P.D 3.4

Total No. of Questions : 8] [Total No.of Pages : 01

III/VI PHARM - D DEGREE EXAMINATIONS, JULY - 2022

Third Year

PHARMACEUTICAL JURISPRUDENCE

Time : **Three Hours** Maximum : **70** Marks

Answer any FIVE Questions. **5x14 = 70 M**

All Questions carry equal marks

1. a) Write about the constitution and functions of State Pharmacy Council of India.
   b) Offences and Penalties under Pharmacy Act.
2. a) Write in detail about sale and packaging of drugs under Drugs and Cosmetics Act 1940.
   b) Write a note on Schedule H, N, X and Y.
3. Give an account on
   a) Qualification and duties of Govt. Analyst.
   b) Constitution and functions of CDL.
4. Write a note on
   a) Ethics to be followed by pharmacist towards a medical profession.
   b) Non prescription products.
5. a) Write an account on manufacture of Ayurvedic and Homeopathic preparation under Medicinal and Toilet Preparations Act 1955.
   b) Write a short note on pricing of essential commodity.
6. Write a brief outline on
   a) Drug Price Control Order.
   b) Prevention of cruelty to Animals Act 1960.
7. Discuss the salient features of Drugs and Magic Remedies Act and its rules.
8. a) Define Patent and discuss the provision for getting the patent rights under Patent Act 1970.
   b) Write a brief outline on Pharmaceutical Legislation in India.

P.D 3.4

Total No. of Questions :08] [Total No. of Pages : 02

III/VI Pharma.D (Regular) DEGREE EXAMINATIONS, APRIL-2019

Third Year

PHARMA-D

PHARMACEUTICAL JURISPRUDENCE

Time: Three Hours **Maximum marks:70**

Answer any FIVE questions.

All questions carry equal marks. **5X14=70M**

1. a) What are the qualifications and duties of the Government Analyst and procedures to be followed?

   b) Briefly write about the procedure to be followed by the Drugs Inspector for taking samples.

2. Bring out the salient features of the Narcotic Drugs and Psychotropic Substances Act. Mention the offences and penalties under the Act.

3. a) What are the objectives of Drugs and Magic Remedies Act? Write about types of advertisements prohibited under this act?

   b) Define the following as per Drugs and Cosmetics Act.

   i) Drug (allopathic) ii) Adulterated drugs iii) Manufacture

4. a) Write the constitution and functions of Pharmacy Council of India.

   b) Write a note on the preparation of First Register of Pharmacist under Pharmacy Act.

5. a) Explain the manufacture in a non-bonded laboratory. What are the advantages and disadvantages of manufacture in non-bonded laboratory?

   b) Write the labeling requirements for Schedule H and Schedule X drugs?

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6. a) Enumerate the salient features of Prevention of Cruelty to Animals Act.

   b) Write the procedure for obtaining Indian patent.

7. a) What are the salient differences between wholesale and retail sale of drugs? What are the licensing requirements for retail sale of drugs?

   b) Write about cGMP in pharmaceutical industry.

8. Write short notes on

   a) Write the code of Pharmaceutical Ethics drafted by PCI.

   b) Explain the procedure for fixing the ceiling price of a formulation as per Drug Price Control Order 2013.

Total No. of Questions :07]

P.D 3.4

[Total No. of Pages : 01

III/VI PHARMA-D (Regular) DEGREE EXAMINATIONS, April-2018

Third Year

PHARMA-D

PHARMACEUTICAL JURISPRUDENCE

Time: Three Hours **Maximum marks:70**

SECTION-A

Answer any FIVE Questions **5X14=70M**

All Questions carry equal marks.

1. a) What are the qualifications and duties of the Drugs Inspector and procedures to be followed for sampling.
   b) Write about Drugs Consultative Committee.
2. a) Write the constitution and functions of Pharmacy Council of India.
   b) Write about joint state pharmacy council.
3. a) What are the salient features of Indian Patents and Design Act 1970 with reference to Drugs and Cosmetics?
   b) Write about Drug Price Control Order 2013.
4. Bring out the salient features of the Narcotic Drugs and Psychotropic Substances Act. Explain the procedure for cultivation and production of opium.
5. a) Match the following

