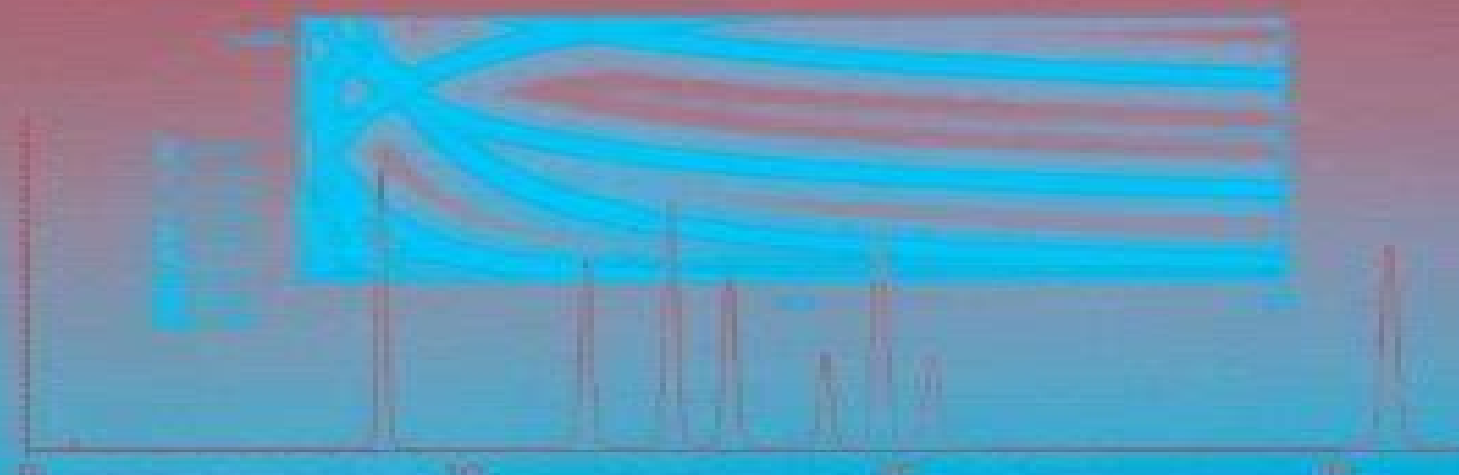




HANDBOOK OF PHARMACEUTICAL ANALYSIS BY HPLC

Edited by
Satinder Ahuja
Michael W. Dong



VOLUME 6

Series Editor Satinder Ahuja



SEPARATION SCIENCE AND TECHNOLOGY

Handbook Of Pharmaceutical Analysis By Hplc Volume 6 Separation Science And Technology

Jicheng Xie



Handbook Of Pharmaceutical Analysis By Hplc Volume 6 Separation Science And Technology:

