

DRUG DISPOSITION DURING DEVELOPMENT



Edited by PAOLO LUCIO MORSELLI



Drug Disposition During Development

Richard Caprioli



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Drug Disposition During Development Paolo Lucio Morselli, 1977 *Drug Disposition During Development* Paolo Lucio Morselli, 1977 **Neonatal and Pediatric Clinical Pharmacology, An Issue of Pediatric Clinics** John N van den Anker, Max J. Coppes, Gideon Koren, 2012-10-28 The need for safe and effective use of medicines in children and WHO's initiative Make Medicines Child Size have boosted research and educational activities in the area of pediatric clinical drug research. This issue focuses on both general and specific aspects of neonatal and pediatric clinical pharmacology including ethics, pharmacogenomics, metabolomics, adverse drug reactions, pain medication, pulmonary hypertension, and several other hot topics. The editors have been able to find outstanding authors for the different parts on neonatal and pediatric pharmacology. **ADME-Enabling Technologies in Drug Design and Development** Donglu Zhang, Sekhar Surapaneni, 2012-04-13 A comprehensive guide to cutting edge tools in ADME research. The last decade has seen tremendous progress in the development of analytical techniques such as mass spectrometry and molecular biology tools resulting in important advances in drug discovery particularly in the area of absorption, distribution, metabolism, and excretion (ADME). ADME Enabling Technologies in Drug Design and Development focuses on the current state of the art in the field, presenting a comprehensive review of the latest tools for generating ADME data in drug discovery. It examines the broadest possible range of available technologies, giving readers the information they need to choose the right tool for a given application, a key requisite for obtaining favorable results in a timely fashion for regulatory filings. With over thirty contributed chapters by an international team of experts, the book provides a thorough examination of current tools covering both electronic/mechanical technologies and biologically based ones. Coverage of applications for each technology including key parameters, optimal conditions for intended results, protocols, and case studies. Detailed discussion of emerging tools and techniques from stem cells and genetically modified animal models to imaging technologies. Numerous figures and diagrams throughout the text. Scientists and researchers in drug metabolism, pharmacology, medicinal chemistry, pharmaceuticals, toxicology, and bioanalytical science will find ADME Enabling Technologies in Drug Design and Development an invaluable guide to the entire drug development process from discovery to regulatory issues. Preclinical Drug Development Mark Rogge, David R. Taft, 2016-04-19 Preclinical Drug Development, Second Edition, discusses the broad and complicated realm of preclinical drug development. Topics range from assessment of pharmacology and toxicology to industry trends and regulatory expectations to requirements that support clinical trials. Highlights of the Second Edition include Pharmacokinetics Modeling and simulation, Formulation and routes of administration, Toxicity evaluations, The assessment of drug absorption and metabolism, Interspecies scaling, Lead molecule selection and optimization via profiling, Screening using in silico and in vitro toxicity evaluations. The book also includes case studies on preclinical pharmacokinetic/pharmacodynamic modeling and simulation in drug development, a review of ICH preclinical guidelines, and experimental methods used to study membrane drug transport.

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Environmental Factors in Human Growth and Development Vilma R. Hunt, M. Kate Smith, Dorothy Worth, 1982

Government Reports Announcements & Index, 1988

Clinical Chemistry Michael L. Bishop, Janet L. Duben-Engelkirk, Edward P. Fody, 1992

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Developmental Pharmacology and Therapeutics, 1990

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