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Roll No. of candidate | | | | | | | | | | |

2019

B.Pharm. 6th Semester End-Term Examination

PHARMACEUTICAL JURISPRUDENCE AND ETHICS

Full Marks – 100 Time – Three hours

Answer question No. 1 and any *six* from the rest :

1. Multiple Choice questions. (10 × 1 = 10)

(i) Person authorized to seize advertisement related documents shall be deemed to be public servants within the meaning of

(a) Sec 4 of Drugs and magic remedies Act

(b) Set 21 of the IPC

(c) Sec 22 of Drugs and Cosmetic Act

(d) Sec 26 of Drug Price Control Order

(ii) Which of the following statement is not true for advertisement

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(a) Advertisements of any type can be exported or imported in accordance with the provisions of Drugs and Magic Remedies Act

(b) The prohibited categories of advertisement within the country are also prohibited for export or import

(c) The custom collector is the authorized person for any seize or detention of advertisement for export or import

(d) Export and import of advertisement is prohibited under section 19 of the Sea Custom Act 1878.

(iii) Provisions of the Poison Act do not apply to

(a) Person involved in import of poison with license granted by Central Government

(b) Authorities empowered to make rules under the Act.

(c) Medical or veterinary practitioners in exercise of their profession.

(d) Government Analyst

(iv) The elected members in the committee governing experimentation on animals are

(a) Two members each of the ICMR and CAR

(b) Two members representing universities granting degrees in medical and veterinary sciences.

(c) Five members actively engaged in the promotion of animal welfare

(d) One member from Lok Sabha and one from Rajya Sabha

(v) The appointment of a pharmacist for maintaining quality of drugs in hospitals was recommended by

(a) Drug Enquiry Committee

(b) Health survey and development committee

(c) Hathi committee

(d) Indian Pharmacopoeial committee

(vi) The termination of pregnancy in Eugenic aspects means

(a) When pregnancy poses grave risk to the life or health of a woman

(b) When pregnancy arises from sexual crime

(c) Pregnancy of lunatic woman

(d) If the child to be born is likely to have deformities

(vii) The current round of negotiation in WTO is known as the

(a) Torquay Round

(b) Doha Development Round

(c) Cancun declarations

(d) Seattle Summit

(viii) Dunkel Draft was the initiative taken in the establishment of

(a) WTO (b) GATT

(c) TRIM (d) TRIPS

(ix) In the Factories Act, Indian standard time is _____ hours ahead of Greenwich Mean Time.

(a) 4.20 (b) 5.0

(c) 5.5 (d) 6.15

(x) Which of the following statement is not true for Rights of patentee

(a) The legal rights in intellectual property secured in one country cannot be enforced in another country.

(b) The legal rights in intellectual property secured in one country also enforced in other countries.

(c) The patentee has the right to prevent third parties of using that process who do not have his consent

(d) The patentee has an exclusive right to make use of the patent within the period of patent, sell or distribute such an article.

2. (a) What are the provisions in the Factories Act to ensure health of the workers in a factory? (5)

(b) Give the important aspects of the poison act including the prescribed penalties. (5)

(c) Discuss the provisions for regulation and control of opium. (5)

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

(b) Give details of the requirements for approval of places for pregnancy termination.

(c) As per the Drugs and Cosmetic Act what are the qualifications for considering a person as qualified person.

(d) Explain the provisions of manufacturing of schedule C and C1 drugs under the Drugs and Cosmetic acts. (2+2+3+8)

4. (a) Write the objectives of the Prevention of Cruelty to Animals Act. What are the requirements of care to animals used for experimental purpose? (4+5+6)

(b) Explain the provisions of import of drugs.

(c) What is bonded and non-bonded manufacturing. Give the required facilities in bonded manufacturing unit.

5. (a) What are three categories of advertisement prohibited?

(b) Give details of the advertisements through the Medical Representatives.

(c) What are various type of excise duty levied on alcohol containing preparations.

(d) What are the prescribed offences and penalties under Narcotic and Psychotropic Substances Act. (3+3+3+6)

6. (a) Give details of members in the composition of pharmacy Council of India (PCI). (5+3+7)

(b) What are the functions of Drug Technical Advisory Board (DTAB)?

(c) How state pharmacy council and joint state pharmacy council differs in their composition and function.

7. (a) What are bulk drug, formulation ceiling price, pre tax return and sale turnover?

(b) Describe how the price of drug formulation is calculated.

(c) Write the purpose and application of loan license and repacking license (5+5+5)

8. (a) Define- drug, adulterated drug, patent and proprietary drug, illicit traffic, cannabis.

(b) Write a note on manufacturing of homeopathic medicine.

(c) Discuss the processing of sample for testing in Central Drug Laboratory. (5+5+5)
