

(LQ 2058)

MARCH 2020

Sub. Code: 2058

**B.PHARM. DEGREE EXAMINATION**  
**PCI Regulation – SEMESTER V**  
**PAPER V – PHARMACEUTICAL JURISPRUDENCE**

*Q.P. Code: 562058*

**Time: Three hours**

**Maximum: 75 Marks**

**I. Elaborate on: Answer any TWO questions.**

**(2 x 10 = 20)**

1. Explain in detail about the constitution and function of state pharmacy council.
2. Describe in detail about Institutional Animal Ethical Committee and CPCSEA guidelines for Animals breeding.
3. Give an account of Import and Export of Narcotic drugs and psychotropic substances.

**II. Write notes on: Answer any SEVEN questions.**

**(7 x 5 = 35)**

1. Education regulation.
2. Code of Ethics of Pharmacist relation to his profession.
3. Registration procedure for Pharmacist after formation of state pharmacy council.
4. Discuss the rule relating to import of drugs as per drugs and cosmetics Act.
5. Conditions for grant of license for wholesale of schedule C and C1 drugs.
6. Classes of prohibited advertisements.
7. Export of alcoholic preparations under bond.
8. Retail price of formulation calculation under drug price control order.
9. Prices of Bulk drugs calculation under drug price Control Order.

**III. Short answers on: Answer ALL questions.**

**(10 x 2 = 20)**

1. Central register.
2. Schedule H.
3. Adulterated drugs.
4. Labeling condition for ophthalmic preparations.
5. State the advisory agency as per drugs and cosmetics Act.
6. Qualification of Drug Inspectors.
7. Misbranded cosmetics.
8. Define Ayurvedic and Unani drugs.
9. Registered Medical Practitioner.
10. Schedule 'X' Drugs.

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THE TAMIL NADU DR. M.G.R. MEDICAL UNIVERSITY

[BPHARM 0921]

SEPTEMBER 2021  
(SEPTEMBER 2020 EXAM SESSION)

Sub. Code: 2058

B.PHARM. DEGREE EXAMINATION  
PCI Regulation 2017 – SEMESTER V  
PAPER V – PHARMACEUTICAL JURISPRUDENCE

*Q.P. Code: 562058*

**Time: Three hours**

**Maximum: 75 Marks**

**I. Elaborate on: Answer any TWO questions. (2 x 10 = 20)**

1. Explain in detail about the Constitutions and functions of Joint State Pharmacy Council.
2. Describe in detail about manufacture in Non bonded laboratory.
3. Give an account on qualification and duties of government analyst.

**II. Write notes on: Answer any SEVEN questions. (7 x 5 = 35)**

1. Labeling and packing of Schedule X drugs.
2. Code of Ethics of Pharmacist relation to his job.
3. Registration of Pharmacist.
4. Repacking Licences and their conditions.
5. Classes of drugs prohibited to sale.
6. Classes of exempted advertisements.
7. Export of duty paid alcoholic preparations.
8. Mention the operations controlled by the central government under Narcotics and psychotropic substances.
9. Salient features of Rights to information Act.

**III. Short answers on: Answer ALL questions. (10 x 2 = 20)**

1. Subsequent registers.
2. Schedule C and C1.
3. Spurious drug.
4. Labeling condition for Schedule G.
5. State the administrative agencies as per Drugs and Cosmetics Act.
6. Define magic remedies.
7. Schedule J.
8. Define Psychotropic substances.
9. Chemists and Druggists.
10. Define Manufacture of drugs.

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**THE TAMIL NADU DR. M.G.R. MEDICAL UNIVERSITY**

**[BPHARM 0122]**

**JANUARY 2022  
(MARCH 2021 EXAM SESSION)**

**Sub. Code: 2058**

**B.PHARMACY DEGREE COURSE (SEMESTER EXAMINATIONS)  
PCI Regulation 2017 – SEMESTER V  
PAPER V – PHARMACEUTICAL JURISPRUDENCE  
Q.P. Code: 562058**

**Time: Three hours**

**Maximum: 75 Marks**

**I. Elaborate on: Answer any TWO questions. (2 x 10 = 20)**

1. Discuss about the conditions of license for manufacture of drugs for test, examination and analysis.
2. Write a note on in bond and outside bond manufacturing.
3. Write in detail about the medical termination of pregnancy act, 1971.

