

**GUJARAT TECHNOLOGICAL UNIVERSITY**  
**B.PHARM – SEMESTER –4 EXAMINATION – SUMMER-2024**

**Subject Code: BP405TT****Date: 18/06/2024****Subject Name: Pharmaceutical Jurisprudence****Time: 10.30 a.m. to 1.30 p.m.****Total Marks: 80****Instructions:**

1. Attempt any five questions.
2. Make suitable assumptions wherever necessary.
3. Figures to the right indicate full marks.

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|------------|---|-----------|
| <b>Q.1</b> | (a) Define and explain: loan license and repacking license.   | <b>06</b> |
|            | (b) Write labelling requirements of Schedule G, Schedule H and Schedule X drugs.                                | <b>05</b> |
|            | (c) Write about offences and Penalties related to Narcotic & Psychotropic substances Act-1985 and Rules.        | <b>05</b> |
| <b>Q.2</b> | (a) Write constitution & functions of Drug Technical Advisory Board   | <b>06</b> |
|            | (b) Explain Qualification and duties of government analyst.   | <b>05</b> |
|            | (c) Write constitution and functions of PCI.  | <b>05</b> |
| <b>Q.3</b> | (a) Define Registered Pharmacist. Discuss the procedure for subsequent registration.                            | <b>06</b> |
|            | (b) Differentiate State & Joint state Pharmacy council.   | <b>05</b> |
|            | (c) Write a note on central drug laboratory.  | <b>05</b> |
| <b>Q.4</b> | (a) Define: Adulterate drug, Spurious drug, Hemp, Spirit, Poppy straw & Medicinal preparation                   | <b>06</b> |
|            | (b) Write difference between bonded and non-bonded laboratory.  | <b>05</b> |
|            | (c) Explain in detail the procedure for the import of drugs.  | <b>05</b> |
| <b>Q.5</b> | (a) Write a short note on Right to Information Act.   | <b>06</b> |
|            | (b) Give labeling conditions of ophthalmic preparations and narcotic preparations.                              | <b>05</b> |
|            | (c) Write a note on building facilities requirement as per schedule "M".  | <b>05</b> |
| <b>Q.6</b> | (a) Write a short note on definitions, objective and prohibited advertisements of drugs and magic remedies act. | <b>06</b> |
|            | (b) What are the provisions of Prevention of Cruelty to Animals act?  | <b>05</b> |
|            | (c) How the retail price of formulation is calculated?  | <b>05</b> |
| <b>Q.7</b> | (a) What are the qualifications for appointment as drug inspector? Discuss their power and duties.              | <b>06</b> |
|            | (b) Write a note on code of Pharmaceutical Ethics   | <b>05</b> |
|            | (c) Explain Medical termination of Pregnancy Act.   | <b>05</b> |

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