

DISSOLUTION

- Dissolution – Mass transfer of solid substance from the solid surface to the liquid phase. It is a dynamic process
- Solubility – Maximum amount of solute dissolved in given solvent. It is a static (lack of movement) process
- Dissolution rate - rate at which the drug released from dosage form under standardized conditions of liquid-solid interface, temperature and media composition
- The actual drug release in the human body can be measured *in-vivo* by measuring the plasma or urine concentrations in the patient.
- Due to impracticalities involved in employing such techniques on a routine basis, leads to the introduction of official *in-vitro* dissolution tests and its principle function involves,
 - Optimization of therapeutic effectiveness during product development and stability assessment.
 - Routine assessment of production quality to ensure uniformity between production lots.
 - Assessment of 'bioequivalence'.
 - Prediction of *in-vivo* availability, i.e. bioavailability.
 - Active ingredient is released in the predicted way

Dissolution Techniques For Evaluation Of Novel Drug

**Murray P. Ducharme, Leon
Shargel, Andrew B. C. Yu**



Dissolution Techniques For Evaluation Of Novel Drug:

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

Regulatory Affairs in the Pharmaceutical Industry Javed Ali, Sanjula Baboota, 2021-11-14
Regulatory Affairs in the Pharmaceutical Industry is a comprehensive reference that compiles all the information available pertaining to regulatory procedures currently followed by the pharmaceutical industry Designed to impart advanced knowledge and skills required to learn the various concepts of regulatory affairs the content covers new drugs generic drugs and their development regulatory filings in different countries different phases of clinical trials and the submission of regulatory documents like IND Investigational New Drug NDA New Drug Application and ANDA Abbreviated New Drug Application Chapters cover documentation in the pharmaceutical industry generic drug development code of Federal Regulation CFR the ANDA regulatory approval process the process and documentation for US registration of foreign drugs the regulation of combination products and medical devices the CTD and ECTD formats and much more Updated reference on drug approval processes in key global markets Provides comprehensive coverage of concepts and regulatory affairs Presents a concise compilation of the regulatory requirements of different countries Introduces the fundamentals of manufacturing controls and their regulatory importance

Drug Delivery Systems, 2019-10-23
Drug Delivery Systems examines the current state of the field within pharmaceutical science and concisely explains the history of drug delivery systems including key developments The book translates the physicochemical properties of drugs into drug delivery systems administered via various routes such as oral parenteral transdermal and inhalational Regulatory and product development topics are also explored Written by experts in the field this volume in the Advances in Pharmaceutical

Product Development and Research series deepens our understanding of drug delivery systems within the pharmaceutical sciences industry and research as well as in chemical engineering Each chapter delves into a particular aspect of this fundamental field to cover the principles methodologies and technologies employed by pharmaceutical scientists This book provides a comprehensive examination that is suitable for researchers and advanced students working in pharmaceuticals cosmetics biotechnologies and related industries Provides up to date information on how to translate the physicochemical properties of drugs into drug delivery systems Explores how drugs are administered via various routes such as oral parenteral transdermal and inhalational Contains extensive references and further reading for course and self study Oral Bioavailability and Drug Delivery Ming Hu,Xiaoling Li,2024-02-13 ORAL BIOAVAILABILITY AND DRUG DELIVERY Improve the performance and viability of newly developed and approved drugs with this crucial guide Bioavailability is the parameter which measures the rate and extent to which a drug reaches a user s circulatory system depending on the method of administration For example intravenous administration produces a bioavailability of 100% since the drugs are injected directly into the circulatory system in the case of oral administration however bioavailability can vary widely based on factors which if not properly understood can result in a failure in drug development adverse effects and other complications The mechanics of oral bioavailability are therefore critical aspects of drug development Oral Bioavailability and Drug Delivery provides a comprehensive coverage of this subject as well as its drug development applications Beginning with basic terminology and fundamental concepts it provides a thorough understanding of the challenges and barriers to oral bioavailability as well as the possibilities for improving this parameter The resulting book is an indispensable tool for drug development research Oral Bioavailability and Drug Delivery readers will also find Discussion questions in many chapters to facilitate comprehension Detailed discussion of topics including dissolution absorption metabolism and more Real world examples of methods in actions throughout Oral Bioavailability and Drug Delivery is ideal for pharmaceutical and biotechnology scientists working in drug discovery and development researchers in chemistry biology pharmacology immunology neuroscience and other related fields and graduate courses in drug development and delivery

