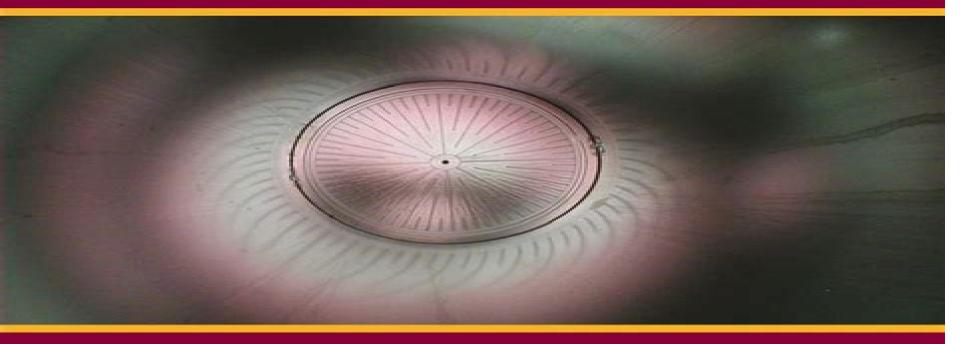


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Donald A.Wellings

A Practical Handbook Of Preparative Hplc

Scott C. Dulebohn

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Quality Control Applications in the Pharmaceutical and Medical Device Manufacturing Industry Carrillo-Cedillo, Eugenia Gabriela, Arredondo-Soto, Karina Cecilia, Palomino-Vizcaino, Kenia, Magaña-Badilla, Héctor Alfonso, 2022-03-18 Quality control in pharmaceutical products and medical devices is vital for users as failing to comply with national and international regulations can lead to accidents that could easily be avoided For this reason manufacturing a quality medical product will support patient safety Microbiologists working in both the pharmaceutical and medical device industries face considerable challenges in keeping abreast of the myriad microbiological references available to them and the continuously evolving regulatory requirements Quality Control Applications in the Pharmaceutical and Medical Device Manufacturing Industry presents the importance of quality control in pharmaceutical products and medical devices which must have very high quality standards to not cause problems to the health of patients It reinforces and updates the knowledge of analytical instrumental and biological methods to demonstrate the correct quality control and good manufacturing practice for pharmaceutical products and medical devices Covering topics such as pharmaceutical nano systems machine learning and software validation this book is an essential resource for managers engineers supervisors pharmacists chemists academicians and researchers Handbook of Process Chromatography Gunter Jagschies, Gail K. Sofer, Lars Hagel, 2007-12-08 This book will update the original edition published in 1997 Since the publication of the first edition the biotechnology and biologics industries have gained extensive knowledge and experience in downstream processing using chromatography and other technologies associated with recovery and purification unit operations. This book will tie that experience together for the next generation of readers Updates include sources and productivity types of products made today experiences in clinical and licensed products economics current status of validation illustrations and tables automated column packing automated systemsNew topics include the use of disposables multiproduct versus dedicated production design principles for chromatography media and filters ultrafiltration principles and optimization risk assessments characterization studies design space platform technologies process analytical technologies PATs biogenerics comparability assessmentsKey Features new approaches to process optimization use of patform technologies applying risk assessment to process design 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 **Supercritical Fluid Chromatography** Gregory K. Webster, 2014-02-04 Analytical chemists in the pharmaceutical industry are always looking for more efficient techniques to meet the analytical challenges of today s pharmaceutical industry One technique that has made steady advances in pharmaceutical analysis is supercritical fluid chromatography SFC SFC is meeting the chromatography needs of the industry by provi

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