UNIVERSITY OF JAFFNA BACHELOR OF PHARMACY

FOURTH YEAR SECOND SEMESTER EXAMINATION MARCH 2019 PHADD 4223 DRUG DISCOVERY AND DEVELOPMENT

Date: 22.03.2019 Time:03 Hours

ANSWER ALL 6 QUESTIONS

	Describe the following as drug targets with examples.	
1.1	Enzyme	(50 Marks)
1.2	Receptor	(50 Marks)
2.1	What is the purpose of the High Throughput Screening (HTS)?	(10 Marks)
2.2	Briefly explain major five components in the HTS?	(30 Marks)
2.3	List five detection methods used in HTS?	(15 Marks)
2.4	Briefly describe three fluorescence techniques that can be used for HTS?	(45 Marks)
3.1	Name the optical biosensor technology with label free principal.	(10 Marks)
3.2	Write the advantages and disadvantages of above mentioned technology	(15 Marks)
3.3	List the information obtained from the above mentioned technology.	(10 Marks)
3.4	Write the principle of the above mentioned technology.	(65 Marks)
4.1	What is the aim of the <i>de novo</i> drug design?	(10 Marks)
4.2	List the advantages of solid phase synthesis.	(30 Marks)
4.3	Write the requirements for the solid phase synthesis	(20 Marks)
4.4	Describe the protection strategy used in the solid phase synthesis	(40 Marks)
Desc	cribe.	(50 Marks)
5.1	Phase I clinical trail.	(50 Marks)
5.2	Phase III clinical trail.	
Writ	te an account on:	
6.1	Approval procedure of clinical trail in Sri Lanka	(25 Marks)
6.2	Cross over studies	(25 Marks)
6.3	Pre-clinical animal studies	(50 Marks)
	1.2 2.1 2.2 2.3 2.4 3.1 3.2 3.3 3.4 4.1 4.2 4.3 4.4 Desc 5.1 5.2 Writ 6.1 6.2	 Enzyme Receptor What is the purpose of the High Throughput Screening (HTS)? Briefly explain major five components in the HTS? List five detection methods used in HTS? Briefly describe three fluorescence techniques that can be used for HTS? Name the optical biosensor technology with label free principal. Write the advantages and disadvantages of above mentioned technology List the information obtained from the above mentioned technology. Write the principle of the above mentioned technology. Write the advantages of solid phase synthesis. Write the requirements for the solid phase synthesis Describe the protection strategy used in the solid phase synthesis Phase I clinical trail. Phase III clinical trail. Approval procedure of clinical trail in Sri Lanka Cross over studies