

QP Code: 525006

Reg. No.....

**Fifth Semester B. Pharm Degree Supplementary Examinations
May 2023**

Pharmaceutical Jurisprudence

(2017 Scheme)

Time: 3 Hours

Max. Marks: 75

- *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

(2x10=20)

1. Explain the qualification, duties and responsibilities of drug inspector. Explain the procedure for taking of samples by drug inspector.
2. Describe the general procedure for obtaining a licence for the manufacture of drugs stating the conditions to be satisfied.

Short Notes

(7x5=35)

3. Restricted licence.
4. Give the constitution of drugs technical advisory board.
5. Explain the conditions for grant of manufacturing licence for Schedule C, C₁ and X drugs.
6. Explain the code of ethics of pharmacists in relation to his Trade.
7. What are the provisions under Medical termination of Pregnancy Act.
8. What are the requirements of a non-bonded laboratory.
9. Explain the recommendations of the Drug Enquiry Committee.

Answer Briefly

(10x2=20)

10. Drugs and Cosmetics Act, 1940.
11. Mention the conditions when names are removed from the first register.
12. Describe the qualification, powers and functions of Licensing Authorities under Drug and Cosmetics Act, 1940.
13. Define Schedule C and C₁ and Schedule X with an example.
14. Constitution of Pharmacy Council of India (PCI).
15. Information that may be refused under Right to Information Act.
16. Define schedule G, schedule H, schedule M and schedule N.
17. Institutional Animal Ethics committee.
18. Give the specimen label for ophthalmic preparation.
19. Define advertisement and magic remedies.
