Drug Discovery Series/5

HANDBOOK OF ASSAY DEVELOPMENT IN DRUG DISCOVERY

EDITED BY LISA K. MINOR



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Patrick Vollmar

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Handbook of Assay Development in Drug Discovery Lisa K. Minor, 2006-01-20 The need to screen targets faster and more efficiently coupled with advances in parallel and multiplex chemical synthesis has contributed to the increasing use of multiwell assays for drug discovery The Handbook of Assay Development in Drug Discovery is a reference that describes the complete armament of tools currently available for performing various assay techniques Featuring contributions from assay developers in the pharmaceutical and vendor communities the book presents descriptions of methods laboratory guidelines and protocols used to perform such methods specific examples of each assay system and troubleshooting tools The handbook describes biochemical assay classes as well as non class specific assay development for cell based assays It covers a wide range of target classes including kinases proteases nuclear receptors and GPCRs and describes currently employed methods and assay types such as radioligand binding assays image analysis assays enzyme fragment complementation and bioluminescent and fluorescent based assays Designed as a guide to running an assay from start to finish the Handbook of Assay Development in Drug Discovery is an ideal bench top companion for discovery researchers laboratory managers academics and other scientists involved in drug discovery screening lead profiling therapeutic target evaluation and assay development and implementation in the pharmaceutical and biotechnology industries Daniel E Levy editor of the Drug Discovery Series is the founder of DEL BioPharma a consulting service for drug discovery programs He also maintains a blog that explores organic chemistry Handbook of Assay Development in Drug Discovery Lisa K. Minor, 2006-01-20 The Handbook of Assay Development in Drug Discovery describes all the tools currently available for performing various assay techniques Featuring troubleshooting advice for common problems from experienced assay developers the vendor community and scientists in the pharmaceutical industry the book presents descriptions of methods laboratory guidelines and protocols used to perform such methods specific examples of each assay system and troubleshooting tools Designed as a quide to running an assay from start to finish this is an ideal bench top companion for scientists involved in drug discovery screening lead profiling therapeutic target evaluation and assay development and implementation GPCR Molecular Pharmacology and Drug Targeting Annette Gilchrist, 2010-12-10 G protein coupled receptors GPCRs are a large protein family of transmembrane receptors vital in dictating cellular responses GPCRs are involved in many diseases but are also the target of around half of all modern medicinal drugs Shifting Paradigms in G Protein Coupled Receptors takes a look at the way GPCRs are examined today how they react how their mutations lead to disease and the many ways in which they can be screened for compounds that modulate them Chemists pharmacologists and biologists will find essential information in this comprehensive reference A Practical Guide to Assay Development and High-Throughput Screening in Drug **Discovery** Taosheng Chen, 2009-12-21 The development of suitable assays the integration of appropriate technology and the effective management of the essential infrastructure are all critical to the success of any high throughput screening HTS

