



# RRB PHARMACIST

MODEL PAPER -17

## 2024

TIME:-  
**9 P.M**



# 40 QUESTIONS

WITH DETAILED EXPLANATION

**PHARMACEUTICAL  
JURISPRUDENCE**

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# **1. Which schedule contains different types of forms in Drugs and Cosmetics Act, 1945**

- (a) Schedule B**
- (b) Schedule C**
- (c) Schedule D**
- (d) Schedule A**

**1. Which schedule contains different types of forms in Drugs and Cosmetics Act, 1945**

- (a) Schedule B**
- (b) Schedule C**
- (c) Schedule D**
- (d) Schedule A**

## SCHEDULES TO THE RULES

- **A-** Performa for application for the licenses, issues and renewal of licenses, for sending memoranda under the act.
- **B-** Rates of fee for test or analysis by the Central Drugs Laboratory or the state drug laboratories.
- **B<sub>1</sub>** - Fee for the test or analysis by the pharmacopeia laboratory for Indian medicine or the govt. analyst.

**2. Which Schedule gives information about fees for test or analysis by the central drugs laboratories or state drugs laboratories**

- (a) Schedule F**
- (b) Schedule Y**
- (c) Schedule K**
- (d) Schedule B**

**2. Which Schedule gives information about fees for test or analysis by the central drugs laboratories or state drugs laboratories**

- (a) Schedule F**
- (b) Schedule Y**
- (c) Schedule K**
- (d) Schedule B**

## SCHEDULES TO THE RULES

- **A-** Performa for application for the licenses, issues and renewal of licenses, for sending memoranda under the act.
- **B-** Rates of fee for test or analysis by the Central Drugs Laboratory or the state drug laboratories.
- **B<sub>1</sub>** - Fee for the test or analysis by the pharmacopeia laboratory for Indian medicine or the govt. analyst.

### **3. Biological and Biotechnological products are included in IMP**

- (a) Schedule A**
- (b) Schedule B**
- (c) Schedule X**
- (d) Schedule C & C1**



### **3. Biological and Biotechnological products are included in**

- (a) Schedule A**
- (b) Schedule B**
- (c) Schedule X**
- (d) Schedule C & C1**

## SCHEDULES TO THE RULES

- **C** - List of biological and other special products whose import, sale, distribution and manufacture are governed by special provision.
- **C<sub>1</sub>** - List of other special products whose import, sale, distribution and mfg are governed by special provision.
- **D** - List of drugs exempted from the provisions to import of drugs.

## **4. Schedule C and C1 of Drugs and Cosmetics**

### **Act deals with**

- (a) List of biological and special products**
- (b) Management of wholesale pharmacy**
- (c) Dyes and pigments in cosmetics**
- (d) Management of retail pharmacy**

## **4. Schedule C and C1 of Drugs and Cosmetics**

**Act deals with**

- (a) List of biological and special products**
- (b) Management of wholesale pharmacy**
- (c) Dyes and pigments in cosmetics**
- (d) Management of retail pharmacy**

## SCHEDULES TO THE RULES

- **C** - List of biological and other special products whose import, sale, distribution and manufacture are governed by special provision.
- **C<sub>1</sub>** - List of other special products whose import, sale, distribution and mfg are governed by special provision.
- **D** - List of drugs exempted from the provisions to import of drugs.

**5. Which one of the following comes under the schedule C of the Drugs and Cosmetics**

- (a) OTC products**
- (b) Drugs from digitalis**
- (c) Biological products**
- (d) Surgical dressings**

**5. Which one of the following comes under the schedule C of the Drugs and Cosmetics**

- (a) OTC products**
- (b) Drugs from digitalis**
- (c) Biological products**
- (d) Surgical dressings**

## SCHEDULES TO THE RULES

- **C** - List of biological and other special products whose import, sale, distribution and manufacture are governed by special provision.
- **C<sub>1</sub>** - List of other special products whose import, sale, distribution and mfg are governed by special provision.
- **D** - List of drugs exempted from the provisions to import of drugs.



