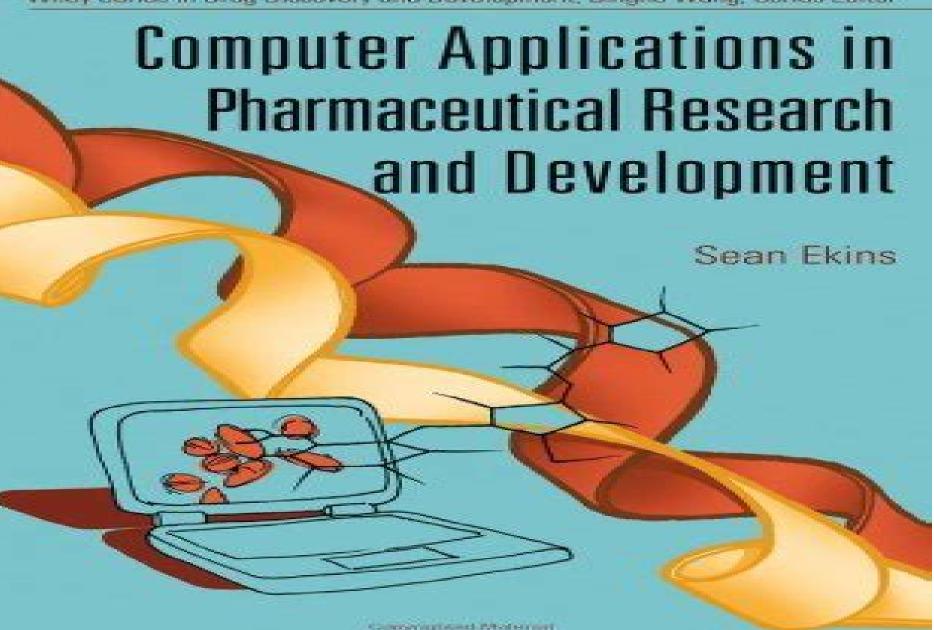


Wiley Series in Drug Discovery and Development, Binghe Wang, Series Editor



Computer Applications In Pharmaceutical Research And Development

Sean Ekins

Computer Applications In Pharmaceutical Research And Development:

Computer Applications in Pharmaceutical Research and Development Sean Ekins, 2006-06-12 A unique holistic approach covering all functions and phases of pharmaceutical research and development While there are a number of texts dedicated to individual aspects of pharmaceutical research and development this unique contributed work takes a holistic and integrative approach to the use of computers in all phases of drug discovery development and marketing It explains how applications are used at various stages including bioinformatics data mining predicting human response to drugs and high throughput screening By providing a comprehensive view the book offers readers a unique framework and systems perspective from which they can devise strategies to thoroughly exploit the use of computers in their organizations during all phases of the discovery and development process Chapters are organized into the following sections Computers in pharmaceutical research and development a general overview Understanding diseases mining complex systems for knowledge Scientific information handling and enhancing productivity Computers in drug discovery Computers in preclinical development Computers in development decision making economics and market analysis Computers in clinical development Future applications and future development Each chapter is written by one or more leading experts in the field and carefully edited to ensure a consistent structure and approach throughout the book Figures are used extensively to illustrate complex concepts and multifaceted processes References are provided in each chapter to enable readers to continue investigating a particular topic in depth Finally tables of software resources are provided in many of the chapters This is essential reading for IT professionals and scientists in the pharmaceutical industry as well as researchers involved in informatics and ADMET drug discovery and technology development The book's cross functional all phases approach provides a unique opportunity for a holistic analysis and assessment of computer applications in pharmaceutics Computer Applications in Pharmaceutical Research and Development. Wiley Series in Drug Discovery and Development Sean Ekins, 2006 Computer Applications in Drug Discovery and Development Ayarivan Puratchikody, Sakthivel Lakshmana Prabu, Appavoo Umamaheswari, 2018-11-23 With more restrictions upon animal experimentations pharmaceutical industries are currently focusing on a new generation of experiments and technologies that are considerably more efficient and less controversial The integration of computational and experimental strategies has led to the identification and development of promising compounds Computer Applications in Drug Discovery and Development is a pivotal reference source that provides innovative research on the application of computers for discovering and designing new drugs in modern molecular biology and medicinal chemistry While highlighting topics such as chemical structure databases and dataset utilization this publication delves into the current panorama of drug discovery where high drug failure rates are a major concern and properly designed virtual screening strategies can be a time saving cost effective and productive alternative This book is ideally designed for chemical engineers pharmacists molecular biologists students researchers and academicians seeking current research on the unexplored avenues and future perspectives of drug design Chemical Engineering in the Pharmaceutical Industry David J. am Ende, Mary T. am Ende, 2019-03-28 A guide to the development and manufacturing of pharmaceutical products written for professionals in the industry revised second edition The revised and updated second edition of Chemical Engineering in the Pharmaceutical Industry is a practical book that highlights chemistry and chemical engineering The book s regulatory quality strategies target the development and manufacturing of pharmaceutically active ingredients of pharmaceutical products The expanded second edition contains revised content with many new case studies and additional example calculations that are of interest to chemical engineers The 2nd Edition is divided into two separate books 1 Active Pharmaceutical Ingredients API s and 2 Drug Product Design Development and Modeling The active pharmaceutical ingredients book puts the focus on the chemistry chemical engineering and unit operations specific to development and manufacturing of the active ingredients of the pharmaceutical product The drug substance operations section includes information on chemical reactions mixing distillations extractions crystallizations filtration drying and wet and dry milling In addition the book includes many applications of process modeling and modern software tools that are geared toward batch scale and continuous drug substance pharmaceutical operations This updated second edition Contains 30new chapters or revised chapters specific to API covering topics including manufacturing quality by design computational approaches continuous manufacturing crystallization and final form process safety Expanded topics of scale up continuous processing applications of thermodynamics and thermodynamic modeling filtration and drying Presents updated and expanded example calculations Includes contributions from noted experts in the field Written for pharmaceutical engineers chemical engineers undergraduate and graduate students and professionals in the field of pharmaceutical sciences and manufacturing the second edition of Chemical Engineering in the Pharmaceutical Industryf ocuses on the development and chemical engineering as well as operations specific to the design formulation and manufacture of drug substance and products