Handbook of Pharmaceutical Analysis by HPLC Satinder Ahuja, Michael Dong, 2005-02-09 High pressure liquid chromatography frequently called high performance liquid chromatography HPLC or LC is the premier analytical technique in pharmaceutical analysis and is predominantly used in the pharmaceutical industry Written by selected experts in their respective fields the Handbook of Pharmaceutical Analysis by HPLC Volume 6 provides a complete yet concise reference guide for utilizing the versatility of HPLC in drug development and quality control Highlighting novel approaches in HPLC and the latest developments in hyphenated techniques the book captures the essence of major pharmaceutical applications assays stability testing impurity testing dissolution testing cleaning validation high throughput screening A complete reference guide to HPLC Describes best practices in HPLC and offers tricks of the trade in HPLC operation and method development Reviews key HPLC pharmaceutical applications and highlights current trends in HPLC ancillary techniques sample preparations and data handling Principles and Practice of Bioanalysis Richard F. Venn, 2008-02-25 In the seven years since the publication of Principles and Practice of Bioanalysis bioanalytical methods have remained the same but their usage patterns have changed This second edition of a bestseller provides an updated guide to the techniques used in developing and running ultra trace analyses for drugs metabolites and other substance Nuclear Magnetic Resonance G A Webb, 2007-10-31 As a spectroscopic method Nuclear Magnetic Resonance NMR has seen spectacular growth over the past two decades both as a technique and in its applications Today the applications of NMR span a wide range of scientific disciplines from physics to biology to medicine Each volume of Nuclear Magnetic Resonance comprises a combination of annual and biennial reports which together provide comprehensive coverage of the literature on this topic For those wanting to become rapidly acquainted with specific areas of NMR this title provides unrivalled scope of coverage Seasoned practitioners of NMR will find this an invaluable source of current methods and applications Each volume of Nuclear Magnetic Resonance comprises a combination of annual and biennial reports which together provide comprehensive coverage of the literature on this topic This Specialist Periodical Report reflects the growing volume of published work involving NMR techniques and applications in particular NMR of natural macromolecules which is covered in two reports NMR of Proteins and Acids and NMR of Carbohydrates Lipids and Membranes For those wanting to become rapidly acquainted with specific areas of NMR this title provides unrivalled scope of coverage Seasoned practitioners of NMR will find this an invaluable source of current methods and applications Specialist Periodical Reports provide systematic and detailed review coverage in major areas of chemical research Compiled by teams of leading authorities in the relevant subject areas the series creates a unique service for the active research chemist with regular in depth accounts of progress in particular fields of chemistry Subject coverage within different volumes of a given title is similar and publication is on an annual or biennial basis

Capillary Electrophoresis Methods for Pharmaceutical Analysis Satinder Ahuja, Mohamed Elias Jimidar, 2011-08-09

Capillary electrophoresis CE is a powerful analytical technique that is widely used in research and development and in quality control of pharmaceuticals. Many reports of highly efficient separations and methods have been published over the past 15 years. CE offers several advantages over high pressure or high performance liquid chromatography (HPLC). These include simplicity, rapid analysis, automation, ruggedness, different mechanisms for selectivity, and low cost. Moreover, CE requires smaller sample size and yet offers higher efficiency and thus greater resolution power over HPLC. These characteristics are very attractive in research and development, even more so in pharmaceutical quality control (QC) and stability monitoring (SM) studies. This book will provide busy pharmaceutical scientists a complete yet concise reference guide for utilizing the versatility of CE in new drug development and quality control. Provides current status and future developments in CE analysis of pharmaceuticals. Explains how to develop and validate methods. Includes major pharmaceutical applications including assays and impurity testing.

Sample Introduction Systems in ICPMS and ICPOES Diane Beauchemin, 2020-03-15. Sample Introduction Systems in ICPMS and ICPOES provides an in depth analysis of sample introduction strategies including flow injection analysis and less common techniques such as arc spark ablation and direct sample insertion. The book critically evaluates what has been accomplished so far along with what can be done to extend the capabilities of the technique for analyses of any type of sample such as aqueous, gaseous or solid. The latest progress made in fields such as FIA, ETV, LC-ICP-MS and CE-ICP-MS is included and critically discussed. The book addresses problems related to the optimization of the system, peak dispersion and calibration and automatization. Provides contributions from recognized experts that give credibility to each chapter as a reference source. Presents a single source providing the big picture for ICPMS and ICPOES. Covers theory, methods, selected applications and discrete sampling techniques. Includes access to core data for practical work, comparison of results and decision making.

Handbook of Isolation and Characterization of Impurities in Pharmaceuticals Satinder Ahuja, Karen Mills Alsante, 2003-07-18. The United States Food and Drug Administration (FDA) and other regulatory bodies around the world require that impurities in drug substance and drug product levels recommended by the International Conference on Harmonisation (ICH) be isolated and characterized. Identifying process related impurities and degradation products also helps us to understand the production of impurities and assists in defining degradation mechanisms. When this process is performed at an early stage, there is ample time to address various aspects of drug development to prevent or control the production of impurities and degradation products well before the regulatory filing and thus assure production of a high quality drug product. This book therefore has been designed to meet the need for a reference text on the complex process of isolation and characterization of process related synthesis and formulation impurities and degradation products to meet critical regulatory requirements. Its objective is to provide guidance on isolating and characterizing impurities of pharmaceuticals such as drug candidates, drug substances and drug products. The book outlines impurity identification processes and will be a key resource document for impurity analysis, isolation

synthesis and characterization Provides valuable information on isolation and characterization of impurities Gives a regulatory perspective on the subject Describes various considerations involved in meeting regulatory requirements Discusses various sources of impurities and degradation products

