Seat No: _____ **Enrollment No: _____**

PARUL UNIVERSITY
FACULTY OF PHARMACY
B. Pharm Winter 2022 - 23 Examination

Semester: 5 **Date: 20/10/2022**
Subject Code: BP505T **Time: 10:00am to 1:00pm**
Subject Name: Pharmaceutical Jurisprudence – Theory **Total Marks: 75**

Instructions:
1. Figures to the right indicate maximum marks.
2. Make suitable assumptions wherever necessary.

Q.1 Multiple Choice Questions (MCQs) (1 Mark Each) **(20)**

1. The members of DTAB hold the office for a period of:
   a) 3 Years b) 5 Years
   c) 4 Years d) 1 Year
2. Good manufacturing practices for Ayurvedic, Siddha and Unani medicines comes under
   a) Schedule T b) Schedule S
   c) Schedule U d) Schedule M
3. Application for grant/renewal of license to manufacture blood products for sale and distribution is applied in
   a) Form 38 b) Form 27E
   c) Form 28 d) Form 32A
4. The appointment of drug inspector is under act of
   a) 51 of IPC b) 22 of IPC
   c) 41 of IPC d) 21 of IPC
5. The _____ are articles meant to be rubbed, poured, sprinkled, sprayed on any part of the human body for cleansing, beautifying, promoting attractiveness or altering appearance.
   a) Drug b) Cosmetic
   c) Adulterated Drug d) Misbranded Drug
6. Drug sample taken by drug inspector for analysis are sent to
   a) Drug Controller b) Government analyst
   c) Chemical analyst d) Testing lab
7. List of coal tar colour permitted to be used in cosmetics come under the schedule
   a) Schedule M b) Schedule V
   c) Schedule P d) Schedule Q
8. Central Register for pharmacist is maintained by
   a) Central Government b) State Government
   c) AICTE d) PCI
9. PCI was constituted in
   a) 1949 b) 1950
   c) 1951 d) 1955
10. Which act provide to levy and collect excise duties on medical and toilet preparations containing alcohol, opium, hemp or other narcotic drugs?
    a) Poisons Act b) Pharmacy Act
    c) Medical and Toilet Preparations Act d) Drugs and Cosmetics Act
11. Government opium factory is situated in
    a) Ghazipur b) Lucknow
    c) Kasauli d) Izatnagar
12. As per narcotic drugs, controlled substance means
    a) Drug notified in central government gazette b) Drug banned by state
    c) Drug included in controlled price list d) Drug dispensed by RMP
13. Which act provides regulations for objectionable advertisement?
    a) Poisons Act b) Drugs and Magic Remedies Act
    c) Medical and Toilet Preparations Act d) Pharmacy Act

14. Prevention of Cruelty to Animals Act was passed in
    a) 1960 b) 1945
    c) 1985 d) 1956
15. Government of India appointed a Drug Enquiry Committee under the chairmanship of
    a) Col. Ramnath N. Chopra b) Jaysukhlal Hathi
    c) Dr. A. L. Mudaliar d) M.L. Schroff
16. The pharmacist should keep in mind the following point while dealing with his medical profession
    a) Limitation of professional activity b) Liaison with public
    c) Clandestine arrangement d) All of the above
17. A 20 weeks pregnancy can be terminated with the consent of
    a) Doctor holding MBBS degree b) Gynecologist
    c) Doctor holding MD degree d) Two registered medical practitioners
18. APIO forward applications for information to
    a) PIO b) First Appellate Authority
    c) The head of the Authority d) Third party
19. Patent is granted for ____ from the filing date of the application.
    a) 20 Years b) 5 Years
    c) 10 Years d) 1 Year
20. The distinctive signs that identify certain goods produced or provided by a specific person or enterprise are known as
    a) Geographical Indications b) Patent
    c) Trade mark d) Copyright

Q.2 Long Answers (any 2 out of 3) (10 Mark Each) **(20)**

1. Discuss in brief about the Drug Technical Advisory Board and Drug consultative committee of Drugs and Cosmetics act.
2. Write a note on Manufacturing in Bonded laboratories as per Medicinal and Toilet Preparation act.
3. Explain briefly about cultivation, production and sale of opium under Narcotic and Psychotropic substances act.

Q.3 Short Answers (any 7 out of 9) (5 Mark Each) **(35)**

1. What are the cautionary labels needed for the following drugs according to D & C act?
   a. Schedule G b. Schedule H c. Veterinary drugs
2. Explain the Qualifications and duties of government analyst.
3. What are the duties and power of a drug inspector?
4. What are the codes of ethics of a pharmacist in relation to his job?
5. Write a note on RTI Act 2005.
6. Write a note on functions of PCI.
7. Give an account on experimentation of animals under Prevention of Cruelty to Animals Act.
8. Define patents. Write the provisions for getting the patents rights quoted in Patents Act.
9. Explain about the Retail price of formulations in detail.