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	
i. Schedule H	
o. Schedule X	
Schedule C	
I. Schedule M	
3. The patent is granted for period of	
a. 20 years	
o. 10 years	
e. 01 years	
l. 05 years	
As per Schedule M (MFR) stands for	
a. Master Formula Receipt	
Manufacture Formula Records	
. Master Finish Records	
I. Master Formula Records	
. Rates of fees charged for analysis by CDL or State Drugs Laboratories are mentioned in	-
chedule	
. Schedule A	
o. Schedule C	
s. Schedule D	
I. Schedule B	
5. In respect of drugs imported into India by rail from Pakistan is through	
. Amritsar Cantonment	
b. Jammu Cantonment	36-
e. Firozpur Cantonment	融
d. Chandigarh Cantonment	100
7. Schedule P states about	10
Life periods of drugs	
b. Pack sizes of the drugs	
E. List of Dyes, Colors and Pigments	
I. Standards for cosmetics in finished form	
B. The manufacturing records, records for raw material and analysis and other operational records	
hould be maintained as per schedule	
. Schedule B	
a. Schedule U	
. Schedule M	
. Schedule R	
	-
The Drugs and Cosmetics Act was passed in	
. 1945	
0. 1955	
2. 1956	
1. 1954	
0. The Indian Drug Regulatory Authority is called as	
. CSIR b. ICMR c. AICTE d. CDSCO	12

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. St. Bhatia b. Rannath Chopra c. Joseph Bhore d. Jatsukhlal 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 1970 b. DPCO 1970 b. DPCO 1975 c. JPCO 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 Council c. Institutional Animal Ethics Committee c. Institutional Animal Pintrusted Committee 17. Flowering or fruit tops of cannabis plant is called a. Hemp b. Charas c. Hashish d. Ganja ners of the proper of the 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 10 years or fine up to 1 lack or both c. Rigorous imprisonment up to 10 -20 years and fine not less than to 1-2 lakh. d. Rigorous imprisonment up to 10 years or fine up to 1 lack or both c. Rigorous imprisonment up to 10 years or fine up to 1 lack or both c. Rigorous imprisonment up to 20 years and fine not less than to 1-2 lakh. d. Rigorous imprisonment up to 10 years or fine up to 10 (0000 or both b. Rigorous imprisonment up to 10 years or fine up to 10 (0000 or both c. Rigorous imprisonment up to 20 years and fine not less than to 1-2 lakh. d. Rigorous imprisonment up to 10-20 years and fine not less than to 1-2 lakh. d. Rigorous imprisonment up to 10-20 years and fine bot less than to 1-2 lakh. d. Can practice medicine only in emergencies d. Can practice medicine only in emergenc	11. First Register is prepared by	
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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.

I Long Answers 2 Q × 1	0 marks = 20) marks
Q1. A. Describe the functions of Pharmacy Council of India. Write a brief no Education Regulations.	ote on	10
B. State the offences and penalties in relation to Cannabis Plant and Cannabi	s prescribed	İ
under The Narcotic Drugs and Psychotropic Substances Act. (5+5=10 marks		
Q2. Explain various phases of clinical trials. Add a note on good manufactur practices.	ing	10
II Short Answers 7 Q >	< 5 marks = 2	5 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	5 5	
and Magic Remedies Act? What do you mean by Non-Bonded manufactory? (3+2=5marks)		
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	