HPLC and UHPLC for Practicing Scientists

Michael W. Dong, 2019-07-23 A concise yet comprehensive reference guide on HPLC UHPLC that focuses on its fundamentals latest developments and best practices in the pharmaceutical and biotechnology industries Written for practitioners by an expert practitioner this new edition of HPLC and UHPLC for Practicing Scientists adds numerous updates to its coverage of high performance liquid chromatography including comprehensive information on UHPLC ultra high pressure liquid chromatography and the continuing migration of HPLC to UHPLC the modern standard platform In addition to introducing readers to HPLC s fundamentals applications and developments the book describes basic theory and terminology for the novice and reviews relevant concepts best practices and modern trends for the experienced practitioner HPLC and UHPLC for Practicing Scientists Second Edition offers three new chapters One is a standalone chapter on UHPLC covering concepts benefits practices and potential issues Another examines liquid chromatography mass spectrometry LC MS The third reviews at the analysis of recombinant biologics particularly monoclonal antibodies mAbs used as therapeutics While all chapters are revised in the new edition five chapters are essentially rewritten HPLC columns instrumentation pharmaceutical analysis method development and regulatory aspects The book also includes problem and answer sections at the end of each chapter Overviews fundamentals of HPLC to UHPLC including theories columns and instruments with an abundance of tables figures and key references Features brand new chapters on UHPLC LC MS and analysis of recombinant biologics Presents updated information on the best practices in method development validation operation troubleshooting and maintaining regulatory compliance for both HPLC and UHPLC Contains major revisions to all chapters of the first edition and substantial rewrites of chapters on HPLC columns instrumentation pharmaceutical analysis method development and regulatory aspects Includes end of chapter quizzes as assessment and learning aids Offers a reference guide to graduate students and practicing scientists in pharmaceutical biotechnology and other industries Filled with intuitive explanations case studies and clear figures HPLC and UHPLC for Practicing Scientists Second Edition is an essential resource for practitioners of all levels who need to understand and utilize this versatile analytical technology It will be a great benefit to every busy laboratory analyst and researcher

Handbook of Modern Pharmaceutical Analysis

Satinder Ahuja, Stephen Scypinski, 2001 This book describes the role modern pharmaceutical analysis plays in the development of new drugs Detailed information is provided as to how the quality of drug products is assured from the point of discovery until the patient uses the drug Coverage includes state of the art topics such as analytics for combinatorial chemistry and high throughput screening formulation development stability studies international regulatory aspects and documentation and future technologies that are likely to impact the field Emphasis is placed on current easy to follow methods that readers can apply in their laboratories No book has

