B Pharm Semester-V
END SEM Examination (Make-Up) January 2023
PMA-BP-505 Pharmaceutical Jurisprudence

Date: 27/1/23 **Duration:** - 3hrs. **Max. Marks:** 75

Instructions: Answer ALL questions.

Question	Ans
1. The regulatory authority of Japan is …….. a. Ministry of Health, Labour and Welfare b. Medicines and Healthcare procedure Regulatory Authority c.Japan Medicines Evaluation Agency d.Ministry and Healthcare products of Japan	
2. List of Psychotropic substances whose import, manufacture and sale, labelling and packaging are governed by special provisions as per schedule…… a. Schedule H b. Schedule X c. Schedule C d. Schedule M	
3. The patent is granted for period of…….. a. 20 years b. 10 years c. 01 years d. 05 years	
4. As per Schedule M (MFR) stands for…….. a. Master Formula Receipt b. Manufacture Formula Records c. Master Finish Records d. Master Formula Records	
5. Rates of fees charged for analysis by CDL or State Drugs Laboratories are mentioned in schedule…… a. Schedule A b. Schedule C c. Schedule D d. Schedule B	
6. In respect of drugs imported into India by rail from Pakistan is through……. a. Amritsar Cantonment b. Jammu Cantonment c. Firozpur Cantonment d. Chandigarh Cantonment	
7. Schedule P states about…… a. Life periods of drugs b. Pack sizes of the drugs c. List of Dyes, Colors and Pigments d. Standards for cosmetics in finished form	
8. The manufacturing records, records for raw material and analysis and other operational records should be maintained as per schedule……. a. Schedule B b. Schedule U c. Schedule M d. Schedule R	
9. The Drugs and Cosmetics Act was passed in …….. a. 1945 b. 1955 c. 1956 d. 1954	
10. The Indian Drug Regulatory Authority is called as…….. a. CSIR b. ICMR c. AICTE d. CDSCO	

11. First Register is prepared by _____ a. State Government b. State Pharmacy Council c. Central Council d. State and Central Council	
12. Which of the following chairman of committee made a comprehensive enquiry into the working of pharmaceutical industry? a. SL Bhatia b. Ramnath Chopra c. Joseph Bhore d. Jaisukhlal Hathi	
13. Which of the following Act was established in 1954? a. Drugs and Magic Remedies Act b. Pharmacy Act c. Medicinal and Toilet Preparations Act d. Essential Commodities Act	
14. Which of the following is the current Drug Price Control Order? a. DPCO 1970 b. DPCO 1976 c. DPCO 1995 d. DPCO 2013	
15. According to Medicinal and Toilet Preparations Act dutiable goods may be kept in a warehouse for a maximum period of _____ a. 1 Years b. 2 Years c. 3 Years d. 5 Year	
16. What is IAEC? a. Institutional Animal Ethics Council b. Institutional Animal Ethics Committee c. Institutional Animal Education Council d. Institutional Animal Entrusted Committee	
17. Flowering or fruit tops of cannabis plant is called _____ a. Hemp b. Charas c. Hashish d. Ganja	
18. Punishment for contravention, where the contravention involves small quantity, in relation to Poppy Straw according to Narcotic Drugs and Psychotropic Substances is _____ a. Rigorous imprisonment up to 6 months or fine up to 10,000 or both b. Rigorous imprisonment up to 10 years or fine up to 1 lakh or both c. Rigorous imprisonment up to 10-20years and fine not less than to 1-2 lakh. d. Rigorous imprisonment up to 20years and fine not less than to 4 lakh.	
19. A registered pharmacist cannot do the following except _____ a. Dispense drugs on humanitarian grounds b. Render first acid in emergencies c. Prescribe medicines in emergencies d. Can practice medicine only in emergencies	
20. Diacetylemorphine is known as _____ a. Ganja b. Charas c. Hashish d. Heroin	

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I Long Answers	**2 Q × 10 marks = 20 marks**
Q1. A. Describe the functions of Pharmacy Council of India. Write a brief note on Education Regulations. B. State the offences and penalties in relation to Cannabis Plant and Cannabis prescribed under The Narcotic Drugs and Psychotropic Substances Act. (5+5=10 marks)	10
Q2. Explain various phases of clinical trials. Add a note on good manufacturing practices.	10

II Short Answers	**7 Q × 5 marks = 35 marks**	
Question	**MARKS**	
Q1. Describe the functions of National Pharmaceutical Pricing Authority.	5	
Q2. What types and classes of advertisements are prohibited under the Drugs and Magic Remedies Act? What do you mean by Non-Bonded manufactory? (3+2=5marks)	5	
Q3. Write short notes Hathi Committees. What ethics a pharmacist should follow with respect to his Profession. (2+3=5marks)	5	
Q4. What are patents and copyright?	5	
Q5. Discuss functions of CDSCO and centres of USFDA.	5	
Q6. Write powers, duties and qualifications of drug inspectors	5	
Q7. Write short note on MTP Act and RTI Act.	5	