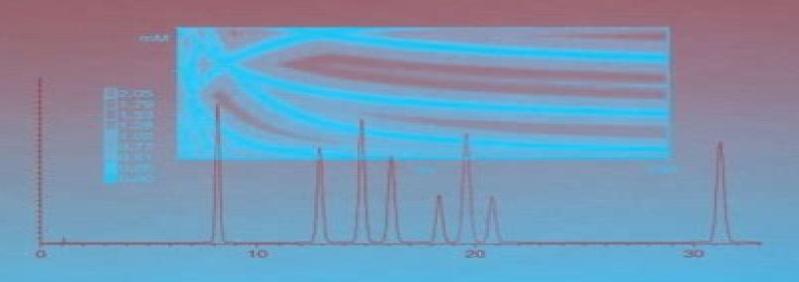


HANDBOOK OF PHARMACEUTICAL ANALYSIS BY HPLC

Edited by Satinder Ahuja Michael W. Dong



VOLUME 6 Series Editor Satinder Ahuja SEPARATION SCIENCE AND TECHNOLOGY

Handbook Of Pharmaceutical Analysis By Hplc

Béla G. Lipták,Kriszta Venczel

Handbook Of Pharmaceutical Analysis By Hplc:

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 guality 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 quide 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 currents trends in HPLC ancillary techniques sample preparations and data handling 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 Handbook of Modern Pharmaceutical Analysis Satinder Ahuja, Stephen Scypinski, 2010-11-11 Handbook of Modern Pharmaceutical Analysis Second Edition synthesizes the complex research and recent changes in the field while covering the techniques and technology required for today s laboratories The work integrates strategy case studies methodologies and implications of new regulatory structures providing complete coverage of quality assurance from the point of discovery to the point of use Treats pharmaceutical analysis PA as an integral partner to the drug development process rather than as a service to it Covers method development validation selection testing modeling and simulation studies combined with advanced exploration of assays impurity testing biomolecules and chiral separations Features detailed coverage of QA ethics and regulatory guidance quality by design good manufacturing practice as well as high tech methodologies and technologies from lab on a chip to LC MS LC NMR and LC NMR MS Modern HPLC for Practicing Scientists Michael W. Dong, 2006-05-19 A comprehesive 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

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everyday laboratory challenges this English edition has been expanded and revised using the feedback received on the successful German precursor Throughout the book Professor Mascher draws on his 30 years of experience and provides abundant practical advice troubleshooting and other hints highlighted in boxes as well as a broad selection of walkthrough case studies Based on a course taught by the author the first part of the book intuitively explains all steps of routine bioanalysis and explains how to set up a robust inexpensive and effi cient method for a given substance In the second part he includes 20 worked example cases that highlight common challenges and how to overcome them With its appendix containing tried and tested analytical methods for 100 clinically relevant substances from the author s own laboratory complete with spectral and MS data as well as literature references and basic pharmacokinetic information this is a life long companion for everyone working in clinical pharmaceutical and biochemical analysis Comments to the German book The book comes to life through its examples showing not only what did work in the author s laboratory but also what didn t ChemieReport Indispensable for novices while even old hands will be able to expand their knowledge A collection of analytical data for ca 100 substances completes the book s offering leaving almost nothing to be desired pharmind

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transition Pharmaceutical Analysis for Small Molecules is a concise yet comprehensive introduction to the drug development process and analysis of chemically synthesized small molecule drugs It features contributions by distinguished experts in the field including editor and author Dr Behnam Davani an analytical chemist with decades of technical management and teaching experience in compendial regulatory and industry This book provides an introduction to pharmaceutical analysis for small molecules non biologics using commonly used techniques for drug characterization and performance tests The driving force for industry to perform pharmaceutical analyses is submission of such data and supporting documents to regulatory bodies for drug approval in order to market their products In addition related required supporting studies including good laboratory documentation practices including analytical instrument qualification are highlighted in this book Topics covered include Drug Approval Process and Regulatory Requirements private standards Pharmacopeias and Compendial Approval Process public standards Common methods in pharmaceutical analysis typically compendial Common Calculations for assays and impurities and other specific tests Analytical Method Validation Verification Transfer Specifications including how to handle out of specification OOS and out of trend OOT Impurities including organic inorganic residual solvents and elemental impurities Good Documentation Practices for regulatory environment Management of Analytical Laboratories Analytical Instrument Qualifications including IQ OQ PQ and VQ Due to global nature of pharmaceutical industry other topics on both regulatory ICH and Compendial harmonization are also highlighted Pharmaceutical Analysis for Small Molecules is a valuable working resource for scientists directly or indirectly involved with the drug development process including analytical chemists pharmaceutical scientists pharmacists and quality control quality assurance professionals It also is an excellent text reference for graduate students in analytical chemistry pharmacy pharmaceutical and regulatory sciences

HPLC Methods for Recently Approved Pharmaceuticals George Lunn, 2005-04-22 An indispensable resource for busy researchers Your time is valuable too valuable to spend hunting through the technical literature in search of the right HPLC assay techniques for your projects With HPLC Methods for Recently Approved Pharmaceuticals you Il quickly identify and replicate the ideal procedures for your project needs without having to refer to original source publications More of your time can then be spent in the lab not the library Covering the relevant world literature through 2003 this book picks up where Dr Lunn's acclaimed HPLC Methods for Pharmaceutical Analysis left off It arms you with established HPLC assay techniques for hundreds of newly approved drugs as well as drugs for which assay methods were only recently developed Combining detailed descriptions of procedures with specially annotated references this practical handbook gives you HPLC methods for 390 commonly prescribed pharmaceutical compounds Various procedures for each drug listed together making it easy to mix and match for customized approaches Methods for drugs in biological fluids and for bulk and formulated drugs Chemical structures molecular weights and formulas and CAS Registry Numbers Cross references to The Merck Index Retention times of other drugs that can be assayed using the same methods

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Ohannesian, Anthony Streeter, 2001-11-09 Exploring the analysis of pharmaceuticals including polymorphic forms this book discusses regulatory requirements in pharmaceutical product development and pharmaceutical testing It covers methods of drug separation and procedures such as capillary electrophoresis for chromatographic separation of molecules Additional topics include drug formulation analysis using vibrational and magnetic resonance spectroscopy and identification of drug metabolites and decomposition products using such techniques as mass spectrometry The book provides more than 300 tables equations drawings and photographs and convenient easy to use indices facilitating quick access to each topic

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