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BPHARM
(SEM V) THEORY EXAMINATION 2025-26
PHARMACEUTICAL JURISPRUDENCE – THEORY

TIME: 3 HRS

M.MARKS: 75

Note: 1. Attempt all Sections. If require any missing data; then choose suitably.

SECTION A

1. Attempt all questions in brief. 10 x 2 = 20

a.	Define misbranded drug as per Drugs and Cosmetics Act, 1940.
b.	State any two functions of the Drugs Technical Advisory Board (DTAB).
c.	What is Schedule H of Drugs and Cosmetics Rules, 1945?
d.	Mention any two objectives of the Pharmacy Act, 1948.
e.	Define Registered Pharmacist.
f.	What is magic remedy under Drugs and Magic Remedies Act?
g.	Define ceiling price under DPCO, 2013.
h.	Define Pharmaceutical ethics?
i.	What is NLEM?
j.	What is meant by loan licence?

SECTION B

2. Attempt any two parts of the following: 2 x 10 = 20

a.	Describe the Drugs and Cosmetics Act, 1940 with respect to: a) Objectives b) Import of drugs c) Prohibition of manufacture and sale d) Offences and penalties
b.	Explain Sale of drugs under Drugs and Cosmetics Rules, 1945 covering: a) Wholesale licence b) Retail licence c) Restricted licence d) Conditions for sale
c.	Discuss the Narcotic Drugs and Psychotropic Substances Act, 1985 with reference to: a) Objectives b) Authorities and officers c) Control and regulation d) Offences and penalties

SECTION C

3. Attempt any five parts of the following: 7 x 5 = 35

a.	Explain conditions for grant and renewal of manufacturing licence for drugs
b.	Describe the constitution and functions of Pharmacy Council of India (PCI) and explain the registration of pharmacists under pharmacy act,1948
c.	Write a short note on Code of Pharmaceutical Ethics.
d.	Explain intellectual property rights (IPR) and Explain the Right to Information (RTI) Act and its relevance to pharmacy profession.
e.	Write a note on Drugs and Magic Remedies (Objectionable Advertisements) Act.
f.	Explain the Prevention of Cruelty to Animals Act, 1960 with reference to animal ethics committee.
g.	Describe Drug Price Control Order (DPCO), 2013 and its objectives.