effectively replaced the very popular text *Pharmaceutical Analysis* that was edited in the 1960s by Tak Higuchi. This book will fill that gap with an up to date treatment that is both handy and authoritative. *Pharmaceutical Stress Testing* Steven W. Baertschi, Karen M. Alsante, Robert A. Reed, 2016-04-19. The second edition of *Pharmaceutical Stress Testing: Predicting Drug Degradation* provides a practical and scientific guide to designing, executing and interpreting stress testing studies for drug substance and drug product. This is the only guide available to tackle this subject in depth. The Second Edition expands coverage from chemical stability. *Supercritical Fluid Chromatography* Colin F. Poole, 2017-02-16. Supercritical fluid chromatography (SFC) is a rapidly developing laboratory technique for the separation and identification of compounds in mixtures. Significant improvements in instrumentation have rekindled interest in SFC in recent years and enhanced its standing in the scientific community. Many scientists are familiar with column liquid chromatography and its strengths and weaknesses but the possibilities brought to the table by SFC are less well known and are underappreciated. *Supercritical Fluid Chromatography* is a thorough and encompassing reference that defines the concept of contemporary practice in SFC and how it should be implemented in laboratory science. Given the changes that have taken place in SFC, this book presents contemporary aspects and applications of the technique and introduces SFC as a natural solution in the larger field of separation science. The focus on state of the art instrumental SFC distinguishes this work as the go to reference work for those interested in implementing the technique at an advanced level. Edited and authored by world leading chromatography experts. Provides comprehensive coverage of SFC in a single source. Extensive referencing facilitates identification of key research developments. More than 200 figures and tables aid in the retention of key concepts. *Handbook of Food Analysis - Two Volume Set* Leo M.L. Nollet, Fidel Toldra, 2015-06-10. Updated to reflect changes in the industry during the last ten years. The *Handbook of Food Analysis* Third Edition covers the new analysis systems, optimization of existing techniques and automation and miniaturization methods. Under the editorial guidance of food science pioneer Leo M L Nollet and new editor Fidel Toldra, the chapters take an in *Sample Preparation of Pharmaceutical Dosage Forms* Beverly Nickerson, 2011-08-05. This book is intended to serve as a resource for analysts in developing and troubleshooting sample preparation methods. These are critical activities in providing accurate and reliable data throughout the lifecycle of a drug product. This book is divided into four parts. Part One covers dosage form and diluent properties that impact sample preparation of pharmaceutical dosage forms and the importance of sampling considerations in generating data representative of the drug product batch. Part Two reviews specific sample preparation techniques typically used with pharmaceutical dosage forms. Part Three discusses sample preparation method development for different types of dosage forms including addressing drug excipient interactions and post extraction considerations as well as method validation and applying Quality by Design (QbD) principles to sample preparation methods. Part Four examines additional topics in sample preparation including automation, investigating aberrant potency results, green chemistry considerations for sample preparation and the ideal case where no sample

preparation is required for sample analysis Encyclopedia of Analytical Science ,2019-04-02 The third edition of the Encyclopedia of Analytical Science Ten Volume Set is a definitive collection of articles covering the latest technologies in application areas such as medicine environmental science food science and geology Meticulously organized clearly written and fully interdisciplinary the Encyclopedia of Analytical Science Ten Volume Set provides foundational knowledge across the scope of modern analytical chemistry linking fundamental topics with the latest methodologies Articles will cover three broad areas analytical techniques e g mass spectrometry liquid chromatography atomic spectrometry areas of application e g forensic environmental and clinical and analytes e g arsenic nucleic acids and polycyclic aromatic hydrocarbons providing a one stop resource for analytical scientists Offers readers a one stop resource with access to information across the entire scope of modern analytical science Presents articles split into three broad areas analytical techniques areas of application and and analytes creating an ideal resource for students researchers and professionals Provides concise and accessible information that is ideal for non specialists and readers from undergraduate levels and higher *Essentials of Pharmaceutical Analysis* Muhammad Sajid Hamid Akash,Kanwal Rehman,2019-12-17 Recent advances in the pharmaceutical sciences and biotechnology have facilitated the production design formulation and use of various types of pharmaceuticals and biopharmaceuticals This book provides detailed information on the background basic principles and components of techniques used for the analysis of pharmaceuticals and biopharmaceuticals Focusing on those analytical techniques that are most frequently used for pharmaceuticals it classifies them into three major sections and 19 chapters each of which discusses a respective technique in detail Chiefly intended for graduate students in the pharmaceutical sciences the book will familiarize them with the components working principles and practical applications of these indispensable analytical techniques Modern HPLC for Practicing Scientists Michael W. Dong,2016-04-06 A comprehensive yet concise guide to Modern HPLC Written for practitioners by a practitioner Modern HPLC for Practicing Scientists is a concise text which presents the most important High Performance Liquid Chromatography HPLC fundamentals applications and developments It describes basic theory and terminology for the novice and reviews relevant concepts best practices and modern trends for the experienced practitioner Moreover the book serves well as an updated reference guide for busy laboratory analysts and researchers Topics covered include HPLC operation Method development Maintenance and troubleshooting Modern trends in HPLC such as quick turnaround and greener methods Regulatory aspects While broad in scope this book focuses particularly on reversed phase HPLC the most common separation mode and on applications for the pharmaceutical industry the largest user segment Accessible to both novice and intermediate HPLC users information is delivered in a straightforward manner illustrated with an abundance of diagrams chromatograms tables and case studies and supported with selected key references and Web resources With intuitive explanations and clear figures Modern HPLC for Practicing Scientists is an essential resource for practitioners of all levels who need to understand and utilize this versatile analytical technology

