

PHARMACEUTICAL JURISPRUDENCE (GUESS PAPER 2025–26)

SECTION A (10 × 2 = 20)

1. Define adulterated, misbranded and spurious drugs.
2. Define Schedule G and Schedule M.
3. What is Drug Technical Advisory Board (DTAB)?
4. Define registered pharmacist.
5. Write objectives of the Pharmacy Act, 1948.
6. What is DPCO? Write its objective.
7. Define narcotic drugs with examples.
8. Write functions of Drug Inspector.
9. What is CPCSEA?
10. Define intellectual property rights (IPR).

SECTION B (Attempt any TWO – 2 × 10 = 20)

1. Explain Schedule M (GMP) under Drugs and Cosmetics Act, 1940.
2. Discuss Right to Information Act (RTI).
3. Describe constitution and functions of Pharmacy Council of India (PCI).
4. Explain Drug Price Control Order (DPCO) with objectives.

SECTION C (Attempt any FIVE – 5 × 7 = 35)

1. Discuss Drug and Cosmetics Act, 1940 with offences and penalties.
2. Explain qualifications, powers and duties of Drug Inspector.
3. Describe Narcotic Drugs and Psychotropic Substances Act, 1985.
4. Explain procedure for registration of pharmacists under Pharmacy Act, 1948.
5. Write a note on Drug & Magic Remedies Act, 1954.
6. Discuss Intellectual Property Rights (IPR) in detail.
7. Explain CPCSEA guidelines for breeding and stocking of animals.

■ QUESTION FREQUENCY ANALYSIS (2019–2025)

Topic	Times Asked	Exam Weight
Drug & Cosmetics Act, 1940	7 / 7	■■■■■
Schedule M (GMP)	7 / 7	■■■■■
Drug Inspector – Powers & Duties	6 / 7	■■■■
Pharmacy Act & PCI	6 / 7	■■■■
DPCO / NLEM	6 / 7	■■■■
NDPS Act, 1985	5 / 7	■■■■
Drug & Magic Remedies Act	5 / 7	■■■■
CPCSEA Guidelines	5 / 7	■■■■
IPR & Patents	4 / 7	■■■