

Analytical Profiles of Drug Substances and Excipients

VOLUME 25



Edited by
Harry G. Brittain

Analytical Pro Of Drug Substances Volume 16

**Priyanka Chhabra, Divya Bajpai
Tripathy, Anjali Gupta, Shruti
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Profiles of Drug Substances, Excipients and Related Methodology, 1987-09-04 Profiles of Drug Substances Excipients and Related Methodology *Pharmaceutical Stress Testing* Steven W. Baertschi, Karen M. Alsante, Robert A. Reed, 2016-04-19 The second edition of *Pharmaceutical Stress Testing Predicting Drug Degradation* provides a practical and scientific guide to designing, executing and interpreting stress testing studies for drug substance and drug product. This is the only guide available to tackle this subject in depth. The Second Edition expands coverage from chemical stability. **HPLC for Pharmaceutical Scientists** Yuri V. Kazakevich, Rosario LoBrutto, 2006-12-13 HPLC for Pharmaceutical Scientists is an excellent book for both novice and experienced pharmaceutical chemists who regularly use HPLC as an analytical tool to solve challenging problems in the pharmaceutical industry. It provides a unified approach to HPLC with an equal and balanced treatment of the theory and practice of HPLC in the pharmaceutical industry. In-depth discussion of retention processes, modern HPLC separation theory, properties of stationary phases and columns are well blended with the practical aspects of fast and effective method development and method validation. Practical and pragmatic approaches and actual examples of effective development of selective and rugged HPLC methods from a physico-chemical point of view are provided. This book elucidates the role of HPLC throughout the entire drug development process from drug candidate inception to marketed drug product and gives detailed specifics of HPLC application in each stage of drug development. The latest advancements and trends in hyphenated and specialized HPLC techniques: LC-MS, LC-NMR, Preparative HPLC, High temperature HPLC, high pressure liquid chromatography are also discussed. *Handbook of Modern Pharmaceutical Analysis* Satinder Ahuja, Stephen Scypinski, 2001 This book describes the role modern pharmaceutical analysis plays in the development of new drugs. Detailed information is provided as to how the quality of drug products is assured from the point of discovery until the patient uses the drug. Coverage includes state-of-the-art topics such as analytics for combinatorial chemistry and high throughput screening, formulation development, stability studies, international regulatory aspects and documentation and future technologies that are likely to impact the field. Emphasis is placed on current, easy-to-follow methods that readers can apply in their laboratories. No book has effectively replaced the very popular text *Pharmaceutical Analysis* that was edited in the 1960s by Tak Higuchi. This book will fill that gap with an up-to-date treatment that is both handy and authoritative. **Pharmaceutical Dosage Forms - Tablets** Larry L. Augsburger, Stephen W. Hoag, 2016-04-19 The ultimate goal of drug product development is to design a system that maximizes the therapeutic potential of the drug substance and facilitates its access to patients. *Pharmaceutical Dosage Forms: Tablets* Third Edition is a comprehensive resource of the design, formulation, manufacture and evaluation of the tablet dosage form. **Formulation and Analytical Development for Low-Dose Oral Drug Products** Jack Zheng, 2009-03-04 There are unique challenges in the formulation, manufacture, analytical chemistry and regulatory requirements of low-dose drugs. This book provides an overview

of this specialized field and combines formulation analytical and regulatory aspects of low dose development into a single reference book It describes analytical methodologies like dissolution testing solid state NMR Raman microscopy and LC MS and presents manufacturing techniques such as granulation compaction and compression Complete with case studies and a discussion of regulatory requirements this is a core reference for pharmaceutical scientists regulators and graduate students

Profiles of Drug Substances, Excipients and Related Methodology Harry G. Brittain, 2007-11-14 Whilst following in the footsteps of previous volumes by presenting comprehensive reviews of drug substances and additional materials this title also heralds a significant expansion of the scope of the series Traditional contributions will now also be augmented by publication of critical review chapters that summarize information related to the characterization of drug substances and excipients This change is required to better meet the needs of the pharmaceutical community and to allow the development of a timely vehicle for publishing review materials on this topic The scope of the Profiles series will encompass review articles and database compilations that fall within one of the following six broad categories Physical profiles of drug substances and excipients Analytical profiles of drug substances and excipients Drug metabolism and pharmacokinetic profiles of drug substances and excipients Methodology related to the characterization of drug substances and excipients Methods of chemical synthesis and Reviews of the uses and applications for individual drug substances classes of drug substances or excipients Presents comprehensive reviews covering all aspects of drug development and formulation of drugs Now encompassing critical review chapters Meets the information needs of the drug development community

