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DRUG INSPECTOR PREVIOUS YEARS PAPERS contains in this book and preparation tipsSYLLABUS FOR POST OF DRUGS INSPECTOR.PAPER-I PHARMACYThere should be 8 units containing the following :Unit-1-FORENSIC PHARMACY1. Drugs and Cosmetic Act, 1940 and Rules thereunder, 1945 with amendments.2. Pharmacy Act, 1948.3. Drug Price Control Order, 1995.4. Medical Termination of Pregnancy Act, 1971.5. Poison Act, 1919 and Dangerous Drugs Act, 1930.6. Drugs and Magic Remedy Act, 1954.7. Medical and Toilet Preparation Act, 1955.8. Prevention of Cruelty to Animal Act.9.

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PHARMACY Handling of prescription, Incompatibility, Storage conditions of drugs, Clinical Pharmacy and its role in Hospital. Unit-8 - ANATOMY, PHYSIOLOGY & HEALTH EDUCATION

1. Elementary knowledge of following systems :- Blood, Digestive system, Respiratory system, Eye, Ear, Reproductive system and Urinary system.
2. Nutrition, First aid, Population Control, Aids Control.

PAPER-II (GENERAL KNOWLEDGE) : The paper in General Knowledge will include knowledge of current events and matters as of everyday observation and experience in their scientific aspects of life as may be expected of an educated person. The paper will also include questions on History of India and Geography of such standard which the candidates should be able to answer without special study. The author of book is 2 times GPAT qualified

1. General Introduction,
2. History of Drug Legislation and Pharmacy Profession in India,
3. Pharmaceutical Ethics,
4. The Pharmacy Act, 1948,
5. The All India Council for Technical Education Act, 1987,
6. The University Grants Commission (U.G.C.) Act, 1956,
7. The Drugs and Magic Remedies (Objectionable Advertisements) Act, 1954 and Rules, 1955,
8. The Drugs and Cosmetics Act, 1940 and Rules, 1945,
9. The Narcotic Drugs and Psychotropic Substances Act, 1985 and Rules, 1985,
10. Medicinal and Toilet Preparations (Excise Duties) Act, 1955 and Rules, 1956,
11. The Industries (Development and Regulations) Act, 1952,
12. The Prevention of Food Adulteration Act, 1954 and Rules, 1955,
13. National Blood Policy,
14. Pharmaceutical Policy-2002,
15. The Drugs (Price Control) Order (DPCO), 1995,
16. WTO,

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GATS and The Indian Patents Act, 1970 with Amendments

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The purpose of this handbook is to assist individuals for the Certified Pharmaceutical Good Manufacturing Practices Professional (CPGP) examination and provide a reference for the practitioner. The second edition reflects the Body of Knowledge which was updated in 2015. This edition has also incorporated additional information including updated references. The updates reflect the current trends and expectations of the evolving pharmaceutical industry driven by consumer expectations and regulatory oversight. This handbook covers compliance with good manufacturing practices (GMPs), as regulated and guided by national and international agencies for the pharmaceutical industry. It covers finished human and veterinary drugs and biologics, and combination devices, as well as their component raw materials (including active pharmaceutical ingredients (APIs) and excipients), and packaging and labeling operations.

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This is an invaluable revision aid for those preparing for multiple choice questions in clinical pharmacy. Questions in this textbook are practice-oriented and are intended to assess students' knowledge of clinical issues, evaluative and analytical skills, and ability to apply their knowledge in clinical practice. The MCQs will be presented as four practice tests and each test should take c3 hours. Each test will consist of 80 MCQs presented in a variety of formats. Main topics include: therapeutics and rational drug use; aetiology of disease states; presentation of conditions; investigations and diagnostic testing; drug therapy including adverse drug reactions; drug interactions; and contra-indications.

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