Name:

Enrolment No:



UNIVERSITY OF PETROLEUM AND ENERGY STUDIES End Semester Examination, May 2022

Course: Biopharmaceutics and Pharmacokinetics Program: B. Pharm. Course Code: BP 604 T Semester: VI Time: 03 h. Max. Marks: 100

Instructions: All the sections are compulsory.

SECTION A						
S. No.	CO			Marks		
		Answer all the questions.		20		
1.	CO1	Which is the major process of absorption for a	nore than 90% of drugs?	1		
		A. Facilitated diffusion	B. Absorption diffusion			
		C. Passive diffusion	D. Pinocytosis			
2.	CO1	Which one of the following is a critical rate-li soluble drugs?	miting step of drug absorption of poorly	1		
		A. Rate of dissolution	B. Rate of permeation			
		C. Size of the drug	D. Rate of drug release			
3.	CO1	Fick's law is used to explain		1		
		A. Dissolution rate	B. Disintegration rate			
		C. Diffusion rate	D. Dissociation rate			
4.	C01	distributed	metabolism r's body due to the presence of uterus, etus increases the volume thus increasing separate compartment for a drug to get	1		
5.	CO1	Which one of the following organ has a high perfusion rate?				
		A. Fats	B. Kidney			
	COA	C. Skin	D. Bone	1		
6.	CO2	Biotransformation of drugs is defined as the conversion from one physical form to another.		1		
		A. True B. False				
7.	CO2	Which enzymes catalyzes the hydrolysis of amides?				
		A. Esterase B.	Amidase			
		C. Amydase D.	Aminodase			
8.	CO2	2 Which one of the following is the principal organ for drug excretion?				
		A. Kidney B.	Liver			
		C. Lung D.	Skin			

9.	CO2 Which of the following is not a factor influencing pulmonary excretion?			
		A. Pulmonary blood flow B. Rate of respiration		
		C. The solubility of volatile substance D. Heart rate	1	
10.	CO2	What will be the elimination rate if the clearance is 130 ml/min and drug concentration is 0.8 g/ml?		
		A. 104 g/min B. 140 g/min		
		C. 130 g/min D. 100 g/min		
11.	CO2	Drugs of molecular range 300-500 Dalton will be excreted out by glomerulus.A.TrueB. False		
12.	CO3	What is the equation to find out the apparent volume of distribution?		
12.		A. Amount of drug in the body/plasma drug concentration		
		 B. Plasma drug concentration/amount of drug in the body C. 1 / plasma drug concentration D. 1 / Amount of drug in the body 	С	
13.	CO3 In which of the model peripheral compartments are connected to a central compartment?			
		A. Compartment model B. Mammillary model		
		C. Caternary model D. Physiological model		
14.	CO3	What is the chemical equivalence?	1	
		 A. Two or more drug products contain the same labeled chemical substance in the same amount B. Two or more drug products contain the same labeled chemical substance in different quantity C. Two or more drug products contain different labeled chemical substance giving the same therapeutic effect D. Two or more drug products contain the same labeled chemical substance giving a different therapeutic effect 		
15.	CO3	The period for which the plasma concentration of drug remains above minimum effective concentration is known as		
		A. Onset of time B. Duration of action		
		C. Onset of action D. Therapeutic range		
16.	CO3	Multiple dose study is better since we can understand the peak, valley, drug blood levels, etc. A. True B. False	1	
17.	CO4	Draw a block diagram for two compartment open model extravascular administration.	1	
18.	CO4	In the given picture, the marking "?" represents the drug concentration of which compartment? A. The central compartment in a two compartment model B. Peripheral compartment in a two compartment model C. The central compartment in a one compartment model D. Drug concentration of the plasma	1	
19.	C05	The characteristic of non-linear pharmacokinetics include A. Area under the curve is proportional to the dose B. Elimination half-life remains constant		

		C. Area under the curve is not proportional to the dose D. Amount of drug excreted through remains constant	
20.	CO5	In Michaelis-Menton Equation, when the value of Km <<< C, rate of the process is equal to maximum rate of reaction.	1
		A. True B. False	
		SECTION B	
iswer	any two	questions of the following.	20
1.	CO5	What are the causes of non-linearity found in pharmacokinetics (ADME) of drug?	
2.	CO3	 If the plasma concentration of vancomycin after IV bolus administration was found to be 20.0 and 11 μg/mL at 4 and 8 hours, respectfully. By assuming one compartment open model, calculate following parameters: a) The elimination rate constant (3 marks) b) half-life of the drug (3 marks) c) Concentration of drug at zero time (4 marks) 	
3.	CO2	Describe any five factors affecting renal excretion with an example.	4+6
		SECTION C	
iswer	any sev	en questions of the following.	35
1.	CO1	Differentiate active and passive transport.	5
2.	CO1	Explain the term 'volume of distribution'.	1+4
3.	CO1	Write a short note on blood-brain barrier.	5
4.	CO2	"Drug interactions can be employed for selective excretion of some drugs". Explain the statement with an example.	5
5.	CO3	 Half-life of a drug in 60 kg patient is 5 hours and Vd is found to be 5 L/kg. a. Determine the total systemic clearance b. Determine the renal clearance if the fraction excreted unchanged in urine is 0.045. 	5
6.	CO3	Explain bioequivalence and therapeutic equivalence.	5
7.	CO2	Explain glucuronidation in biotransformation.	
8.	CO4	How body tissues are classified considering two compartment model? In which compartment the brain should be classified.	
9.	CO4	Why the rapid decline in initial concentration followed by slower decline is observed in plasma levels of the drug that follows two compartment model?	5
		Total	75