

# Pharmaceutical Particulate Carriers

## Therapeutic Applications



edited by  
Alain Rolland

# Pharmaceutical Particulate Carriers Therapeutic Applications

**Dilip M. Parikh**

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## **Pharmaceutical Particulate Carriers Therapeutic Applications:**

**Pharmaceutical Particulate Carriers** Alain Rolland, 1993-07-19 Offers a comprehensive overview of the medical applications of pharmaceutical particulate carriers discussing the various routes of administration and controlled delivery of sites of action Intravenous liposomal therapy for retroviral infections especially AIDS is discussed **Pharmaceutical Applications of Dendrimers** Abhay Singh Chauhan, Hitesh Kulhari, 2019-11-15 **Pharmaceutical Applications of Dendrimers** explores the applications of dendrimers in the solubilization of hydrophobic active ingredients drug delivery gene delivery imaging diagnosis and photodynamic therapy The book discusses the diagnostic applications of dendrimers including their use as MRI contrast agents and in the imaging of diseased areas In addition the anti inflammatory antimicrobial and antiviral properties of PPI and PAMAM are also covered along with a discussion on photosensitizers such as rose Bengal and protoporphyrin IX that have been delivered using PAMAM and PPI dendrimers for the treatment of cancer This book is an important research reference for those who want to learn more about the development of dendrimer based solutions for drug delivery Explores the role of dendrimers in the design of dendritic nanoplateforms for targeted drug and gene delivery systems Discusses the potential of dendrimers in preformulation and formulation development Addresses both clinical and regulatory challenges in the development of dendrimer based formulations **Handbook of Pharmaceutical Granulation Technology** Dilip M. Parikh, 1997-06-17 Integrating the basic principles and industrial practices of pharmaceutical granulation production this book discusses technologies and demonstrates cost effective approaches to manufacturing solid dosage forms with content uniformity and consistent physical properties while complying with regulatory requirements Specialists from pharmaceutical companies academia and the U S Drug Regulatory Affairs agency address current and changing practices in industrial drug granulation production Text charts figures and photographs illustrate the pros and cons of diverse methods and technologies for accurately achieving strong bonding of particles in tablets and capsules **The Pharmaceutical Regulatory Process** Ira R. Berry, Robert P. Martin, 2008-12-02 This Second Edition examines the mechanisms and means to establish regulatory compliance for pharmaceutical products and company practices It focuses on major legislative revisions that impact requirements for drug safety reviews product regulatory approvals and marketing practices Written by top industry professionals practicing attorneys an **Pharmaceutical Gene Delivery Systems** Alain Rolland, 2003-04-11 This volume examines the advantages and limitations of the major gene delivery systems and offers guidelines to select the most appropriate viral or synthetic delivery system for specific therapeutic applications It discusses advances in the design optimization and adaptation of gene delivery systems for the treatment of cancerous cardiovascular pulmonary genetic and infectious diseases **Nanoparticulate Drug Delivery Systems** Deepak Thassu, Michel Deleers, Yashwant Vishnupant Pathak, 2007-03-30 With the advent of analytical techniques and capabilities to measure particle sizes in nanometer ranges there has been tremendous interest in the use of nanoparticles for more efficient methods

