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## Pharmaceutical Jurisprudence and Ethics Short Answers - 02 Marks:

- Q 1. Add a note on Drugs Enquiry Committee.
- Q 2. What is Hathi Committee?
- Q 3. What is Mudaliar Committee?
- Q 4. Explain role of pharmacist in relation to his trade.
- Q 5. Explain role of pharmacist in relation to his job.
- Q 6. Explain role of pharmacist in relation to his profession.
- Q 7. Define the term Drug under D & C Act.
- Q 8. Define the term Adulterated drug under D & C Act.
- Q 9. Define the term Spurious drug under D & C Act.
- Q 10. Define the term Misbranded drug under D & C Act.
- Q 11. Define Schedule X and give 2 examples of drugs.
- Q 12. Define Schedule H and give 2 examples of drugs.
- Q 13. Write the objectives of Medicinal and Toilet Preparations Act.
- Q 14. Define i.) Spirit ii.) Denatured Alcohol under M & TP Act.
- Q 14. Define i.) Lunatic; ii.) RMP under MTP Act.
- Q 15. Write the conditions for termination of Pregnancy.
- Q 16. Write the objectives of pharmaceutical policy 2002.
- Q 17. Define the term MAPE.
- Q 18. Write the formula for calculating retail price of a formulation.
- Q 19. Add a note on NLEM.
- Q 20. Write the salient features of Hatch Waxman Act 1984.
- Q 21. Write a note on generic drugs with two examples.
- Q 22. Write the constitution of IAEC.
- m Q 23. Write the functions of IAEC.  $m \square$
- Q 24. List the conditions required for breeding animals for experimentation.
- Q 25. Add a note on NDA.
- Q 26. Add a note on ANDA
- Q 27. Revocation of Patents
- Q 28. Write about the recommendations of DEC.
- Q 29. Write a note on Joint State Pharmacy Council.
- Q 30. Reproduce Pharmacist's Oath.
- Q 31. What are the labeling requirements for i) Colored Preparations ii.) Medicine for External use



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## **Pharmaceutical Jurisprudence and Ethics** <u>Short Essay - 05 Marks:</u>

Q1. Write about the constitution and functions of PCI

Q 2. Explain about ER. Write the salient features of ER 91.

Q 3. Write the constitution and functions of state pharmacy council.

Q 4. How is the first register prepared and maintained?

Q 5. Explain in brief about registration of pharmacist.

Q 6. Write in brief about the layout of a bonded laboratory.

Q 7. Write a note on warehousing of alcoholic preparations.

Q 8. Discuss the provisions relating to manufacture of ayurvedic and homeopathic medicines containing alcohol

Q 9. Explain in brief about manufacturing alcoholic preparations under M & TP Act.

Q 10. Write the operations controlled by Central and State Government under NDPS Act.

Q 11. Discuss about cultivation of opium under NDPS Act.

Q 12. Explain in brief about offences and penalties under NDPS Act.

Q 13. Explain in brief about import and export of Narcotic drugs and Psychotropic substances under NDPS Act.

Q 14. Discuss the advertisements exempted under Drugs and Magic Remedies Act.

Q 15. Discuss the advertisements prohibited under Drugs and Magic Remedies Act.

Q 16. Write the general procedures with time lines for obtaining a patent.

Q 17. What are the qualifications required for appointing a Government Analyst? What are the functions of the Government Analyst?

Q 18. Write the labeling requirements of medicines for internal use with a model label.

Q 19. Explain in detail about Schedule N.

Q 20. Write a note on loan license and repackaging license.

Q 21. What is subsequent register and how is it prepared?

Q 22. What are the grounds on which the name of a pharmacist can be removed from the Central Register? Write about the rights of pharmacist and procedure to appeal against it?

Q 23. Write about Central Drugs Laboratory and its role.

Q 24. Write about the power of entry, search and seizure without warrant under NDPS Act.





## **Pharmaceutical Jurisprudence and Ethics** Long Essay - 10 Marks:

**Q1.** Describe the constitution and functions of a)DTAB b) DCC

**Q** 2. What are the conditions for the grant of a license to manufacture drugs other than those specified in Schedule C and C1 of D and C Rules 1945?

**Q** 3. Explain the different licenses issued for the sale of drugs. Describe the general procedure for a license stating the conditions to be satisfied.

**Q 4.** Explain in detail about Schedule M (GMP) under D & C Act.

**Q 5.** Explain in detail about Schedule Y under D & C Act.

**Q 6.** What are the qualifications for appointment as a drugs inspector? Describe the powers and duties of drugs inspector appointed under D & C Act?

**Q** 7. Enumerate in detail classes of drugs totally prohibited and permitted to be imported under license or permission under D & C Act.

**Q** 8. Discuss the penalty for manufacturing and sale of drugs in contravention of the D & C Act 1940.

Q 9. Explain in detail about manufacturing, import and sale of ayurvedic / homoeopathic medicines.