

Colonic Drug Absorption and Metabolism



120 min



150 min



180 min



210 min

edited by
Peter R. Bieck

Colonic Drug Absorption And Metabolism

**Alexander Taylor Florence, Jürgen
Siepmann**



Colonic Drug Absorption And Metabolism:

Colonic Drug Absorption and Metabolism Peter Bieck,1993-06-29 An integrated series of chapters on the anatomical and physiological basis of drug absorption and metabolism in the colon as an aid to achieving the optimum delivery of drugs to the colon Among the topics are colon targeted osmotic systems therapeutic peptides and proteins drug induced injuries

Oral Colon-Specific Drug Delivery David R. Friend,1992-07-27 Oral Colon Specific Drug Delivery covers approaches used to deliver a variety of drugs to the colon Anatomy and physiology of the gastrointestinal tract as it affects colonic drug delivery and pharmacokinetics are reviewed as well as drug absorption from the colon The book presents valuable information on a variety of topics including oral peptide protein delivery dextran based delivery systems glycoside glycosidase based delivery azo bond prodrugs hydroxypropyl methacrylamide copolymers for colonic delivery and matrices for colonic drug delivery Special emphasis is placed on delivery systems especially biochemical approaches to delivery such as the use of degradable polymers and both low and high molecular weight prodrugs Oral Colon Specific Drug Delivery will provide a valuable reference resource for gastroenterologists pharmaceutical scientists and other researchers working with drug delivery to the colon

Applications of Polymers in Drug Delivery Ambikanandan Misra, Aliasgar Shahiwala,2020-10-02 Applications of Polymers in Drug Delivery Second Edition provides a comprehensive resource for anyone looking to understand how polymeric materials can be applied to current new and emerging drug delivery applications Polymers play a crucial role in modulating drug delivery and have been fundamental in the successful development of many novel drug delivery systems This book describes the development of polymeric systems ranging from conventional dosage forms to the most recent smart systems Regulatory and intellectual property aspects as well as the clinical applicability of polymeric drug delivery systems are also discussed The chapters are organized by specific delivery route offering methodical and detailed coverage throughout This second edition has been thoroughly revised to include the latest developments in the field This is an essential book for researchers scientists and advanced students in polymer science drug delivery pharmacology pharmaceuticals materials science tissue engineering nanomedicine chemistry and biology In industry this book supports scientists R D and other professionals working on polymers for drug delivery applications Explains how polymers can be prepared and utilized for all major drug delivery routes Presents the latest advances including drug targeting polymeric micelles and polymersomes and the delivery of biologicals and nucleic acid therapeutics Includes appendices with in depth information on pharmaceutical properties of polymers and regulatory aspects

Oral Controlled Release Formulation Design and Drug Delivery Hong Wen, Kinam Park,2010-10-19 This book describes the theories applications and challenges for different oral controlled release formulations This book differs from most in its focus on oral controlled release formulation design and process development It also covers the related areas like preformulation biopharmaceutics in vitro in vivo correlations IVIVC quality by design QbD and regulatory issues

Oral Drug Delivery for Modified Release

Formulations Edmund S. Kostewicz, Maria Vertzoni, Heather A. E. Benson, Michael S. Roberts, 2022-04-26 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 pharmaceutics pharmaceutical sciences biomedical engineering polymer and materials science and chemical and biochemical engineering A Biopharmaceutics Design Model for Oral Delivery Xuanqiang Yu, 1996 *Developing Solid Oral Dosage Forms* Yihong Qiu, Yisheng Chen, Geoff G.Z. Zhang, Lawrence Yu, Rao V. Mantri, 2016-11-08 *Developing Solid Oral Dosage Forms* Pharmaceutical Theory and Practice Second Edition illustrates how to develop high quality safe and effective pharmaceutical products by discussing the latest techniques tools and scientific advances in preformulation investigation formulation process design characterization scale up and production operations This book covers the essential principles of physical pharmacy biopharmaceutics and industrial pharmacy and their application to the research and development process of oral dosage forms Chapters have been added combined deleted and completely revised as necessary to produce a comprehensive well organized valuable reference for industry professionals and academics engaged in all aspects of the development process New and important topics include spray drying amorphous solid dispersion using hot melt extrusion modeling and simulation bioequivalence of complex modified released dosage forms biowaivers and much more Written and edited by an international team of leading experts with experience and knowledge across industry academia and regulatory settings Includes new chapters covering the pharmaceutical applications of surface

phenomenon predictive biopharmaceutics and pharmacokinetics the development of formulations for drug discovery support and much more Presents new case studies throughout and a section completely devoted to regulatory aspects including global product regulation and international perspectives Novel Drug Delivery and Its Therapeutic Application L. F. Prescott, W. S. Nimmo, 1989 International research specialists discuss their work with pharmaceuticals in this text focusing on the mechanisms and assessment of drug absorption and delivery The book also explores the ways in which a drug should be administered to provide self regulating and programmed delivery **Handbook of Biodegradable Polymeric Materials and Their Applications** Surya Mallapragada, Balaji Narasimhan, 2006 *Mechanistic Studies on the Enhanced Mucosal Transport of Human Growth Hormone by Certain Amino Acid Derivatives* Sy-Juen Wu, 1999 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 **Modern Pharmaceutics, Fourth Edition Revised and Expanded** Gilbert S. Banker, Christopher T. Rhodes, 1996-06-15 An up to date sequenced approach to drug dosage formulation design and evaluation This edition offers new chapters on regulatory aspects of the pharmaceutical industry in the European Union the pharmaceutical needs of special populations target oriented drug delivery systems and more *Good Laboratory Practice Regulations* Sandy Weinberg, 1995 This work covers all aspects of the Food and Drug Administration's Good Laboratory Practice regulations and techniques for implementation This edition includes general knowledge on computer system validation details on implementing GIPs in an automated laboratory a forecast of the flexibility and effectiveness of GLPs in the changing laboratory environment and a contemporary bibliography with new references **Pharmaceutical Process Validation, Second Edition** Ira R. Berry, Robert A. Nash, 1993-01-29 The second edition of this text has been updated and enlarged to reflect current good manufacturing practice CGMP regulations and the increased interest in and applicability of process validation Pharmaceutical Process Validation offers up to the minute coverage of regulations and validation sterile process validation organization in validation processes solid dosage forms validation raw material validation analytical methods validation and prospective and retrospective validation Providing the contributions of leading experts in the field the text also supplies examinations of current concepts in validation and new topics such as validation of cleaning systems and computer systems equipment and water systems validation and lyophilized and aerosol product validation *The Cumulative Book Index*, 1994 A world list of books in the English language **Scandinavian Journal of Gastroenterology**, 1995 **Biopharmaceutics and Clinical Pharmacokinetics** Milo Gibaldi, 1991 This updated introduction to the clinical applications of pharmacokinetics looks at gastrointestinal absorption prolonged release medication and drug disposition The effects of disease weight age sex and

genetic factors on pharmacokinetic variability and drug response are detailed Bioequivalence and regulatory considerations for generic drug

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 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 *Kirk-Othmer Encyclopedia of Chemical Technology, Composites Materials to Detergency* Raymond Eller Kirk, 1993-07-20 Contains a library of information for the chemical industry The 4th edition has undergone a complete revision with the inclusion of many new subjects which reflect the growth in chemical technology through the 1990s The book includes expanded coverage of biotechnology and materials science

Modern Pharmaceuticals Alexander Taylor Florence, Jürgen Siepmann, 2009 Modern Pharmaceuticals 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 Pharmaceuticals is the must have current reference text for pharmaceuticals and drug delivery

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