

Total No. of Questions : 6]

SEAT No. :

P3186

[Total No. of Pages : 3

[5245]-805

Final Year B.Pharmacy (Semester - VIII)

PHARMACOLOGY - V (INCLUDING BIostatISTICS)

(2013 Pattern)

Time : 3 Hours]

[Max. Marks :70

Instructions to the candidates:

- 1) *All questions are compulsory.*
- 2) *Answers to the two sections should be written in separate books.*
- 3) *Neat diagrams must be drawn wherever necessary.*
- 4) *Figures to the right indicate full marks.*

SECTION - I

Q1) Classify drug Interaction. Explain in Brief drug interaction during absorption and distribution with suitable example. **[10]**

OR

Define pharmacovigilance. Write in brief about monitoring, detection and reporting of adverse drug reactions (ADR).

Q2) Solve any five : **[15]**

- a) Justify the use of penicillin and probenecid as beneficial drug interaction.
- b) Write the variables affecting drug abuse and misuse.
- c) Explain the laboratory tests for liver dysfunction.
- d) Discuss drug - food interaction with example
- e) Explain the types of hypersensitivity reactions.
- f) Write the applications of stem cell therapy.
- g) Explain the effect of change in pH during drug excretion with example.

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Q3) Write notes on (any two) [10]

- a) Advantages and applications of TDM.
- b) Pharmacodynamic drug interaction.
- c) Drug Induced Diseases.
- d) Carcinogenicity

SECTION - II

Q4) Explain clinical research. Write in brief about different phases of clinical trials. [10]

OR

Write the composition, objectives and process of pharmacy and therapeutic committee in hospital pharmacy.

Q5) Solve any five : [15]

- a) Write about inclusion and exclusion criterion in subject selection.
- b) Define cross over trial, control group and double blinded study.
- c) Explain unit dose drug distribution system with its advantages.
- d) Write about role and responsibilities of contract. Research organization (CRO)
- e) Write the importance of data handling and record keeping in clinical trials.
- f) Explain the role of hospital pharmacist in practice of rational drug therapy.
- g) Ethical issues in nuremberg code.

Q6) Write short notes on (any 2)

[10]

- a) Drug development process
- b) Hospital formulary.
- c) Composition and process of - IRB
- d) Principles of ICH GCP guidelines.

