QP Code: 525006 **Reg. No……..**

Fifth Semester B. Pharm Degree Regular/Supplementary Examinations January 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 administrative bodies of Drugs and Cosmetics Act, 1940 and Rules 1945. Write a note on the qualification, powers and duties of drug inspectors.
2. Describe the formation and circumstances which lead to the formation of Drug Enquiry Committee (Chopra Committee) and explain its objectives and final recommendations to the government.

Short Notes **(7x5=35)**

3. Describe the Drug Price Control Order,2013.
4. What are the duties and procedures adopted by government analysts.
5. What are the provisions under Medical Termination of Pregnancy Act.
6. Give the class of drugs and cosmetics prohibited from import.
7. Explain the constitution and function of state pharmacy council.
8. Define patent and proprietary medicines. Explain the labelling and packaging requirements of these medicines for export.
9. Procedure for the maintenance, transfer, acquisition of animals for experiment under Prevention of Cruelty to Animal Act, 1960.

Answer Briefly **(10x2=20)**

10. Explain schedule X.
11. Name the controlling authorities under Drugs and Cosmetics Act,1940.
12. Name the office for the control and regulation of opium cultivation.
13. Explain the power to suspend or revoke registration under Prevention of Cruelty to Animals Act, 1960.
14. Explain the conditions required for obtaining an import licence for drugs under Drugs and Cosmetics Act, 1940.
15. Differentiate between State and joint state pharmacy council.
16. Give few lines on prohibition of certain advertisements.
17. Define restricted licence.
18. Give the national list of essential medicines.
19. Draw the specimen label for schedule H drug

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