REV-01
BPH/113/38/43

2023/12

SET B

B. PHARM.
FIFTH SEMESTER
PHARMACEUTICAL JURISPRUDENCE
BP505T

[USE OMR FOR OBJECTIVE PART]

Duration : 3 hrs. **Full Marks : 75**

PART-A: Objective

Time : 30 min. **Marks : 20**

Choose the correct answer from the following: ***1×20=20***

1. Which of the following is/are non-patentable?
   a. A new medical device for diagnosis
   b. Admixture of two known substances to give a new effect
   c. Algorithms
   d. A genetically modified microorganism

2. Plant of the species of Papaver, from which opium or any phenanthrene alkaloid can be extracted, is called..
   a. opium poppy
   b. coca
   c. heroin
   d. Cannabis sativum

3. Who has the power to fix the ceiling price of scheduled formulations?
   a. State Government
   b. Central Government
   c. Lok Sabha
   d. Rajya Sabha

4. NDPS Act provide licensing system to regulate.....of narcotic and psychotropic substances
   a. Manufacturing
   b. Cultivation
   c. Sale and export
   d. All of the above

5. Manufacture of Opium can be made only by the Central Government at its two factories situated at..
   a. Ghazipur and Neemuch
   b. Kolkata and Delhi
   c. Mumbai and Lucknow
   d. None

6. The non bonded laboratory shall be inspected by the proper Excise officer..
   a. Once in two months
   b. Once in a month
   c. Once in a year
   d. Once in six months

7. The first Pharmacy Council of India was constituted in the year..
   a. 1945
   b. 1955
   c. 1948
   d. 1959

8. The Pharmacy Act extends to the whole of India except..
   a. Karnataka
   b. Jammu and Kashmir
   c. Maharashtra
   d. None of the above

9. The schedule in D&C Act that deals with the standard for disinfectant fluid is
   a. Schedule B
   b. Schedule F
   c. Schedule O
   d. Schedule M

10. Requirements of factory premises for manufacture of medical devices is dealt under...
    a. Schedule M1
    b. Schedule M2
    c. Schedule M3
    d. Schedule N

11. Blood Bank comes under the schedule
    a. Schedule B
    b. Schedule D
    c. Schedule F
    d. Schedule G

12. If the drug contains in filthy, putrid or decomposed substance then is known as
    a. Spurious Drug
    b. Adulterated drug
    c. Misbranded drug
    d. None of the above

13. "Schedule F3" is related with
    a. Standard for surgical dressing
    b. Standard for sterilized umbilical tapes
    c. Standard for ophthalmic preparation
    d. Standard for production of sera

14. 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 President of Medical Council of India
    d. The Director General of Health Services

15. Which amongst the following is the duty of a Drug Inspector ?
    a. Seizure of stocks
    b. Take samples of any drug or cosmetics
    c. Inspection
    d. All of the above

16. Central drug laboratory is located at..
    a. Mumbai
    b. Calcutta
    c. Delhi
    d. Lucknow

17. Schedule P is related to..
    a. Packing of drugs
    b. Toilet preparation
    c. Ophthalmic preparation
    d. Biologicals

18. The premises for manufacturing of drug must fulfil the condition according..
    a. Schedule N.
    b. Schedule M
    c. Schedule O
    d. Schedule X

19. The Narcotic Drugs and Psychotropic Substances Act was passed in the year..
    a. 1940
    b. 1955.
    c. 1985
    d. 2000

20. What is the definition of a "Magic Remedy?

a. Talisman, mantra, kavacha or any other charm or any substance alleged to possess miraculous powers powers to diagnose, cure, mitigate, treat or prevent a disease in humans or animals

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. Any substance, whether processed, partially processed or unprocessed, which is intended for human consumption.

d. Ethyl alcohol of any strength and purity having the chemical composition C2H5OH

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PART-B: Descriptive

Time : 2 hrs. 30 min. Marks : 35

[Answer any seven (7) questions]

1. Write a short note on Prohibition of import of certain Drugs or cosmetics. 5
2. Define the term Misbranded, Adulterated and Spurious drugs according to the drugs and cosmetics act. 5
3. Write down the offences and penalties relating to import of Drugs. 5
4. Write down the duties of Drug Inspector. 5
5. Write a short note on Drug Technical Advisory Board (DTAB) 5
6. Write down the documents required for obtaining Retail Drug license. 5
7. Write about the fixation of ceiling and retail prices of scheduled formulation under DPCO act, 1995. 5
8. Discuss the provision of the Medical termination of pregnancy for the termination of pregnancy 5
9. Discuss the advertisements which are prohibited under objectionable advertisement. 5

PART-C : Long type questions

[Answer any two (2) questions]

1. Elaborately discuss loan license and Repacking license 5+5=10
2. Elaborately discuss the Good Manufacturing Practises (GMP) required for factory premises. 10
3. Write the constitution of IAEC and describe the CPSCEA guidelines as per the prevention of cruelty to Animals act, 1960 10

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