

B. PHARM.
(SEM-III) THEORY EXAMINATION 2019-20
PHARMACEUTICAL JURISPRUDENCE AND ETHICS

Time: 3 Hours**Total Marks: 100****Note:** Attempt all Sections. If require any missing data; then choose suitably.**SECTION A****1. Attempt all questions in brief. 2 x 10 = 20**

a.	Define Adulterated Drugs.
b.	Explain steps for manufacturing of new drug.
c.	Define Schedule J.
d.	What do you mean by Drug Analysts?
e.	Define Patent.
f.	What are the objectives of Trademark Act?
g.	Define Ayurvedic & Homeopathic Medicines.
h.	What do you mean by Suspend Registration?
i.	Write the qualifications of Registered Pharmacist.
j.	Discuss about Enquiry Committee.

SECTION B**2. Attempt any three of the following: 10x3=30**

a.	Describe Drugs and Cosmetics Act 1940 with reference to Schedule C,C1 and X.
b.	Give detail study of schedule G,M and Y.
c.	What is the constitution and functions of Narcotic & Psychotropic Consultative Committee?
d.	What is difference between Retail Price and Ceiling price of scheduled formulations?
e.	Discuss the terms and conditions of Medical Termination of Pregnancy Act.

SECTION C**3. Attempt any one part of the following: 10x1=10**

a.	Discuss in detail about Offence and Penalties under Import of Drugs.
b.	Classify the drugs and cosmetics prohibited from import.

4. Attempt any one part of the following: 10x1=10

a.	Write in detail on Drug Technical Advisory Board (DTAB).
b.	What do you mean by Labeling and Packing of Drugs?

5. Attempt any one part of the following: 10x1=10

a.	Write a note on Manufacture, Sale and Export of Opium.
b.	Discuss the objectives and main functions of PCI.

6. Attempt any one part of the following: 10x1=10

a.	Write a detail note on Drugs and Magic Remedies Act.
b.	Explain the role of Drug Price Control Order (DPCO).

7. Attempt any one part of the following: 10x1=10

a.	What is role and responsibilities of AICTE in Pharmaceutical Jurisprudence?
b.	Write a detail note on Pharmaceutical Legislations.