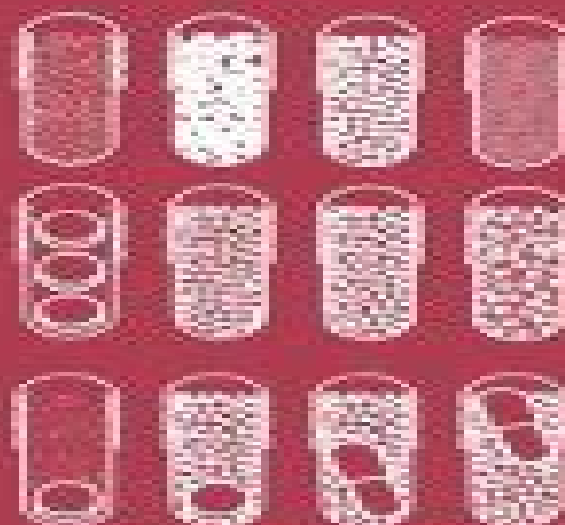


# Multiparticulate Oral Drug Delivery



edited by  
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# Multiparticulate Oral Drug Delivery

**Clive G. Wilson, Patrick J. Crowley**



## **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

*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

Matrix- and Reservoir-type Oral Multiparticulate Drug Delivery Systems Joana Mota, 2010 Drug Delivery

Any 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 Presents an in depth analysis of the opportunities and obstacles afforded by the application of nanotechnologies for drug delivery and targeting Includes a revised and expanded section on the major epithelial routes of drug delivery currently under investigation Describes the most recent emerging and innovative technologies of drug delivery Provides real life examples of the clinical translation of drug delivery technologies through the use of case studies Discusses the pertinent regulatory hurdles and safety issues of drug delivery and targeting systems crucial considerations in order to achieve licensing approval for these new technologies *The Design and Characterisation of Multiparticulate Lipidic Systems for Oral Drug Delivery* Gemma Coombs, 2013 Pharmaceutical Dosage Forms Larry L. Augsburger, Stephen W. Hoag, 2017-10-30 *Pharmaceutical Dosage Forms* Capsules covers the development composition and manufacture of capsules Despite the important role that capsules play in drug delivery and product development few comprehensive texts on the science and technology of capsules have been available for the research and academic environments This text addresses this gap discussing how capsules provide unique capabilities and options for dosage form design and formulation **Emulsions and Nanosuspensions for the Formulation of Poorly Soluble Drugs** Rainer H. Müller, Simon Benita, Bernhard H. L. Böhm, 1998 Explore possible new approaches for overcoming poorly soluble drugs a challenge to drug formulation work and an increasing problem Many newly developed drugs are poorly soluble very often simultaneously in aqueous and in organic media Emulsions and Nanosuspensions for the Formulation of Poorly Soluble Drugs aims to review the possibilities limitations and future perspectives of emulsions as drug carriers considering technology from other than the pharmaceutical industry i e food industry show the production technology of nanosuspensions explain the special dissolution properties i e increased saturation solubility and increased dissolution velocity theory and cover the possible applications present the theory of high pressure homogenization and high pressure extrusion in dispersion techniques including examples of applications and size measurements in concentrated dispersions *The ADME Encyclopedia* Alan Talevi, 2022-06-14 The ADME Encyclopedia covers pharmacokinetic phenomena Absorption Distribution Metabolism and Excretion processes and their relationship with the design of pharmaceutical carriers and the success of drug therapies It covers both basic and advanced knowledge serving as introductory material for students of biomedical careers and also as reference updated material for graduates and professionals working in any field related to pharmaceutical sciences medicine pharmaceutical technology materials science medicinal chemistry Structured as

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**in Drug Development** Rajesh Krishna, 2006-05 This reference provides a concise overview of the key principles in dose selection and optimization and demonstrates applicability to recent successful new drug applications Compiling key issues and current research on safety efficacy and clinical pharmacology and PK PD this volume critically highlights the multidisciplinary nature of drug development and spans the fields of pharmacokinetics clinical pharmacology biostatistics and experimental medicine **Good Laboratory Practice Regulations** Sandy Weinberg, 1995 This work covers all aspects of the Food and Drug Administration's Good Laboratory Practice regulations and techniques for implementation This edition includes general knowledge on computer system validation details on implementing GIPs in an automated laboratory a forecast of the flexibility and effectiveness of GLPs in the changing laboratory environment and a contemporary bibliography with new references

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**Good Manufacturing Practices for Pharmaceuticals** Sidney H. Willig, James R. Stoker, 1997 Revised to ensure GMP compliance this text examines US laws affecting domestic and multinational pharmaceutical manufacturing It recommends practical ways to interpret and comply with FDA cGMP regulations while meeting the goals of a comprehensive controls system to preserve product integrity

**Endotoxins** Kevin L. Williams, 2007-02-23 This source expertly examines the discovery biological structure control and continued clarification of endotoxin from a parenteral manufacturing

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**Handbook of Pharmaceutical Excipients** Raymond C. Rowe,Paul J. Sheskey,Paul J. Weller,2003 Describes the chemical and physical properties of pharmaceutical excipients Each monograph contains nonproprietary names synonyms chemical name and CAS registry number empirical formula and molecular weight structural formula functional category applications in pharmaceutical formulation or technology description pharmacopeial specifications typical properties stability and storage conditions incompatibilities method of manufacture safety handling precautions regulatory status pharmacopeias related substances comments specific references general references and authors Modern Pharmaceutics Alexander Taylor Florence,Jürgen Siepmann,2009 Modern Pharmaceutics examines the impact of pharmaceutical biotechnology cell therapy pharmacogenomics biotherapeutics and nanotechnology on current practice and the potential for personalized medicines and implications for pediatric and geriatric formulations Reflecting the shift away from physical pharmacy Modern Pharmaceutics is the must have current reference text for pharmaceutics and drug delivery

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