

Drug Delivery to the Oral Cavity Molecules to Market



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Tapash K. Ghosh
William R. Pfister

Drug Delivery To The Oral Cavity Molecules To Market

**Donald C. Singer, Raluca-Ioana
Stefan, Jacobus F. van Staden**



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Drug Delivery to the Oral Cavity Tapash K. Ghosh, William R. Pfister, 2005-02-28 With contributions from recognized authorities in industry academia and government this reference presents the state of the art in the testing formulation and clinical evaluation of intraoral drug delivery products summarizing intraoral dosage forms in various stages of research as well as products currently on the market

Modified-Release Drug Delivery Technology Michael Rathbone, Jonathan Hadgraft, Michael S. Roberts, Majella E. Lane, 2008-05-28 This two volume Second Edition describes the anatomical physiological pharmaceutical and technological aspects of delivery routes found in areas like Oral Ocular Dermal and transdermal Vaginal Colonic Oral mucosal Nasal Pulmonary Providing insight and critical assessment of the many available and emerging modified release drug delivery systems for

Oral Mucosal Drug Delivery and Therapy Michael J. Rathbone, Sevda Senel, Indiran Pather, 2015-03-13 This volume provides a comprehensive overview of the current issues facing scientists working on delivering drugs locally and systemically via the membranes that line the mouth The book describes the anatomical and physiological challenges of this route for drug delivery and how they impact the design of oral mucosal drug delivery systems It also provides a detailed description of current oral mucosal drug delivery technologies that overcome these challenges alongside research development and assessment methods In 11 authoritative chapters the book affords an in depth evaluation of the major issues associated with this route of administration namely the retention of the drug product at the site of administration and increasing drug permeability through the oral mucosa The book provides insights into the in vitro and in vivo methods available to assess drug permeability and retention offers solutions on how to improve the permeation of the drugs through the oral mucosa and explores approaches to prolong drug product retention at the site of administration It also indicates future directions in research and product development Oral Mucosal Drug Delivery and Therapy is a key resource for those wishing to extend their knowledge of this field

Environmental Monitoring for Cleanrooms and Controlled Environments Anne Marie Dixon, 2016-04-19 A critical technology in the science of contamination control environmental monitoring is a technique that provides important data on the quality of a process processing environment and final product which can aid scientists in identifying and eliminating potential sources of contamination in cleanrooms and controlled environments In response

Microbial Contamination Control in the Pharmaceutical Industry Luis Jimenez, 2004-08-30 This authoritative reference presents an up to date review of the testing methods emerging technologies and analytical systems and procedures used to prevent the microbial contamination of pharmaceutical processes products and environments It identifies new tools for sample analysis and evaluation and the impact of these advancements on the co

Novel Drug Delivery Systems Mehrdad Hamidi, Mahsa Sayed Tabatabaei, 2025-01-29 This book serves as a unique resource on the field of novel drug delivery systems NDDSs catering to both academic audiences researchers teachers students and industry professionals pharmaceutical policymakers managers R D regulatory business

development and marketing involved in NDDSs The evolving nature of diseases and the emergence of new health complications have driven a significant shift in drug therapy Coupled with changes in human lifestyles and economic conditions these factors have compelled the pharmaceutical industry to develop novel efficient and affordable drug products The rise of promising technologies aimed at enhancing therapeutic performance further underscores the importance of novel drug delivery systems NDDSs making this field one of the most dynamic in pharmaceutical sciences today This book offers a comprehensive exploration of NDDS applications catering to both academic and industry professionals It is designed to be both user friendly and thorough meeting the needs of diverse readers from those seeking practical insights to those delving deeply into the field s various sectors The content has been meticulously compiled organized and analyzed by a team of experts to ensure accuracy relevance and currency The book provides up to date information suitable for a broad audience reflecting the wide scope of the NDDS field The first section covers the foundational principles of NDDS including their scientific and therapeutic basis as well as the epidemiological and economic trends driving their development Subsequent sections explore market oriented aspects such as global trends and projections Chapters 3 to 12 present a taxonomy of NDDS categorized by their routes of administration The book concludes with a forward looking report on the potential future directions of NDDS Written in clear concise language the book is accessible to non native English readers ensuring broad usability Original figures created by experts with scientific and graphic design expertise enhance the reading experience and aid in understanding complex concepts Audience A wide variety of professional readers from universities and other research institutions to health industries industrial pharmaceutical and related to the relevant high tech startups to health and pharma policymakers and managers will be among the audiences of this book

