

Roll No

PY-802

B.Pharm. VIII Semester

Examination, December 2016

Pharmaceutics - XI
(Pharmaceutical Jurisprudence)

Time : Three Hours

Maximum Marks : 70

Note: i) Answer five questions. In each question part A, B, C is compulsory and D part has internal choice.

ii) All parts of each question are to be attempted at one place.

iii) All questions carry equal marks, out of which part A and B (Max.50 words) carry 2 marks, part C (Max.100 words) carry 3 marks, part D (Max.400 words) carry 7 marks.

iv) Except numericals, Derivation, Design and Drawing etc.

1. a) What is the scope of Forensic Pharmacy?
- b) What do you understand by "first register" and "Subsequent registers" under pharmacy Act.
- c) What are the functions of state pharmacy council?
- d) Discuss the constitution of pharmacy council of India. Discuss in detail education regulation.

OR

Give an account of Drug technical advisory board and drug inspectors.

2. a) Name three drugs each from Schedule C and C₁.
- b) Give the definition of Spurious drug.
- c) Define the terms Drug and standards of quality.
- d) List out the schedules and respective descriptions as per the D and C Act.

OR

What are the minimum requirement of drug manufacturing units as per schedule M under drug and cosmetic Act 1940 and rules there under?

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3. a) Mention the general labeling requirements of Ophthalmic.
- b) Give labeling requirement of preparations containing Schedule X drugs.
- c) What are the objectives of NDPS Act?
- d) Give the salient features of Medical Termination of Pregnancy Act.

OR

Define the term cocoa leaf, opium, hemp, manufactured drugs and Narcotic drugs and psychotropic substances. Discuss the conditions for manufacture and sale of opium as per Narcotic and Psychotropic substances Act 1985.

4. a) What is provisional specification?
- b) Define patent and invention.
- c) Discuss the objectives of prevention of cruelty to animals Act.
- d) Discuss and differentiate between "product patent" and "process patent"? Give the documents required to file Indian patent.

OR

How are the shops dealing in drugs and medicines exempted from the provision of shops and establishments Act?

5. a) Give the storage conditions for schedule H drugs.
- b) Give formula for calculation of retail price of formulations as under DPCO 1995.
- c) Define the term advertisement.
- d) Describe the provisions of the Factories Act relating to the approval, licensing and registration of Factories.

OR

Distinguish bonded manufactory from Non-Bonded manufactory. Write a detail note on the manufacture in Bonded Laboratory.

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