

End Semester/ Reappear (Semester V) Examination Dec 2022

Programme: B. Pharm
Course: Pharmaceutical Jurisprudence
Course Code: BP505T
Enrollment No: _____

Full Marks: 75
Time: 3 Hrs.

Section I

1. Objective type questions. Answer all questions.

20 x 1 = 20

- i. The Pharmacy Council of India is constituted
 - (a) Every 3 years
 - (b) Every 2 years
 - (c) Every 5 years
 - (d) Every 6 years
- ii. How many ex officio members are there in DTAB
 - (a) 6
 - (b) 5
 - (c) 8
 - (d) 4
- iii. Biological and Biotechnological products are listed under
 - (a) Schedule C & C₁
 - (b) Schedule A
 - (c) Schedule X
 - (d) Schedule B
- iv. Schedule J is related to
 - (a) GMP
 - (b) Curable disease
 - (c) Pack size of drug
 - (d) List of disease and ailments which drug cannot claim
- v. One of the following is a magic remedy
 - (a) Talisman
 - (b) Kavacha
 - (c) Mantra
 - (d) All of the above
- vi. Analysis and test of vaccine samples is carried out at
 - (a) Central Research Institute, Kaushali
 - (b) Pasteur Institute of India, Coonoor
 - (c) Central Drug Testing Laboratories, Thane
 - (d) Central Indian Pharmacopeia Laboratory, Ghaziabad
- vii. Cosmetic means any article intended to
 - (a) Affect the structure of human body
 - (b) Destruct the vermin of insect
 - (c) Alter the appearance of human body
 - (d) All of the above
- viii. The requirement of factory for manufacture of medical device is dealt under
 - (a) Schedule M1
 - (b) Schedule M2
 - (c) Schedule M3
 - (d) Schedule N
- ix. Any cosmetic preparation containing Hexachlorophene or Mercury are from import
 - (a) Exempted
 - (b) Prohibited
 - (c) Allowed
 - (d) Provided special provision
- x. The President of PCI is elected by
 - (a) Pharmacy Council Members
 - (b) Nominated members of PCI
 - (c) Elected members of PCI
 - (d) Ex-Officio members of PCI
- xi. A drug that is imported under a name which belong to another drug
 - (a) Spurious
 - (b) Misbranded
 - (c) Adulterated
 - (d) None
- xii Opium derivatives includes all the following except
 - (a) Prepared Opium
 - (b) Charas
 - (c) Heroin
 - (d) Medicinal Opium
- xiii The Prevention of Cruelty to Animal Act is enacted in
 - (a) 1940
 - (b) 1950
 - (c) 1960
 - (d) 1970
- xiv. Publication of advertisement that prohibited that refer to the use of any drug for the treatment of disease comes under
 - (a) Schedule P
 - (b) Schedule M
 - (c) Schedule J
 - (d) Schedule X

- xv. License issued for bonded and non-bonded laboratory is by
 (a) Excise Commission (b) Central Government
 (c) State Government (d) Director of Health Services
- xvi. NDPS committee consist of maximum -----number
 (a) 20 (b) 25 (c) 15 (d) 30
- xvii. A person is called Registered Pharmacist
 (a) Holding diploma in pharmacyI (b) Having sufficient experience in pharmacy profession
 (c) Holding pharmacy degree (d) Having his name entered in the state register pharmacist
- xviii. The Pharmacy Act came to force in
 (a) 1958 (b)1968 (c) 1948 (d)1938
- xix. The constitution of PCI consist of following members except
 (a) Elected members (b) Nominated members
 (c) Selected members (d) Ex-Officio members.
- xx. License issued for bonded and non-bonded laboratory is by
 (a) Excise Commission (b) Central Government
 (c) State Government (d) Director of health service

Section II

2. Short Answer type questions. Answer any five.

5 x 7 = 35

- a. Write the pharmacist oath.
- b. Discuss the advertisements that can be made without any prohibition.
- c. What is repacking license? Discuss condition to get the repacking license.
- d. Write the experience or training required for RMP according to Medical Termination of Pregnancy Act
- e. What are the duties of Drug Inspector?
- f. Write a note on interstate transport of alcoholic goods.
- g. As per The Prevention of Cruelty to Animals Act what all includes in “treating animals cruelly”.

Section III

Long Answer type questions. Answer any two.

2 x 10 = 20

3. Discuss the labelling and packaging of drugs.
4. What are the particulars of register for the entry of registered pharmacist? Explain the First Register including qualification to entry on First Register.
5. Discuss the procedure for sale prices of bulk drugs and the retail price of formulation.