Drug Products for Clinical Trials Donald Monkhouse, Charles F. Carney, Jim Clark, Peter Brun, 2005-11-09 Addressing issues at the forefront of interest for the Clinical Trial Materials Professional CTMP this Second Edition highlights the most critical concepts related to the planning manufacturing packaging labeling distribution reconciliation and quality and regulatory control of clinical trial materials offering an authoritative selection of chapters on the current and evolving state of clinical supplies operations by esteemed researchers and consultants in industry

Aqueous Polymeric Coatings for Pharmaceutical Dosage Forms, Third Edition Linda A. Felton, James W. McGinity, 2008-01-09 Thoroughly updated and expanded this new Third Edition provides the latest information on dosage forms film defects and polymer characterization Written by renowned leaders in the field Aqueous Polymeric Coatings for Pharmaceutical Dosage Forms is easily the most comprehensive book available on the market today New to the Third Edition the interaction of drugs with functional polymers the influence of processing parameters on coating quality the stabilization of polymeric film coats plasticizers and their applications in pharmaceutical coatings adhesion of polymeric films to solid substrates basic properties of latex and pseudolatex colloidal dispersions Key topics included polymer interactions with drugs and excipients physical aging of polymeric films a complete overview and in depth analysis of recent

advances in the field which includes information on the latest equipment used to apply polymers to a pharmaceutical system illustrated examples explaining the appropriate steps to be taken in order to solve formulation processing and stability problems to achieve an optimized dosage form *Pharmaceutical Photostability and Stabilization Technology* Joseph T. Piechocki, Karl Thoma, 2006-09-18 Based on a training course developed by Dr Joseph T Piechocki and other experts in this field whose contributions appear in this book for two International Meetings on the Photostability of Drugs and Drug Products this text clarifies the guidelines set by the International Conference on Harmonization ICH and provides a comprehensive background *Filtration and Purification in the Biopharmaceutical Industry* Maik J. Jornitz, Maik W. Jornitz, Theodore H. Meltzer, 2007-11-28 Filtration and Purification in the Biopharmaceutical Industry First Edition greatly expands its focus with extensive new material on the critical role of purification and the significant advances in filtration science and technology This new edition provides state of the science information on all aspects of filtration and purification in *Dermal Absorption and Toxicity Assessment* Michael S. Roberts, 2007-12-14 The source Dermal Absorption and Toxicity Assessment supplies a state of the art overview of the dermal absorption process and is divided into six well organized sections Written by internationally recognized experts in the field this Second Edition is a complete revised and updated text covering the wide range of methods used to assess skin ab **Pharmacogenomics** Werner Kalow, Urs B. Meyer, Rachel F. Tyndale, 2005-09-02 Reflecting the shift from genetics to genomics in the pharmaceutical sphere this Second Edition traces the evolution of the science of pharmacogenetics and gathers research from the forefront of the field spanning the most influential breakthroughs in molecular diagnostics metabonomics proteomics bioinformatics disease mapping pharmacodynamic *Microencapsulation* Simon Benita, 2005-11-01 Presenting breakthrough research pertinent to scientists in a wide range of disciplines from medicine and biotechnology to cosmetics and pharmacy this Second Edition provides practical approaches to complex formulation problems encountered in the development of particulate delivery systems at the micro and nano size level Completely revised and e *Injectable Dispersed Systems* Diane J. Burgess, 2005-05-23 This authoritative guide will serve as the most current source on the design and manufacturing of parenteral dispersed systems showcasing the utility of dispersed systems in drug delivery drug targeting and pharmaceutical engineering *Laboratory Auditing for Quality and Regulatory Compliance* Donald C. Singer, Raluca-Ioana Stefan, Jacobus F. van Staden, 2005-07-25 Identifying current tools techniques and approaches for the evaluation of laboratory operations this reference reviews the latest regulatory standards and auditing practices to test laboratory safety quality and performance *Good Laboratory Practice Regulations* Sandy Weinberg, 2007-01-19 Recent changes in the interpretation and enforcement of 21 CFR Part 11 have shifted the focus of Good Laboratory Practice GLP regulations to concentrate on the acceptance of electronic signatures the archiving of data the security of electronic documents and the automation of laboratory procedures This all encompassing Fourth Edition addresse *Endotoxins* Kevin L. Williams, 2007-02-23 This

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