Retrosynthetic Analysis and Synthesis of Natural Products 1 Olivier Piva, 2019-12-12 For chemists attempting to mimic nature by synthesizing complex natural products from raw material is a challenge that is fraught with pitfalls To tackle this unique but potentially rewarding task researchers can rely on well established reactions and methods of practice or apply their own synthesis methods to verify their potential Whatever the goal and its complexity there are multiple ways of achieving it We must now establish a strategic and effective plan that requires the minimum number of steps but lends itself to widespread use This book is structured around the study of a dozen target products butyrolactone macrolide indole compound cyclobutanic terpene spiro and polycyclic derivatives etc For each product the different disconnections are presented and the associated syntheses are analyzed step by step The key reactions are described explicitly followed by diagrams showing the range of impact of certain transformations This set of data alone is conducive to understanding syntheses and indulging in this difficult but worthwhile activity

Advances in Analytical Techniques for Forensic Investigation Priyanka Chhabra, Divya Bajpai Tripathy, Anjali Gupta, Shruti Shukla, Rajeev Kumar, Kajol Bhati, 2024-08-06 This book is essential for anyone seeking to understand and apply the latest analytical techniques in forensic investigation saving time materials energy and manpower by providing guidance on the most appropriate techniques for different types of investigations *Advances in Analytical Techniques for Forensic Investigation* is aimed to describe the applicability of different types of

analytical techniques used for the forensic investigation including FT IR chromatography mass spectroscopy NMR spectroscopy atomic absorption spectroscopy UV vis spectroscopy etc This book will focus on current and emerging developments in the latest analytical techniques and methods used in the forensic investigation and sample analysis of various physical chemical and biological samples in order to facilitate the smooth conduction of justice *Analytical Chemistry* Harold H. Trimm, 2011-04-15 This title includes a number of Open Access chapters This collection presents a broad selection of recent research on analytical chemistry including methods of determination and analysis as applied to plants pharmaceuticals foods proteins and more Analytical chemistry is the study of what chemicals are present and in what amount in nature

New Psychoactive Substances Hans H. Maurer, Simon D. Brandt, 2019-02-20 This volume is designed to feature the pharmacology of new psychoactive substances legislative aspects information exchange including epidemiology and clinical forensic and analytical toxicology in order to facilitate the understanding of this complex and rapidly developing phenomenon

Helix: The Capstone Research Journal of Ivy Collegiate School Joon Kim, Chaeho Shin, Hayden Lee, Han Na Shin, Tae Hwan Kim, Yeongung Baek, Seokbin Yoon, Yoon-Seo Ha, Dawn Oh, Hyounghbin Koh, Sia Chin, Seoyun Jang, Gabin Kim, Nahee Kwon, Stacy Kwon, Sunghwan Kim, Joshua Lee, Yoonsoo Choi, Jiyong Sim, Sejin Tennant, Jihoon Choi, Woojoo Kim, Daniel Cho, Junwon Lee, Sejoong Park, Sarang Park, Brian Choi, Eddie Koh, 2022-09-22 Helix is the Capstone Research journal of Ivy Collegiate School Student researchers conduct annual research projects in an area of their choosing Selected presentations and papers are published in this journal Research topics vary greatly in approach and field Topics range from questions in the life sciences to studies in musicology

Advances in Neuropharmacology Md. Sahab Uddin, Mamunur Rashid, 2020-01-31 Here is a comprehensive overview of the drugs that act on the central and peripheral nervous systems This volume thoroughly describes the diseases associated with the nervous system and the drugs used for their treatment while also looking at the current status of these drugs and their future potential and challenges Divided into three sections the book first focuses on the drugs that affect the functions of the autonomic nervous system to produce therapeutic effects These drugs may act presynaptically by manipulating the genesis storage and secretion and by blocking the action of neurotransmitters Some drugs may trigger or impede postsynaptic receptors Section 2 focuses on drugs that affect the central nervous system including antianxiety drugs sedative and hypnotic drugs antidepressant drugs antipsychotic drugs antiepileptic drugs and many more It covers the pharmacological management of various diseases including Alzheimer's Parkinson's Huntington's and others The last section offers explanations of neurochemical interactions with the aim to develop drugs that have beneficial effects on neurochemical imbalances This section demonstrates models to assess the transport of drugs across the blood brain barrier and nanomedicine to treat brain disorders This rich compilation provides thorough and extensive research updates on the important advances in neuropharmacological drugs and drug therapy from experienced and eminent academicians researchers and scientists from throughout the world

