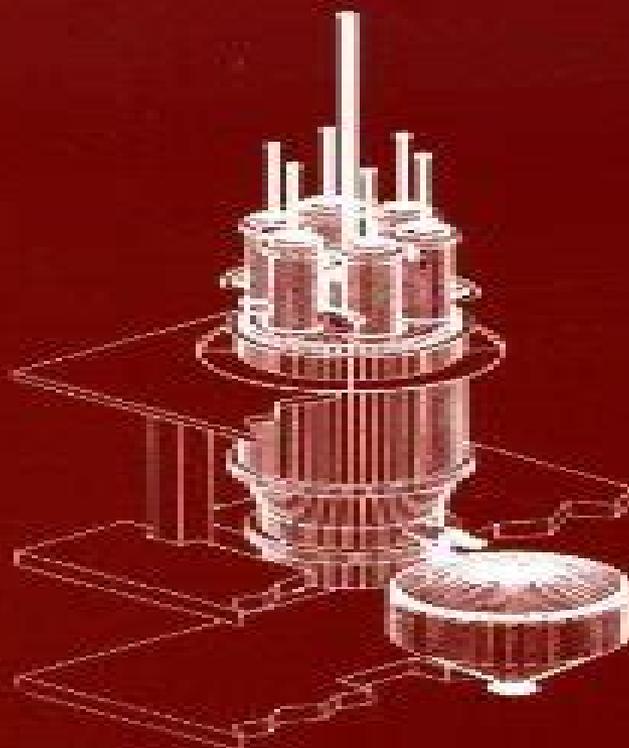


Handbook of Pharmaceutical Granulation Technology



edited by
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Handbook Of Pharmaceutical Granulation Technology

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Handbook of Pharmaceutical Granulation Technology Dilip M. Parikh, 2021-05-11 This fully revised edition of Handbook of Pharmaceutical Granulation Technology covers the rapid advances in the science of agglomeration process control process modelling scale up emerging particle engineering technologies along with current regulatory changes presented by some of the prominent scientist and subject matter experts around the globe Learn from more than 50 global subject matter experts who share their years of experience in areas ranging from drug delivery and pharmaceutical technology to advances in nanotechnology Every pharmaceutical scientist should own a copy of this fourth edition resource Key Features Theoretical discussions covering granulation and engineering perspectives Covers new advances in expert systems process modelling and bioavailability Chapters on emerging technologies in particle engineering Updated Current research and developments in granulation technologies

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well as academics regulatory officials and equipment vendors who provide PAT tools and granulation equipment

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 Handbook of Encapsulation and Controlled Release Munmaya Mishra, 2015-12-01 The field of encapsulation especially microencapsulation is a rapidly growing area of research and product development The Handbook of Encapsulation and Controlled Release covers the entire field presenting the fundamental processes involved and exploring how to use those processes for different applications in industry Written at a level comp **Water-Insoluble Drug Formulation** Ron Liu, 2018-03-12 Properties and Formulation From Theory to Real World Application Scientists have attributed more than 40 percent of the failures in new drug development to poor biopharmaceutical properties particularly water insolubility Issues surrounding water insolubility can postpone or completely derail important new drug development Even the much needed reformulation of currently marketed products can be significantly affected by these challenges More recently it was reported that the percentage increased to 90% for the candidates of new chemical entities in the discovery stage and 75% for compounds under development In the most comprehensive resource on the topic this third edition of Water Insoluble Drug Formulation brings together a distinguished team of experts to provide the scientific background and step by step guidance needed to deal with solubility issues in drug development Twenty three chapters systematically describe the detailed discussion on solubility theories solubility prediction models the aspects of preformulation biopharmaceutics pharmacokinetics regulatory and discovery support of water insoluble drugs to various techniques used in developing delivery systems for water insoluble drugs This book includes more than 15 water insoluble drug delivery systems or technologies illustrated with case studies and featuring oral and parenteral applications Highlighting the most current information and data available this seminal volume reflects the significant progress that has been made in nearly all aspects of this field The aim of this book is to provide a handy reference for pharmaceutical scientists in the handling of formulation issues related to water insoluble drugs In addition this book may be useful to pharmacy and chemistry undergraduate students and pharmaceutical and biopharmaceutical graduate students to enhance their knowledge in the techniques of drug solubilization and dissolution enhancement **Handbook of Drug Screening, Second Edition** Ramakrishna Seethala, Litao Zhang, 2009-06-24 A presentation of screening techniques modern technologies and high capacity instrumentation for increased productivity in the development and discovery of new drugs

chemical compounds and targeted delivery of pharmaceuticals It contains practical applications and examples of strategies in cell based and cell free screens as well as homogeneous fluorescence chemiluminescence and radioactive based technologies

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

Ansel's Pharmaceutical Dosage Forms and Drug Delivery Systems Loyd V. Allen, Nicholas G. Popovich, Howard C. Ansel, 2011 Thanks to its comprehensive coverage clear explanations and logical organization Ansel's Pharmaceutical Dosage Forms and Drug Delivery Systems has been a core pharmaceuticals text in the pharmacy curriculum for more than 40 years As you progress through this thoroughly updated Ninth Edition you'll master all the principles practices and technologies essential for the preparation of pharmaceutical dosage forms and drug delivery systems The text's integrated approach will help you understand the interrelationships among pharmaceutical and biopharmaceutical principles product design formulation manufacturing compounding and the clinical application of dosage forms for effective patient care Book Jacket

Guidelines for Safe Handling of Powders and Bulk Solids Stanley S. Grossel, 2005 Powders and bulk solids handled widely in the chemical pharmaceutical agriculture smelting and other industries present unique fire explosion and toxicity hazards Indeed substances which are practically inert in consolidated form may become quite hazardous when converted to powders and granules The U S Chemical Safety and Hazard Investigation Board is currently investigating dust explosions that occurred in 2003 at WestPharma CTA Acoustics and Hayes Lemmerz and is likely to recommend that companies that handle powders or whose operations produce dust pay more attention to understanding the hazards that may exist at their facility This new CCPS guidelines book will discuss the types of hazards that can occur in a wide range of process equipment and with a wide range of substances and will present measures to address these hazards

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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

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

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

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

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

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 amebocyte lysate LAL such as assay development automation 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

Effects of Solid-state Structure on the Tableting Properties of Pharmaceuticals Yushi Feng,2006

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