

**BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI
(END SEMESTER EXAMINATION)**

**CLASS: B.PHARM
BRANCH: PHARMACY**

**SEMESTER: V
SESSION: MO 2025**

SUBJECT: BP505T PHARMACEUTICAL JURISPRUDENCE

TIME: 3.00 Hours

FULL MARK: 75

INSTRUCTIONS:

1. The missing data, if any, may be assumed suitably.
2. Before attempting the question paper, be sure that you have got the correct question paper.
3. Tables/Data handbook/Graph paper etc. to be supplied to the candidates in the examination hall.
4. This question paper consists of (03) three parts. Read the part wise instructions before attempting the questions.

PART-I

Objective type questions (Instruction: Answer all questions)

Q1. (10 x 2 = 20 Marks)

- A. Define Proprietary medicine.
- B. What is Schedule C & C1 of Drugs and cosmetics Rules 1945?
- C. What is Schedule H of Drugs and cosmetics Rules 1945?
- D. Define spurious cosmetics.
- E. What is expanded form of CPCSEA, and under which act CPCSEA is provisioned?
- F. Enlist the role of Pharmacy council of India?
- G. What is the significance of Drugs and magic Remedies (Objectionable Advt.) Acts and Rules?
- H. Define NLEM.
- I. Enumerate the differences between Pharmacy and Drug store?
- J. What is Schedule P of Drugs and Cosmetics rules?

PART-II

Short Answers

(Instruction: Answer seven out of nine questions)

(7 x 5 = 35 Marks)

- Q2. Discuss in brief about price application and ceiling price, as per DPCO.
- Q3. Explain the role and use of alcohol in drug formulation industry with reference to Medicinal and Toilet preparation (Excise Duty) Acts and Rules 1955.
- Q4. Write in brief about Narcotics and Psychotropic substances Acts and Rules.
- Q5. State the qualifications required for appointment as a Drugs Inspector.
- Q6. Describe the composition of the Drug Technical Advisory Board (DTAB).
- Q7. Describe the composition of the Pharmacy Council of India (PCI).
- Q8. Illustrate the objectives of Pharmacy Act 1948 and Drug Price Control Order 2003.
- Q9. Enumerate the functions designated to Animals Welfare Board of India.
- Q10. Explain the Medical Termination of Pregnancy Act.

PART-III

Long Answers

(Instruction: Answer two out of three questions)

(2 x 10 = 20 marks)

- Q11. Discuss the Rule 96 manner of labelling for formulated products.
- Q12. Explain the requirements for obtaining drug licence for drug manufacturing unit.
- Q13. Discuss cGMP of Schedule M of Drugs and Cosmetics Rules 1945.

:::25/11/2025:::M