

Pharmaceutical Biotechnology • Volume 5

Stability and Characterization of Protein and Peptide Drugs

Case Histories

**Edited by
Y. John Wang
and
Rodney Pearlman**

Stability And Characterization Of Protein And Peptide Drugs

Vincent Lee



Stability And Characterization Of Protein And Peptide Drugs:

Stability and Characterization of Protein and Peptide Drugs Y. John Wang, Rodney Pearlman, 1993-05-31 This is the first volume to make available specific case histories of therapeutic proteins and peptides that have been marketed or are currently under clinical testing The editors have selected a wide range of molecules derived from monoclonal antibodies recombinant DNA and natural and chemical sources to provide formulation scientists with practical examples of the development of pharmaceutical products *Therapeutic Peptides and Proteins* Ajay K. Banga, 2005-09-14 Upon publication of the first edition of *Therapeutic Peptides and Proteins* ten years ago there were only 19 biotechnology medicines on the market Currently there are more than 100 with at least 400 more in various stages of development That alone would be grounds for a new edition Add to that the fact that it is still difficult to find up *Pharmaceutical Biotechnology* Daan J. A. Crommelin, Robert D. Sindelar, Bernd Meibohm, 2019-04-13 This introductory text explains both the basic science and the applications of biotechnology derived pharmaceuticals with special emphasis on their clinical use It serves as a complete one stop source for undergraduate graduate pharmacists pharmaceutical science students and for those in the pharmaceutical industry The Fifth Edition completely updates the previous edition and also includes additional coverage on the newer approaches such as oligonucleotides siRNA gene therapy and nanotech and enzyme replacement therapy Parenteral Medications, Fourth Edition Sandeep Nema, John D. Ludwig, 2019-07-19 Parenteral Medications is an authoritative comprehensive reference work on the formulation and manufacturing of parenteral dosage forms effectively balancing theoretical considerations with practical aspects of their development Previously published as a three volume set all volumes have been combined into one comprehensive publication that addresses the plethora of changes in the science and considerable advances in the technology associated with these products and routes of administration Key Features Provides a comprehensive reference work on the formulation and manufacturing of parenteral dosage forms Addresses changes in the science and advances in the technology associated with parenteral medications and routes of administration Includes 13 new chapters and updated chapters throughout Contains the contributors of leading researchers in the field of parenteral medications Uses full color detailed illustrations enhancing the learning process The fourth edition not only reflects enhanced content in all the chapters but also highlights the rapidly advancing formulation processing manufacturing parenteral technology including advanced delivery and cell therapies The book is divided into seven sections Section 1 Parenteral Drug Administration and Delivery Devices Section 2 Formulation Design and Development Section 3 Specialized Drug Delivery Systems Section 4 Primary Packaging and Container Closure Integrity Section 5 Facility Design and Environmental Control Section 6 Sterilization and Pharmaceutical Processing Section 7 Quality Testing and Regulatory Requirements **Physical Methods to Characterize Pharmaceutical Proteins** James N. Herron, Wim Jiskoot, Daan J.A. Crommelin, 2013-11-21 Proteins are still gaining importance in the pharmaceutical world where they are used to improve our

arsenal of therapeutic drugs and vaccines and as diagnostic tools Proteins are different from traditional low molecular weight drugs As a group they exhibit a number of biopharmaceutical and formulation problems These problems have drawn considerable interest from both industrial and academic environments forcing pharmaceutical scientists to explore a domain previously examined only by peptide and protein chemists Biopharmaceutical aspects of proteins e g low oral bioavailability have been extensively investigated Although all possible conventional routes of administration have been examined for proteins no real generally applicable alternative to parenteral administration in order to achieve systemic effects has yet been discovered Several of these biopharmaceutical options have been discussed in Volume 4 of this series Biological Barriers to Protein Delivery Proteins are composed of many amino acids several of which are notorious for their chemical instability Rational design of formulations that optimize the native structure and or bioactivity of a protein is therefore of great importance when long shelf life is required as it is for pharmaceutical products This issue has also been examined in two prior volumes of this series Volume 2 Stability of Protein Pharmaceuticals Part A and Volume 5 Stability and Characterization of Protein and Peptide Drugs

Pharmaceutical Dosage Forms - Parenteral Medications Sandeep Nema, John D. Ludwig, 2016-04-19 This three volume set of *Pharmaceutical Dosage Forms Parenteral Medications* is an authoritative comprehensive reference work on the formulation and manufacture of parenteral dosage forms effectively balancing theoretical considerations with the practical aspects of their development As such it is recommended for scientists and engineers in the

Pharmaceutical Dosage Forms Sandeep Nema, John D. Ludwig, 2010-08-26 *Pharmaceutical Dosage Forms Parenteral Medications* explores the administration of medications through other than the enteral route First published in 1984 as two volumes and then last revised in 1993 this three volume set presents the plethora of changes in the science and considerable advances in the technology associated with these products

