20 Marks)
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0.22		(30 Marks)
6.3		(30 Marks)
	5.2	 5.2 Illustrate the mechanism of inhibition of proton pump by Omeprazole with the help of curly arrows. 6.1 Give two examples with chemical structure of 6.1.1 ester based local anaesthetics. 6.1.2 amide based local anaesthetics. 6.2 Illustrate the structural characteristics of a local anaesthetic using a diagram.