## **6. Schedule D in Drugs and Cosmetics Act refers to**

- (a) List of poisonous substances**
- (b) Standards for surgical dressing**
- (c) List of drugs exempted from certain manufacturing provisions**
- (d) List of drugs exempted from import provision**

## **6. Schedule D in Drugs and Cosmetics Act refers to**

- (a) List of poisonous substances**
- (b) Standards for surgical dressing**
- (c) List of drugs exempted from certain manufacturing provisions**
- (d) List of drugs exempted from import provision**

## SCHEDULES TO THE RULES

- **C** - List of biological and other special products whose import, sale, distribution and manufacture are governed by special provision.
- **C<sub>1</sub>** - List of other special products whose import, sale, distribution and mfg are governed by special provision.
- **D** - List of drugs exempted from the provisions to import of drugs.

## **7. Poisonous drugs are covered in**

- (a) Schedule E1**
- (b) Schedule Y**
- (c) Schedule F**
- (d) Schedule X**

## **7. Poisonous drugs are covered in**

**(a) Schedule E1**

**(b) Schedule Y**

**(c) Schedule F**

**(d) Schedule X**

## SCHEDULES TO THE RULES

- **E<sub>1</sub>** - List of poisonous substances under the Ayurvedic (including Siddha) and Unani systems of medicine.
- **F** - Requirement for the functioning and operation of the blood bank and/or for preparation of blood components.
- **F<sub>1</sub>** -
  - **Part I** - Provisions applicable to the production of all bacterial and viral vaccine.
  - **Part II** - Provisions applicable to the production of all sera from living animal of blood components.
  - **Part III** - Provisions applicable to the manufacture and standardization of diagnostic agents (Bacterial origin)

## **8. Schedule F means**

- (a) Regulations and standards for surgical dressings**
- (b) Regulations and standards for umbilical tapes**
- (c) Regulations and standards for vaccines**
- (d) Regulations and standards for running a blood bank**

## **8. Schedule F means**

- (a) Regulations and standards for surgical dressings**
- (b) Regulations and standards for umbilical tapes**
- (c) Regulations and standards for vaccines**
- (d) Regulations and standards for running a blood bank**



## SCHEDULES TO THE RULES

- **E<sub>1</sub>** - List of poisonous substances under the Ayurvedic (including Siddha) and Unani systems of medicine.
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  - **Part I** - Provisions applicable to the production of all bacterial and viral vaccine.
  - **Part II** - Provisions applicable to the production of all sera from living animal of blood components.
  - **Part III** - Provisions applicable to the manufacture and standardization of diagnostic agents (Bacterial origin)

## **9. Which schedule is standards for surgical dressing**

- (a) Schedule D**
- (b) Schedule F1**
- (c) Schedule F2**
- (d) Schedule F3**

## 9. Which schedule is standards for surgical dressing

- (a) Schedule D
- (b) Schedule F1
- (c) Schedule F2**
- (d) Schedule F3

- **F<sub>2</sub>** - Standards for surgical dressings.
- **F<sub>3</sub>** - Standards for sterilized umbilical tapes.
- **FF** - Standards of ophthalmic preparations.

**10. Under the Drugs and Cosmetics Rules 1945, Schedule \_\_\_ gives standards for umbilical tapes**

- (a) M II**
- (b) M III**
- (c) F II**
- (d) F III**

**10. Under the Drugs and Cosmetics Rules 1945, Schedule \_\_\_ gives standards for umbilical tapes**

- (a) M II**
- (b) M III**
- (c) F II**
- (d) F III**

- **F<sub>2</sub>** - Standards for surgical dressings.
- **F<sub>3</sub>** - Standards for sterilized umbilical tapes.
- **FF** - Standards of ophthalmic preparations.



