

Drug Stability for Pharmaceutical Scientists

Thersteinn Loffsson



Drug Stability For Pharmaceutical Scientists

Nita K. Pandit



Drug Stability For Pharmaceutical Scientists:

Drug Stability for Pharmaceutical Scientists Thorsteinn Loftsson, 2014-01-25 Drug Stability for Pharmaceutical Scientists is a clear and easy to follow guide on drug degradation in pharmaceutical formulation This book features valuable content on both aqueous and solid drug solutions the stability of proteins and peptides acid base catalyzed and solvent catalyzed reactions how drug formulation can influence drug stability the influence of external factors on reaction rates and much more Full of examples of real life formulation problems and step by step calculations this book is the ideal resource for graduate students as well as scientists in the pharmaceutical and related industries Illustrates important theoretical concepts with numerous examples figures calculations learning problems and questions for self study and retention of material Provides answers and explanations to test your knowledge Enables you to better understand key concepts such as rate and order of reaction reaction equilibrium complex reaction mechanisms and more Includes an in depth discussion of both aqueous and solid drug solutions and contains the latest international regulatory requirements on drug stability

Drug Stability Jens Thurø Carstensen, 1995 Combining basic theory current industrial practice and useful regulatory aspects in an original overview of pharmaceutical stability this thoroughly rewritten and enlarged reference text examines data analysis of the packaged drug s stability experimental methods for achieving stable marketed products and the stability principles of drugs in dissolved dispersed and solid states

Drug Stability Jens Thurø Carstensen, Christopher T. Rhodes, 2000

Martin's Physical Pharmacy and Pharmaceutical Sciences Patrick J. Sinko, 2023-03-01 Consistently revised and updated for more than 60 years to reflect the most current research and practice Martin s Physical Pharmacy and Pharmaceutical Sciences 8th Edition is the original and most comprehensive text available on the physical chemical and biological principles that underlie pharmacology and the pharmaceutical sciences An ideal resource for PharmD and pharmacy students worldwide teachers researchers or industrial pharmaceutical scientists this 8th Edition has been thoroughly revised enhanced and reorganized to provide readers with a clear consistent learning experience that puts essential principles and concepts in a practical approachable context Updated content reflects the latest developments and perspectives across the full spectrum of physical pharmacy and a new full color design makes it easier than ever to discover distinguish and understand information providing users the most robust support available for applying the elements of biology physics and chemistry in work or study

Pharmaceutical Stability Testing to Support Global Markets Kim Huynh-Ba, 2009-12-04 The International Conference of Harmonization ICH has worked on harmonizing the stability regulations in the US Europe and Japan since the early 1990s Even though the Stability Guidelines Q1A R2 was issued over a decade ago issues surrounding this arena continue to surface as the principles described in the guideline are applied to different technical concentrations As a result the stability community has continued to discuss concerns and find ways of harmonizing regulatory requirements streamlining practices improving processes in order to bring safe and effective medical supplies to

the patients around the world In 2007 the American Association of Pharmaceutical Scientists AAPS Stability Focus Group organized two workshops the Stability Workshop and the Degradation Mechanism Workshop These meetings attracted many industry scientists as well as representatives from several regulatory agencies in the world to discuss important topics related to pharmaceutical stability practices Recognizing the importance of documenting these discussions and with the permission of AAPS I have worked with speakers to assemble a collection of 30 articles from presentations given at these two meetings mainly the Stability Workshop I trust that this book will be beneficial to all of you in providing guidance and up to date information for building quality stability programs v Freedom of our mind is Mother of all inventions HPLC for Pharmaceutical Scientists Yuri V. Kazakevich, Rosario LoBrutto, 2006-12-13 HPLC for Pharmaceutical Scientists is an excellent book for both novice and experienced pharmaceutical chemists who regularly use HPLC as an analytical tool to solve challenging problems in the pharmaceutical industry It provides a unified approach to HPLC with an equal and balanced treatment of the theory and practice of HPLC in the pharmaceutical industry In depth discussion of retention processes modern HPLC separation theory properties of stationary phases and columns are well blended with the practical aspects of fast and effective method development and method validation Practical and pragmatic approaches and actual examples of effective development of selective and rugged HPLC methods from a physico chemical point of view are provided This book elucidates the role of HPLC throughout the entire drug development process from drug candidate inception to marketed drug product and gives detailed specifics of HPLC application in each stage of drug development The latest advancements and trends in hyphenated and specialized HPLC techniques LC MS LC NMR Preparative HPLC High temperature HPLC high pressure liquid chromatography are also discussed *Introduction to the Pharmaceutical Sciences* Nita K. Pandit, 2007 This unique textbook provides an introductory yet comprehensive overview of the pharmaceutical sciences It is the first text of its kind to pursue an interdisciplinary approach in this area of study Readers are introduced to basic concepts related to the specific disciplines in the pharmaceutical sciences including pharmacology pharmaceutics pharmacokinetics and medicinal chemistry In an easy to read writing style the book provides readers with up to date information on pharmacogenomics and includes comprehensive coverage of industrial drug development and regulatory approval processes Each chapter includes chapter outlines and critical thinking exercises as well as numerous tables and graphs More than 160 illustrations complement the text **Developing Solid Oral Dosage Forms** Yihong Qiu, Yisheng Chen, Geoff G.Z. Zhang, Lirong Liu, William Porter, 2009-03-10 Developing Solid Oral Dosage Forms is intended for pharmaceutical professionals engaged in research and development of oral dosage forms It covers essential principles of physical pharmacy biopharmaceutics and industrial pharmacy as well as various aspects of state of the art techniques and approaches in pharmaceutical sciences and technologies along with examples and or case studies in product development The objective of this book is to offer updated or current knowledge and skills required for rational oral product design and