of drug delivery Nanoparticulate Drug Delivery Systems addresses the scientific methodologies formulation processing applications recent trends and e  *Microparticulate Systems for the Delivery of Proteins and Vaccines Smadar Cohen,Howard Bernstein,2020-07-24* This practical guide offers concise coverage of the scientific and pharmaceutical aspects of protein delivery from controlled release microparticulate systems emphasizing protein stability during encapsulation and release **Good Manufacturing Practices for Pharmaceuticals** Joseph D. Nally,2016-04-19 With global harmonization of regulatory requirements and quality standards and national and global business consolidations ongoing at a fast pace pharmaceutical manufacturers suppliers contractors and distributors are impacted by continual change Offering a wide assortment of policy and guidance document references and interpretations this Sixth Edition is significantly expanded to reflect the increase of information and changing practices in CGMP regulation and pharmaceutical manufacturing and control practices worldwide An essential companion for every pharmaceutical professional this guide is updated and expanded by a team of industry experts each member with extensive experience in industry or academic settings  *Physics and Chemistry Basis of Biotechnology* M. de Cuyper,Jeff W.M. Bulte,2006-04-11 At the end of the 20th century a tremendous progress was made in biotechnology in its widest sense This progress was largely possible as a result of joint efforts of top academic researchers in both pure fundamental sciences and applied research The surplus value of such interdisciplinary approaches was clearly highlighted during the 9th European Congress on Biotechnology that was held in Brussels Belgium 11 15 July 1999 The present volume in the Focus on Biotechnology series entitled Physics and Chemistry Basis for Biotechnology contains selected presentations from this meeting A collection of experts has made serious efforts to present some of the latest developments in various scientific fields and to unveil prospective evolutions on the threshold of the new millenium In all contributions the emphasis is on emerging new areas of research in which physicochemical principles form the foundation In reading the different chapters it appears that more than ever significant advances in biotechnology very often depend on breakthroughs in the biotechnology itself e g Oral Lipid-Based Formulations David J. Hauss,2007-06-08 Oral lipid based formulations are attracting considerable attention due to their capacity to facilitate gastrointestinal absorption and reduce or eliminate the effect of food on the absorption of poorly water soluble lipophilic drugs Despite the obvious and demonstrated utility of these formulations for addressing a persistent and growing problem **Handbook of Pharmaceutical Analysis** Lena Ohannesian,Anthony Streeter,2001-11-09 Exploring the analysis of pharmaceuticals including polymorphic forms this book discusses regulatory requirements in pharmaceutical product development and pharmaceutical testing It covers methods of drug separation and procedures such as capillary electrophoresis for chromatographic separation of molecules Additional topics include drug formulation analysis using vibrational and magnetic resonance spectroscopy and identification of drug metabolites and decomposition products using such techniques as mass spectrometry The book provides more than 300 tables equations drawings and photographs and

convenient easy to use indices facilitating quick access to each topic

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New Drug Development Chandrahas Sahajwalla, 2004-05-24 Highlighting key points from the latest regulatory requirements New Drug Development helps those new to the world of pharmaceutical development understand regulatory steps reduce cost by avoiding unnecessary trials and attain guidance through each step of the drug approval process This volume acquaints readers with procedures that determine the

**Nanotherapeutics** Alf Lamprecht, 2016-04-19 This timely book provides an overview of possible therapeutic applications The first part of the book highlights general properties of and phenomena observed with nanoparticles and the subsequent consequences for applications in drug delivery The second part focuses on the therapeutic approaches that are possible through the use of nanoparticles

Affinity Capillary Electrophoresis in Pharmaceutics and Biopharmaceutics Reinhard H. H. Neubert, Hans-Hermann Ruttinger, 2003-01-14 This reference presents the most recent breakthroughs and techniques in affinity capillary electrophoresis ACE to measure and determine the physicochemical and thermodynamic parameters of drug compounds The authors offer strategies to explore and characterize interactions between drugs drug vehicles and biological membranes to facilitate devel

Pharmacogenomics Werner Kalow, Urs B. Meyer, Rachel F. Tyndale, 2001-05-08 Summarizes the history of and available applications techniques and approaches to pharmacogenomics improving efficacy and tolerance to pharmaceutical compounds evaluating pharmacogenomics on clinical trial design and increasing the chances of successful clinical trials and patient outcomes Compares pharmacogenomics to pharmacogenetics focusing on the essential components that incorporate individual genetic variations to drug regimens resulting in different responses to therapeutics

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**Mechanisms of Transdermal Drug Delivery** Russel O. Potts, 1997-07-15 Provides an up to date and critical examination of biophysical techniques used in the analysis of molecular mechanisms underlying transdermal drug delivery as well as a physical and chemical evaluation of the stratum corneum necessary for the enhancement of percutaneous drug transport Reflects the hands on experience of established and novel researchers in the field

**Polymeric Drug Delivery Systems** Glen S. Kwon, 2005-04-12 Emphasizing four major classes of polymers for drug delivery water soluble polymers hydrogels biodegradable polymers and polymer assemblies this reference surveys efforts to adapt modify and tailor polymers for challenging molecules such as poorly water soluble compounds peptides proteins and plasmid DNA

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