BRAINWARE UNIVERSITY

Term End Examination 2020 - 21

Programme – Bachelor of Pharmacy

Course Name – Pharmaceutical Jurisprudence

Course Code - BP505T

Semester / Year - Semester V

Time allotted : 90 Minutes | Full Marks : 75

[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 75=75

1. *(Answer any Seventy five)*

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

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

(ii) The total area required for the manufacturer of cosmetic aerosole as per the Schedule M II of Drug and Cosmetic Act is-

a) 15 sq. mt
b) 25 sq. mt
c) 30 sq. mt
d) 35 sq. mt

(iii) The term of patent for invention of the process of manufacture of a substance used as drug or as a food is-

a) 2 years from the date of patent
b) 5 years from the date of sealing
c) 7 years from the date of patent
d) b. 5 years from the date of sealing or 7 years from the date of patent whichever is earlier

(iv) Repacking of drugs means

a) Formulation of drugs in bulk and packing in bulk and packing in small units
b) Breaking up of any drug from a bulk container into small packs and labeling them for sale

c) Packing, dispensing or formulation of drugs in retail sale

d) Compounding of drugs in wholesale business

(v) VDRL antigen is to be tested and analyzed by the

a) Drug Inspector

b) Excise Commissioner

c) Serologist and Chemical Examiner

d) Drug Controller of India

(vi) An example of artificial colour is

a) Titanium dioxide

b) Caramel

c) Cochineal

d) Curcumin

(vii) All adults employed for work in a shop or other establishment should not be made to works daily for more than

a) 8 hours

b) 10 hours

c) 6 hours

d) 12 hours

(viii) Insulin should be taken under the supervision of

a) RMP

b) Pharmacist

c) Nurse

d) All of these

(ix) Drugs and Chemists shops are exempted from the provision of

a) Opening time

b) Closing time

c) a. Opening time & b. Closing time

d) Lunch time

(x) 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

(xi) All the statements regarding CDL are true except

a) To carry analysis of sample of drugs and cosmetics sent by Court or Custom Collector

b) Biological or microbiological testing carried out by CDL, Kolkata

c) To carry all duties suggested by Central or State Govt

d) Analytical report with all protocol of test supplied by the Director

(xii) "Drugs standard" as per the provision of Drugs and Cosmetics Act includes-

a) Drugs complying with standard of official Pharmacopoeia

b) Drugs complying with standard of Drugs and Cosmetics Act

c) Drugs with international standard

d) Drugs complying with standard of all Pharmacopoeia

(xiii) Major amendment in Drugs and Cosmetics Act was made in-

a) 1975

b) 1982

c) 1985

d) 1986

(xiv) Patent and proprietary medicines can be imported to some extent in

a) Crude conditions

b) Multi-dose vials

c) Bulk form

d) Unit containers

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

a) E

b) F and F1

c) E1

d) H

(xvi) Drug retail sale licenses are issued by

a) Drugs Controller of India

b) Union Health Minister

c) Drug Control Authorities Of the States

d) Director of Health Services

(xvii) Establishment which have a qualified person and engage in compounding of drugs is

a) Drug Store
b) Chemists and Druggists
c) Pharmacy
d) Testing Laboratory

(xviii) Ergot and its preparations belong to Schedule

a) P
b) Q
c) C1
d) L

(xix) Digitalis belongs to Schedule

a) E
b) X
c) G
d) H

(xx) Schedule X drug is

a) Amphetamine
b) Cyclobarbital
c) Glutethimide
d) All

(xxi) Competent technical staff engaged in manufacturing of drug should be a

a) Pharmacy graduate with 18 months practical experience
b) Science graduate with chemistry and microbiology with 3 years practical experience
c) Graduate in medicine with 3 years' experience in manufacturing drug
d) Any of these

(xxii) Antisera and toxoids are tested at

a) Chennai
b) Lucknow
c) Izatnagar
d) Delhi

(xxiii) A list of allopathic poison are given in Schedule

a) E
b) F
c) G
d) H

(xxiv) A drug sample taken by the drug inspector for analysis is sent to

a) Drug Controller
b) Drug Inspector
c) Excise Commissioner
d) Government Analyst

