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GPAT/DI PREVIOUS YEARS PAPERS UPTO 2019 AVAILABLE IN THIS BOOKGPAT/NIPER-JEE/ DRUG

INSPECTOR PREPARATION TIPS AS PER VARIOUS SUBJECTSCracking GPAT was easier when it used to be conducted with GATE by the IITs earlier. It was very much templatized and there were certain set pieces of questions which were repeated every year. However, now the level of unpredictability has increased in the exam. That is why students should study all topics thoroughly. The preparation tips candidates can use for each section are given below:PharmacognosyThis is a very theoretical section. Applicants must read the chapters comprehensively and make micro notes based on it. These micro notes can help in last minute revision.From the analysis of previous years' exam papers, it can be seen that the most asked drugs from this section are Senna, Morphine and Vinca rosa . Individuals must study all the concepts of these drugs; understand their mechanisms

and their effects & uses. It is best to devote recurrent time periods to Pharmacognosy so that candidates do not forget the concepts and details.Important topics in this section are-*Glycosides*Alkaloids*Volatile Oils*Resins*Tannins*Carbohydrates*Tissue culture*Herbal drugsRecommended books for Pharmacognosy:*Pharmacognosy by C K Kokate*The complete guide for Pharmacognosy by Trease and EvansPharmacologyMany students find this section very interesting. Topics from this section carry a lot of weightage in GPAT. Aspirants must study drug interaction and mechanisms as well as unique side effects of drugs (not general side effects like body pain). For the chapter of Classification of drugs, candidates can use acronyms to remember the names. For eg: 1.Semisynthetic derivatives: Atropine Methonitrate, Homatropine, Hyoscine butyl

bromide, Ipratropium bromide, Tiotropium Bromide (AHITH)2.Quaternary compounds: Propantheline, Clidimium, Oxyphenonium, Pipenzolate methyl bromide, Isopropamide, Glycopyrrolate (PCO PIG)Important topics in this section are:*Oncology (drugs for chemotherapy, upcoming technologies, etc.)*Neuropharmacology and rare diseases*Cardiovascular and Blood productsRecommended books:*Essential of Medical Pharmacology by K D TripathiThe author of this book is 2 times GPAT qualified

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. Trademark Registration Act.10. Pharmaceutical Ethics.Unit-2- MANUFACTURING PHARMACY1. Tablet and Tablet coating.2. Capsule.3. Emulsion, Suspension, Ointment and Cream.4. Ophthalmic Solutions.5. Blood Fluid and Electrolytes.6. Parenteral preparation and Quality Control.7. Surgical Dressing.8. Biological preparation ... (Sera, Vaccine and Anti-Sera)9. Biopharmaceutics.Unit-3- PHARMACEUTICAL ANALYSIS1. Limit Test.2. Bio-Assay.3. Sterility Test.4. Pyrogen Test.5. Theory & Application of

Colorimeter, Florimeter, Nephlometer and Turbidometer, U.V.Visilile Spectrophotometer.6. Kerl Fischer Titration.7. Alcohol determination.8. Microbiological Assay of Vitamins, Antibiotics and Vaccine Preparation.Unit-4- MEDICINAL CHEMISTRYStructure, Storage, Preparation & Brand names of the Following Classes (Definition, Classification etc.) :1. Steroids2. Sedatives and Hypnotics.3. Psycho-therapeutic Agents.4. Antihistaminic Agents.5. Analgesics (narcotic, non-narcotic and NSAID)6. Cardiovascular Agents.Unit-5 -PHARMACOGNOSYSource, Chemical constituents, uses and adulteration of the following classes of natural drugs, Rauwolfia, Ipecacuahna, Belladona, Cinchona, Cinnamon, Digitalis, Senna, Aloe, Noxvomica, Opium, Kurchi, Brahmi, Tulsi, Bael and Ephedra.Unit-6- PHARMACOLOGY &

