

DRUGS AND THE PHARMACEUTICAL SCIENCES

VOLUME 197

PHARMACEUTICAL POWDER COMPACTION TECHNOLOGY

SECOND EDITION



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Sidney H. Willig, James R. Stoker



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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 is 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

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Pharmaceutical Technology James Swarbrick,2013-07-01 Presenting authoritative and engaging articles on all aspects of drug development dosage manufacturing and regulation this Third Edition enables the pharmaceutical specialist and novice alike to keep abreast of developments in this rapidly evolving and highly competitive field A dependable reference tool and constant companion for years to com *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 **Aulton's Pharmaceutics E-Book** Kevin M.G. Taylor, Michael E. Aulton,2021-04-23 The essential pharmaceutics textbook One of the world s best known texts on pharmaceutics Aulton s Pharmaceutics offers a complete course in one book for students in all years of undergraduate pharmacy and pharmaceutical sciences degrees Thoroughly revised updated and extended by experts in their fields and edited by Professors Kevin Taylor and Michael Aulton this new edition includes the science of formulation pharmaceutical manufacturing and drug delivery All aspects of pharmaceutics are covered in a clear and readily accessible way and extensively illustrated throughout providing an essential companion to the entire pharmaceutics curriculum from day one until the end of the course Fully updated throughout with the addition of new chapters to reflect advances in formulation and drug delivery science pharmaceutical manufacturing and medicines regulation Designed and written for newcomers to the design and manufacture of dosage forms Relevant pharmaceutical science covered throughout Includes the science of formulation and drug delivery Reflects current practices and future applications of formulation and drug delivery science to small drug molecules biotechnology products and nanomedicines Key points boxes throughout Over 400 online multiple choice questions **Novel Belt Compaction Technology for Pharmaceutical Powder Dry Granulation Processing** Stephanie Casilagan,2023 **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 **Parts 1-5** ,2005 Process Modeling in Powder

Metallurgy & Particulate Materials, 2002

Pharmaceutical Statistics Practical And Clinical Applications, Third

Edition Bolton Sanford, 1997-01-17 This edition offers new and expanded information on recent developments in stability data analysis concepts of statistical outliers bioequivalence studies problems in sampling and devising limits for product release covariance analysis and tolerance intervals multiple endpoints and clinical data analysis and more student price which is available upon request from Marcel Dekker *Freeze-drying/lyophilization of Pharmaceutical and Biological Products* Louis Rey, Joan Christine May, 1999 Highlights the application of freeze drying to pharmaceuticals illustrating practical industry tested methods of preserving reactivating delicate biologicals biochemicals Discusses the basic principles engineering aspects of lyophilization also the role of bulking agents additives cryoprotectants antioxidants free radicals other products that protect the biological integrity of active substances during freezing drying storage **Agglomeration**

Processes Wolfgang Pietsch, 2002 Agglomeration is integral to the processes of modification of powders production of composites and creation of new materials which are required in pharmaceuticals foods chemicals fertilizers and agrochemicals minerals ceramics metallurgy and all material producing industries The binding mechanisms and the particle behavior as well as the characteristics of the processes and the resulting agglomerates are the same whether they are occurring in the ultra clean pharmaceutical or food industries or in dirty minerals or waste processing plants The book introduces the interdisciplinary approach to the development of new concepts and the solution of problems It is a complete and up to date practical guide describing the various agglomeration phenomena and industrial techniques for size enlargement In addition to introducing the properties of agglomerates and the characteristics of the different methods descriptions of the machinery and discussions of specific equipment features are the main topics The detailed evaluation of the subject is based on the authors experience as student researcher teacher developer designer vendor and user as well as expert and consultant in the field of agglomeration its technologies and products and is complemented by the know how of colleagues who are active in specific areas and information from vendors It is intended for everybody working in industries that process and handle particulate solids as it aims to help understand and control unwanted agglomeration as well as use improve and develop methods for the beneficial size enlargement by agglomeration **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

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 *Biosystems Engineering* ,2009 **Modern Pharmaceutics, Fourth Edition**
Revised and Expanded Gilbert S. Banker, Christopher T. Rhodes, 1996-06-15 An up to date sequenced approach to drug
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