Printed Pages:01 **Sub Code: BP505T**

Paper Id: 231622 **Roll No.** | | | | | | | | | | |

B PHARM

(SEM -V) THEORY EXAMINATION 2022-23

PHARMACEUTICAL JURISPRUDENCE

Time: 3 Hours ***Total Marks: 75***

Note: 1. Attempt all Sections.

SECTION A

1. Attempt *all* questions in brief. **10 x 2 = 20**

a. Define adulterated drugs.

b. What are schedules J and M?

c. What do you mean by the forensic pharmacy?

d. Define the term "retail sale".

e. Define the term "restricted license"

f. Write the objectives of the medicinal and toilet preparation act.

g. Define magic remedy.

h. What are CPCSEA guidelines?

i. Give a short note on the patent.

j. Write the pharmacist's oath.

SECTION B

2. Attempt any *two* parts of the following: **2 x 10 = 20**

a. Discuss in detail about right to information act (RTI).

b. What is the national list of essential medicines (NLEM)? Write objectives of drug price control order (DPCO).

c. Write the constitution and functions of the narcotic & psychotropic consultative committee.

SECTION C

3. Attempt any *five* parts of the following: **7 x 5 = 35**

a. Explain the qualifications, powers, and duties of drug inspectors.

b. Write offense and penalties under the sale of drugs.

c. Discuss the classes of drugs and cosmetics prohibited from import.

d. What do you understand by loan license and repacking license?

e. Write terms and conditions for the cultivation and collection of the opium poppy.

f. What do you mean by intellectual property rights (IPR)?

g. Define advertisement; What do you mean by exempted advertisement?

Subject Code: BP505T

PAPER ID-411016

Roll No: | | | | | | | | | | | | |

B PHARM
(SEM-V) THEORY EXAMINATION 2021-22
PHARMACEUTICAL JURISPRUDENCE

Time: 3 Hours **Total Marks: 75**

Note: 1. Attempt all Sections.

SECTION A

1. Attempt *all* questions in brief. **10 x 2 = 20**

a. What do you mean by misbranded drug?
b. What are the objectives of Patent Act?
c. What is Hathi Committee?
d. Define Pharmaceutical Jurisprudence.
e. Give examples of any 04 Narcotic drugs.
f. Write in brief about code of Pharmaceutical ethics.
g. Write the full form of CPCSEA and IAEC
h. Define the term Poison.
i. What are schedule G and N?
j. What are the objectives of Trademark Act?

SECTION B

2. Attempt any *two* parts of the following: **2 x 10 = 20**

a. Write in detail about Schedule M (GMP)under D and C Act1940.
b. Describe the classes of drug which can be import, export and transshipment under narcotic drugs and psychotropic substances.
c. Write in detail the constitution and functions PCI.

SECTION C

3. Attempt any *five* parts of the following: **7 x 5 = 35**

a. Write in brief the constitution and function of DTAB.
b. Write in brief the qualification, powers and duties of drug Inspector.
c. Give the specimen label for Schedule H and schedule X drug with suitable example?
d. Write a note on drug and cosmetic act 1940 and explain the requirement for obtaining a retail license.
e. Write a note on registration of pharmacists.
f. Write short notes on:
   (i) Pharmacist's role as a member of health care team.
   (ii) Spurious drugs.
g. Define the term advertisement. Mention the objective of Drug and Magic Remedies Act .

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Subject Code: BP505T

B PHARM
(SEM-V) THEORY EXAMINATION 2020-21
PHARMACEUTICAL JURISPRUDENCE

Time: 3 Hours *Total Marks: 75*

Note: 1. Attempt all Sections. If require any missing data; then choose suitably.

SECTION A

1. Attempt *all* questions in brief. **10 x 2 = 20**

a.	What is Schedule G?
b.	Define spurious drug
c.	Define psychotropic substance
d.	Define 'registered pharmacist' and 'medical practitioner'
e.	What is Schedule M?
f.	Write the functions of Drug inspector
g.	Define adulterated drug
h.	Define the terms 'absolute alcohol' and 'denatured alcohol'
i.	Write the objective of Right to Information Act
j.	Write the objective of Medicinal and Toilet preparation Act-1955.

SECTION B

2. Attempt any *two* parts of the following: **2 x 10 = 20**

a.	Explain in detail about Medical Termination of Pregnancy Act
b.	Write a note on licenses required for wholesale of drugs under the provisions of Drug and Cosmetic Act.
c.	Discuss in brief about CPCSEA guidelines for Breeding and Stocking of Animals

SECTION C

3. Attempt any *five* parts of the following: **7 x 5 = 35**

a.	Write a short note on Intellectual Property Rights (IPR).
b.	Discuss the GMP guidelines mentioned under the Schedule M
c.	Describe the procedure for obtaining a patent. Write a note on opposition to grant of patent.
d.	Give a detail account on labelling and packing of drug
e.	Write a note on constitution and functions of Pharmacy Council of India.
f.	Describe the provisions for registration and removal of names as notified in the pharmacy act 1948.
g.	Discuss the rules relating to the export, import and transshipment of Narcotic Drugs

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B PHARM
(SEM-V) THEORY EXAMINATION 2019-20
PHARMACEUTICAL JURISPRUDENCE

Time: 3 Hours ***Total Marks: 75***

Note: 1. Attempt all Sections. If require any missing data; then choose suitably.