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## **11. Under Drugs and Cosmetics Act, Schedule FF contains the list of the following**

- (a) Drugs which can be marked under generic names only**
- (b) Drugs which are habit forming**
- (c) Standards for ophthalmic preparation**
- (d) Drugs which are exempt from certain provisions applicable to manufacturing**

## **11. Under Drugs and Cosmetics Act, Schedule FF contains the list of the following**

- (a) Drugs which can be marked under generic names only**
- (b) Drugs which are habit forming**
- (c) Standards for ophthalmic preparation**
- (d) Drugs which are exempt from certain provisions applicable to manufacturing**

- **F<sub>2</sub>** - Standards for surgical dressings.
- **F<sub>3</sub>** - Standards for sterilized umbilical tapes.
- **FF** - Standards of ophthalmic preparations.

## **12. Schedule for medications under supervision of RMP**

- (a) T**
- (b) H**
- (c) S**
- (d) G**

## 12. Schedule for medications under supervision of RMP

(a) T

**(b) H**

(c) S

(d) G



- **G** - List of substances that are required to be used only under medical supervision and which are to be labeled accordingly.
- **H** - List of prescription drugs.
- **J** - Disease or ailments which a drug may not purport to prevent or cure.
- **K** - Drugs exempted from certain provision relating to manufacture of drugs.
- **L -1** Good laboratory practice (GLP) and requirement of premises and equipment.
- **M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

**13. The words "It is dangerous to take this preparation except under medical supervision" can be found on label of drugs belonging to**

- (a) Schedule X**
- (b) Schedule H**
- (c) Schedule C**
- (d) Schedule G**

**13. The words "It is dangerous to take this preparation except under medical supervision" can be found on label of drugs belonging to**

- (a) Schedule X**
- (b) Schedule H**
- (c) Schedule C**
- (d) Schedule G**

- **G** - List of substances that are required to be used only under medical supervision and which are to be labeled accordingly.
- **H** - List of prescription drugs.
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- **K** - Drugs exempted from certain provision relating to manufacture of drugs.
- **L -1** Good laboratory practice (GLP) and requirement of premises and equipment.
- **M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

## **14. Prescription drugs are given under which schedule of Drugs and Cosmetics Rules**

- (a) Schedule K**
- (b) Schedule H**
- (c) Schedule R**
- (d) Schedule M**

## 14. Prescription drugs are given under which schedule of Drugs and Cosmetics Rules

- (a) Schedule K
- (b) Schedule H**
- (c) Schedule R
- (d) Schedule M

- **G** - List of substances that are required to be used only under medical supervision and which are to be labeled accordingly.
- **H** - List of prescription drugs.
- **J** - Disease or ailments which a drug may not purport to prevent or cure.
- **K** - Drugs exempted from certain provision relating to manufacture of drugs.
- **L -1** Good laboratory practice (GLP) and requirement of premises and equipment.
- **M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

**15. Which of the following drug must to be sold by retail only after producing a prescription by a registered medical practitioner**

- (a) Schedule H**
- (b) Schedule G**
- (c) Schedule X**
- (d) Schedule M**



**15. Which of the following drug must to be sold by retail only after producing a prescription by a registered medical practitioner**

- (a) Schedule H**
- (b) Schedule G**
- (c) Schedule X**
- (d) Schedule M**

- **G** - List of substances that are required to be used only under medical supervision and which are to be labeled accordingly.
- **H** - List of prescription drugs.
- **J** - Disease or ailments which a drug may not purport to prevent or cure.
- **K** - Drugs exempted from certain provision relating to manufacture of drugs.
- **L -1** Good laboratory practice (GLP) and requirement of premises and equipment.
- **M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

**16. List of diseases and ailments which a drug may not purport to prevent or cure or make claims to prevent or cure under the Drugs and Cosmetics Rule 1945 is given under**

- (a) Schedule J**
- (b) Schedule K**
- (c) Schedule M**
- (d) Schedule P**

**16. List of diseases and ailments which a drug may not purport to prevent or cure or make claims to prevent or cure under the Drugs and Cosmetics Rule 1945 is given under**