High Performance Liquid Chromatography Omar Al Sayed Omar, Moustafa A. Khalifa, 2022-02-21 During the past decade modern high performance liquid chromatography HPLC utilization has expanded greatly especially in the quality control of pharmaceutical products in drug quality control laboratories This book provides an extensive collection of technical information about HPLC Columns physicochemical properties and chromatographic characteristics from various manufacturers and helps analysts to decide on the ideal approach for their analysis according to the requirements of drug manufacturers specifications and the desired Pharmacopeia In addition the authors give practical advice on how to prepare mobile phases choose a suitable detector and set up an HPLC analysis This book is comprehensive for the average professional or technician who plans to work with modern HPLC This book is useful for most Drug Quality Control Laboratories where modern HPLC is utilized Following a hands on approach the book gives key insights into the pharmaceutical applications of HPLC and the latest requirements of the major regulatory agencies such as ICH FDA or USP

GeNeDis 2020 Panayiotis Vlamos, 2022-01-12 The 4th World Congress on Genetics Geriatrics and Neurodegenerative Diseases Research GeNeDis 2020 focuses on the latest major challenges in scientific research new drug targets the development of novel biomarkers new imaging techniques novel protocols for early diagnosis of neurodegenerative diseases and several other scientific advances with the aim of better safer and healthier aging The relation between genetics and its effect on several diseases are thoroughly examined in this volume This volume focuses on the sessions from the conference on Genetics and Neurodegenerative Diseases

TEXT BOOK OF MODERN PHARMACEUTICAL ANALYTICAL TECHNIQUES Dr. MOHD JUNAID ,Dr. Pawan Singh ,Dr. Manoj Shrawan Charde ,Dr. Prithu Pathak ,Prof. Sushil K. Kashaw, 2025-01-07 The Textbook of Modern Pharmaceutical Analytical Techniques is a comprehensive resource designed for students researchers and professionals in pharmaceutical sciences It provides an in depth exploration of advanced analytical methodologies critical to drug development quality control and research

- 1 UV Visible Spectroscopy Covers fundamental principles laws instrumentation solvent effects and versatile applications in pharmaceutical analysis
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- 8 Electrophoresis Explores diverse techniques their instrumentation and roles in pharmaceutical separation processes
- 9 X ray Crystallography Examines diffraction methods Bragg's law and their importance in structural determination of compounds
- 10 Immunological Assays Details RIA ELISA and bioluminescence techniques pivotal in drug

and disease research The textbook emphasizes both theoretical foundations and practical applications bridging the gap between academic learning and industrial practice Rich in diagrams examples and technical insights it s an essential guide for mastering modern analytical techniques Ewing's Analytical Instrumentation Handbook, Fourth Edition Nelu Grinberg, Sonia Rodriguez, 2019-02-21 This handbook is a guide for workers in analytical chemistry who need a starting place for information about a specific instrumental technique It gives a basic introduction to the techniques and provides leading references on the theory and methodology for an instrumental technique This edition thoroughly expands and updates the chapters to include concepts applications and key references from recent literature It also contains a new chapter on process analytical technology **Subject Guide to Books in Print** ,1996

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