BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI (END SEMESTER EXAMINATION)

CLASS: B.PHARM SEMESTER: VIII BRANCH: PHARMACY SESSION: SP/19

SUBJECT: PS8407 CLINICAL PHARMACY & DRUG INTERACTIONS

TIME: 3 Hours FULL MARKS: 60

INSTRUCTIONS:

- 1. The question paper contains 7 questions each of 12 marks and total 84 marks.
- 2. Candidates may attempt any 5 questions maximum of 60 marks.
- 3. The missing data, if any, may be assumed suitably.
- 4. Before attempting the question paper, be sure that you have got the correct question paper.
- 5. Tables/Data hand book/Graph paper etc. to be supplied to the candidates in the examination hall.

Q.1(a) Q.1(b) Q.1(c)	Define clinical pharmacy and mention it's importance. Define ADR and give a detailed classification of ADR according to Wills and Brown. Elaborate the mnemonics WWHAM, ENCORE, AS METTHOD. Mention the steps involved in the pharmaceutical care and consultation process.	[2] [4] [6]
Q.2(a) Q.2(b) Q.2(c)	Define TDM. Enlist the health professionals involved in TDM services and what are their individual roles. Discuss the criteria for conducting TDM. Mention 5 classes of drugs with examples for which TDM is essential.	[2] [4] [6]
Q.3(a) Q.3(b) Q.3(c)	What is cirrhosis? Classify the types of hepatitis. Enumerate the causes of liver cirrhosis. Elaborate on the symptoms, complications and treatment of cirrhosis.	[2] [4] [6]
Q.4(a) Q.4(b)	Define ulcerative colitis. Illustrate the symptoms and diagnostic procedures which differentiates it from other inflammatory conditions.	[2] [4]
Q.4(c)	Discuss the various treatment regimens of ulcerative colitis.	[6]
Q.5(a) Q.5(b) Q.5(c)	Enumerate the methods of drug transport across membranes. What is the significance of Henderson-Hasselbach equation in drug distribution? Write a note on excretion of drugs with examples.	[2] [4] [6]
Q.6(a) Q.6(b) Q.6(c)	What is bioequivalence? Discuss the different drugs used in Alzheimer's disease. Illustrate the pathogenesis of Alzheimer's disease.	[2] [4] [6]
Q.7(a) Q.7(b) Q.7(c)	Define bioavailability. Why do we need to bother about pharmacokinetics in drug discovery? Illustrate the absorption of drug via different routes of administration.	[2] [4] [6]

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