TOXICOLOGYIntroduction and General Principle-Mode of action, Drug receptor interaction, Drug, antagonist, Absorption, distribution, metabolism and excretion of drugs, Routes of administration, Bioavailability, Drug dependence and addiction, Drug abuse and toxicity, Adverse drug reaction, Drug allergy, Biostatistics.Unit-7- HOSPITAL & CLINICAL PHARMACYHandling of prescription, Incompatibility, Storage conditions of drugs, ClinicalPharmacy and its role in Hospital.Unit-8 - ANATOMY, PHYSIOLOGY & HEALTH EDUCATION1. Elementary knowledge of following systems :-Blood, Digestive system, Respiratory system, Eye, Ear, Reproductive system andUrinary system.2. Nutrition, First aid, Population Control, Aids Control.PAPER-II (GENERAL KNOWLEDGE) : The paper in General Knowledge willinclude

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

A collection of recommended procedures for analysis and specifications for the determination of pharmaceutical substances, excipients and dosage forms intended to serve as source material for reference by any WHO member state.

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),

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superbly written. It starts with never-before focus points that dissect the whole 2020 paper and explains how to structure and answer each question of the 2020 All Subjects (Mathematics, Science, Social Science, English, Hindi A & Hindi B Exam efficiently. Extra value items added in this book: Utilising 15 minute reading time just before the exam (by CBSE topper) Focus points at the beginning (6 pages) Structuring your Maths Exam 3 hours smartly (by CBSE Markers) 2020 marking scheme points (value points) underlined in all papers (CBSE markers look for these key points to allot full marks) Self-assessments are included to help you practice without the temptation of checking the answers at the back and thus strain your memory further to get to the answer. This book provides the right recipe to practice for the English 2020 board exam. Take our word for it :) And of course we are here should you have any

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The process of user-centered innovation: how it can benefit both users and manufacturers and how its emergence will bring changes in business models and in public policy. Innovation is rapidly becoming democratized. Users, aided by improvements in computer and communications technology, increasingly can develop their own new products and services. These innovating users—both individuals and firms—often freely share their innovations with others, creating user-innovation communities and a rich intellectual commons. In Democratizing Innovation, Eric von Hippel looks closely at this emerging system of user-centered innovation. He explains why and when users find it profitable to develop new

products and services for themselves, and why it often pays users to reveal their innovations freely for the use of all.The trend toward democratized innovation can be seen in software and information products—most notably in the free and open-source software movement—but also in physical products. Von Hippel's many examples of user innovation in action range from surgical equipment to surfboards to software security features. He shows that product and service development is concentrated among "lead users," who are ahead on marketplace trends and whose innovations are often commercially attractive. Von Hippel argues that manufacturers should redesign their innovation processes and that they should systematically seek out innovations developed by users. He points to businesses—the custom semiconductor industry is one example—that have learned to assist user-innovators by providing

them with toolkits for developing new products. User innovation has a positive impact on social welfare, and von Hippel proposes that government policies, including R&D subsidies and tax credits, should be realigned to eliminate biases against it. The goal of a democratized user-centered innovation system, says von Hippel, is well worth striving for. An electronic version of this book is available under a Creative Commons license.

Available now to FDA-regulated organizations, this manual allows facility managers to look at their operation's regulatory compliance through the eyes of the government. Because this is the primary reference manual used by FDA personnel to conduct field investigation activities, you can feel confident you are preparing appropriate planning or action. This manual includes revised

instructions regarding the release of information and covers FDA's policies and expectations on a comprehensive range of topics: FDA's authority to enter and inspect, inspection notification, detailed inspection procedures, recall monitoring, inspecting import procedures, computerized data requests, federal/state inspection relationships, discussions with management regarding privileged information, seizure and prosecution, HACCP, bioengineered food, dietary supplements, cosmetics, bioterrorism, and product disposition. The manual also includes a directory of Office of Regulatory Affairs offices and divisions.

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, GATS and The Indian Patents Act, 1970 with Amendments