9789815223026 Dilpreet Singh, Prashant Tiwari,2024-03-25 Software and Programming Tools in Pharmaceutical Research is a detailed primer on the use for computer programs in the design and development of new drugs Chapters offer information about different programs and computational techniques in pharmacology The book will help readers to harness computer technologies in pharmaceutical investigations Readers will also appreciate the pivotal role that software applications and programming tools play in revolutionizing the pharmaceutical industry The book includes nine structured chapters each addressing a critical aspect of pharmaceutical research and software utilization From an introduction to pharmaceutical informatics and computational chemistry to advanced topics like molecular modeling data mining and high throughput screening this book covers a wide range of topics Key Features Practical Insights Presents practical knowledge

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Annual Reports of Shionogi Research Laboratory, 1996 **Remington** David B. Troy, Paul Beringer, 2006 For over 100 years Remington has been the definitive textbook and reference on the science and practice of pharmacy This Twenty First Edition keeps pace with recent changes in the pharmacy curriculum and professional pharmacy practice More than 95 new contributors and 5 new section editors provide fresh perspectives on the field New chapters include pharmacogenomics application of ethical principles to practice dilemmas technology and automation professional communication medication errors re engineering pharmacy practice management of special risk medicines specialization in pharmacy practice disease state management emergency patient care and wound care Purchasers of this textbook are entitled to a new fully indexed Bonus CD ROM affording instant access to the full content of Remington in a convenient and portable format

New Scientist, 2001 **Shargel and Yu's Applied Biopharmaceutics & Pharmacokinetics, 8th Edition** Murray P. Ducharme, Leon Shargel, Andrew B. C. Yu, 2022-01-24 The authoritative textbook on the principles and practical applications of biopharmaceutics and pharmacokinetics Shargel Yu s Applied Biopharmaceutics Pharmacokinetics has been the standard textbook in its field for over 40 years This eighth edition includes recent scientific developments in the field and embodies the collective contribution of experts with deep knowledge and experience in the selected subject areas Shargel Yu s Applied Biopharmaceutics Pharmacokinetics Eighth Edition provides the reader with a fundamental understanding of biopharmaceutics and pharmacokinetics principles that can be applied to patient drug therapy and rational drug product development Shargel Yu s Applied Biopharmaceutics Pharmacokinetics Eighth Edition has been expanded and revised to include advancements in biopharmaceutics and pharmacokinetics The chapter sequence has been reorganized into four main sections providing a more logical sequence for students The textbook starts with fundamental concepts followed by application of these principles to optimize drug therapy and to the rational development of drug products Each chapter includes theoretical concepts with practical examples and clinical applications Frequently asked questions provide a discussion of overall concepts Features Expanded and revised chapters to include scientific advances in biopharmaceutics and pharmacokinetics Four main sections providing a natural buildup of knowledge introduction to biopharmaceutics and pharmacokinetics fundamentals of biopharmaceutics pharmacokinetic calculations

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Drug Product Development Additional information on drug therapy drug product performance and other related topics
Frequently asked questions practice problems clinical examples and learning questions **The Development and Control
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by the U S Army Medical Library Library & Information Science Abstracts ,2003 British Medical Journal ,1930
Drug Intelligence & Clinical Pharmacy ,1972 *Applied Biopharmaceutics & Pharmacokinetics, Fifth Edition* Leon
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of concepts Basic theoretical discussions of the principles of biopharmaceutics and pharmacokinetics are provided along with
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Hybrids and Composites Vol. 12 Sooraj Hussain Nandyala, 2016-11-01 *Nanomaterials Advances in Biomedical and
Environmental Applications* Special topic volume with invited peer reviewed papers only Applied Biopharmaceutics &
Pharmacokinetics Leon Shargel, Andrew B. C. Yu, 1999 This work emphasizes the application and understanding of core areas
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