BRAINWARE UNIVERSITY

Term End Examination 2022
Programme – B.Pharm-2018/B.Pharm-2019/B.Pharm-2020
Course Name – Pharmaceutical Jurisprudence
Course Code - BP505T
(Semester V)

Full Marks : 75 **Time : 3:0 Hours**

[The figure in the margin indicates full marks. Candidates are required to give their answers in their own words as far as practicable.]

Group-A
(Multiple Choice Type Question) 1 x 20=20

1. *Choose the correct alternative from the following :*

(i) A list of allopathic poisen are given in Schedule
a) E b) F
c) G d) H

(ii) Tests on oral polio vaccine are to be carried out at the
a) National Institute of Communicable Disease, Delhi b) Indian Pharmacopoeial Laboratory, Ghaziabad
c) Central Drug Laboratory, Kolkata d) Central Drug Research Institute, Lucknow

(iii) The Pharmacy Act was passed in
a) 1945 b) 1948
c) 1947 d) 1951

(iv) Government opium factory is situated at
a) Delhi b) Mumbai
c) Hyderabad d) Neemuch

(v) Intellectual Property Rights (IPR) protect the use of information and ideas that are of
a) Ethical value b) Moral value
c) Social value d) Commercial value

(vi) Medical Termination of Pregnancy Act was established in:
a) 1971 b) 1989
c) 1999 d) 1994

(vii) Drug Inspector is appointed under the section of the D & C Act
a) 19 b) 42
c) 21 d) 30

(viii) Patent and proprietary medicines can be imported to some extent in
a) Crude conditions b) Multi-dose vials
c) Bulk form d) Unit containers

(ix) The term "Intellectual Property Rights" covers
a) Copyrights b) Know-how
c) Trade dress d) All of these

(x) The RTI Act gives right to seek information from the following:

a) private companies
b) MNCs
c) public authorities
d) associations

(xi) List of substances that are required to be used only under medical supervision and which are labeled accordingly are included in Schedule

a) R
b) G
c) FF
d) C and C1

(xii) Trade mark

a) is represented graphically
b) is capable of distinguishing the goods or services of one person from those of others
c) may include shapes of goods or combination of colours
d) All of these

(xiii) Biological and microbiological tests are conducted at

a) Mumbai
b) Kolkata
c) Chennai
d) Kasauli

(xiv) The manufacturing blood products or to operate blood bank license is issued in the form no.

a) 28A
b) 28
c) 28B
d) 28C

(xv) Section-14 of the Factory Act states that

a) Effective measures should be taken by the employers to keep workrooms free from dust and fame
b) A factory must be fenced by safeguards of substantial construction
c) Adequate arrangement should be made for running canteens and dispensary for employees
d) Aseptic environment should be maintain around the manufacture process

(xvi) The Inspector general of Forest,Government of India is a member of _____.

a) Drug Consultative Committee
b) Drug Technical Advisory Board
c) Pharmacy Council of India
d) Animal Board of India

(xvii) Drug samples from magistrates are sent for analysis to

a) State Drug Control Laboratory
b) Pharmacy Council of India
c) Central Drug Laboratory
d) Indefinite period

(xviii) Manufactured drugs can be imported into India only under the authorization of

a) Customs Collector
b) State Government
c) Narcotic Commissioner
d) Excise Commissioner

(xix) List of Ayurvedic and Unani poisons is given in schedule

a) E
b) F and F1
c) H
d) E1

(xx) Example of Narcotic drug is

a) Coca
b) Opium
c) Charas
d) All

Group-B

(Short Answer Type Questions) 5 x 7=35

2. Write a short on NLEM and its uses as per DPCO Act. (5)

OR

Justify the details of the schedule A, C, C (1), F, G, H, M, N, X, Y. (5)

3. Describe what type of Drugs and Cosmetics are prohibited for the provision of import under the Drugs and Cosmetics Act, 1940? (5)

OR

Define Register Medical Practitioner and Register Pharmacist. (5)

4. Describe functions of PCI (5)

OR

Discuss about CDL and its function (5)

5. Briefly Explain about duties of government analyst and licensing authority. (5)

OR

Differentiate between Register Medical Practitioner and Register Pharmacist. (5)

6. Explain the functions of DTAB & DCC (5)

OR

Briefly explain the Composition of PCI. (5)

7. Write down about State Pharmacy Council and its composition (5)

OR

Explain the Preparation of first register of pharmacists. (5)

8. Illustrate the prohibition, control & regulation of Opium Act. (5)

OR

Explain the prohibition, control & regulation of Drug and Magic remedies act. (5)

Group-C

(Long Answer Type Questions) 10 x 2=20

9. Briefly explain the functions of Analytical Administrative members. (10)

OR

Explain the power & duties of Drug Inspectors (10)

10. Justify in brief CPCSEA guidelines as per the Prevention of Cruelty to Animals Act, 1960 (10)

OR

Summarize the details on Administrative Authority and its Act & laws under the Drugs and Cosmetic Act, 1940. (10)

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