

Stability-Indicating HPLC Methods for Drug Analysis

SECOND EDITION



Quanyun A. Xu
Lawrence A. Trissel



Stability Indicating Hplc Methods For Drug Analysis

Guodong Chen



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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

Stability-indicating HPLC methods for drug analysis, 2008
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Thin Layer Chromatography in Drug Analysis Lukasz Komsta, Monika Waksmundzka-Hajnos, Joseph Sherma, 2013-12-20 Used routinely in drug control laboratories forensic laboratories and as a research tool thin layer chromatography TLC plays an important role in pharmaceutical drug analyses It requires less complicated or expensive equipment than other techniques and has the ability to be performed under field conditions Filling the need for an up to date

Development of Novel Stability Indicating Methods Using Liquid Chromatography Mukesh Maithani, Parveen Bansal, 2019-08-07 Reversed phase high performance liquid chromatography RP HPLC has become the most widely used method for pharmaceutical analysis as it ensures accuracy specificity and reproducibility for the quantification of drugs while avoiding interference from any of the excipients that are normally present in pharmaceutical dosage forms This book presents a simple methodology for developing stability indicating methods and offers a how to guide to creating novel stability indicating methods using liquid chromatography It provides the detailed information needed to devise a stability indicating method for drug substances and drug products that comply with international regulatory guidelines As such it is a must read for anyone engaged in analytical and bioanalytical chemistry

professionals at reference test and control laboratories students and academics at research laboratories and scientists working for chemical pharmaceutical and biotechnology companies Analytical Methods for Drug Development Dr. Sumithra M, Dr. Sharath Chandra Seelam, Mrs. Ramya Nagabathula, Dr. Harikrishnan Hariharan, Mrs. Vinny Therissa Mangam, Dr. Syed Ansar Ahmed, 2025-04-16 This textbook offers a practical approach to understanding analytical methods in drug development Written for students researchers and industry professionals it bridges fundamental concepts with real world applications The book covers essential techniques from early stage drug discovery through manufacturing incorporating current regulatory standards and industry practices Each chapter builds analytical knowledge through practical examples case studies and detailed protocols Whether you re studying pharmacy working in quality control or conducting research this guide provides the tools needed to master modern pharmaceutical analysis and implement effective analytical strategies in drug development Characterization of Impurities and Degradants Using Mass Spectrometry Guodong Chen, 2011-04-27 The book highlights the current practices and future trends in structural characterization of impurities and degradants It begins with an overview of mass spectrometry techniques as related to the analysis of impurities and degradants followed by studies involving characterization of process related impurities including potential genotoxic impurities and excipient related impurities in formulated products Both general practitioners in pharmaceutical research and specialists in analytical chemistry field will benefit from this book that will detail step by step approaches and new strategies to solve challenging problems related to pharmaceutical research **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

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