



Automated Process Control System In Pharmacy

Thomas Bernold

Automated Process Control System In Pharmacy:

A Textbook of Pharmaceutical Engineering Prof. (Dr.) Sreedhar Ranjan Das, Dr Kaushik Santara, Dr. Neeti Srivastav, Dr. Dipali M. Dhoke, Mrs Mayuri Gupta, 2025-09-16 This textbook is designed specifically to meet the requirements of the Pharmacy Council of India PCI prescribed syllabus for BP 304T Pharmaceutical Engineering in the Bachelor of Pharmacy B Pharm curriculum. It provides a comprehensive and structured overview of the fundamental engineering principles that are essential in pharmaceutical manufacturing and processing. Each chapter integrates theoretical concepts with practical applications relevant to the pharmaceutical industry. Special emphasis is given to equipment design, process efficiency, materials of construction, and corrosion control. Critical elements in ensuring product quality and safety in pharmaceutical settings. Complex topics are broken down into simplified explanations, making it suitable not only for academic study but also for competitive examinations and industrial training. This book serves as a bridge between pharmaceutical sciences and process engineering. It is an invaluable resource for pharmacy undergraduates, educators, and professionals aiming to build a strong foundation in pharmaceutical engineering and develop a sound understanding of the manufacturing processes that underpin drug development and delivery.

An Industrial IoT Approach for Pharmaceutical Industry Growth Valentina Emilia Balas, Vijender Kumar Solanki, Raghvendra Kumar, 2020-05-15 An Industrial IoT Approach for Pharmaceutical Industry Growth Volume Two uses an innovative approach to explore how the Internet of Things (IoT) and big data can improve approaches and make discoveries. Rapid growth of the IoT has encouraged many companies in the manufacturing sector to make use of this technology to unlock its potential. Using clear language and real world case studies, this book discusses systems level from both a human factors point of view and the perspective of networking databases, privacy, and anti-spoofing. The wide variety in topics presented offers multiple perspectives on how to integrate the Internet of Things into pharmaceutical manufacturing. This book represents a useful resource for researchers in pharmaceutical sciences, information and communication technologies, and those who specialize in healthcare and pharmacovigilance. Emphasizes efficiency in pharmaceutical manufacturing through an IoT Big Data approach. Explores cutting edge technologies through sensor enabled environments in the pharmaceutical industry. Discusses system levels from both a human factors point of view and the perspective of networking databases, privacy, and anti-spoofing.

Microfluidics for Pharmaceutical Applications Hélder A. Santos, Dongfei Liu, Hongbo Zhang, 2018-10-12 Microfluidics for Pharmaceutical Applications From Nano Micro Systems Fabrication to Controlled Drug Delivery is a concept orientated reference that features case studies on utilizing microfluidics for drug delivery applications. It is a valuable learning reference on microfluidics for drug delivery applications and assists practitioners developing novel drug delivery platforms using microfluidics. It explores advances in microfluidics for drug delivery applications from different perspectives covering device fabrication, fluid dynamics, cutting edge microfluidic technology in the global drug delivery industry, lab on chip, nano micro fabrication, and drug encapsulation.

cell encapsulation and delivery and cell drug interaction screening These microfluidic platforms have revolutionized the drug delivery field but also show great potential for industrial applications Presents detailed coverage on the fabrication of novel drug delivery systems with desired characteristics such as uniform size Janus particles and particular or combined responsiveness Includes a variety of case studies that explain principles Focuses on commercialization cost safety society and educational issues of microfluidic applications showing how microfluidics is used in the real world Comprehensive Quality by Design for Pharmaceutical Product Development and Manufacture Gintaras V. Reklaitis,Christine Seymour,Salvador García-Munoz,2017-10-09 Covers a widespread view of Quality by Design QbD encompassing the many stages involved in the development of a new drug product The book provides a broad view of Quality by Design QbD and shows how QbD concepts and analysis facilitate the development and manufacture of high quality products QbD is seen as a framework for building process understanding for implementing robust and effective manufacturing processes and provides the underpinnings for a science based regulation of the pharmaceutical industry Edited by the three renowned researchers in the field Comprehensive Quality by Design for Pharmaceutical Product Development and Manufacture guides pharmaceutical engineers and scientists involved in product and process development as well as teachers on how to utilize QbD practices and applications effectively while complying with government regulations The material is divided into three main sections the first six chapters address the role of key technologies including process modeling process analytical technology automated process control and statistical methodology in supporting QbD and establishing the associated design space The second section consisting of seven chapters present a range of thoroughly developed case studies in which the tools and methodologies discussed in the first section are used to support specific drug substance and drug product QbD related developments The last section discussed the needs for integrated tools and reviews the status of information technology tools available for systematic data and knowledge management to support QbD and related activities Highlights Demonstrates Quality by Design QbD concepts through concrete detailed industrial case studies involving of the use of best practices and assessment of regulatory implications Chapters are devoted to applications of QbD methodology in three main processing sectors drug substance process development oral drug product manufacture parenteral product processing and solid liquid processing Reviews the spectrum of process model types and their relevance the range of state of the art real time monitoring tools and chemometrics and alternative automatic process control strategies and methods for both batch and continuous processes The role of the design space is demonstrated through specific examples and the importance of understanding the risk management aspects of design space definition is highlighted Comprehensive Quality by Design for Pharmaceutical Product Development and Manufacture is an ideal book for practitioners researchers and graduate students involved in the development research or studying of a new drug and its associated manufacturing process **Chemical Engineering in the Pharmaceutical Industry** David J. am Ende,Mary T. am Ende,2019-03-28 A guide to the development

and manufacturing of pharmaceutical products written for professionals in the industry revised second edition The revised and updated second edition of Chemical Engineering in the Pharmaceutical Industry is a practical book that highlights chemistry and chemical engineering The book s regulatory quality strategies target the development and manufacturing of pharmaceutically active ingredients of pharmaceutical products The expanded second edition contains revised content with many new case studies and additional example calculations that are of interest to chemical engineers The 2nd Edition is divided into two separate books 1 Active Pharmaceutical Ingredients API s and 2 Drug Product Design Development and Modeling The active pharmaceutical ingredients book puts the focus on the chemistry chemical engineering and unit operations specific to development and manufacturing of the active ingredients of the pharmaceutical product The drug substance operations section includes information on chemical reactions mixing distillations extractions crystallizations filtration drying and wet and dry milling In addition the book includes many applications of process modeling and modern software tools that are geared toward batch scale and continuous drug substance pharmaceutical operations This updated second edition Contains 30new chapters or revised chapters specific to API covering topics including manufacturing quality by design computational approaches continuous manufacturing crystallization and final form process safety Expanded topics of scale up continuous processing applications of thermodynamics and thermodynamic modeling filtration and drying Presents updated and expanded example calculations Includes contributions from noted experts in the field Written for pharmaceutical engineers chemical engineers undergraduate and graduate students and professionals in the field of pharmaceutical sciences and manufacturing the second edition of Chemical Engineering in the Pharmaceutical Industryf ocuses on the development and chemical engineering as well as operations specific to the design formulation and manufacture of drug substance and products

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