**(a) Schedule J**

**(b) Schedule K**

**(c) Schedule M**

**(d) Schedule P**

- **G** - List of substances that are required to be used only under medical supervision and which are to be labeled accordingly.
- **H** - List of prescription drugs.
- **J** - Disease or ailments which a drug may not purport to prevent or cure.
- **K** - Drugs exempted from certain provision relating to manufacture of drugs.
- **L -1** Good laboratory practice (GLP) and requirement of premises and equipment.
- **M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

**17. Good laboratory practices and requirements of premises and equipment is specified in**

- (a) Schedule M**
- (b) Schedule N**
- (c) Schedule L-1**
- (d) Schedule M-1**

**17. Good laboratory practices and requirements of premises and equipment is specified in**

- (a) Schedule M**
- (b) Schedule N**
- (c) Schedule L-1**
- (d) Schedule M-1**

- **G** - List of substances that are required to be used only under medical supervision and which are to be labeled accordingly.
- **H** - List of prescription drugs.
- **J** - Disease or ailments which a drug may not purport to prevent or cure.
- **K** - Drugs exempted from certain provision relating to manufacture of drugs.
- **L-1** Good laboratory practice (GLP) and requirement of premises and equipment.
- **M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.



**18. In the Drugs and Cosmetics Act and Rules, the schedule relating to GMP is**

- (a) Schedule X**
- (b) Schedule P**
- (c) Schedule M**
- (d) Schedule Y**

**18. In the Drugs and Cosmetics Act and Rules, the schedule relating to GMP is**

- (a) Schedule X**
- (b) Schedule P**
- (c) Schedule M**
- (d) Schedule Y**

**M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

- **M<sub>1</sub>** - Requirement of factory premises etc. for manufacture of homoeopathic preparation
- **M<sub>2</sub>** - Requirement of factory premises etc. for manufacture of cosmetics.
- **M<sub>3</sub>** - Requirements of factory premises for the manufacture of medical devices.

**19. The schedule for good manufacturing practices and requirements of factory premises for the manufacture of belongs to**

- (a) Schedule X**
- (b) Schedule W**
- (c) Schedule L**
- (d) Schedule M**

**19. The schedule for good manufacturing practices and requirements of factory premises for the manufacture of belongs to**

- (a) Schedule X**
- (b) Schedule W**
- (c) Schedule L**
- (d) Schedule M**

**M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

- **M<sub>1</sub>** - Requirement of factory premises etc. for manufacture of homoeopathic preparation
- **M<sub>2</sub>** - Requirement of factory premises etc. for manufacture of cosmetics.
- **M<sub>3</sub>** - Requirements of factory premises for the manufacture of medical devices.

## **20. Schedule M1 of Drugs and Cosmetics Act 1940 & Rules 1945 deals with**

- (a) Allopathic medicines**
- (b) Homeopathic medicines**
- (c) Medical devices**
- (d) Cosmetics**

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- (a) Allopathic medicines**
- (b) Homeopathic medicines**
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**M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

- **M<sub>1</sub>** - Requirement of factory premises etc. for manufacture of homoeopathic preparation
- **M<sub>2</sub>** - Requirement of factory premises etc. for manufacture of cosmetics.
- **M<sub>3</sub>** - Requirements of factory premises for the manufacture of medical devices.



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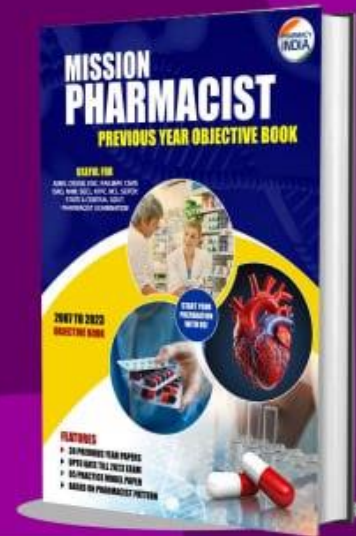


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**21. The version of GMP in India that describe requirement of factory premises for manufacture of cosmetics**

- (a) Schedule-M**
- (b) Schedule-M III**
- (c) Schedule-M II**
- (d) Schedule -MI**

**21. The version of GMP in India that describe requirement of factory premises for manufacture of cosmetics**

- (a) Schedule-M**
- (b) Schedule-M III**
- (c) Schedule-M II**
- (d) Schedule -MI**

**M** - Good manufacturing practice (GMP) requirement of factory premises, plants and equipment.

- **M<sub>1</sub>** - Requirement of factory premises etc. for manufacture of homoeopathic preparation
- **M<sub>2</sub>** - Requirement of factory premises etc. for manufacture of cosmetics.
- **M<sub>3</sub>** - Requirements of factory premises for the manufacture of medical devices.



