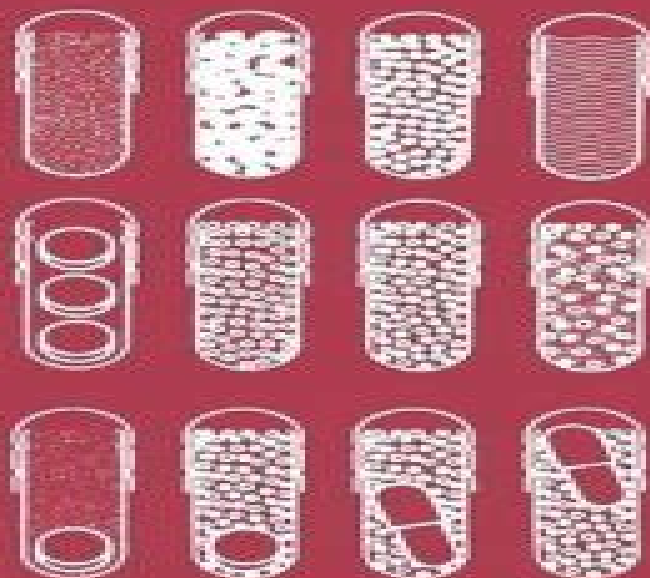


Multiparticulate Oral Drug Delivery



edited by
Isaac Ghebre-Sellassie

Multiparticulate Oral Drug Delivery

**Gordon L. Amidon, Ping I. Lee, Elizabeth
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Multiparticulate Oral Drug Delivery:

Multiparticulate Oral Drug Delivery Isaac Ghebre-Selassie,1994-05-19 This definitive reference explores the various aspects of multiparticulate dosage form development assessing the in vivo behavior and performance of multiparticulates as well as comparing their market position to other dosage forms Discussing for the first time in a comprehensive manner alternative pelletization techniques such as balling spherical agglomeration spray congealing and cryopelletization Multiparticulate Oral Drug Delivery describes formulation and processing variables that affect the formation integrity and performance of coatings derived from molten materials and polymeric solutions and dispersions analyzes the biopharmaceutical aspects and in vivo performance of multiparticulate drug delivery systems details the idiosyncrasies of the manufacturing process of hard shell gelatin capsules explains the different kinds of packaging materials and machinery and the importance of packaging during the development phase and beyond previews the marketing considerations of multiparticulate systems in the years to come presents practical solutions to problems encountered in the area of film coating processes and more *Multiparticulate Oral Drug Delivery Systems* Christoph Schmidt,2000 *Multiparticulate Drug Delivery* Ali R. Rajabi-Siahboomi,2017-05-26 Authored by leading experts from academia users and manufacturers this book provides an authoritative account of the science and technology involved in multiparticulate drug delivery systems which offer superior clinical and technical advantages over many other specialized approaches in drug delivery The book will cover market trends potential benefits and formulation challenges for various types of multiparticulate systems Drug solubility dose chemistry and therapeutic indications as well as excipient suitability coupled with manufacturing methods will be fully covered Key approaches for taste masking delayed release and extended release of multiparticulates systems are of significant interest especially their in vivo and in vitro performance In addition the principles of scale up QbD and regulatory aspects of common materials used in this technology will be explained as well as recent advances in materials and equipment enabling robust flexible and cost effective manufacture Case studies illustrating best practices will also make the book a valuable resource to pharmaceutical scientists in industry and academia **Oral Drug Delivery for Modified Release Formulations** Edmund S. Kostewicz,Maria Vertzoni,Heather A. E. Benson,Michael S. Roberts,2022-04-04 ORAL DRUG DELIVERY FOR MODIFIED RELEASE FORMULATIONS Provides pharmaceutical development scientists with a detailed reference guide for the development of MR formulations Oral Drug Delivery for Modified Release Formulations is an up to date review of the key aspects of oral absorption from modified release MR dosage forms This edited volume provides in depth coverage of the physiological factors that influence drug release and of the design and evaluation of MR formulations Divided into three sections the book begins by describing the gastrointestinal tract GIT and detailing the conditions and absorption processes occurring in the GIT that determine a formulation s oral bioavailability The second section explores the design of modified release formulations covering early drug substance testing the biopharmaceutics classification system an

