



SECOND EDITION

Developing Solid Oral Dosage Forms

Pharmaceutical Theory & Practice

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Lawrence Tu, and Rao K. Mestri



Developing Solid Oral Dosage Forms Pharmaceutical Theory Practice

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Zhang, Lirong Liu, William Porter**

Developing Solid Oral Dosage Forms Pharmaceutical Theory Practice:

Developing Solid Oral Dosage Forms Yihong Qiu, Yisheng Chen, Geoff G.Z. Zhang, Lirong Liu, William Porter, 2009-03-10 *Developing Solid Oral Dosage Forms* is intended for pharmaceutical professionals engaged in research and development of oral dosage forms. It covers essential principles of physical pharmacy, biopharmaceutics, and industrial pharmacy, as well as various aspects of state-of-the-art techniques and approaches in pharmaceutical sciences and technologies, along with examples and/or case studies in product development. The objective of this book is to offer updated or current knowledge and skills required for rational oral product design and development. The specific goals are to provide readers with Basics of modern theories of physical pharmacy, biopharmaceutics, and industrial pharmacy and their applications throughout the entire process of research and development of oral dosage forms. Tools and approaches of preformulation, investigation, formulation, process design, characterization, and scale-up in pharmaceutical sciences and technologies. New developments, challenges, trends, opportunities, intellectual property issues, and regulations in solid product development. The first book ever that provides comprehensive and in-depth coverage of what's required for developing high-quality pharmaceutical products to meet international standards. It covers a broad scope of topics that encompass the entire spectrum of solid dosage form development for the global market, including the most updated science and technologies, practice applications, regulation, intellectual property protection, and new development trends, with case studies in every chapter. A strong team of more than 50 well-established authors, co-authors of diverse background, knowledge, skills, and experience from industry, academia, and regulatory agencies.

Physico-Chemical Aspects of Dosage Forms and Biopharmaceutics Amit Kumar Nayak, Kalyan Kumar Sen, 2024-08-09 *Physico-Chemical Aspects of Dosage Forms and Biopharmaceutics: Recent and Future Trends in Pharmaceutics, Volume Two* explores aspects of pharmaceutics with an original approach that focuses on technology, novelties, and future trends. The field of pharmaceutics is highly dynamic and rapidly expanding day by day, so it demands a variety of amplified efforts for designing and developing pharmaceutical processes and formulation strategies. Readers will find practical information for conducting research in pharmaceutics that is ideal for researchers in academia and industry, as well as advanced graduate students in pharmaceutics. In addition, the book discusses the most recent developments in biopharmaceutics, including important and exciting areas such as solubility of drugs, pharmaceutical granulation, routes of drug administration, drug absorption, bioavailability, and bioequivalence. Provides extensive details on the most recent developments in biopharmaceutics. Contains contributions from leading experts from academia, research, industry, and regulatory agencies. Includes high-quality illustrations, flow charts, and tables for easier understanding of the concepts. Discusses practical examples and research case studies.

How to Develop Robust Solid Oral Dosage Forms Bhavishya Mittal, 2016-10-05 *How to Develop Robust Solid Oral Dosage Forms from Conception to Post Approval* uses a practical and hands-on approach to cover the development process of solid oral dosage forms in one single

source The book details all of the necessary steps from formulation through the post approval phase and contains industry case studies real world advice and troubleshooting tips By merging the latest scientific information with practical instructions this book provides pharmaceutical scientists in formulation research and development with a concrete look at the key aspects in the development of solid oral dosage forms Focuses on important topics such as robustness bioavailability formulation design continuous processing stability tests modified release dosage forms international guidelines process scale up and much more Part of the Expertise in Pharmaceutical Process Technology series edited by Michael Levin Discusses common real world problems and offers both theoretical and practical solutions to these everyday issues

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

ADME Processes in Pharmaceutical Sciences Alan Talevi, Pablo A. M. Quiroga, 2018-11-30 Absorption Distribution Metabolism and Excretion ADME processes and their relationship with the design of dosage forms and the success of pharmacotherapy form the basis of this upper level undergraduate graduate textbook As an introduction oriented to pharmacy students it is also written for scientist from different fields outside of pharmaceuticals e.g. material scientist material engineers medicinal chemists who

might be working in a positions in pharmaceutical companies or whose work might benefit from basic training in the ADME concepts and some biological background Pedagogical features such as objectives keywords discussion questions summaries and case studies add valuable teaching tools This book will provide not only general knowledge on ADME processes but also an updated insight on some hot topics such as drug transporters multi drug resistance related to pharmacokinetic phenomena last generation pharmaceutical carriers nanopharmaceuticals in vitro and in vivo bioequivalence studies biopharmaceuticals pharmacogenomics drug drug and food drug interactions and in silico and in vitro prediction of ADME properties In comparison with other similar textbooks around half of the volume would be focused on the relationship between expanding scientific fields and ADME processes Each of these burgeoning fields has a separate chapter in the second part of the volume and was written with leading experts on the correspondent topic including scientists and academics from USA and UK Duquesne University School of Pharmacy Indiana University School of Medicine University of Utah College of Pharmacy University of Maryland University of Bath Additionally each of the initial chapters dealing with the generalities of drug absorption distribution metabolism and excretion would include relevant classic examples related to each topic with appropriate illustrations e g importance of active absorption of levodopa implications in levodopa administration drug drug interactions and food drug interactions emerging from the active uptake intoxication with paracetamol as a result of glutathione depletion CYP induction and its relationship with acute liver failure caused by paracetamol etc ADME Processes and Pharmaceutical Sciences is written as a core textbook for ADME processes pharmacy pharmacokinetics drug delivery biopharmaceutics drug disposition drug design and medicinal chemistry courses *Aulton's Pharmaceutics* Michael E. Aulton, Kevin Taylor, 2013 Pharmaceutics is the art of pharmaceutical preparations It encompasses design of drugs their manufacture and the elimination of micro organisms from the products This book encompasses all of these areas Provided by publisher **Pharmaceutical Dosage Forms and Drug Delivery** Ram I. Mahato, Ajit S. Narang, Virender Kumar, 2024-11-22 Completely revised and updated this fourth edition elucidates the principles of pharmaceutics biopharmaceutics dosage form design and drug delivery including emerging new biotechnology based treatment modalities The authors integrate aspects of physical pharmacy chemistry biology and biopharmaceutics into drug delivery With the expiration of older patents and generic competition the biopharmaceutical industry is evolving faster than ever Consequently this edition of the book emphasizes the heightened focus that the recent remarkable progress in gene editing immunotherapy and nanotechnology has brought to the design of new drugs and diagnostic approaches along with novel dosage forms Apart from new chapters this edition highlights the emerging emphasis on the role of artificial intelligence AI in drug discovery mRNA and antibody based therapies genome editing immunotherapy chemical kinetics and the stability of drug products Features Includes new chapters on antibody therapeutics gene editing and immunotherapy Explains newer approaches and future methods and the significance of artificial intelligence AI in drug discovery Updated sections on pharmacy mathematics

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