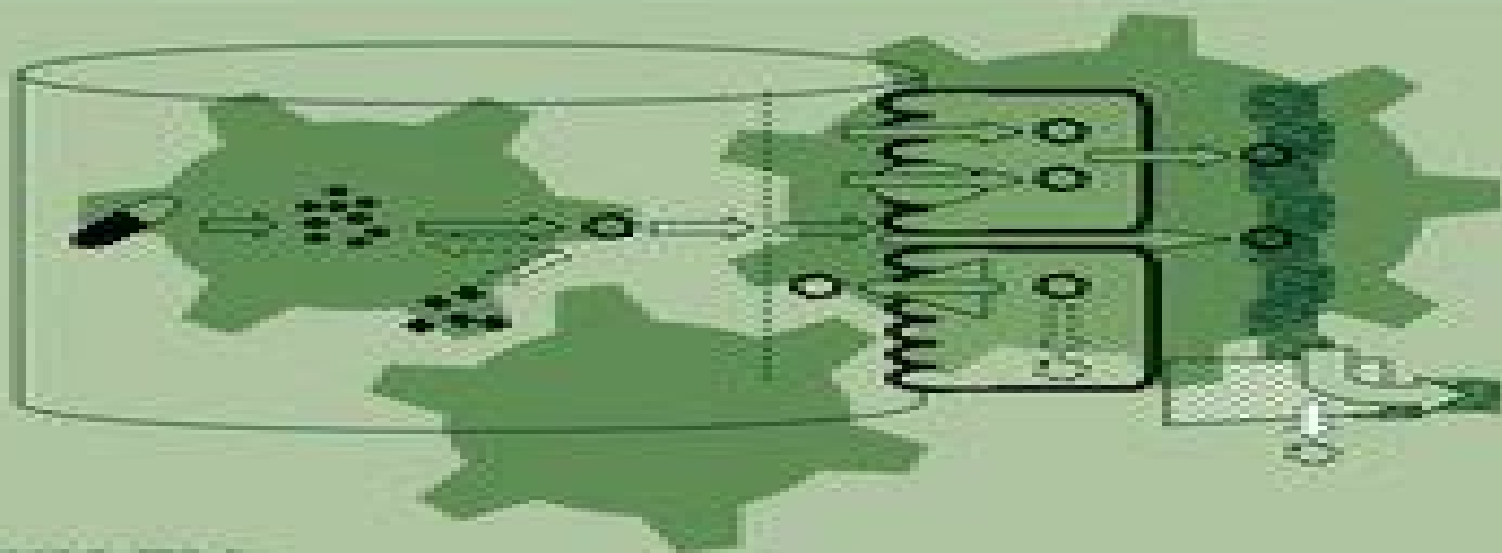


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Biopharmaceutics and Pharmacokinetics Considerations, 2021-07-07 Biopharmaceutics and Pharmacokinetics Considerations examines the history of biopharmaceutics and pharmacokinetics The book provides a biopharmaceutics and pharmacokinetics approach to addressing issues in formulation development and ethical considerations in handling animals Written by experts in the field this volume within the Advances in Pharmaceutical Product Development and Research series deepens understanding of biopharmaceutics and pharmacokinetics within drug discovery and drug development Each chapter delves into a particular aspect of this fundamental field to cover the principles methodologies and technologies employed by pharmaceutical scientists researchers and pharmaceutical industries to study the chemical and physical properties of drugs and the biological effects they produce Examines the most recent developments in biopharmaceutics and pharmacokinetics for pharmaceutical sciences Covers the principles methodologies and technologies of biopharmaceutics and pharmacokinetics Focuses on the pharmaceutical sciences but also encompasses aspects of toxicology neuroscience environmental sciences and nanotechnology

ADME Processes in Pharmaceutical Sciences Alan Talevi, Pablo A. M. Quiroga, 2018-11-30 Absorption Distribution Metabolism and Excretion ADME processes and their relationship with the design of dosage forms and the success of pharmacotherapy form the basis of this upper level undergraduate graduate textbook As an introduction oriented to pharmacy students it is also written for scientist from different fields outside of pharmaceutics e g material scientist material engineers medicinal chemists who might be working in a positions in pharmaceutical companies or whose work might benefit from basic training in the ADME concepts and some biological background Pedagogical features such as objectives keywords discussion questions summaries and case studies add valuable teaching tools This book will provide not only general knowledge on ADME processes but also an updated insight on some hot topics such as drug transporters multi drug resistance related to pharmacokinetic phenomena last generation pharmaceutical carriers nanopharmaceuticals in vitro and in vivo bioequivalence studies biopharmaceuticals pharmacogenomics drug drug and food drug interactions and in silico and in vitro prediction of ADME properties In comparison with other similar textbooks around half of the volume would be focused on the relationship between expanding scientific fields and ADME processes Each of these burgeoning fields has a separate chapter in the second part of the volume and was written with leading experts on the correspondent topic including scientists and academics from USA and UK