array of formulation technologies that can be used for MR dosage forms and more The final section focuses on in vitro in silico and in vivo evaluation and regulatory considerations for MR formulations Topics include biorelevant dissolution testing preclinical evaluation and physiologically based pharmacokinetic modelling PBPK of in vivo behaviour Featuring contributions from leading researchers with expertise in the different aspects of MR formulations this volume Provides authoritative coverage of physiology physicochemical determinants and in vitro in vivo correlation IVIVC Explains the different types of MR formulations and defines the key terms used in the field Discusses the present status of MR technologies and identifies current gaps in research Includes a summary of regulatory guidelines from both the US and the EU Shares industrial experiences and perspectives on the evaluation of MR dosage formulations Oral Drug Delivery for Modified Release Formulations is an invaluable reference and guide for researchers industrial scientists and graduate students in general areas of drug delivery including pharmaceuticals pharmaceutical sciences biomedical engineering polymer and materials science and chemical and biochemical engineering Pharmaceutical Extrusion Technology Isaac Ghebre-Selassie, Charles Martin, 2003-05-14 Pharmaceutical Extrusion Technology is the only resource to provide in depth descriptions and analyses of the key parameters of extruders and extrusion processes The book highlights the applicability of melt extrusion in pharmaceutical drug development and product manufacturing including controlled release dissolution rate and bioavailability enhancement and granulation technology It brings together the technical information necessary to develop and market pharmaceutical dosage forms that meet current quality and regulatory requirements and details extruder hardware and controls process definition and troubleshooting of single and twin screw extrusion processes and more

Controlled Release in Oral Drug Delivery Clive G. Wilson, Patrick J. Crowley, 2011-09-22 Controlled Release in Oral Drug Delivery provides focus on specific topics complementing other books in the initial CRS series Each chapter sets the context for the inventions described and describe the latitude that the inventions allow In order to provide some similar look to each chapter the coverage includes the historical overview candidate drugs factors influencing design and development formulation and manufacturing and delivery system design This volume was written along three main sections the relevant anatomy and physiology a discussion on candidates for oral drug delivery and the major three groups of controlled release systems diffusion control swelling and inert matrices environmental control pH sensitive coatings time control enzymatic control pressure control and finally lipidic systems **Drug Delivery** Anya Hillery, 2016-09-15 This book provides a comprehensive introduction to advanced drug delivery and targeting covering their principles current applications and potential future developments This edition has been updated to reflect significant trends and cutting edge advances that have occurred since the first edition was published All the original chapters have been retained but the material therein has been updated Eight new chapters have been added that deal with entirely new technologies and approaches Features Offers a comprehensive introduction to the fundamental concepts and underlying scientific principles of drug delivery and targeting

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Matrix- and Reservoir-type Oral Multiparticulate Drug Delivery Systems Joana Mota,2010 Gibaldi's Drug Delivery Systems in Pharmaceutical Care Mary Lee,Archana Desai,2007 Tying together concepts of traditional pharmaceuticals in a way this text focuses on the selection of appropriate dosage forms as an integral part of drug therapy Pharmaceutical Inhalation Aerosol Technology, Second Edition Anthony J. Hickey,2003-09-03 This thoroughly revised and expanded reference provides authoritative discussions on the physiologic pharmacologic metabolic molecular cellular and physicochemical factors influencing the efficacy and utilization of pharmaceutical aerosol It analyzes the latest science and developments in the generation administration and characterization of these compounds showcasing current clinical applications the efficiency and limitations of major aerosol products and emerging aerosol therapies impacting the field