**22. List of minimum equipment for efficient running of a pharmacy is given in Schedule**

- (a) M**
- (b) N**
- (c) Q**
- (d) R**

**22. List of minimum equipment for efficient running of a pharmacy is given in Schedule**

**(a) M**

**(b) N**

**(c) Q**

**(d) R**

- **N** - List of minimum equipment for efficient running of a pharmacy.
- **O** - Standard for disinfectant fluid.
- **P** - Life period of drug.
- **P<sub>1</sub>** - Pack sizes of drugs.

**23. Which of the following schedules to the Drug and Cosmetics Rules refers to requirements for a pharmacy**

- (a) Schedule M**
- (b) Schedule N**
- (c) Schedule O**
- (d) Schedule P**

**23. Which of the following schedules to the Drug and Cosmetics Rules refers to requirements for a pharmacy**

- (a) Schedule M**
- (b) Schedule N**
- (c) Schedule O**
- (d) Schedule P**

- **N** - List of minimum equipment for efficient running of a pharmacy.
- **O** - Standard for disinfectant fluid.
- **P** - Life period of drug.
- **P<sub>1</sub>** - Pack sizes of drugs.

**24. A modern drug store should fulfil all minimum requirements as specified in the Drugs and Cosmetics Act 1940 and Rules 1945**

- (a) Schedule M**
- (b) Schedule N**
- (c) Schedule P**
- (d) Schedule S**

**24. A modern drug store should fulfil all minimum requirements as specified in the Drugs and Cosmetics Act 1940 and Rules 1945**

**(a) Schedule M**

**(b) Schedule N**

**(c) Schedule P**

**(d) Schedule S**



- **N** - List of minimum equipment for efficient running of a pharmacy.
- **O** - Standard for disinfectant fluid.
- **P** - Life period of drug.
- **P<sub>1</sub>** - Pack sizes of drugs.

## **25. Schedule 0 is related to**

- (a) Standard for ophthalmic preparation**
- (b) Standard for disinfectant fluids**
- (c) For external use only**
- (d) Standard for cosmetics**

## **25. Schedule 0 is related to**

- (a) Standard for ophthalmic preparation**
- (b) Standard for disinfectant fluids**
- (c) For external use only**
- (d) Standard for cosmetics**

- **N** - List of minimum equipment for efficient running of a pharmacy.
- **O** - Standard for disinfectant fluid.
- **P** - Life period of drug.
- **P<sub>1</sub>** - Pack sizes of drugs.

## **26. Schedule P of Drugs and Cosmetics Act refers to**

- (a) Provisions for disinfectants**
- (b) Life period of drugs**
- (c) Pack sizes of drugs**
- (d) Standards for contraceptives**

## **26. Schedule P of Drugs and Cosmetics Act refers to**

- (a) Provisions for disinfectants**
- (b) Life period of drugs**
- (c) Pack sizes of drugs**
- (d) Standards for contraceptives**

- **N** - List of minimum equipment for efficient running of a pharmacy.
- **O** - Standard for disinfectant fluid.
- **P** - Life period of drug.
- **P<sub>1</sub>** - Pack sizes of drugs.

**27. The schedule in D and C act that deals with the pack sizes of drugs is**

- (a) Schedule P1**
- (b) Schedule K**
- (c) Schedule R**
- (d) Schedule S**



**27. The schedule in D and C act that deals with the pack sizes of drugs is**

**(a) Schedule P1**

**(b) Schedule K**

**(c) Schedule R**

**(d) Schedule S**

- **N** - List of minimum equipment for efficient running of a pharmacy.
- **O** - Standard for disinfectant fluid.
- **P** - Life period of drug.
- **P<sub>1</sub>** - Pack sizes of drugs.