development The specific goals are to provide readers with Basics of modern theories of physical pharmacy biopharmaceutics and industrial pharmacy and their applications throughout the entire process of research and development of oral dosage forms Tools and approaches of preformulation investigation formulation process design characterization and scale up in pharmaceutical sciences and technologies New developments challenges trends opportunities intellectual property issues and regulations in solid product development The first book ever that provides comprehensive and in depth coverage of what is required for developing high quality pharmaceutical products to meet international standards It covers a broad scope of topics that encompass the entire spectrum of solid dosage form development for the global market including the most updated science and technologies practice applications regulation intellectual property protection and new development trends with case studies in every chapter A strong team of more than 50 well established authors co authors of diverse background knowledge skills and experience from industry academia and regulatory agencies

Drug Stability and Chemical Kinetics Muhammad Sajid Hamid Akash, Kanwal Rehman, 2020-11-01 This book comprehensively reviews drug stability and chemical kinetics how external factors can influence the stability of drugs and the reaction rates that trigger these effects Explaining the important theoretical concepts of drug stability and chemical kinetics and providing numerous examples in the form of illustrations tables and calculations the book helps readers gain a better understanding of the rates of reactions order of reactions types of degradation and how to prevent it as well as types of stability studies It also offers insights into the importance of the rate at which the drug is degraded and or decomposed under various external and internal conditions including temperature pH humidity and light This book is intended for researchers PhD students and scientists working in the field of pharmacy pharmacology pharmaceutical chemistry medicinal chemistry and biopharmaceutics

Chemical Stability of Pharmaceuticals Kenneth A. Connors, Gordon L. Amidon, Valentino J. Stella, 1986-10-13 Provides a sound theoretical basis for understanding chemical kinetics and its uses in studying drug stability Treats the calculations approximations and estimates that are useful to the pharmacist in professional practice and presents a collection of selected drug stability data from the pharmaceutical literature This Handbook makes accessible to the pharmacist much of the information necessary to make pharmaceutical decisions about drug stability Changes in this edition include thorough revision of the chapter on oxidation addition of a new chapter on solid state stability and a tripling of the number of stability monographs All monographs figures have been redrawn most of them from published data and all sources are cited

Drug Stability, Third Edition, Revised, and Expanded Jens T. Carstensen, Christopher Rhodes, 2000-07-28 Addressing concerns for patient welfare while protecting producer reputation and providing a database for formulation of other products this multiauthored reference blends fundamental theory and practical advice on drug product stability in scientific technical and regulatory environments covering development of indicating assays computer use clinical trial materials strategic planning and packaging Describing the documentation required to minimize the changes of

regulatory citations the book lists manufacturers of photostability testing chambers stability system software and laboratory information management systems for pharmaceutical applications **Journal of Pharmaceutical Sciences** ,1965

Stability of Drugs and Dosage Forms Sumie Yoshioka,Valentino J. Stella,2007-05-08 Drug products are complex mixtures of drugs and excipients and as such their chemical and physical stability kinetics are complex This book discusses the stability of these dosage forms with preformulation studies through to the studies on the final products The book is intended for graduate students researchers and professionals in the field of Pharmaceutics and Pharmaceutical Chemistry **Journal of Pharmaceutical Sciences** ,1912 Vols for 1912 45 include proceedings of the association s annual meeting

Handbook of Stability Testing in Pharmaceutical Development Kim Huynh-Ba,2008-11-16 This handbook is the first to cover all aspects of stability testing in pharmaceutical development Written by a group of international experts the book presents a scientific understanding of regulations and balances methodologies and best practices *Pharmaceutical Science : Its Past and Its Future* ,1985 **ICH Quality Guidelines** Andrew Teasdale,David Elder,Raymond W. Nims,2017-10-09

Examining the implications and practical implementation of multi disciplinary International Conference on Harmonization ICH topics this book gives an integrated view of how the guidelines inform drug development strategic planning and decision making Addresses a consistent need for interpretation training and implementation examples of ICH guidelines via case studies Offers a primary reference point for practitioners addressing the dual challenge of interpretation and practical implementation of ICH guidelines Uses case studies to help readers understand and apply ICH guidelines Provides valuable insights into guidelines development with chapters by authors involved in generating or with experience implementing the guidelines Includes coverage of stability testing analytical method validation impurities biotechnology drugs and products and good manufacturing practice GMP **Remington's Pharmaceutical Sciences** Joseph Price Remington,Alfonso R. Gennaro,1990 **Pharmaceutical Practice** Diana M. Collett,Michael E. Aulton,1990 This edition of Pharmaceutical Practice replaces the 12th edition of Cooper and Gunn s Dispensing for Pharmaceutical Students and has a redesigned and updated content Written by specialists in pharmacy education and practice it aims to provide a sound base for all aspects of the work **Pharmaceutical Sciences** Joseph Price Remington,1975

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