

# Advanced Pharmaceutical Solids



Jens T. Carstensen

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**Rajesh Krishna**



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operations involved in the manufacturing process of solid and liquid dosage forms along with the scale up of each operation This book is a valuable resource for professionals working in the pharmaceutical industry and researchers seeking to develop a comprehensive understanding of the various aspects of the manufacturing process The book is divided into four sections covering a range of topics Section I provide readers with a comprehensive understanding of the basic principles behind the manufacturing process of solid and liquid dosage forms Section II covers the different unit operations involved in the production of solid dosage forms including mixing granulation drying compression coating and size reduction This section includes case studies to provide readers with practical insights into the scale up principles involved in the manufacturing process Section III focuses on the manufacturing and scale up of liquid formulations covering topics such as mixing filtration and scale up of liquid mixing process This section offers a comprehensive understanding of the various aspects of the manufacturing process including the challenges and opportunities associated with the scale up of liquid formulations Finally Section IV includes two chapters that describe the manufacturing and scale up of advanced drug delivery systems including the manufacturing and scale up of nanoparticles and biotechnology derived products This section provides readers with insights into the development of innovative drug delivery systems and the challenges involved in their scale up Overall the book is an essential guide for professionals and researchers seeking a deeper understanding of the manufacturing process The case studies and practical examples offer valuable insights into the challenges and opportunities involved in the scale up process making it an indispensable resource for those involved in the pharmaceutical industry Only book that is dedicated to pharmaceutical process engineering and scale up Contain numerous case studies for easy reference Covers solid liquid and advanced dosage forms

**Modern Pharmaceuticals, Two Volume Set** Alexander T. Florence, Juergen Siepmann, 2016-04-19 This new edition brings you up to date on the role of pharmaceuticals and its future paradigms in the design of medicines Contributions from over 30 international thought leaders cover the core disciplines of pharmaceuticals and the impact of biotechnology gene therapy and cell therapy on current findings Modern Pharmaceuticals helps you stay current

Generic Drug Product Development Leon Shargel, Isadore Kanfer, 2013-10-24 In this era of increased pharmaceutical industry competition success for generic drug companies is dependent on their ability to manufacture therapeutic equivalent drug products in an economical and timely manner while also being cognizant of patent infringement and other legal and regulatory concerns Generic Drug Product Development Solid Oral

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**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.

*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 perspective with in depth discussion of state of the art technologies involving Limulus amoebocyte lysate (LAL) such as assay development, automation, and depyrogenation. Completely revised and expanded, this Third Edition contains the knowledge necessary to apply endotoxin testing in the increasingly complex pharmaceutical environment, featuring sections detailing the latest information regarding clinical advances, regulation standards, and validation procedures for computerized kinetic tests.

**Polysaccharides for Drug Delivery and Pharmaceutical Applications** R. H. Marchessault, François Ravenelle, American Chemical Society. Meeting, 2006. Cellulose and starch are dry binders for tablets. They are essential for drug formulation and for slow release to maintain drug concentration in the blood. Unusual excipient uses are described such as protein separation and purification, gene delivery, and iron content in human physiology.

Polymorphism in Pharmaceutical Solids, Second Edition H. G. Brittain, 2009-07-27. Using clear and practical examples, this volume presents a comprehensive examination of polymorphic behavior in pharmaceutical development. The book is ideal for pharmaceutical development scientists and graduate students in pharmaceutical science. This second edition focuses on the pharmaceutical aspects of polymorphism and solvatomorphism while systematically explaining their thermodynamic and structural origins.

Topics in this edition include computational methodologies high throughput screening methods pharmaceutical cocrystals and thermally induced and solution mediated phase transformations The book also examines amorphous solids the amorphous state and crystallography preparation and characterization

**Dose Optimization 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

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