REV-01 BPH/05/10

## B. PHARM. FIFTH SEMESTER PHARMACEUTICAL JURISPRUDENCE BP505T [REPEAT] JUSE OMR FOR OBJECTIVE PART]

SET

2023/12

JUSE OMR FOR OBJECTIVE PARTI

Duration: 3 hrs.

PART-A: Objective

Time: 30 min.

Choose the correct answer from the following:

Full Marks: 75

Marks: 20 1×20=20

- 1. 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
- Requirements of factory premises for manufacture of medical devices is dealt under...
  - a. Schedule M1

- b. Schedule M2
- c. Schedule M3 d. Schedule N
- 3. Blood Bank comes under the schedule
  - a. Schedule B

b. Schedule D

c. Schedule F

- d. Schedule G
- 4. If the drug contains in filthy, putrid or decomposed substance then is known as
  - a. Spurious Drug

- b. Adulterated drugd. None of the above
- c. Misbranded drug
- "Schedule F3" is related with
  - a. Standard for surgical dressing
- b. Standard for sterilized umbilical
- c. Standard for ophthalmic preparation
- d. Standard for production of sera
- 6. Who among the following is the Chairman of Drug Technical Advisory Board?
  - a. The Drugs Controller of India
- The President of Pharmacy Council of India
- c. The President of Medical Council of India
- The Director General of Health Services
- 7. Which amongst the following is the duty of a Drug Inspector?
  - a. Seizure of stocks

Take samples of any drug or cosmetics

c. Inspection

- d. All of the above
- 8. Central drug laboratory is located at ...
  - a. Mumbai

b. Calcutta

c. Delhi

- d. Lucknow
- 9. Schedule P is related to..
  - a. Packing of drugs

- b. Toilet preparation
- c. Ophthalmic preparation
- d. Biologicals

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USTM/COE/R-01

	a. Once in two months c. Once in a year	b.	Once in a month Once in six months
17.	Manufacture of Opium can be made only by factories situated at  a. Ghazipur and Neemuch c. Mumbai and Lucknow The non bonded laboratory shall be inspect	b. d.	Kolkata and Delhi None
16.	NDPS Act provide licensing system to regul substances a. Manufacturing c. Sale and export	b.	Cultivation  All of the above
15.	Who has the power to fix the ceiling price of a. State Government c. Lok Sabha	b. d.	Central Government Rajya Sabha
14.	Plant of the species of Papaver, from which be extracted, is called a. opium poppy c. heroin	b.	ium or any phenanthrene alkaloid can coca Cannabis sativum
	c. Algorithms		to give a new effect A genetically modified microorganism
13.	Which of the following is/are non-patentable a. A new medical device for diagnosis		Admixture of two known substances
	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
12.	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 of any salt or preparation of such substance or material, included in the list of psychotropic substances specified in the
11.	The Narcotic Drugs and Psychotropic Substa a. 1940 c. 1985	b.	es Act was passed in the year 1955. 2000
10.	The premises for manufacturing of drug mu a. Schedule N. c. Schedule O	b.	Schedule X

19.	The first Pharmacy Council of India was constituted in the year					
	a. 1945	b. 1955				
	c. 1948	d. 1959				
20.	The Pharmacy Act extends to the whole of India except					
	a. Karnataka	b. Jammu and Kashmi				
	c. Maharashtra	d. None of the above				

## PART-B: Descriptive

Time: 2 hrs. 30 min. Marks: 35 [Answer any seven (7) questions] 1. 5 Write a short note on Prohibition of import of certain Drugs or cosmetics. Define the term Misbranded, Adulterated and Spurious drugs 5 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. Write a short note on Drug Technical Advisory Board (DTAB) 5 5. Write down the documents required for obtaining Retail Drug 5 license. 7. Write about the fixation of ceiling and retail prices of scheduled 5 formulation under DPCO act, 1995. Discuss the provision of the Medical termination of pregnancy for 5 the termination of pregnancy 9. prohibited 5 Discuss the advertisements which are objectionable advertisement.

## (PART-C: Long type questions)

## [ Answer any two (2) questions ]

1.	Elaborately discuss loan license and Repacking license				
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			