**28. "Rule 134 of Drugs and Cosmetics Rules" stipulates that no cosmetic shall be imported which contains coal tar colour other than the one prescribed in Schedule**

- (a) Q**
- (b) N**
- (c) M**
- (d) C**

**28. "Rule 134 of Drugs and Cosmetics Rules" stipulates that no cosmetic shall be imported which contains coal tar colour other than the one prescribed in Schedule**

**(a) Q**

**(b) N**

**(c) M**

**(d) C**

- **Q** - List of coals tar colors permitted to be used in cosmetics.
- **R** - Standard for condoms made of rubber latex and other mechanical contraceptives.
- **R<sub>1</sub>** - Standard for mechanical contraceptive.
- **S**-Standard for cosmetics.
- **T**- Good manufacturing practice for Ayurvedic Siddha, Unani medicines.

## **29. Schedule R of Drugs and Cosmetics Act deals with**

- (a) Disinfectants**
- (b) Antiseptics**
- (c) Mechanical Contraceptives**
- (d) Dyes, Colours & Pigments**

## **29. Schedule R of Drugs and Cosmetics Act deals with**

- (a) Disinfectants**
- (b) Antiseptics**
- (c) Mechanical Contraceptives**
- (d) Dyes, Colours & Pigments**

- **Q** - List of coals tar colors permitted to be used in cosmetics.
- **R** - Standard for condoms made of rubber latex and other mechanical contraceptives.
- **R<sub>1</sub>** - Standard for mechanical contraceptive.
- **S**-Standard for cosmetics.
- **T**- Good manufacturing practice for Ayurvedic Siddha, Unani medicines.



## **30. Schedule "S" in Drugs and Cosmetics Act to the rules stands for**

- (a) Standard for surgical dressing**
- (b) Standard for sterilized umbilical tapes**
- (c) Standard for cosmetics**
- (d) Standard for disinfectant fluids**

## **30. Schedule "S" in Drugs and Cosmetics Act to the rules stands for**

- (a) Standard for surgical dressing**
- (b) Standard for sterilized umbilical tapes**
- (c) Standard for cosmetics**
- (d) Standard for disinfectant fluids**

- **Q** - List of coals tar colors permitted to be used in cosmetics.
- **R** - Standard for condoms made of rubber latex and other mechanical contraceptives.
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**31. Manufacturer of cosmetics to conform to the latest Bureau of Indian Standards requirements is specified in**

- (a) Schedule-A**
- (b) Schedule-S**
- (c) Schedule-F**
- (d) Schedule T**



**31. Manufacturer of cosmetics to conform to the latest Bureau of Indian Standards requirements is specified in**

- (a) Schedule-A**
- (b) Schedule-S**
- (c) Schedule-F**
- (d) Schedule T**

- **Q** - List of coals tar colors permitted to be used in cosmetics.
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- **R<sub>1</sub>** - Standard for mechanical contraceptive.
- **S**-Standard for cosmetics.
- **T**- Good manufacturing practice for Ayurvedic Siddha, Unani medicines.

**32. Under which Schedule "Good Manufacturing practices of Ayurveda, Siddha and for all Unani drugs" are described**

- (a) Schedule -T**
- (b) Schedule -S**
- (c) Schedule M**
- (d) Schedule -Z**

**32. Under which Schedule "Good Manufacturing practices of Ayurveda, Siddha and for all Unani drugs" are described**

- (a) Schedule -T**
- (b) Schedule -S**
- (c) Schedule M**
- (d) Schedule -Z**

- **Q** - List of coals tar colors permitted to be used in cosmetics.
- **R** - Standard for condoms made of rubber latex and other mechanical contraceptives.
- **R<sub>1</sub>** - Standard for mechanical contraceptive.
- **S**-Standard for cosmetics.
- **T**- Good manufacturing practice for Ayurvedic Siddha, Unani medicines.

