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undergraduate (B. Sc. First year) students. It is based on UGC model curriculum and as per revised syllabus of Dr. Babasaheb Ambedkar Marathwada University, Aurangabad (w.e.f. June 2013). Present book covers the syllabus of Organic chemistry and Inorganic chemistry papers prescribed for first semester followed by Physical and Inorganic chemistry papers of second semester. The prime objective behind writing this book is to facilitate our dear students for grasping better knowledge of chemistry in an easy, lucid and understandable language. Each topic in the text is provided with point-wise description and elaborated figures. Furthermore, separate Question Bank comprising of long answer questions which are frequently asked in the university examinations with lot of multiple choice questions have been provided at the end of each chapter which will help students to face successfully not only the university examinations but also competitive exams like GATE, SET, NET/JRF, IIT, PET etc. through this platform.

The textbook of Pharmaceutical Chemistry has been written for students of diploma in pharmacy first-year students keeping in mind specific requirements of the Pharmacy Council of India (PCI), Education Regulation - 2020. This is a bilingual book in both English and Hindi for easy understanding to students. This book is covering

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the entire syllabus as per new PCI norms including practicals and previous year questions. This book containing thirteen chapters covering pharmaceutical inorganic chemistry and medicinal chemistry topics. Chapter 1 is introduction to pharmaceutical chemistry containing limit tests, error in analysis, scope, significant figures and quality control methods. Chapter 2 is volumetric analysis containing fundamentals, acid and base theories and titrations. Chapter 3 is related to inorganic pharmaceuticals comprise of hematinics and antacids. Chapter 4 belongs to heterocyclic compounds and their nomenclature. Chapter 5-13 belongs to synthesis and classification of medicinal drugs and their chemistry used in the treatment of several disorder.

Medicinal Chemistry–III provides information pertinent to the fundamental aspects of medicinal chemistry. This book discusses the mechanism of action of drugs at the molecular level. Organized into eight chapters, this book begins with an overview of the systems for protein biosynthesis that are classified according to their mechanism and the selective action of drugs on their reactions. This text then examines the key structure in the biosynthesis of the peptidoglycans of bacterial cell walls. Other chapters consider the enzymes that transcribe DNA synthesizing RNA, which have structural differences in

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eukaryotic and prokaryotic cells. This book discusses as well the concepts for drug screening as they have been emerged from neurochemical research. The final chapter deals with the significance of molecular orbital theory in the hands of the biologist or chemists, which lies in its ability to derive information regarding the properties of molecules. This book is a valuable resource for organic, physical, and biological chemists.

1 Mass transfer 2 Drying 3 Heat transfer 4 Evaporation 5
Crystallization 6 Flow of fluids 7 Distillation 8 Corrosion

Draws parallels between different religious faiths by presenting side-by-side comparisons of four leaders' teachings on topics such as knowledge, suffering, death, and liberation, along with commentaries for each topic.

Drug Regulatory Affairs is a compilation of fundamental concepts pertaining to pharmaceutical drug regulation. Governments protect public health by strictly controlling the safety and efficacy of human medicine, veterinary medicines, medical devices, cosmetics and

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complementary medicines. Companies responsible for the discovery, testing, manufacture and marketing of these products must ensure sale of safe and effective products. Drug Regulatory Affairs refers to fulfilment of all aspects of drug regulations within the pharmaceutical companies from development process to finished product marketing. The book provides a sound basis on understanding of international drug regulatory guidelines controlling the quality, safety and purity of marketed drugs. The content of this book covers the syllabus of pharmacy undergraduate and postgraduate course content of drug regulatory affairs. The text focuses on delivering updated and reviewed up-to-date information on current global regulatory guidelines. The book contains information that is substantial to a comprehensive understanding of regulatory affairs and the practice in pharmaceutical industry.

I-Dispensing Pharmacy - II-Dispensed Medications - a-Monophasic Liquid Dosage Forms - b-Biphasic Liquid Dosage Forms - c- Semi-solid Dosage Forms - III - Sterile Dosage Forms

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