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Prodrugs As Novel Drug Delivery Systems Acs Symposium Series 14

D. Robert Lu, Svein Øie



Prodrugs As Novel Drug Delivery Systems Acs Symposium Series 14:

Prodrugs Valentino Stella, Ronald Borchardt, Michael Hageman, Reza Oliyai, Hans Maag, Jefferson Tilley, 2007-08-26

Prodrugs are substances administered in an inactive form that is then metabolized in the body in vivo into the active compound. The rationale behind administering prodrugs is to optimize absorption, distribution, metabolism, and excretion of these drugs. Since first described in the 1950s, prodrugs continue to be a fertile area of research. There are a number of small pharmaceutical biotech companies dedicated to using prodrugs for the delivery of older but problematic drugs as well as to developing broad based prodrug technologies for application to new and future drugs. These volumes represent a comprehensive guide to prodrugs and will guide the reader through the current status of the prodrug concept and its many applications and to highlight its many successes in overcoming formulation and delivery of problematic drugs. **Pro-drugs**

as Novel Drug Delivery Systems Takeru Higuchi, Valentino J. Stella, American Chemical Society, Division of Medicinal Chemistry, 1975

Optimizing the "Drug-Like" Properties of Leads in Drug Discovery Ronald Borchardt, Edward Kerns, Michael Hageman, Dhrien Thakker, James Stevens, 2007-12-31. This book arises from a workshop organized by the American Association of Pharmaceutical Scientists entitled Optimizing the Drug Like Properties of Leads in Drug Discovery which took place in Parsippany NJ in September 2004. The workshop focused on the optimization of the drug like properties of leads in drug discovery. The volume outlines strategies and methodologies designed to guide pharmaceutical and biotechnology companies through the drug discovery and development process. **Topical Drug Bioavailability,**

Bioequivalence, and Penetration Vinod P. Shah, Howard I. Maibach, 2013-06-29. This volume presents the state of the art of measuring percutaneous penetration and determining biological relevance in dermal and transdermal drug delivery. Both in vivo and in vitro models and methods are discussed in detail to provide pharmaceutical drug developers with an invaluable guide and reference. *Novel Approaches to the Treatment of Alzheimer's Disease* E. Meyer, 2013-03-09. Alzheimer's disease

afflicts up to 1 in 5 people over the age of 65 years and causes untold suffering of the patient and their family. The cause of this disease is unknown; indeed, evidence increasingly suggests that there may be multiple Alzheimer type syndromes with different etiologies analogous to different types of psychosis. Currently, there are no means to prevent the disease, slow its progress, or reverse its neurodegenerative consequences. With few exceptions, clinical trials of a variety of compounds have resulted in patient responses that are disappointing with respect to both the proportion of responders and the magnitude of the responses. Novel approaches to the treatment of Alzheimer's disease are clearly warranted. For this reason, we organized the First Suncoast Workshop on the Neurobiology of Aging in St. Petersburg, Florida, which took place from February 26 to March 1, 1989. This workshop focused on novel treatments and models for Alzheimer's disease and represented a cooperative venture among academia, government, and industry, both in its participants and sponsorship. The Center for the Neurobiology of Aging at the University of Florida, the National Institute on Aging, and Taiho Pharmaceutical Corporation in Japan.

sponsored the workshop in which scientists from the North America Europe Japan and other parts of Asia participated

National Library of Medicine Current Catalog National Library of Medicine (U.S.),1971 First multi year cumulation covers six years 1965 70 *Drug Design* BAREHAM BAREHAM J,2014-11-26 *Drug Design* Volume IX examines various aspects of drug design and covers topics ranging from the consequences of the Hansch paradigm for the pharmaceutical industry to the Masca model of pharmacochimistry A physicochemical basis for the design of orally active prodrugs is also considered along with the use of interactive graphics in medicinal chemistry Comprised of seven chapters this volume begins with a discussion on efforts to avoid toxicity not only of drugs pesticides and food additives but also of chemicals in general The reader is then introduced to various aspects of the development of bioactive agents including the optimization of existing agents by the design of more efficient prodrugs Other chapters focus on Hansch s paradigm and its application to industrial practice the application of multivariate statistics to pharmacochimistry a logico structural approach to computer assisted drug design and spatial arrangements in bioactive molecules This book will be of interest to pharmacologists chemists and those involved in drug design **Ocular Therapeutics and Drug Delivery** Indra K. Reddy,1995-12-12 Since ocular therapeutics and drug delivery is a subject of interest to specialists from various disciplines such as chemical biochemical medical pharmaceutical and toxicological it truly presents a unique situation requiring a multi disciplinary approach in understanding and addressing various problems Yet we see that scientists associated with these areas are working by and large independently of one another thus limiting dissemination of knowledge experience and ideas that would greatly enhance the overall progress in this area of research Regrettably the information currently available in ocular therapeutics and drug delivery though extensive is still fragmented into various disciplines making meaningful synthesis difficult Several books are available that cover one or two aspects of the multi disciplinary fields on an individual basis however none is available that covers all of them The chapters contained in this book are specific to various interrelated areas of ocular therapeutics and drug delivery and are written by acknowledged experts from both academia and industry The book itself is divided into five parts namely I overview basic principles and methodology II pathopharmacology and clinical applications III chemical biochemical approaches to ocular drug delivery IV formulation and drug delivery considerations and V industrial and regulatory considerations Individual chapters in each section apart from presenting a concise text entail an extensive listing of references Analytical Profiles of Drug Substances and Excipients ,1993-11-17 Although the official compendia define a drug substance as to identity purity strength and quality they normally do not provide other physical or chemical data nor do they list methods of synthesis or pathways of physical or biological degradation and metabolism Such information is scattered throughout the scientific literature and the files of pharmaceutical laboratories Analytical Profiles of Drug Substances and Excipients brings this information together into one source The series has recently been expanded to include profiles of excipient materials **Cellular Drug Delivery** D. Robert Lu,Svein Øie,2004-02-25 An authoritative and up to

date survey of the fundamental principles and practice of drug delivery at the cellular level On the principles side the authors discuss the broad spectrum of cellular delivery ranging from coverage of cell mediated immunity gene delivery and protein targeting to cellular drug transport cellular drug permeability and a variety of carrier system related to targeted drug delivery On the practice side the authors focus on technological developments in cellular drug delivery including novel formulations for the delivery of DNA and antisense oligonucleotides as well as drug targeting with immunoglobulin formulations and antibody mediated approaches

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or multiplying out the rhs or substituting may be implied by a 2 or d 4 comparing coefficients of x 3 a 2 b1 comparing coefficients of x 2 b a b 2 0 4 b1 b1 comparing coefficients of x