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

CLASS: B.PHARM **SEMESTER: V**
BRANCH: PHARMACY **SESSION: MO/2019**

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 hand book/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 types questions (Instruction: Answer all questions)

Q1. (10 x 2 = 20 Marks)

A. What is Section 42 of Pharmacy Act?

B. What do you mean by PCI?

C. What do you mean by DPCO?

D. What is CDSCO?

E. What is Trade mark registration?

F. What is Sch. C & C1 Drugs?

G. What is Ceiling Price?

H. What is objective of Pharmacy Act?

I. What is objective of Narcotics and Psychotropic Substances Acts & Rules 1985?

J. What is objective of Drugs and Magic (Objec. Advt) Acts and Rules 1955?

PART-II
Short Answers
(Instruction: Answer seven out of nine questions)

(7 x 5 = 35 Marks)

Q2. Write the qualifications of Drugs inspector.
Q3. What do you mean by Bonded Laboratory?
Q4. Define Medical Devices.
Q5. Define New Drugs.
Q6. What do you mean by Banning of Drugs?
Q7. Write the composition of DTAB.
Q8. How would you file Patent Application?
Q9. How would you became Patent agent?
Q10. Elaborate the following schedule of Drugs & Cosmetics Rules i. A ii. G , iii K iv N v R.

PART-III
Long Answers
(Instruction: Answer two out of three questions)

(2 x 10 = 20 marks)

Q11. Discuss Schedule M applicable to drug Formulation Industry.
Q12. How would you get Drug Licence for setting **drug Formulation Industry?**
Q13. As a Technical manager how you would label the Cefitixime 200 mg capsule.

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