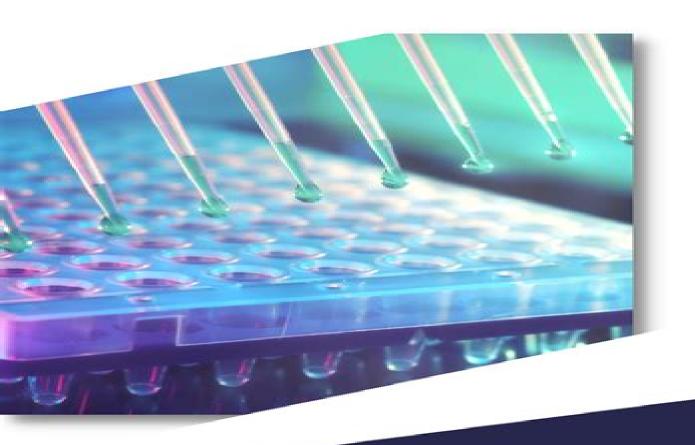
Pharmaceutical Technology EXCELLENCE AWARDS







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Sarfaraz K. Niazi

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Henrique, Jamil, Liliane C., 2021-10-22 Post COVID 19 pandemic researchers have been evaluating the healthcare system for improvements that can be made Understanding global healthcare systems operations is essential to preventative measures to be taken for the next global health crisis A key part to bettering healthcare is the implementation of information management and One Health The Handbook of Research on Essential Information Approaches to Aiding Global Health in the One Health Context evaluates the concepts in global health and the application of essential information management in healthcare organizational strategic contexts This text promotes understanding in how evaluation health and information management are decisive for health planning management and implementation of the One Health concept Covering topics like development partnerships global health and the nature of pandemics this text is essential for health administrators policymakers government officials public health officials information systems experts data scientists analysts health information science and global health scholars researchers practitioners doctors students and academicians

Approaching China's Pharmaceutical Market Ming Q. Lu, 2015-07-30 This authoritative volume examines the major laws regulations and guidelines related to pharmaceutical product development in China With a focus on patent clinical and registration strategies the book helps Western companies introduce their clinical drugs to the Chinese market determine a strategic path and bridge the gap for regulatory and legal differences between China and the Western world For a better understanding of the drug registration process it explores the differences between the China Food and Drug Administration CFDA including its regulations and registration procedures and those of the Western world The volume discusses disparities between China's application requirements compared to Western standards to make it easier for companies to prepare their application packages It also provides detailed commentary on CFDA guidelines in reference to clinical trial IND and market application NDA requirements Overall this book offers guidance for Western companies aspiring to expand into China's pharmaceutical market in hopes that they may gain a fundamental understanding of its rules and complexities in order to ensure a smooth transition and prevent future issues Handbook of Pharmaceutical Manufacturing Formulations Sarfaraz K. Niazi, 2016-04-19 While liquid drugs do not share the compression problems of solid dosage forms the filling problems of powder dosage forms or the consistency problems of semisolid dosage forms they do have their own set of considerations in the formulation and manufacturing stages Highlights from Liquid Products Volume Three include practical details invo Handbook of Pharmacogenomics and Stratified Medicine Sandosh Padmanabhan, 2014-04-28 Handbook of Pharmacogenomics and Stratified Medicine is a comprehensive resource to understand this rapidly advancing field aiming to deliver the right drug at the right dose to the right patient at the right time It is designed to provide a detailed but accessible review of the entire field from basic principles to applications in various diseases The chapters are written by international experts to allow readers from a wide variety of backgrounds clinical and non clinical basic geneticists pharmacologists clinicians trialists industry personnel ethicists to understand the principles underpinning the progress in this area the

successes failures and the challenges ahead To be accessible to the widest range of readers the clinical application section introduces the disease process existing therapies followed by pharmacogenomics and stratified medicine details Medicine is the cornerstone of modern therapeutics prescribed on the basis that its benefit should outweigh its risk It is well known that people respond differently to medications and in many cases the risk benefit ratio for a particular drug may be a gray area The last decade has seen a revolution in genomics both in terms of technological innovation and discovering genetic markers associated with disease In parallel there has been steady progress in trying to make medicines safer and tailored to the individual This has occurred across the whole spectrum of medicine some more than others In addition there is burgeoning interest from the pharmaceutical industry to leverage pharmacogenomics for more effective and efficient clinical drug development Provides clinical and non clinical researchers with practical information normally beyond their usual areas of research or expertise Includes an basic principles section explaining concepts of basic genetic epidemiology bioinformatics pharmacokinetics and pharmacodynamics Covers newer technologies next generation sequencing proteomics metabolomics Provides information on animal models lymphoblastoid cell lines stem cells Provides detailed chapters on a wide range of disease conditions implementation and regulatory issues Includes chapters on the global implications of pharmacogenomics Handbook of Preformulation Sarfaraz K. Niazi, 2019-03-22 Preformulation studies are the physical chemical and biological studies needed to characterize a drug substance for enabling the proper design of a drug product whereas the effectiveness of a drug product is determined during the formulation studies phase Though the two disciplines overlap in practice each is a significantly distinct phase of new drug development Entirely focused on preformulation principles this fully revised and updated Handbook of Preformulation Chemical Biological and Botanical Drugs Second Edition provides detailed descriptions of preformulation methodologies gives a state of the art description of each technique and lists the currently available tools useful in providing a comprehensive characterization of a new drug entity Features Addresses the preformulation studies of three different types of new active entities chemical biological and botanical which is the latest established class of active ingredient classified by the FDA Illustrates the activities comprised in preformulation studies and establishes a method of tasking for drug development projects Includes extensive flow charts for characterization decision making Gives extensive theoretical treatment of principles important for testing dissolution solubility stability and solid state characterization Includes over 50% new material **Directory of Business Information Resources** Laura Mars-Proietti.2008 Resources in Education ,2001-10 **Clinical Informatics Study Guide** John T. Finnell, Brian E. Dixon, 2022-04-22 This completely updated study guide textbook is written to support the formal training required to become certified in clinical informatics. The content has been extensively overhauled to introduce and define key concepts using examples drawn from real world experiences in order to impress upon the reader the core content from the field of clinical informatics. The book groups chapters based on the major foci of the core content health care delivery and policy clinical

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