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Total No. of Printed Pages: [02]

Total No. of Questions: [13]

**B. Pharmacy (Semester: 5<sup>th</sup>)**  
**PHARMACEUTICAL JURISPRUDENCE (Theory)**  
**Subject Code: BP505T**  
**Paper ID: [17170127]**

**Time: 03 Hours**

**Maximum Marks: 75**

**Instruction for candidates:**

1. Section A is compulsory. It consists of 10 parts of two marks each.
2. Section B consist of 9 questions of 5 marks each. The student has to attempt any 7 questions out of it.
3. Section C consist of 3 questions of 10 marks each. The student has to attempt any 2 questions.

**Section – A**

**(2 marks each)**

Q1. Attempt the following:

- a) Define 'Import License' under the Drugs and Cosmetics Act, 1940.
- b) What is Schedule X? Provide two examples of drugs listed under this schedule.
- c) Explain the main objective of the Pharmacy Act, 1948.
- d) List two offenses under the Narcotic Drugs and Psychotropic Substances Act, 1985.
- e) What is the role of the Central Drugs Laboratory?
- f) Define 'In-bond' manufacture under the Medicinal and Toilet Preparations Act, 1955.
- g) What is meant by 'Ceiling Price' as per the Drugs Price Control Order (DPCO), 2013?
- h) Mention any two classes of exempted advertisements under the Drugs and Magic Remedies Act.
- i) State the purpose of the Code of Pharmaceutical Ethics.
- j) Explain the concept of Intellectual Property Rights (IPR) in brief.

**Section – B**

**(5 marks each)**

- Q2. Describe the legal definitions of different schedules in the Drugs and Cosmetics Act, 1940, and explain their significance.
- Q3. Explain the process of registration of pharmacists under the Pharmacy Act, 1948.
- Q4. Outline the conditions for the sale and labeling of Schedule H drugs.
- Q5. Discuss the composition and functions of the Drugs Technical Advisory Board (DTAB) under the Drugs and Cosmetics Act.
- Q6. Describe the penalties for violations under the Narcotic Drugs and Psychotropic Substances Act, 1985.
- Q7. Explain the major provisions of the Prevention of Cruelty to Animals Act, 1960, concerning pharmaceutical research.
- Q8. Discuss the major objectives of the National Pharmaceutical Pricing Authority and the DPCO, 2013.
- Q9. Explain the ethical responsibilities of a pharmacist in relation to their profession, as per the Code of Pharmaceutical Ethics.
- Q10. Outline the functions of the Institutional Animal Ethics Committee as per CPCSEA guidelines.

**Section – C**

**(10 marks each)**

- Q11. Elaborate on the Drugs and Cosmetics Act, 1940, particularly focusing on prohibited drugs and conditions for granting manufacturing licenses. Include a discussion on penalties for offenses.
- Q12. Provide a detailed account of the Medicinal and Toilet Preparations Act, 1955, including licensing requirements and conditions for export and manufacture of Ayurvedic preparations.
- Q13. Describe the role of various committees (Drugs Enquiry Committee, Health Survey Committee, Hathi Committee, and Mudaliar Committee) in shaping pharmaceutical legislation in India.