**33. Every drug manufacturer shall maintain manufacturing and analytical records to the Drugs and Cosmetics Rules, 1945 as per**

- (a) Schedule M**
- (b) Schedule U**
- (c) Schedule V**
- (d) Schedule T**

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**(a) Schedule M**

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**(d) Schedule T**

- **U** - Particulars to be shown in manufacturing records.
- **U<sub>1</sub>** - Particulars to be shown in manufacturing, raw material and analytical records of cosmetic
- **V** - Standard for patent or proprietary medicines.
- **W** - List of drug which is to be marketed under generic names only.
- **X** - List of drugs whose import, manufacture and sale, labeling and packaging are governed by special provision.
- **Y** - Requirement and guideline for permission to import and manufacture of new drugs for sale or to undertake clinical trial



- 34. What does Schedule W of Drugs and Cosmetics Act 1940 and Rules 1945 contain**
- (a) List of drugs which shall be marketed under generic name only**
  - (b) Life period of drugs**
  - (c) Standards for cosmetics**
  - (d) Biological and special products**

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## **35. Habit forming drugs are concerned with**

- (a) Schedule FF**
- (b) Schedule K**
- (c) Schedule X**
- (d) Schedule D**

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## **36. List of narcotic drugs and psychotropic substances are listed in**

- (a) Schedule U**
- (b) Schedule R**
- (c) Schedule W**
- (d) Schedule X**

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**37. The schedule in Drugs and Cosmetics Act that deals with requirement and guidelines the clinical trial import and manufacture of new drug is**

- (a) Schedule O**
- (b) Schedule M**
- (c) Schedule F**
- (d) Schedule Y**

**37. The schedule in Drugs and Cosmetics Act that deals with requirement and guidelines the clinical trial import and manufacture of new drug is**

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## 38. Ibuprofen tablet belongs to schedule

- (a) X
- (b) G
- (c) H
- (d) B

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- (a) X
- (b) G
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- (d) B

## Schedule H Drugs

- Acebutol HCl
- Acyclovir
- Ibuprofen
- Imipramine
- Ketamine HCl
- Ketoprofen
- Metronidazole etc.

**39. Which of the following is a schedule X drug**

- (a) Thalidomide**
- (b) Amoxycillin**
- (c) Glutethimide**
- (d) Halothane**



**39. Which of the following is a schedule X drug**

**(a) Thalidomide**

**(b) Amoxycillin**

**(c) Glutethimide**

**(d) Halothane**

## Schedule H Drugs

- Amobarbital
- Amphetamine
- Barbital
- Dexamphetamine
- Glutethimide
- Methylphenidate
- Phencyclidine etc.

**40. Which of the following is a Schedule G drug**

- (a) Heparin**
- (b) Labetalol HCl**
- (c) Barbituric acid, its salts and derivatives**
- (d) Hydantoin, its salts; its derivatives and their salts**

**40. Which of the following is a Schedule G drug**

**(a) Heparin**

**(b) Labetalol HCl**

**(c) Barbituric acid, its salts and derivatives**

**(d) Hydantoin, its salts; its derivatives and their salts**

## Schedule G Drugs

- Bleomycin
- Chlorpropamide
- Cisplatin
- Cyclophosphamide & its salts
- Doxorubicin HCl
- Ethosuximide
- Hydantoin, its salts, its derivatives etc



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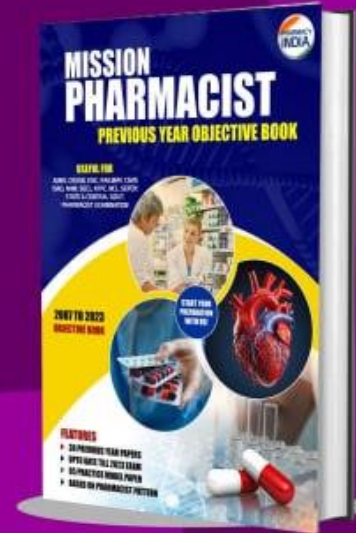


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