Annual (Year II) Examination June 2022

Programme: D. Pharm
Subject: Pharmaceutical Jurisprudence
Time: 3Hrs.

Subject Code: 2BD204

Enrollment No:

Section I

1. Objective type questions. Answer all questions.

 $1 \times 20 = 20$

- i. The First Schedule to the Drugs and Cosmetics Act, 1940 prescribes
 - a. Authoritative books of Ayurvedic, Siddha and Unani Tibb system
 - b. Standards for cosmetics
 - c. Standards for medical devices
 - d. Standards of the drugs to be complied with by imported drugs
- ii. Which of the following is a 'drug' as per the law?
 - a. Energy drinks and candy bars
- c. Mouth freshening spearmint chewing gum.
- b. Empty gelatin capsules, bandages and insect repellent creams
- d. Fish and Chicken
- iii. Who among the following is the Chairman of Drug Technical Advisory Board?
 - a. The Drugs Controller of India
- b. The President of Pharmacy Council of India
- c. The Director General of Health Services
- d. The President of Medical Council of India
- iv. The Narcotic Drugs and Psychotropic Substances Act was passed in the year
 - a. 1940
- b. 1955
- c. 2000
- d. 1985

- v. Prepared opium
 - a. is an extract suitable for smoking
- b. is any medicine containing opium
- c. is also called as hemp
- d. has undergone the processes to adapt it for medicinal use
- vi. Under the Medicinal and Toilet Preparations (Excise Duties) Act, manufacture of any dutiable goods without a valid license is punishable with
 - a. imprisonment up to six years or fine of twenty thousand rupees or both
 - b. imprisonment up to six months or fine of two thousand rupees or both
 - c. imprisonment up to six months or fine of twenty thousand rupees or both
 - d. imprisonment up to six years or fine of two thousand rupees or both
- vii. What is the definition of a "Magic Remedy"?
 - a. Any substance, whether processed, partially processed or unprocessed, which is intended for human consumption.
 - b. Any substance natural or synthetic or any salt or preparation of such substance or material, included in the list of psychotropic substances specified in the
 - c. talisman, mantra, kavacha or any other charm or any substance alleged to possess miraculous powers to diagnose, cure, mitigate, treat or prevent a disease in humans or animals
 - d. Ethyl alcohol of any strength and purity having the chemical composition C_2H_5OH

viii. Schedule K pertains to

- a. List of diseases that a drug cannot claim to cure
- b. List of drugs exempted from certain provisions governing import of drugs
- c. List of drugs exempted from certain provisions governing manufacture and sale of drugs
- d. List of drugs that can be used with caution under medical supervision
- ix. Which Act applies to evolving educational standard and regulations for the course in pharmacy through education regulations?
 - a. Pharmacy Act 1948

- b. Education Act 2009
- c. Drugs and Cosmetics Act 1940
- d. Indian Patent Act 1970

| х. | Qualification in pharmacy granted outside India | | |
|---|--|---|--|
| | a. UGC B. PCI C. DT. | | |
| xi. | The order of executive committee directing remo | oval of a name from register should be | |
| | confirmed by | | |
| | a. State Government | b. Central Pharmacy Council | |
| | c. State Pharmacy Council | d. Central Government | |
| xiiare provided for the analysis of drugs and cosmetics. | | | |
| a. State Drug Laboratory b. Central Drug Laboratory c. Control Laboratory d. Both A and B | | | |
| xiii. | In relation to ayurvedic, siddha or unani drug Go | | |
| | a. Section 33-F b. Section 33-G | | |
| xiv. As per Drugs and Cosmetic Act 1940, 'standard drug' is one | | | |
| a. That Compiles with standards of official pharmacopeia | | | |
| b. That Complies with the standard of Drug and Cosmetic Act 1940 | | | |
| | c. That compiles with the standard of national and International pharmacopeias | | |
| | d. That Compiles with International standards | | |
| XV. | Analysis and test of samples of vaccines are carrie | | |
| | a. Central Indian Pharmacopeia Laboratory, Gha | | |
| | b. Central Research Institute, Kasauli | d. Central Drug Testing Laboratories, Thane | |
| xvi. | As per Schedule M, the permitted limit of solid c | <u> </u> | |
| | a. 10 ppm b. 100 ppm c. 0.1 | | |
| xvii. A person is called as "Registered Pharmacist", if he is | | | |
| a. Having his name entered in the state register of pharmacists | | | |
| | b. Having sufficient experience in pharmacy profession | | |
| c. Holding diploma in pharmacy d. Holding degree in pharmacy | | | |
| xviii. Cosmetics means any article intended to | | | |
| | a. Affect the structure of the human body | b. Alter the appearance of the human body | |
| c. Destruct vermin of insects d.All of the above | | | |
| xix. | Medical Termination of Pregnancy Act was passed | | |
| | a. 1871 b.1980 c. 1971 | d.1949 | |
| XX. | Drugs consisting of filthy and putrid substances | | |
| | a. Proprietary b. Misbranded | c. Spurious d. Adulterated | |
| | | | |
| Section II | | | |
| 2. Short Answer type questions. Answer any four. $4 \times 5 = 20$ | | | |
| a. Discuss any three offenses and penalties under Pharmacy Act, 1948 for falsely claiming to | | | |
| be Registered Pharmacist: | | | |
| b. Define "Adulterated Drug" as per the Drugs and Cosmetics Act 1940. | | | |
| c. How retail price of a formulation is calculated as per the Drugs (Price Control) Order? | | | |
| d. Write the powers of a drug inspector. | | | |
| e. Define 'cocoa derivative' as per the NDPS Act 1985. | | | |
| f. Discuss the composition of DTAB. | | | |
| Section III | | | |
| Long Answer type questions. Answer any four. $4 \times 10 = 40$ | | | |
| 3. What is manufactured in Bond? Give two points to differentiate it from manufacture outside Bond as per | | | |
| Medicinal & Toilet prep. (ED) Act. | | | |
| 4 Define following terms according to D & C Act 1940: a Misbranded drugs b Spurious drug | | | |

- 5. How drugs are imported for personal use as per D & C Act 1940.
- 6. Discuss the classes of advertisement exempted under the Drugs and Magic Remedies Act 1954.
- 7. Give the composition of Pharmacy Council of India.
- 8. Give the procedure to be followed by a drugs Inspector while collecting samples from a manufacturing premises as per Drugs & cosmetic rules 1945.