SECTION A

1. Attempt *all* questions in brief. **10 x 2 = 20**

a.	What do you mean by jurisprudence?
b.	What are the objectives of drug and cosmetics Act 1940?
c.	What do mean by import of the drugs?
d.	Define coca leaf.
e.	Define prepared opium.
f.	Define bonded laboratory.
g.	Write the full form of CPCSEA and NLEM.
h.	Define advertisement.
i.	Define misbranded drug.
j.	What are the objectives of RTI Act?

SECTION B

2. Attempt any *two* parts of the following: **2 x 10 = 20**

a.	Write in detail about import of the drug and classes of the drug prohibited from import.
b.	Write in detail about PCI and its constitution with its functions.
c.	Write in detail about pharmaceutical legislations.

SECTION C

3. Attempt any *five* parts of the following: **7 x 5 = 35**

a.	Write in brief about Schedule-M.
b.	Write in brief about wholesale and retail sale.
c.	What are the prohibited advertisements?
d.	Write in brief about objectives of prevention to cruelty to animals Act 1960.
e.	Write a note on registration of pharmacists.
f.	Write a note on retail price of formulations.
g.	Write in brief about manufacturing outside bond.

PAPER ID-410773

Roll No: | | | | | | | | | | | | |

BPHARM
(SEM V) THEORY EXAMINATION 2023-24
PHARMACEUTICAL JURISPRUDENCE THEORY

TIME: 3 HRS **M.MARKS: 75**

Note: 1. Attempt all Sections. If require any missing data; then choose suitably.

SECTION A

1. Attempt *all* questions in brief. **10 x 2 = 20**

a.	List of two offences & penalties of the Cruelty to Animal Act 1961.
b.	Write in detail about schedule M.
c.	What are the objective of the RTI Act?
d.	What are adulterated and spurious drugs?
e.	Write the full form of CPCSEA, IAEC & NLEM.
f.	Write the objective of the Narcotics Drug and Psychotropic Substance Act-1985.
g.	Write the formula for calculation of retail price of drug.
h.	Give the 4 functions of the Pharmacy Council of India.
i.	Write the function of the Drug technical advisory board.
j.	Copyright Act, Poison Act, Trademark Act and Medical Termination of Pregnancy Act passed in which year.

SECTION B

2. Attempt any *two* parts of the following: **2 x 10 = 20**

a.	Write about Pharmaceutical Education Regulation under the Pharmacy Act.
b.	Write a note on Qualification of a medical practitioner and place where the pregnancy may be terminated.
c.	Discuss in brief about CPCSEA guidelines for breeding and stocking of animal.

SECTION C

3. Attempt any *five* parts of the following: **7 x 5 = 35**

a.	Write the requirement for obtaining a retail license under D&C Act of 1940.
b.	Write a note on the State & Joint State Pharmacy Council and their function.
c.	Write a brief note on the cultivation and production of opium.
d.	What is the prohibited advertisement under Drug & Magic Remedies Act?
e.	Write a brief review of Pharmaceutical Legislation.
f.	Define the role of the Pharmacist and give the procedure for pharmacist registration.
g.	Write in brief the qualifications, power and duties of a drug inspector.

PAPER ID-311093

Subject Code: BP505T

Roll No: | | | | | | | | | | | | | |

BPHARM

(SEM V) THEORY EXAMINATION 2023-24

PHARMACEUTICAL JURISPRUDENCE – THEORY

TIME: 3 HRS **M.MARKS: 75**

Note: 1. Attempt all Sections. If require any missing data; then choose suitably.

SECTION A

1. Attempt *all* questions in brief. **10 x 2 = 20**

a.	State the objective of the Drug and Cosmetic Act,1940 and its rules 1945
b.	Differentiate between misbranded drugs and adulterated drugs.
c.	Recall Schedule G and H
d.	Write the role of Central Drugs Laboratory and Drugs Inspectors.
e.	Define the term absolute alcohol and psychotropic substances.
f.	Give the composition of the Institutional Animal Ethics Committee
g.	Define patent.
h.	List the offenses and penalties under Narcotic Drugs and Psychotropic Substances Act.
i.	State the places where pregnancy may be terminated.
j.	Define NLEM and write the formula of the retail price of any drug formulation.

SECTION B

2. Attempt any *two* parts of the following: **2 x 10 = 20**

a.	Explain in detail the classes of drugs that can be imported under license, the procedure for application, conditions to be fulfilled, and suspension and cancellation of Import license under the Drug and Cosmetic Act.
b.	Discuss the Right to Information Act.
c.	Explain Education Regulations and registration of pharmacists under the Pharmacy Act.

SECTION C

3. Attempt any *five* parts of the following: **7 x 5 = 35**

a.	Discuss i."Manufacturing of drugs for examination, test or analysis"ii. Loan license.
b.	Illustrate the Sale of Drugs under the Drug and Cosmetic Act.
c.	Explain the procedure for obtaining the license to manufacture Medicinal and Toilet preparations containing alcohol or other narcotic substances.
d.	Enumerate the retail price and the ceiling price of scheduled formulations.
e.	Write note on CPCSEA guidelines for Breeding and Stocking of Animals.
f.	Explain various types of Intellectual Property Rights.
g.	Describe the functions of the Narcotic & Psychotropic Consultative Committee.