Duquesne University School of Pharmacy Indiana University School of Medicine University of Utah College of Pharmacy University of Maryland University of Bath Additionally each of the initial chapters dealing with the generalities of drug absorption distribution metabolism and excretion would include relevant classic examples related to each topic with appropriate illustrations e g importance of active absorption of levodopa implications in levodopa administration drug drug interactions and food drug interactions emerging from the active uptake intoxication with paracetamol as a result of glutathione depletion CYP induction and its relationship with acute liver failure caused by paracetamol etc ADME Processes and Pharmaceutical Sciences is written as a core textbook for ADME processes pharmacy pharmacokinetics drug delivery biopharmaceutics drug disposition drug design and medicinal chemistry courses

Revising Oral Pharmacokinetics, Bioavailability and Bioequivalence Based on the Finite Absorption Time Concept Panos Macheras, Athanasios A. Tsekouras, 2023-01-01 This book casts new light on the field of oral drug absorption It outlines both the concept of the past and the novel concept of Finite Absorption Time FAT In addition the authors explore the correlated need for re definition of bioavailability bioequivalence providing a plethora of experimental data Accordingly this book is intended for academics students or scientists working in pharmaceutical industries regulatory agencies and contract research organizations It can be used for teaching purposes in under and post graduate courses dealing with biopharmaceutics pharmacokinetics and biomedical engineering

Polymorphism in the Pharmaceutical Industry Rolf Hilfiker, Markus von Raumer, 2019-01-04 Polymorphism in the Pharmaceutical Industry Solid Form and Drug Development highlights the relevance of polymorphism in modern pharmaceutical chemistry with a focus on quality by design QbD concepts It covers all important issues by way of case studies ranging from properties and crystallization via thermodynamics analytics and theoretical modelling right up to patent issues As such the book underscores the importance of solid state chemistry within chemical and pharmaceutical development It emphasizes why solid state issues are important the approaches needed to avoid problems and the opportunities offered by solid state properties The authors include true polymorphs as well as solvates and hydrates while providing information on physicochemical properties crystallization thermodynamics quantum mechanical modelling and up scaling Important analytical tools to characterize solid state forms and to quantify mixtures are summarized and case studies on solid state development processes in industry are also provided Written by acknowledged experts in the field this is a high quality reference for researchers project managers and quality assurance managers in pharmaceutical agrochemical and fine chemical companies as well as for academics and newcomers to organic solid state chemistry

Molecular Dynamics of Nanostructures and Nanoionics Junko Habasaki, 2020-11-30 Nanostructured materials with multiple components and complex structures are the current focus of research and are expected to develop further for material designs in many applications in electrochemical colloidal medical pharmaceutical and several other fields This book discusses complex nanostructured systems exemplified by nanoporous silicates spontaneously formed gels from silica nanocolloidal solutions

and related systems and examines them using molecular dynamics simulations Nanoporous materials nanocolloidal systems and gels are useful in many applications and can be used in electric devices and storage and for gas ion and drug delivery The book gives an overview of the history current status and frontiers of the field It also discusses the fundamental aspects related to the common behaviors of some of these systems and common analytical methods to treat them *Predictive ADMET* Jianling Wang,Laszlo Urban,2014-02-28 This book helps readers integrate in silico in vitro and in vivo ADMET absorption distribution metabolism elimination and toxicity and PK pharmacokinetics data with routine testing applications so that pharmaceutical scientists can diagnose ADMET problems and present appropriate recommendations to move drug discovery programs forward The book introduces the current clinical practice for drug discovery and development along with the impact on early risk assessment consolidates the tools and models to intelligently integrate existing in silico in vitro and in vivo ADMET data and demonstrates successful cases and lessons learned from real drug discovery and development In short it is a book aimed to provide a practical road map for drug discovery and development scientists to generate efficacious and safe drugs for unmet medical needs **Pharmacokinetic-Pharmacodynamic Modeling and Simulation** Peter L. Bonate,2006-05-14 A natural hierarchy exists in pharmacokinetic pharmacodynamic modeling culminating in population pharmacokinetic models which are a specific type of nonlinear mixed effects model The purpose of this book is to present through theory and example how to develop pharmacokinetic models both at an individual and population level In order to do so however one must first understand linear models and then build to nonlinear models followed by linear mixed effects models and then ultimately nonlinear mixed effects models This book develops in that manner each chapter builds upon previous chapters by first presenting the theory and then illustrating the theory using published data sets and actual data sets that were used in the development of new chemical entities collected by the author during his years in industry A key feature of the book is the process of modeling Most books and manuscripts often present the final model never showing how the model evolved In this book all examples are presented in an evolutionary manner

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