1)	Schedule H	a) Particulars to be shown in manufacturing records
2)	Schedule P	b) List of minimum equipment required to run pharmacy
3)	Schedule N	c) List of prescription drugs
4)	Schedule U	d) Life period of drugs

   b) Mention the labelling requirements of Schedule X and G drugs with suitable examples.
   c) Define adulterated and misbranded drugs.
6. a) Explain the construction and working of a bonded laboratory.
   b) Mention the objectives of Drugs and Magic Remedies Act? Write about prohibited advertisements with suitable examples under this Act.
7. a) Explain the procedure for obtaining the license for the manufacture of drugs. What are conditions for grant of license and conditions of license?
   b) Write about the significance of Schedule M
8. Write about the following
   a) Pharmacist ethics with respect to trade and profession
   b) Prevention of cruelty to animals.

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P.D 3.4

Total No. of Questions :08]

[Total No. of Pages : 02

III/VI PHARMA.D (Regular) DEGREE EXAMINATIONS, JULY- 2017

THIRD YEAR

PHARMACEUTICAL JURISPRUDENCE

Time: Three Hours **Maximum marks:70**

Answer any FIVE questions.

All questions carry equal marks. **5X14=70M**

1. a) Write the constitution and functions of State Pharmacy Council.
   b) What are the objectives and duties of Pharmacy Council of India.
2. a) What different bodies are constituted under Drugs and Cosmetics Act for its administration. Give the constitution and functions of Drug Technical Advisory Board.
   b) Write the qualifications of goverment analyst. Write the procedure to be followed for analysis of drugs.
3. a) What classes of drugs are prohibited for import? Explain the procedure for import of drugs meant for different purposes.
   b) What are the objectives of Drugs and Cosmetics Act 1940? What are the subsequent modifications that are made to the Act?
4. a) What are the salient differences between manufacturing inside and outside bond. Write about the procedure for obtaining spirit form distillery.
   b) Explain the procedure for obtaining patent according to Indian Patent Act.
5. a) What is the significance of 'Code of Ethics'? Write about the Code of Ethics of Pharmacists in relation to his profession.
   b) Write about the Prevention of Cruelty to Animals Act 1960.
6. a) Discuss the operations that are permitted and prohibited under the Narcotics and Psychotropic Substances Act and Rules.
   b) Explain the cultivation, production and sale of opium.

7. a) What different types of licenses are issued for the sale of drugs? Bring out the salient differences between them and the necessary conditions for grant of these licenses.

b) Write about Schedule N of Drugs and Cosmetics Act.

8. Write about the following

a) Drug price control order

b) Drugs and Magic Remedies Act.

Total No. of Questions :08]

P.D 3.4

[Total No. of Pages : 01

III/VI Pharm.D (Regular) DEGREE EXAMINATIONS, JULY/AUG-2016

Paper IV- PHARMACEUTICAL JURISPRUDENCE

Time: Three Hours

Maximum marks:70

Answer any FIVE questions.

All questions carry equal marks.

1. a) What are the qualifications and duties of the Government Analyst and procedures to be followed?
   b) Briefly write about the procedure to be followed by the Drugs Inspector for taking samples.
2. Bring out the salient features of the Narcotic Drugs and Psychotropic Substances Act. Mention the offences and penalties under the Act.
3. a) What are the objectives of Drugs and Magic Remedies Act? Which types of advertisements are prohibited under this act?
   b) Define the following as per Drugs and Cosmetics Act.
      i) Drug (allopathic) ii) Adulterated drugs iii) Manufacture iv) Spurious drugs
4. a) Write the constitution and functions of Pharmacy Council of India.
   b) Write a note on the preparation of First Register of Pharmacist under Pharmacy Act.
5. a) Explain the manufacture in a non-bonded laboratory. What are the advantages and disadvantages of manufacture in non-bonded laboratory?
   b) What are the labeling requirements for Schedule H and Schedule X drugs?
6. a) Enumerate the salient features of Prevention of Cruelty to Animals Act.
   b) Explain the significance and features of Indian Patent Act.
7. a) What are the salient differences between wholesale and retail sale of drugs? What are the licensing requirements for retail sale of drugs?
   b) Write about CGMP in pharmaceutical industry.
8. Write short notes on
   a) Professional ethics in relation to trade and profession
   b) What are the reasons for controlling the prices of drugs? Explain the fixation of prices as per DPCO for formulations.