Carbon Monoxide in Drug Discovery Binghe Wang, Leo E. Otterbein, 2022-06-01 CARBON MONOXIDE IN DRUG DISCOVERY An insightful reference for the latest physiological and therapeutic studies of carbon monoxide In Carbon Monoxide in Drug Discovery Basics Pharmacology and Therapeutic Potential a team of distinguished authors delivers foundational knowledge the latest research and remaining challenges regarding the physiological roles and therapeutic efficacy of carbon monoxide CO The editors have included a broad selection of resources from leading experts in the field that discuss the background and physiological roles of CO a variety of delivery forms including CO prodrugs using benign carriers CO sensing therapeutic applications and clinical trials Organized by topic to allow each chapter to be read individually the book covers a wide range of topics from physiological and patho physiological mechanisms at the molecular level to clinical applications for multiple disease processes The editors of Carbon Monoxide in Drug Discovery have created a compelling argument for shifting the accepted understanding of CO from poison to bioactive molecule with enormous clinical

benefits Readers will also benefit from A thorough introduction to the background and physiological actions of carbon monoxide including endogenous CO production in sickness and in health Comprehensive explorations of CO delivery forms including non carrier formulations metal carbonyl complexes and organic CO donors Practical discussions of carbon monoxide sensing and scavenging including fluorescent probes for intracellular carbon monoxide detection In depth examinations of the therapeutic applications of CO including CO in solid organ transplantation Perfect for professors graduate students and postdocs in the fields of biology pharmacology immunology medicinal chemistry toxicology and drug delivery Carbon Monoxide in Drug Discovery Basics Pharmacology and Therapeutic Potential is also an invaluable resource for industrial scientists in these areas Plant Bioactives and Drug Discovery Valdir Cechinel-Filho, 2012-04-11 An in depth exploration of the applications of plant bioactive metabolites in drug research and development Highlighting the complexity and applications of plant bioactive metabolites in organic and medicinal chemistry Plant Bioactives and Drug Discovery Principles Practice and Perspectives provides an in depth overview of the ways in which plants can inform drug research and development An edited volume featuring multidisciplinary international contributions from acclaimed scientists researching bioactive natural products the book provides an incisive overview of one of the most important topics in pharmaceutical studies today With coverage of strategic methods of natural compound isolation structural manipulation natural products in clinical trials quality control and more and featuring case studies on medicinal plants the book serves as a definitive guide to the field of plant biodiversity as it relates to medicine In addition chapters on using natural products as drugs that target specific disease areas including neurological disorders inflammation infectious diseases and cancer illustrate the myriad possibilities for therapeutic applications Wide ranging and comprehensive Plant Bioactives and Drug Discovery also includes important information on marketing regulations intellectual property rights and academic industry collaboration as they relate to plant based drug research making it an essential resource for advanced students and academic and industry professionals working in biochemical pharmaceutical and related fields **Drug Discovery Research** Ziwei Huang, 2007-05-23 Post Genomics Drug Discovery and Research explores and discusses some of the most important topics in post genomics life and biopharmaceutical sciences It provides an introduction to the field outlining examples of many techniques currently used as well as those still under development which are important for the research of biopharmaceutical discovery in the post genomics era Integrates several developing and cutting edge technologies and methods like bioinformatics experimental therapeutics and molecular recognition Includes discussion on topics such as computer aided ligand design peptide and protein chemistry and synthesis synthesis of active natural products and the use of emerging technologies like proteomics nanotechnology or bioengineering Guide to the Library of Congress Classification Lois Mai Chan, Sheila S. Intner, Jean Weihs, 2016-09-26 Like earlier editions this thoroughly updated sixth edition of the classic textbook provides readers with a basic understanding of the Library of Congress Classification system and its