(xxv) The records for the drugs having date of expiry should be preserved for a period of at least

a) 5 years
b) 2 years
c) 1 year
d) 3 years

(xxvi) If a drug is not labeled in prescribed manner than it is known as

a) Spurious
b) Misbranded
c) Adulterated
d) Genuine

(xxvii) If the drug has been prepared, packed and stored under insanitary conditions where it may have been rendered injurious to health, it may be termed as

a) Toxic drug
b) Adulterated drug
c) Misbranded drug
d) Spurious drug

(xxviii) Schedule M (GMP) and Y were introduced in Drugs and Cosmetics Act in

a) 1976
b) 1982
c) 1988
d) 1980

(xxix) Drugs by air can be imported into India through

a) Ahmadabad
b) Delhi
c) Chennai
d) All

(xxx) The maximum punishment of imprisonment for the manufacture of spuriousdrug which is likely to cause death is

a) 10 years
b) Life time
c) 5 years
d) 15 years

(xxxi) The front of a pharmacy shall bear an inscription

a) Chemist
b) Druggist
c) Pharmacy
d) Chemist and Druggist

(xxxii) The area of the section to be used as dispensing department shall be not less than

a) 4 sq. mt
b) 6 sq. mt
c) 10 sq. mt
d) 8 sq. mt

(xxxiii) The left hand top corner of the label of schedule 'X' drugs contains symbol

a) XRx
b) Rx
c) RMP
d) TDS

(xxxiv) The tablet or capsule drugs specified in Schedule 'X' can be marketed in packing not exceeding

a) 200 units doses
b) 150 units doses
c) 100 units doses
d) 250 units doses

(xxxv) The period in hours of training to be under taken by a student pharmacist in a hospital is

a) 500
b) 750
c) 600
d) 800

(xxxvi) Education Regulations are approved by

a) State Government
b) Central Government
c) Tribunal
d) Educational Institutions

(xxxvii) A State Council may appoint a Registrar who may act as

a) Secretary
b) Treasurer
c) Vice President
d) Executive member

(xxxviii) The Education Regulations are laid down by

a) Central Government
b) State Government
c) Pharmacy Council of India
d) Ayurveda Council of India

(xxxix) The committee that advises the DTAB and various governments is

a) DCC
b) DEC
c) SPC
d) PCI

(xl) Names from the register can removed only by an order of the-

a) PCI
b) Registration tribunal
c) EC of the State PCI
d) Director of Health Services

(xli) The Essential Commodity Act came into force in-

a) 1945
b) 1950
c) 1955
d) 1960

(xlii) Government opium factory is situated at

a) Delhi
b) Mumbai
c) Hyderabad
d) Neemuch

(xliii) All of the followings are psychotropic agents except

a) Amobarbital
b) Mebrobamate
c) Barbital
d) Doxapram

(xliv) Cannabis sativa yields

a) Opium
b) Heroin
c) Morphine
d) Hemp

(xlv) The Excise-Officer-in-Charge of Bonded Laboratory may permit to take a sample from each batch of finished preparation free of duty up to a maximum amount of

a) 250 ml
b) 150 ml
c) 100 ml
d) 50 ml

(xlvi) From a bonded laboratory the spirituous preparations are sent to the following for the determination of the alcohol strength

a) Government Analyst
b) Excise Commissioner
c) Chemical Examiner
d) Drug Inspector

(xlvii) Manufacture without bond licenses are issued by

a) Excise Commissioner
b) Drug Inspector
c) Government Analyst
d) Registrar

(xlviii) The Medicinal and Toilet Preparations (Excise Duties) Act was enacted for levy and collection of excise duties on……..

a) drugs and cosmetics
b) soaps and detergents
c) Alcohol , Opium, Indian Hemp and other Narcotic drugs
d) Foreign Liquor

(xlix) In accordance with the provisions of the Medicinal and Toilet Preparations (Excise Duties)Act , if manufacturing has mandatorily to be done in presence of the Excise Officer/staff ,the manufactory is called….

a) Bonded
b) Non-bonded
c) unlicensed
d) Liberalised

(l) Medical and Toilet preparation Act was extended in which of the following state(s)?

