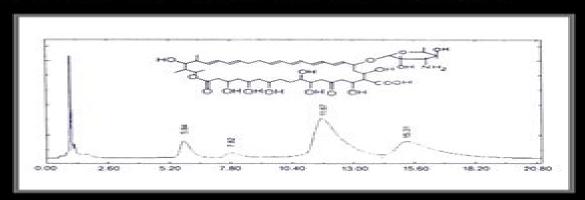
Chromatographic Analysis of Pharmaceuticals

Second Edition, Revised and Expanded



edited by John A. Adamovics

Andras Guttman, László Hajba

Chromatographic Analysis of Pharmaceuticals John A. Adamovics, 2017-09-29 Updated and revised throughout Second Edition explores the chromatographic methods used for the measurement of drugs impurities and excipients in pharmaceutical preparations such as tablets ointments and injectables Contains a 148 page table listing the chromatographic data of over 1300 drugs and related substances including sample matrix analyzed sample handling procedures column packings mobile phase mode of detection and more Chromatographic Analysis of Pharmaceuticals, Second Edition John A. Adamovics, 1996-10-11 Updated and revised throughout Second Edition explores the chromatographic methods used for the measurement of drugs impurities and excipients in pharmaceutical preparations such as tablets ointments and injectables Contains a 148 page table listing the chromatographic data of over 1300 drugs and related substances including sample matrix analyzed sample handling procedures column packings mobile phase mode of detection and more National Library of Medicine Current Catalog National Library of Medicine (U.S.), **Generics and Bioequivalence** Andre J. Jackson, 2019-06-13 Published in 1994 This text focuses on the determination of bioequivalence between formulations that are pharmaceutically equivalent and manufactured using acceptable chemistry manufacturing and controls and in accordance with Good Manufacturing Practices Handbook of Isolation and Characterization of Impurities in Pharmaceuticals Satinder Ahuja, Karen Mills Alsante, 2003-06-26 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 It's 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 degredation products Current Catalog National Library of Medicine (U.S.), First multi year cumulation covers six years 1965 70 Analysis of Drug Impurities Richard J. Smith, Michael L. Webb, 2008-04-15 A key component of the overall quality of a pharmaceutical is control of impurities as their presence even in small amounts may affect drug

safety and efficacy The identification and quantification of impurities to acceptable standards presents a significant challenge to the analytical chemist Analytical science is developing rapidly and provides increasing opportunity to identify the structure and therefore the origin and safety implications of these impurities and the challenges of their measurement drives the development of modern quantitative methods Written for both practicing and student analytical chemists Analysis of Drug Impurities provides a detailed overview of the challenges and the techniques available to permit accurate identification and quantification of drug impurities American Book Publishing Record, 1997 Capillary Gel Electrophoresis Andras Guttman, László Hajba, 2021-12-04 Capillary Gel Electrophoresis and Related Microseparation Techniques covers all theoretical and practical aspects of capillary gel electrophoresis It also provides an excellent overview of the key application areas of nucleic acid protein and complex carbohydrate analysis affinity based methodologies micropreparative aspects and related microseparation methods It not only gives readers a better understanding of how to utilize this technology but also provides insights into how to determine which method will provide the best technical solutions to particular problems This book can also serve as a textbook for undergraduate and graduate courses in analytical chemistry analytical biochemistry molecular biology and biotechnology courses Covers all theoretical and practical aspects of capillary gel electrophoresis Excellent overview of the key applications of nucleic acid protein and complex carbohydrate analysis affinity based methodologies micropreparative aspects and related microseparation methods Teaches readers how to use the technology and select methods that are ideal for fundamental problems Can serve as a textbook for undergraduate and graduate courses in analytical chemistry analytical biochemistry molecular biology and biotechnology courses A Century of Separation Science Haleem J. Issag, 2001-12-19 A Century of Separation Science presents an extensive overview of the critical developments in separation science since 1900 covering recent advances in chromatography electrophoresis field flow fractionation countercurrent chromatography and supercritical fluid chromatography for high speed and high throughput analysis Journal of Chromatography, 2002 **500 Referenced Review Questions in Toxicology** Robert M. The Chemical Bulletin of the Polytechnic Institute "Traian Vuia" of Timisoara, 1990 White,2002 Capillary Electrophoresis Patrick Camilleri, 1997-11-25 The importance of capillary electrophoresis CE as an analytical tool has increased dramatically over the last ten years It has changed from being an exploratory technique mainly of academic interest to one that is applied to solve real analytical problems CE is easily adapted to its various modes of operation often requiring little more than a change of the buffer solution and is quickly becoming the preferred technique when analyzing minute amounts of available material Featuring new chapters on CE analysis of inorganic ions and carbohydrates the new edition of Capillary Electrophoresis not only presents this method as an academic tool but also provides applications for solving real world analytical problems This updated Second Edition reflects the increasing use of CE over the last 10 years how it is being applied and the basic theoretical aspects of the separation and detection methodology of CE Capillary

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