

# **2010 Scheme**

**QP CODE: 305006**

**Reg. No: .....**

## **Third Year B.Pharm Degree Supplementary Examinations March 2025 Pharmaceutical Jurisprudence**

**Time: 3 Hours**

**Total Marks: 100**

- *Answer all questions to the point neatly and legibly • Do not leave any blank pages between answers • Indicate the question number correctly for the answer in the margin space*
- *Answer all parts of a single question together • Leave sufficient space between answers*
- *Draw table/diagrams/flow charts wherever necessary*

### **Essays**

**(3x10=30)**

1. Discuss in detail, the pharmaceutical legislation in India along with historical developments.
2. Explain Narcotic Drugs and Psychotropic Substances Act and Rules and write in detail about prohibition, control and regulation of the operations
3. Discuss the general requirements of Good Manufacturing Practices (GMP) for premises and materials.

### **Short notes**

**(14x5=70)**

4. Describe the role of pharmacist in drug treatment and drug usage.
5. Briefly describe about the Drugs Technical Advisory Board (DTAB).
6. Differentiate between loan license and repacking license.
7. Explain the general labelling requirements for drugs and cosmetics.
8. Formula used to calculate retail price of a formulation.
9. Explain the constitution and the role of Pharmacy Council of India (PCI).
10. Elaborate on Patent Act 1970.
11. The Central Register of Pharmacist.
12. What are the qualification of a registered pharmacist to be entered in first register.
13. Describe the powers of drug inspector and procedure for inspection.
14. Write a note on the category of drugs that can be imported.
15. Explain about Poisons Act.
16. Describe and differentiate between bonded and non-bonded manufactory.
17. Discuss about prohibited and exempted advertisements.

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