Pharmaceutical Process Scale-Up Michael Levin,2001-12-12 Focusing on scientific and practical aspects of process scale up this resource details the theory and practice of transferring pharmaceutical processes from laboratory scale to the pilot plant and production scale It covers parenteral and nonparenteral liquids and semi solids products derived from biotechnology dry blending and powder handling

Containment in the Pharmaceutical Industry James P. Wood,2020-03-26 Delivering an encompassing overview of the factors varieties and applications determining product containment this concise reference provides authoritative information on containment processes It reviews the historical context definition evolution and application of containment technology analyzes a variety of containment techniques in new *Good Manufacturing Practices for Pharmaceuticals* D. Nally Joseph,2000-10-12 Highlighting key issues and differences among GMPs of Europe Canada and the WHO this reference examines US law and governmental policy affecting domestic and multinational pharmaceutical manufacturing The book recommend pragmatic ways to interpret and comply with FDA CGMP regulation and related criteria They focus on geographical redistribution of manufacturing facilities accommodation of a diversity of regulatory and statutory governance adaptation to disparate human resources and new growth areas of manufacture and distribution of homeopathic remedies and dietary supplements in addition to the greater quality control required of pharmacists and other authorized dispensers

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 Transport Processes in Pharmaceutical Systems Gordon L. Amidon, Ping I. Lee, Elizabeth M. Topp, 1999-11-24 This cutting edge reference clearly explains pharmaceutical transport phenomena demonstrating applications ranging from drug or nutrient uptake into vesicle or cell suspensions drug dissolution and absorption across biological membranes whole body kinetics and drug release from polymer reservoirs and matrices to heat and mass transport in freeze drying and hygroscopicity Focuses on practical applications of drug delivery from a physical and mechanistic perspective highlighting biological systems Written by more than 30 international authorities in the field *Transport Processes in Pharmaceutical Systems* discusses the crucial relationship between the transport process and thermodynamic factors analyzes the dynamics of diffusion at liquid liquid liquid solid and liquid cultured cell interfaces covers prodrug design for improving membrane transport addresses the effects of external stimuli in altering some natural and synthetic polymer matrices examines properties of hydrogels including synthesis swelling degree swelling kinetics permeability biocompatibility and biodegradability presents mass transfer of drugs and pharmacokinetics based on mass balance descriptions and more Containing over 1000 references and more than 1100 equations drawings photographs micrographs and tables *Transport Processes in Pharmaceutical Systems* is a must read resource for research pharmacists pharmaceutical scientists and chemists chemical engineers physical chemists and upper level undergraduate and graduate students in these disciplines *Affinity Capillary Electrophoresis in Pharmaceuticals and Biopharmaceutics* Reinhard H. H. Neubert, Hans-Hermann Rüttinger, 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 development Pharmaceutical Product Development Chilukuri Dakshina Murthy, Gangadhar Sunkara, David Young, 2007-02-12 During the last two decades the pharmaceutical industry has been under pressure to reduce development costs and the time needed to bring drugs to market in order to maximize return on investment and bring treatments to patients sooner To meet these ends pharmaceutical scientists working in the differing areas of pharmacy pharmaceutics and pharmaceutical science Microencapsulation Simon Benita, 2005-11-01 Presenting breakthrough research pertinent to scientists in a wide range of disciplines from medicine and biotechnology to cosmetics and pharmacy this Second Edition provides practical approaches to complex formulation problems encountered in the development of particulate delivery systems at the micro and nano size level Completely revised and expanded *Percutaneous Absorption* Robert L. Bronaugh, Howard I. Maibach, 1999-05-28 Since publication of the Second Edition in 1989 numerous innovations have occurred that affect the way scientists look at issues in the field of percutaneous absorption Focusing on recent advances as well as updating and expanding the scope of topics covered in the previous edition *Percutaneous Absorption* Third Edition

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