2. What is Drug Enquiry committee ? Enumerate the aspects and recommendation of this committee. 2 +4+4=10

3. Elaborate the constitution and functions of PCI. 5+5=10

III. Short answers (Answer seven out of nine questions) 5x7=35

1. What is restricted licence? Describe the conditions of restricted licence to be fulfilled by the licensee. 5
2. What are the classes of drugs which are prohibited for its import in India? Mention the places through which drugs can be imported to India. 3+2=5
3. Write in details about the constitution and functions of Drugs Technical Advisory Board. 5
4. Who was the chairman of "Bhore committee"? Enumerate the recommendation of the committee. When it was set-up? 1+3+1=5
5. Write down the applicability of RTI act. 5
6. Explain the functions of Drug Price Control Order (DPCO). 5
7. Discuss in detail about manufacture in bonded laboratory. 5
8. Give a detailed note on salient features of Drugs and Magic Remedies Act 5
9. Write a short note on "Bhatia committee" 5

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UG Odd Semester (CBCS) Examination, 2022
held in March 2023

PHARMACEUTICAL SCIENCES

(5th Semester)

Course No: BP 505T

(Pharmaceutical Jurisprudence-Theory)

Full Marks: 75

Time: 3 Hours

The figures in the margin indicate full marks for the questions

I (A). Multiple choice questions **1x10=10**

1. Which of the following is NOT the ex-offcio member of PCI.
   a) Director General of health service
   b) Drug controller of India
   c) Chief Administrative Medical Officer
   d) Director of Central Drug Laboratory
2. Under the Medicinal and Toilet Preparations (Excise Duties) Act, manufacture of any dutiable goods without a valid licence 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

3. The Drugs and Magic Remedies (Objectionable Advertisements), Rules were introduced in which year?
   a) 1935 b) 1954
   c) 1955 d) 1965

4. 212 recommendations were implemented by
   a) BHORE COMMITEE
   b) BHATIA COMMITTEE
   c) MUDALIAR COMMITEE
   d) HATHI COMMITTEE

5. Drug shall be deemed to be adulterated except if it-
   a) Is manufactured under insanitary conditions.
   b) Contains different colors other than those prescribed.
   c) Is imported under a name that belongs to another drug.
   d) Contains filthy, putrid or decomposed substances.

6. Implementation of the NDPS Act 1985 was carried out by
   a) NCB b) PCB
   c) PCI d) MCI

7. Chopra committee report published in
   a) 1931 b) 1930
   c) 1935 d) 1934

8. In pharmacy, a separate cupboard should be provided for the storage of-
   a) Cosmetics
   b) Patent and proprietary medicine
   c) Ayurvedic, Siddha and Unani drugs
   d) Poisons

9. Government Analyst is appointed by Central Government or State Government under section_____ in relation to Ayurvedic, Siddha and Unani systems of medicine. Fill in the blank.
   a) 20 b) 21
   c) 33F d) 21A

10. Who among the following is not an Ex-Officio member of Drug Technical Advisory Board?
   a) The President, Pharmacy Council of India.
   b) Director, Indian Institute of Chemical Technology, Hyderabad
   c) The President, Medical Council of India.
   d) Director, Central Drugs Research Institute, Lucknow

I (B). Objective type (Answer the following in brief)

2x5=10

1. Define drug according to Drug and Cosmetic act 1940.
2. What qualifications will be entitled to a person to be appointed as a Government Analyst?
3. What is Trademark?
4. Define RTI.
5. What is denatured sprit?

II. Long answers (Answer two out of three questions)

10x2=20

1. What qualifications are required for a person to be appointed as Drug Inspector? What procedure should be followed by Drug inspectors while sending the samples for test or analysis. 3+7=10