PH-II/6 (N)

2024

PHARMACY LAW & ETHICS

Time Allowed: 3.00 Hours *Full Marks: 80*

1. **Choose the correct alternative** *1X10=10*

i) When did the Pharmacy Act 1948, come into force-

a) 01 April 1949 b) 01 May 1949 c) 04 March 1948 d) 23 April 1948

ii) Joint State Pharmacy Councils are constituted for-

a) 2 years b) 3 years c) 5 years d) Indefinite period as per agreement

iii) What is the life of a patented drug?

a) 10 years b) 15 years c) 20 years d) 25 years

iv) Premises licensed for sale of drug having qualified person but drugs are not compounded in it, called:

a) Pharmacy b) Chemist & Druggist c) Drug vendor d) Chemist Shop

v) Medical devices are used for which purpose?

a) Diagnosis b) Prevention c) Monitoring d) All of these

vi) Which phase is also known as "Confirmation Phase"?

a) Phase 2 b) Phase 4 c) Phase 3 d) Phase 0

vii) In a purple container which waste is collected?

a) Syringes b) Hazardous waste c) Cytotoxic and cytostatic waste d) None of the above

viii) DTAB contains _____ Ex- Officio members-

a) Five b) Six c) Seven d) Eight

ix) _____ is an example of natural poison.

a) Carbolic Acid b) Chloroform c) Digitalis d) Mercury oxide

x) The chairman of DTAB is the-

a) Director of Central Drug Laboratory b) Director of Central Drug Research Laboratory

c) Director of General Health Services d) Drug Controller of India

2. **Answer the following questions.** *1X10=10*

i) Define Biomedical Waste.

ii) The latest Drug Price Control Order (DPCO) comes into force in _____

iii) What is the full form of ICMR?

iv) The full form of DTAB is _____

v) What is EUA?

vi) Insulin injection comes under schedule _____

vii) Radioactive waste is covered under _____

viii) FSSAI means _____

ix) Schedule N deals with _____

x) The full form of cGMP is _____

3. Answer the following questions (any TEN) *3X10*

i) Discuss details on NLEM.
ii) What are the general requirements of GRP?
iii) Define code of ethics. Explain receiving and handling of prescriptions by Pharmacists.
iv) Explain in detail "The Drug Price Control Order Act 1995".
v) Define the patent as per Patent Act. Which inventions are not patentable under this act.
vi) What are the objectives of 'Narcotic Drug and Psychotropic Substances Act 1985'.
vii) Write a brief note on Clinical Establishment Act.
viii) Define: (a) Magic Remedy, (b) Advertisement, (c) Registered Pharmacist.
ix) What is a blood bank? Write the criteria for blood donation.
x) What are the functions of pharma regulatory bodies?
xi) Mention the vision, mission and objectives of Indian Pharmacopoeia Commission.
xii) Explain in detail about State Pharmacy Councils.

4. Answer the following questions (any six) *5X6=30*

i) Mention the qualification, power of a Drug Inspector.
ii) Discuss the constitution and functions of PCI.
iii) Define magic remedies. Give the constitution and functions of the Institutional Animal Ethical Committe
iv) What are the objectives of the Consumer Protection Act?
v) Explain the various licenses issued under Drugs and Cosmetics Act 1940.
vi) What are the minimum qualifications for appointment as a Government Analyst?
vii) Discuss under what circumstances termination of pregnancy is legal? Who is the person eligible for terminating a pregnancy?