Genetic Analysis of Animal

Models to Understand the Genomic Architecture of Substance Use Disorders Gary T. Hardiman, Roberto Ciccocioppo, Peter Kalivas, Amy C. Lossie, 2022-11-03

Chemical Analysis of Non-antimicrobial Veterinary Drug Residues in Food Jack F. Kay, James D. MacNeil, Jian Wang, 2016-11-21 Provides a single source reference for readers interested in the development of analytical methods for analyzing non antimicrobial veterinary drug residues in food Provides a comprehensive set of information in the area of consumer food safety and international trade Covers general issues related to analytical quality control and quality assurance measurement uncertainty screening and confirmatory methods Details many techniques including nanotechnology and aptamer based assays covering current and potential applications for non antimicrobial veterinary drugs Provides guidance for analysis of banned drugs including natural and synthetic steroids Resorcylic acid lactones and Beta agonists

Early Drug Development Mitchell N. Cayen, 2011-02-25 The focus of early drug development has been the submission of an Investigational New Drug application to regulatory agencies Early Drug Development Strategies and Routes to First in Human Trials guides drug development organizations in preparing and submitting an Investigational New Drug IND application By explaining the nuts and bolts of preclinical development activities and their interplay in effectively identifying successful clinical candidates the book helps pharmaceutical scientists determine what types of discovery and preclinical research studies are needed in order to support a submission to regulatory agencies

Cumulated Index Medicus , 1991

Practical Pharmaceutics Paul Le Brun, Sylvie Crauste-Manciet, Irene Krämer, Julian Smith, Herman Woerdenbag, 2023-06-15 Practical Pharmaceutics contains essential knowledge on the preparation quality control logistics dispensing and use of medicines It features chapters written by experienced pharmacists and scientists working in hospitals academia and industry throughout Europe including practical examples as well as information on current GMP and GMP based guidelines and EU legislation In this second edition all chapters have been updated with numerous new as well as didactically revised illustrations and tables A completely new chapter about therapeutic proteins and Advanced Therapy Medicinal Products was added From prescription to production from usage instructions to procurement and the impact of medicines on the environment the book provides step by step coverage that will help a wide range of readers students as well as professionals It offers product knowledge for all pharmacists working directly with patients and it will enable them to make the required medicine available to store medicines properly to adapt medicines if necessary and to dispense medicines with the appropriate information for patients as well as caregivers about product care and how to maintain the quality of the product The basic knowledge presented in the book will also be valuable for industrial pharmacists to remind and focus them on the application of the medicines manufactured The basic and practical knowledge on the design preparation and quality management of medicines can directly be applied by the pharmacists whose main duty is production in community and hospital pharmacies and in industry Undergraduate as well as graduate pharmacy students will find knowledge presented in a coherent way and fully supported with relevant examples

Practical Pharmaceutics has become a reliable and recognised source for the acquisition of pharmaceutical technological knowledge. The book is used in the curriculum of a number of international universities and schools of Pharmacy. *National Library of Medicine Current Catalog* National Library of Medicine (U.S.), 1993. Toxicity and Drug Testing Bill Acree, 2012-02-10. Modern drug design and testing involves experimental in vivo and in vitro measurement of the drug candidate's ADMET: adsorption, distribution, metabolism, elimination, and toxicity properties in the early stages of drug discovery. Only a small percentage of the proposed drug candidates receive government approval and reach the market place. Unfavorable pharmacokinetic properties, poor bioavailability, and efficacy, low solubility, adverse side effects, and toxicity concerns account for many of the drug failures encountered in the pharmaceutical industry. Authors from several countries have contributed chapters detailing regulatory policies, pharmaceutical concerns, and clinical practices in their respective countries, with the expectation that the open exchange of scientific results and ideas presented in this book will lead to improved pharmaceutical products.

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