**II. Write notes on: Answer any SEVEN questions. (7 x 5 = 35)**

1. What are the conditions for the issue of import license?
2. Discuss about the compositions of Drugs Technical Advisory Board.
3. Explain the functions of state and joint state pharmacy council.
4. Explain the procedure to sale and export of opium.
5. What are the offences and penalties of drugs and magic remedies act.
6. Write about the sales prices of bulk drugs fixed as per DPCO.
7. Write a brief review on Hathi committee.
8. Give a note on Pharmacist oath.
9. Discuss the powers and functions of information commission.

**III. Short answers on: Answer ALL questions. (10 x 2 = 20)**

1. Define the term misbranded and adulterated drug.
2. Explain the term new drug.
3. What are functions of drugs consultative committee?
4. What is central register?
5. What are Exempted advertisements?
6. Give the main objectives of DPCO 2013.
7. Write the scope of pharmaceutical legislations.
8. Define magic remedies.
9. What is third party license?
10. Define trademark.

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THE TAMIL NADU DR. M.G.R. MEDICAL UNIVERSITY

[BPHARM 0522]

MAY 2022  
(SEPTEMBER 2021 EXAM SESSION)

Sub. Code: 2058

B.PHARM. DEGREE EXAMINATION  
PCI Regulation 2017 – SEMESTER V  
PAPER V – PHARMACEUTICAL JURISPRUDENCE

*Q.P. Code: 562058*

**Time: Three hours**

**Maximum: 75 Marks**

**I. Elaborate on: Answer any TWO questions. (2 x 10 = 20)**

1. Explain the constitutions and functions of state pharmacy council.
2. Discuss the rule belong to sale of drugs as per drugs and cosmetic act.
3. Discuss about the Drugs Price control order.

**II. Write notes on: Answer any SEVEN questions. (7 x 5 = 35)**

1. Discuss about the loan and repacking license.
2. Discuss about the Drugs Technical Advisory Board.
3. Give a note on Pharmacist Oath.
4. Explain the procedure for license to export of alcoholic preparations.
5. Write a short note on Pharmacist in relation to his job.
6. Write a note on Schedule M.
7. Explain about the Medical Termination of pregnancy Act.
8. Write the duties of public authorities.
9. Write the procedure for getting patents and trademark.

**III. Short answers on: Answer ALL questions. (10 x 2 = 20)**

1. Define the term misbranded and adulterated drugs.
2. What are controlling authorities?
3. Define opium derivatives.
4. Define Drugs and magic remedies.
5. Define CPCSEA and IAEC.
6. Define the Nonscheduled formulation.
7. Write the scope of pharmaceutical formulation.
8. Define Schedule C.
9. Define the right to information act.
10. Define registered as medical practitioner?

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**THE TAMIL NADU DR. M.G.R. MEDICAL UNIVERSITY**

**[BPHARM 1022]**

**OCTOBER 2022  
(MARCH 2022 EXAM SESSION)**

**Sub. Code: 2058**

**B.PHARMACY DEGREE COURSE (SEMESTER EXAMINATIONS)**

**PCI Regulation 2017 – SEMESTER V**

**PAPER V – PHARMACEUTICAL JURISPRUDENCE**

***Q.P. Code: 562058***

**Time: Three hours**

**Maximum: 75 Marks**

**I. Elaborate on: Answer any TWO questions. (2 x 10 = 20)**

1. Explain in details about the conditions of license for manufacture of drugs under Drugs and Cosmetics Act, 1940 and its rules 1945.
2. Discuss about the qualification, powers and duties of drugs inspector.
3. Discuss about the salient features of drugs and magic remedies act and its rules.

**II. Write notes on: Answer any SEVEN questions. (7 x 5 = 35)**

1. Describe the Classes of drugs and cosmetics prohibited from import.
2. Loan license.
3. Export under claim for rebate of duty or under bond as per Medicinal and Toilet Preparations Act, 1955.
4. Discuss the legal requirement of cultivation and production of Opium.
5. CPCSEA guidelines for breeding and stocking of animals.
6. The Drugs (Price Control) Order (DPCO), 1995.
7. First and Subsequent Registration of Pharmacist.
8. Medical Termination of Pregnancy Act 1971.
9. Schedule U.

**III. Short answers on: Answer ALL questions. (10 x 2 = 20)**

1. Define Misbranded Drugs.
2. Master Formula Records.
3. What is Schedule P?
4. Non-Bonded manufactory.
5. Manufacture of opium.
6. Functions of DTAB.
7. Offences and penalties under Pharmacy Act 1948.
8. Define Rectified spirit.
9. Hathi committee.
10. Health survey and development committee.

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