PHB Education

PHB Sample Paper - 2

D. Pharm IInd Year, THIRD SESSIONAL EXAMINATION, 2022 - 23

(PHARMACY LAW & ETHICS)

Time: 02 Hrs **Maximum Marks: 40**

Instructions:

1. Write the Roll no. on your question paper.
2. Candidate should ensure that the question paper supplied to them is complete in all respects. Complaints in this regards, if any, should be made to the invigilatory staff on the duty in the exam centre within 15 minutes of Commencement of the exam. No complaint shall be entertained thereafter.

(Section: A)

Q. A Multiple-Choice Questions: **(5)**

1. The poison act was passed on……..

a. 13thSeptember1919 b. 3rdSeptember1919

c. 3rdSeptember1918 d. 23rdSeptember1919

2. Which class of drug poorly absorbed by orally administration?

a. Class-I b. Class-II

c. Class-III d. Class-IV

3. Which one is not the principle of bioethics……

a. Justice b. Dependency

c. Non-maleficence d. Beneficence

4. Drug and magic remedy act is also called as :

a. Objectionable Advertisements b. Wrong Advertisements

c. Magic Advertisements d. All of these

5. RMP should have experience in gynaecology and obstetrics for atleast……..

a. 1-year b. 3-year

c. 5-year d. 2-year

Q. B Define the some terminology: **[5]**

1. Generic Drug

2. Clinical Trials

3. Narcotic Drugs

4. Cosmetics

5. CDSCO

(Section: B)

Q. C Short Answer Type Questions: (Attempt any five) (5 × 3 = 15)

1. Give the objective of the Drug and Cosmetics Act,1940 and rules .
2. Write short note on:
   a. Role of FSSAI
   b. Function of FSSAI
3. Explain the function of CPCSEA.
4. Write a short note on duties of a Drug Inspector.
5. Explain MTP amendment act 2021and its significances.
6. Write short note on history and scope of pharmaceutical legislations.

(Section: C)

Q. D Long Answer Type Questions: (Attempt any three) (5 × 3 = 15)

1. Write a detail note on code of pharmaceutical ethics.
2. Explain in detail Drug Inspector including qualification criteria for DI, Power of DI and Duties of Inspectors.
3. Write a note on NDA and ANDA.
4. What are the requirements for import of medical devices in India?