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following World War I that the administration of toxic chemicals such as nitrogen mustards in a controlled manner could shrink malignant tumor masses for relatively substantial periods of time gave great impetus to the search for molecules that would be lethal to specific cancer cells Weare still actively engaged in that search today The question is how to discover these anticancer molecules Anticancer Drug Development Guide Preclinical Screening Clinical Trials and Approval Second Edition describes the evolution to the present of preclinical screening methods The National Cancer Institute's high throughput in vitro disease specific screen with 60 or more human tumor cell lines is used to search for molecules with novel mechanisms of action or activity against specific phenotypes The Human Tumor Colony Forming Assay HTCA uses fresh tumor biopsies as sources of cells that more nearly resemble the human disease There is no doubt that the greatest successes of traditional chemotherapy have been in the leukemias and lymphomas Since the earliest widely used in vivo drug screening models were the murine L 1210 and P388 leukemias the community came to assume that these murine tumor models were appropriate to the discovery of antileukemia agents but that other tumor models would be needed to discover drugs active against solid tumors A Comprehensive Guide to Toxicology in Preclinical Drug Development Ali S. Fagi, 2012-10-18 A Comprehensive Guide to Toxicology in Preclinical Drug Development is a resource for toxicologists in industry and regulatory settings as well as directors working in contract resource organizations who need a thorough understanding of the drug development process Incorporating real life case studies and examples the book is a practical guide that outlines day to day activities and experiences in preclinical toxicology This multi contributed reference provides a detailed picture of the complex and highly interrelated activities of preclinical toxicology in both small molecules and biologics The book discusses discovery toxicology and the international guidelines for safety evaluation and presents traditional and nontraditional toxicology models Chapters cover development of vaccines oncology drugs botanic drugs monoclonal antibodies and more as well as study development and personnel the role of imaging in preclinical evaluation and supporting materials for IND applications By incorporating the latest research in this area and featuring practical scenarios this reference is a complete and actionable guide to all aspects of preclinical drug testing Chapters written by world renowned contributors who are experts in their fields Includes the latest research in preclinical drug testing and international guidelines Covers preclinical toxicology in small molecules and biologics in one single source Drua Discovery and Development Ramarao Poduri, 2021-02-15 This book describes the processes that are involved in the development of new drugs The authors discuss the history role of natural products and concept of receptor interactions with regard to the initial stages of drug discovery In a single highly readable volume it outlines the basics of pharmacological screening drug target identification and genetics involved in early drug discovery The final chapters introduce readers to stem therapeutics pharmacokinetics pharmacovigilance and toxicological testing Given its scope the book will enable research scholars professionals and young scientists to understand the key fundamentals of drug discovery including

stereochemistry pharmacokinetics clinical trials statistics and toxicology **Methods in Microarray Normalization** Phillip Stafford, 2008-01-31 This organized text compiles for the first time the most useful normalization methods developed for interpreting microarray data Experts examine the mathematical processes that are important in normalizing data and avoiding inherent systematic biases They also review modern software including discussions on key algorithms comparative data and download locations The book contains the latest microarray innovations from companies such as Agilent Affymetrix and GeneGo as well as new readily adaptable normalization methods for expression and CGH arrays It also lists of open source molecular profiling normalization algorithms available and where to access them Microarray Innovations Gary Hardiman, 2009-04-02 In recent years high density DNA microarrays have revolutionized biomedical research and drug discovery efforts by the pharmaceutical industry Their efficacy in identifying and prioritizing drug targets based on their ability to confirm a large number of gene expression measurements in parallel has become a key element in drug discovery Handbook of Bioequivalence Testing Sarfaraz K. Niazi, 2014-10-29 As the generic pharmaceutical industry Microarr continues to grow and thrive so does the need to conduct adequate efficient bioequivalence studies In recent years there have been significant changes to the statistical models for evaluating bioequivalence In addition advances in the analytical technology used to detect drug and metabolite levels have m Preclinical Development Handbook Shayne Cox Gad, 2008-03-11 A clear straightforward resource to guide you through preclinical drug development Following this book s step by step guidance you can successfully initiate and complete critical phases of preclinical drug development The book serves as a basic comprehensive reference to prioritizing and optimizing leads toxicity pharmacogenomics modeling and regulations This single definitive easy to use resource discusses all the issues that need consideration and provides detailed instructions for current methods and techniques Each chapter was written by one or more leading experts in the field These authors representing the many disciplines involved in preclinical toxicology screening and testing give you the tools needed to apply an effective multidisciplinary approach The editor with more than thirty years experience working with pharmaceutical and biotechnology companies carefully reviewed all the chapters to ensure that each one is thorough accurate and clear Among the key topics covered are In vitro mammalian cytogenetics tests Phototoxicity Carcinogenicity studies The pharmacogenomics of personalized medicine Bridging studies Toxicogenomics and toxicoproteomics Each chapter offers a full exploration of problems that may be encountered and their solutions. The authors also set forth the limitations of various methods and techniques used in determining the safety and efficacy of a drug during the preclinical stage This is a hands on guide for pharmaceutical scientists involved in preclinical testing enabling them to perform and document preclinical safety tests to meet all FDA requirements before clinical trials may begin

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