**QP Code: 525006** **Reg. No……**

## Fifth Semester B. Pharm Degree Supplementary Examinations
## January 2021
## 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 conditions of import licence and classes of drugs that prohibited to be imported into India.
2. Explain in detail the manufacture of alcoholic preparation in bonded and non-bonded manufactory.

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

3. Explain about Drugs Technical Advisory Board (DTAB).
4. Loan licence
5. Explain the conditions for grant of licence for retail sale of drugs.
6. Explain the code of ethics of pharmacists in relation to his Job
7. List out offences and penalties under Narcotic Drugs and Psychotropic Substances Act, 1985.
8. Describe the procedure and conditions for obtaining a licence for the manufacture of drugs specified in schedule C, $C_1$ and X.
9. How are experimental animals required to be handled during and after experiments

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

10. Give the specimen label for Schedule X.
11. Conditions for removal names from the first register.
12. Define the following • Schedule T • Schedule M2 • Schedule J • Schedule H.
13. Explain the duties of Govt. Analyst.
14. Define Repacking Licence.
15. Exempted advertisements from the provisions of Drugs and Magic Remedies Act, 1955.
16. Define misbranded, adulterated and spurious drugs under Drug and Cosmetics Act and Rules, 1940.
17. Give the specimen label for ophthalmic preparation.
18. Storage conditions for Schedule X and Veterinary drugs.
19. Functions of Drugs Consultative Committee.

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