

2010 Scheme

QP CODE: 305006

Reg. No:

Third Year B.Pharm Degree Supplementary Examinations August 2023 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. Elaborate on Current Good Manufacturing Practices
2. State the offences and penalties in relation to a) Cannabis plant and Cannabis b) Manufactured Drugs and Preparations prescribed under The Narcotic Drugs and Psychotropic Substances Act. In what way penalties under NDPS Act are different from other Acts.
3. Explain general and restricted license for sale of drugs. Write a note on drugs for personal use and why India is favourable manufacturing destination for many pharmaceutical companies.

Short notes

(14x5=70)

4. Discuss the salient features of Minimum Wages Act 1948.
5. Elaborate the powers of Drug Inspector as per Drugs and Cosmetics Act.
6. Discuss about Prohibited advertisements.
7. Describe the salient features of Patent Act 1970.
8. Discuss Schedule H in detail.
9. Enlist the various types of price calculations as specified under DPCO 2013. Describe the calculation of anyone type.
10. What are the classes of advertisements exempted under the Drugs and Magic Remedies (Objectionable Advertisements) Act.
11. Explain the recommendations put forwarded by Drug Enquiry Committee.
12. Describe the conditions for Termination of Pregnancy as per MTP Act.
13. Discuss the Poisons Act.
14. What are the qualifications and duties of a drug inspector.
15. Define drug, cosmetic, new drug, repacking licenses and Manufacture as per Drugs and Cosmetics Act.
16. Explain various phases of clinical trials.
17. What ethics a pharmacist should follow with respect to his profession. Explain.