Journal of Pharmacy and Pharmacology, 1995

Controlled Drug Delivery Kinam Park, 1997 This volume discusses the challenges of creating controlled release dosage forms that will deliver new therapeutic agents based on high molecular weight molecules It examines strategies for delivering drugs through resistant biological barriers and surveys a variety of topics including drug targeting self regulated drug delivery protein drug delivery biosensors cell and tissue engineering new biomaterials modeling methods pharmacokinetics and U S federal regulations

Novel Therapeutics from Modern Biotechnology Dale L. Oxender, Leonard E. Post, 1999-07-23 While addressing the particular problems associated with several classes of biotechnology products this book also demonstrates that the principles are the same as in the development of small new chemical entities It begins by studying FDA regulatory expectations for biotech products before moving on to discuss general issues common to each class of biotech drug such as proteins peptides and nucleic acids The text deals with specific biotech drugs that have successfully made it into clinical trials and each review is written by a renowned expert in the relevant fields

Mechanistic Studies on the Enhanced Mucosal Transport of Human Growth Hormone by Certain Amino Acid Derivatives

Sy-Juen Wu,1999 *Biologija* ,2002 *Vaccine Design* Jessica R. Burdman,1995 Immunological and formulation design considerations for subunit vaccines Public health implications of emerging vaccine technologies Preclinical safety assessment considerations in vaccine development Regulatory considerations in vaccine design Clinical considerations in vaccine trials with special reference to candidate HIV vaccines Laboratory empiricism clinical design and social value the rough road toward vaccine development A compedium of vaccine adjuvants and excipients Adjuvant properties of aluminum and calcium compounds Structure and properties of aluminum containing adjuvants MF59 design and evaluation of a safe and potent adjuvant for human vaccines Development of vaccines based on formulations containing nonionic block copolymers Development of an emulsion based muramyl dipeptide adjuvant formulation for vaccines Liposomal presentation of antigens for human vaccines Liposome design and vaccine development Lipid matrix based vaccines for mucosal and systemic immunization Polymer microspheres for vaccine delivery Vehicle for oral immunization Design and production of single immunization vaccines using polylactide polyglycolide microsphere systems Nanoparticles as adjuvants for vaccines Water soluble phosphazene polymers for parenteral and mucosal vaccine delivery Monophosphoryl lipid A as an adjuvant past experiences and new directions Structural and immunological characterization of the vaccine adjuvant QS 21 A novel generation of viral vaccines based on the ISCOM matrix Vaccine adjuvants based on gamma inulin A new approach to vaccine adjuvants immunopotential by intracellular T helper like signals transmitted by loxoribine Stearyl tyrosine an organic equivalent of aluminum based immunoadjuvants Cytokines as vaccine adjuvants current status and potential applications Cytokines as immunological adjuvants Cytokine containing liposomes as adjuvants for subunit vaccines Haemophilus influenzae type b conjugate vaccines Pneumococcal conjugate vaccines Lyme vaccine enhancement N terminal acylation of a protein antigen and inclusion of a saponin adjuvant Vaccine research and development for the prevention of filarial nematode infections Retrovirus and retrotransposon particles as antigen presentation and delivery systems Rationale and approaches to constructing preerythrocytic malaria vaccines The MAP system a flexible and unambiguous vaccine design of branched peptides Design of experimental synthetic peptide immunogens for prevention of HIV 1 and HTLV I retroviral infections Design and testing of peptide based cytotoxic T cell mediated immunotherapeutics to treat infectious diseases and cancer Development of active specific immunotherapeutic agents based on cancer associated mucins Synthetic peptide vaccines for schistosomiasis Synthetic hormone growth factor subunit vaccine with application to antifertility and cancer

Therapeutic Protein and Peptide Formulation and Delivery Zahra Shahrokh,1997 This volume reviews protein stability and the analytical and biophysical characterization of proteins It emphasizes drug delivery approaches especially local delivery through the skin Including both academic and industrial perspectives from such companies as Genentech Amgen and Merck the book also discusses novel drug delivery polymers and the development of pharmaceutical protein formulations

Development of an Oil/water TIRF-RET Instrument for the Study of Interfacial Protein Adsorption, Packing,

and Conformational Change Mark Ronald Payne, 2004 **American Book Publishing Record** ,1996 Bibliography of Agriculture ,1995 **Transactions** Biochemical Society (Great Britain),1995 *Formulation and Delivery of Proteins and Peptides* Jeffrey L. Cleland, Robert S. Langer, 1994 Presents an updated review of the recent research in formulation and delivery of proteins and peptides Examines measurement and characterization of degradation products formed during storage degradation mechanisms and model systems designed to predict formulation stability Discusses stabilization of proteins and peptides through lyophilization and focuses on the use of FTIR as a tool for understanding the role of excipients during lyophilization Describes potential new routes and methods of delivery such as depot systems based on biodegradable polymers and aerosol systems for pulmonary delivery The Cumulative Book Index ,1997 A world list of books in the English language

Whispering the Strategies of Language: An Mental Quest through **Stability And Characterization Of Protein And Peptide Drugs**

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