National Board of Examinations

Question Paper Name :	DNB Pharmacology Paper1
Subject Name :	DNB Pharmacology Paper1
Creation Date :	2024-10-17 16:12:22
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DNB Pharmacology Paper1

Group Number :	1
Group Id :	3271872822
Group Maximum Duration :	0
Group Minimum Duration :	180
Show Attended Group? :	No
Edit Attended Group? :	No
Break time :	0
Group Marks :	100

DNB Pharmacology Paper1

Section Id :	3271872825
Section Number :	1
Section type :	Offline
Mandatory or Optional :	Mandatory
Number of Questions :	10
Number of Questions to be attempted :	10
Section Marks :	100
Maximum Instruction Time :	0
Sub-Section Number :	1
Sub-Section Id :	3271872829
Question Shuffling Allowed :	No

Question Number : 1 Question Id : 32718734294 Question Type : SUBJECTIVE Consider As Subjective : Yes Correct Marks : 10

Question Number	Answer to be attempted within	Question Number	Answer to be attempted within
Q. 1	Page 1-5	Q. 6	Page 26-30
Q. 2	Page 6-10	Q. 7	Page 31-35
Q. 3	Page 11-15	Q. 8	Page 36-40
Q. 4	Page 16-20	Q. 9	Page 41-45
Q. 5	Page 21-25	Q. 10	Page 46-50

Please write your answers in the answer booklet within the allotted pages as follows:-

1. Define Pharmacovigilance. Describe Pharmacovigilance Programme of India. Discuss various challenges faced by this Programme. [2+5+3]

Question Number : 2 Question Id : 32718734295 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

Define medication errors. What are the various types of medication errors? Discuss the strategies to prevent medication errors. [2+4+4]

Question Number : 3 Question Id : 32718734296 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

What is plasma half-life of a drug? Discuss clinical importance of knowing plasma half-life of a drug. Write the differences between plasma half-life and biological half-life of a drug. [2+4+4]

Question Number : 4 Question Id : 32718734297 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

Define the terms Bioavailability & Bioequivalence respectively. What are the guidelines for conduct of Bioavailability & Bioequivalence study of Investigational New Drug (IND) as per the "New Drugs & Clinical trials rules 2019"? [(2+2)+6]

Question Number : 5 Question Id : 32718734298 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

Define Biosimilars. How are they different from Biological drugs? Describe the challenges for their use. [2+4+4]

Question Number : 6 Question Id : 32718734299 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

a) Periodic safety update report (PSUR). [5]

b) Receptor regulation. [5]

Question Number : 7 Question Id : 32718734300 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

Define the phenomenon of genetic polymorphism. Describe its role in clinical practice by giving suitable examples. [2+8]

Question Number : 8 Question Id : 32718734301 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

What do you understand by Individualization of drug therapy? Enumerate the factors which influence action of various drugs. Discuss how pathological conditions affect the drug action. [3+3+4]

Question Number : 9 Question Id : 32718734302 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

Compare & contrast between the following:

- a. Autoreceptors versus Heteroreceptors. [5]
- b. Drug specificity versus Drug selectivity. [5]

Question Number : 10 Question Id : 32718734303 Question Type : SUBJECTIVE Consider As Subjective : Yes

Correct Marks : 10

Define translational pharmacology. Describe the need of translational pharmacology. Discuss the challenges in translational research. [2+5+3]