Total No. of printed pages = 4

2.4

23208/22

Roll No. of candidate

2022

D.Pharm. 2nd Year Part-II End-Term Examination

PHARMACEUTICAL JURISPRUDENCE

Full Marks – 80 Time – Three hours

The figures in the margin indicate full marks for the questions.

1. Answer all the questions: (20 × 1 = 20)

(i) 'Drug Enquiry Committee' is also known as ———— committee.

(a) Mudaliar Committee (b) Srivasatava Committee

(c) Chopra Committee (d) None of the above

(ii) The 1st edition of Indian Pharmacopoeia was published in the year

(a) 1940 (b) 1955

(c) 1965 (d) 1985

(iii) Excise Commissioner of a state is responsible for enforcement of law regarding;

(a) Cocoa Derivatives (b) Psychotropic substance

(c) Toilet preparation (d) Opium derivatives

(iv) Lisence for wholesale of drugs specified in Schedule C and C_1 are issued in the forms

(a) 20 A (b) 20 B

(c) 21 A (d) 21 B

(v) The Drug (Price Control) Order was passed in the year

(a) 1955 (b) 1948

(c) 1970 (d) 1995

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(vi) Denatured alcohol refers to an alcohol of any strength which has been ——

(a) Rendered not fit for storage
(b) Rendered not fit for sale
(c) Rendered fit for human consumption
(d) Rendered unfit for human consumption

(vii) If any substance has been mixed with the drug so as to reduce its quality or strength, then the drug is considered as ——————.

(a) Adulterated (b) Misbranded
(c) Spurious (d) All of the above

(viii) As per the rules, the manufacturing of opium can only take place at ——— and

(a) Jaipur and Ghazipur (b) Meerut and Ghazipur
(c) Ghazipur and Neemuch (d) Meerut and Neemuch

(ix) Who is NOT the ex-Officio member of DTAE?

(a) Drug controller of India (b) Government analyst
(c) President PCI (d) Director CDRI

(x) In medical termination of pregnancy act (1971), a registered medical practitioners can terminate a pregnancy only if:

(a) It is more than 06 weeks but not more than 10 weeks
(b) It is more than 12 weeks but not more than 20 weeks
(c) It is more than 18 weeks but not more than 30 weeks
(d) None of the above

(xi) The 'Poison Act' was passed on:

(a) 1919 (b) 1985
(c) 1954 (d) 1971

(xii) Which class of advertisement is prohibited?

(a) Procurement of miscarriage of conception in woman
(b) Menstrual disorder
(c) Improvement of sexual Pleasure
(d) All of the above

(xiii) Lisence for wholesale of drugs specified in Schedule C and C_1, are issued in the forms

(a) 20 A (b) 20 B
(c) 21 A (d) 21 B

(xiv) In case of oral liquid preparations, the drug specified in schedule X can be marketed in packing not exceeding:

(a) 100 ml (b) 250 ml
(c) 300 ml (d) 500 ml

(xv) As per Drug (Price Control) Order Act, 'Retail Price' means:

(a) Price fixed by the Government for scheduled formulations
(b) Price after bringing into India from a place outside India
(c) Price fixed in accordance with provisions of the act including ceiling price
(d) Profit before payment of income tax

(xvi) Which of the following is true about Ganja?

(a) It is composed of resin extracted from cannabis plant
(b) It is composed of flowering or fruiting tops of cannabis plant
(c) It is composed of leaf and seed parts of cannabis plant
(d) All of the above

(xvii) All opium products should be delivered to ————————.

(a) Narcotic commissioner
(b) District opium officer
(c) Central government
(d) State government

(xviii) MAPE should not exceed ———————— % in the case of category I formulation and ———————— % in the case of category II formulations.

(a) 70% and 100%
(b) 50% and 70%
(c) 60% and 80%
(d) 100% and 70%

(xix) Minimum age for registered pharmacist is ————————. (fill in the blank)

(xx) DTAB constituted in every ———————— years. (fill in the blank).

2. Short Answer (Answer 10 out of 11) (10 × 3 = 30)

(a) Define the following terms:

(i) Magic Remedy
(ii) Registered pharmacists
(iii) Advertisement

(b) Match the following (6 × 0.5 = 3)

Group A		Group B	
(i)	Schedule O	(p)	Guidelines for clinical trials
(ii)	Schedule H	(q)	Standard for disinfectant fluids
(iii)	Schedule F	(r)	Standards for cosmetics
(iv)	Schedule Y	(s)	Provision applicable to vaccines, toxins, antigens
(v)	Schedule S	(t)	Standard for ophthalmic preparations
(vi)	Schedule FF	(u)	List of Prescription drugs

(c) Write the composition of Pharmacy Council of India.

(d) Explain design and construction of bonded laboratory.

(e) Describe in detail the classes of prohibited advertisement under Drugs and Magic Remedies Act, 1954.

(f) What are the training and experience prescribed in the act for a RMP for termination of pregnancies?

(g) What are the qualifications needed for appointment as a Drug Inspector?

(h) Define ceiling price. Give the calculation of Retail Price of formulation as per Drug (Price Control) Order, 2013.

(i) How Joint State Pharmacy Council is Constituted?

(j) Explain in details what do you mean by Illicit Traffic' under NDPS Act and rules.

(k) Give the specific mandatory requirements of labelling for Schedule H drugs.

3. Long Answer Questions (Answer 6 out of 7): $(6 \times 5 = 30)$

(a) Describe in details the code of ethics a pharmacist needs to follow in relation to his job.

(b) What do you mean by eugenic aspect as per Medical Termination of Pregnancy Act, 1971'? Under what conditions can a pregnancy be terminated medically under the act? (2+3)

(c) Define narcotic drugs. Briefly explain the procedure of entry, search, seizure and arrest under the Narcotic Drugs and Psychotropic Substances Act. (1+4)

(d) Give the composition and function of Drug Technical Advisory Board (DTAB).

(e) Give a brief account of historical development of pharmaceutical legislations.

(f) Explain the provisions of manufacturing license of Schedule C and C_1 drugs.

(g) Write short notes on : (2.5 + 2.5)

(i) Poison Act

(ii) Drug Enquiry Committee.