a) Jhammmu and Kashmir
b) West Bengal
c) Chennai
d) All of these

(li) Any person aggrieved be any notification issued regarding Drugs (Price Control) Order, may apply to Govt. for a review within _____of its publication in the official Gazette

a) 10 days
b) 15 days
c) 30 days
d) 6 months

(lii) Price of formulation sold to retailer in case of Schedule Drug as per Drugs (Price Control) Order should be

a) Retail price minus 5%
b) Retail price minus 10%
c) Retail price minus 16%
d) Retail price and 15% local tax

(liii) List of ailments and diseases that a drug should not claim to cure is given in schedule

a) L
b) J
c) C
d) H

(liv) The D.P.C.O.,2013 is made under

a) The Environment Protection Act
b) The Essential Commodities Act
c) The Drugs and Magic Remedies (Obj. Adv.) Act
d) The Drugs and Cosmetics Act

(lv) When Prevention of Cruelty to Animal Act was established?

a) 1945
b) 1960
c) 1971
d) 1948

(lvi) The Animal Board of India is constituted under the section_____ ofPrevention of Cruelty to Animal Act, 1960.

a) 5A
b) 5B
c) 5C
d) 5D

(lvii) Medical Termination of Pregnancy Act was established in:

a) 1971
b) 1989
c) 1999
d) 1994

(lviii) The Maternity act was established in..

a) 1961 b) 1976

c) 1989 d) 1970

(lix) Name of the Chairman of the Drugs Enquiry Committee is

a) Col.R.N.Chopra b) Maj. Gen. S.L.Bhatia

c) Col. A.L.Chopra d) Dr.A.L. Mudaliar

(lx) The Drugs Enquiry Committee is also known as

a) The Chopra Committee b) The Bhore Committee

c) The Hathi Committee d) The Bhatia Committee

(lxi) The Bhore Committee was set up in the year of …..

a) 1930 b) 1953

c) 1946 d) 1943

(lxii) The Chopra Committee was set up by the….

a) President of India b) Government of India

c) Governor of India d) Prime Minister of India

(lxiii) The first Pharmaceutical industry in India is…

a) the Bengal Chemical and Pharmaceutical Works b) the East India Pharmaceutical Works

c) the Alembic Chemical Works d) the Glaxo

(lxiv) Who is considered as the Father of Pharmacy in India?

a) Mahadev Lal Schroff b) R.N. Chopra

c) J.S. Hathi d) S.L. Bhatia

(lxv) 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

(lxvi) The following cannot be exploited by assigning or by licensing the rights to others.

a) Patents
b) Designs
c) Trademark
d) All of these

(lxvii) In 'quid-pro-quo', quo stands for

a) knowledge disclosed to the public
b) monopoly granted for the term of the patent
c) exclusive privilege of making, selling and using the invention
d) none of these

(lxviii) Design does not include

a) features of shape
b) composition of lines or colours
c) mode or principle of construction
d) none of these

(lxix) The agreement that is enforceable by law is known as

a) Valid agreement
b) Void agreement
c) Illegal agreement
d) Unenforceable agreement

(lxx) RTI Act 2005 came into force on_____.

a) 38637
b) 38579
c) 38518
d) 38657

(lxxi) If the interests of a third party are involved in information sought for, the maximum time limit to get the information will be

a) 30 days
b) 40 days
c) 45 days
d) 60 days

(lxxii) If information sought has been supplied by third party or is treated as confidential by that third party, the third party must be given a representation before the PIO in reply to the notice issued to him within ------ days from the date of receipt of such notice.

a) 5 days
b) 10 days
c) 15 days
d) 30 days

(lxxiii) What is the fee for getting information under RTI Act?

a) Rs. 50 /-
b) Rs. 10/-
c) Rs. 100/-
d) Rs. 1000/-

(lxxiv) First appeal to the first appellate authority can be preferred by the applicant within -------- days from the expiry of the prescribed time limit or from the receipt of the decision from the PIO

a) 30 days
b) 45 days
c) 90 days
d) 60 days

(lxxv) The long title of the RTI Act seeks to promote the following qualities in the working of everypublic authority:

a) Transparency
b) Punctuality
c) Efficiency
d) Reputation