Sub Code:BP505T

Paper Id: 150720 *Roll No:*

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-310673

Printed Page: 1 of 1

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

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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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 meanby 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 .

Printed Pages:01 SubCode:BP505T

Paper Id: 231622 Roll No.

B PHARM

(SEM -V) THEORY EXAMINATION 2022-23

PHARMACEUTICALJURISPRUDENCE

Time: 3Hours ***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 detailabout right toinformationact (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.

b. Explain the qualifications, powers, and duties ofdrug inspectors.

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

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

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

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

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

Define advertisement; What do you mean by exempted advertisement?

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. https://www.aktuonline.com

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.

PAPER ID-310801

Roll No:

BPHARM
(SEM V) THEORY EXAMINATION 2024-25
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.	
b.	Define the termAdulterated, Misbranded & Spurious drugs according to D&CAct.
c.	What is the objective of the Pharmacy Act?
d.	Write the formula for calculation of retail price of drug.
e.	Write a note on Repacking License.
f.	Describe the first register and Central register.
g.	Write the function of Drugs Technical Advisory Board.
h.	Copyright Act, Poison Act, Trademark Act and Drug & Magic Remedies Act passed in which year.
i.	Write the difference b/w Charas, Ganja and Bhang.
j.	List of two offences & penalties of the Drug and Magic Remedies Act.
	Write about Schedule M & Schedule G.

SECTION B

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

a.	
b.	Write in brief the constitution and function of the Drug Technical Advisory Board.
c.	Discuss in brief CPCSEA guidelines for the breeding and stocking of animals.
	Write in detail about Import of the drugs & classes of the drug prohibited from import.

SECTION C

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

a.	
b.	Whatis the Hathi Committee and Bhore Committee?
c.	Discuss drug and magic remedies Act 1954 with offences and penalties.
d.	Describe the composition and functions of the Pharmacy Council of India.
e.	Give the qualification and function of Drug Inspector & Government Analyst.
f.	Write a note on Narcotic & Psychotropic substances act 1985 and Rules.
g.	Write in detail about the Drugs Price Control Order 1995.
	Define the role of the Pharmacist and give the procedure for pharmacist registration.

PAPER ID-310888

Roll No:

BPHARM
(SEM V) THEORY EXAMINATION 2025-26
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.	Define misbranded drug as per Drugs and Cosmetics Act, 1940
b.	State any two functions of the Drugs Technical Advisory Board (DTAB).
c.	What is Schedule H of Drugs and Cosmetics Rules, 1945?
d.	Mention any two objectives of the Pharmacy Act, 1948.
e.	Define Registered Pharmacist.
f.	What is magic remedy under Drugs and Magic Remedies Act?
g.	Define ceiling price under DPCO, 2013.
h.	Define Pharmaceutical ethics?
i.	What is NLEM?
j.	What is meant by loan licence?

SECTION B

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

a.	Describe the Drugs and Cosmetics Act, 1940 with respect to: a) Objectives b) Import of drugs c) Prohibition of manufacture and sale d) Offences and penalties
b.	Explain Sale of drugs under Drugs and Cosmetics Rules, 1945 covering: a) Wholesale licence b) Retail licence c) Restricted licence d) Conditions for sale
c.	Discuss the Narcotic Drugs and Psychotropic Substances Act, 1985 with reference to: a) Objectives b) Authorities and officers c) Control and regulation d) Offences and penalties

SECTION C

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

a.	Explain conditions for grant and renewal of manufacturing licence for drugs
b.	Describe the constitution and functions of Pharmacy Council of India (PCI) and explain the registration of pharmacists under pharmacy act,1948
c.	Write a short note on Code of Pharmaceutical Ethics.
d.	Explain intellectual property rights (IPR) and Explain the Right to Information (RTI) Act and its relevance to pharmacy profession.
e.	Write a note on Drugs and Magic Remedies (Objectionable Advertisements) Act.
f.	Explain the Prevention of Cruelty to Animals Act, 1960 with reference to animal ethics committee.
g.	Describe Drug Price Control Order (DPCO), 2013 and its objectives.

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