applications The Library of Congress Classification system is used in academic legal medical and research libraries throughout North America as well as worldwide accordingly catalogers and librarians in these settings all need to be able to use it The established gold standard text for Library of Congress Classification LCC the sixth edition of Guide to the Library of Congress Classification updates and complements the classic textbook s coverage of cataloging in academic and research libraries Clear and easy to understand the text describes the reasoning behind assigning subject headings and subheadings including use of tables explains the principles structure and format of LCC details notation tables assigning class numbers and individual classes and covers classification of special types of library materials. The last chapter of this perennially useful resource addresses the potential role of classification in libraries of the future HIV-1 Integrase Nouri Neamati, 2011-08-10 This book comprehensively covers the mechanisms of action and inhibitor design for HIV 1 integrase It serves as a resource for scientists facing challenging drug design issues and researchers in antiviral drug discovery Despite numerous review articles and isolated book chapters dealing with HIV 1 integrase there has not been a single source for those working to devise anti AIDS drugs against this promising target But this book fills that gap and offers a valuable introduction to the field for the interdisciplinary scientists who will need to work together to design drugs that target HIV 1 integrase **Design Strategies** David J. Livingstone, Andrew M. Davis, 2012 This book brings together drug design practitioners all leaders in their field who are actively advancing the field of quantitative methods to guide drug discovery from structure based design to empirical statistical models from rule based approaches to toxicology to the fields of bioinformatics and systems biology. The aim of the book is to show how various facets of the drug discovery process can be addressed in a quantitative fashion ie numerical analysis to enable robust predictions to be made Each chapter includes a brief review of the topic showing the historical development of quantitative approaches a survey summary of the current state of the art a selection of well chosen examples with some worked through and an appreciation of what problems remain to be overcome as well as an indication of how the field may develop After an overview of quantitative approaches to drug design the book describes the development of concepts of drug like properties of quantitative structure activity relationships and molecular modelling and in particular structure based design approaches to guide lead optimisation How to manage and describe chemical structures underpins all quantitative approaches to drug design and these are described in the following chapters The next chapter covers the value of a quantitative approach and also the challenge which is to describe the confidence in any prediction and methods to assess predictive model quality The later chapters describe the application of quantitative approaches to describing and optimising potency selectivity drug metabolism and pharmacokinetic properties and toxicology and the design of chemical libraries to feed the screening approaches to lead generation that underpin modern drug discovery Finally the book describes the impact of bioinformatics current status of predicting ligand affinity direct from the protein structure and the application of quantitative approaches to predicting environmental risk The book provides a

summary of the current state of the art in quantitative approaches to drug design and future opportunities but it also provides inspiration to drug design practitioners to apply careful design to make best use of the quantitative methods that are available while continuing to improve them Drug discovery still relies heavily on random screening and empirical screening cascades to identify leads and drugs and the process has many failures to deliver only a small handful of drugs With the rapidly escalating costs of drug discovery and development together with spiralling delivery quantitative approaches hold the promise of shifting the balance of success to enable drug discovery to maintain its economic viability

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