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Handbook of Dissolution Testing William A. Hanson, 1982 **Handbook of Bioequivalence Testing** Sarfaraz K. Niazi, 2014-10-29 As the generic pharmaceutical industry continues to grow and thrive so does the need to conduct adequate efficient bioequivalence studies In recent years there have been significant changes to the statistical models for evaluating bioequivalence In addition advances in the analytical technology used to detect drug and metabolite levels have m

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

Biopharmaceutics and Pharmacokinetics Dr.S.Valarmathi, Mrs. P. Parameswari, Mr. Obaidurrahman, Mr. Radheshyam Samanta, Mr. Pushpendu Gaurav, *Pharmaceutical Solid Dispersion Technology* Muhammad J. Habib, 2000-10-05 There has not until now been a single up to date volume to provide those in drug R D with practical information on all aspects of solid dispersion technology for drugs This forthcoming volume finally provides such a guide and reference The unified presentation by a team of specialists in this field is designed for practical application Theoretical concepts are covered for a fuller understanding of current techniques All significant recent developments are included

Physico-Chemical Aspects of Dosage Forms and Biopharmaceutics Amit Kumar Nayak, Kalyan Kumar Sen, 2024-08-09 Physico Chemical Aspects of Dosage Forms and Biopharmaceutics Recent and Future Trends in Pharmaceutics Volume Two explores aspects of pharmaceutics with an original approach that focuses on technology novelties and future trends The field of pharmaceutics is highly dynamic and rapidly expanding day by day so it demands a variety of amplified efforts for designing and developing pharmaceutical processes and formulation strategies Readers will find practical information for conducting research in pharmaceutics that is ideal for researchers in academia and industry as

well as advanced graduate students in pharmaceuticals. In addition, the book discusses the most recent developments in biopharmaceutics, including important and exciting areas such as solubility of drugs, pharmaceutical granulation, routes of drug administration, drug absorption, bioavailability, and bioequivalence. Provides extensive details on the most recent developments in biopharmaceutics. Contains contributions from leading experts from academia, research industry, and regulatory agencies. Includes high quality illustrations, flow charts, and tables for easier understanding of the concepts. Discusses practical examples and research case studies.

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.

The United States Pharmacopeia, 2012. *USP, NF.*, 2006. **Technical Report Series**, 1984. **Principles and Perspectives in Drug Bioavailability** James Blanchard, Ronald J. Sawchuk, Bernard B. Brodie, 1979. **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, clinical pharmacokinetics and pharmacodynamics, and biopharmaceutics and pharmacokinetics.

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