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September 20-21, 2022

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TECHNOLOGY'S CONNECTION IDEAS

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PROGRAM - DAY 1

SEPTEMBER 20, 2022

* JOIN US FOR THE PRE-SUMMIT SUPPLY CHAIN DISRUPTIONS PANEL AND PANEL 4 AT 10 PM - 11 PM PM EDT

10:00 AM - 10:30 AM

REGULATE REGISTRATION AND NETWORKING BREAKFAST

10:30 AM - 11:00 AM

CHAIR'S WELCOME AND OPENING REMARKS

EXTERNAL MANUFACTURING

Ampho

AMPHO
Head of North America Outsourcing
and

EXTERNAL QUALITY

Qflow

Qflow
Head of North America Outsourcing
and

EXTERNAL SUPPLY

Ampho

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and

11:00 AM - 11:30 AM

OPENING REMARKS

Ampho

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THAT'S RELATIONSHIP AND DATA DEPENDENT CONNECTIONS FOR SUPPLIER INTERACTION WITH UPSTREAM AND DOWNSTREAM

- What's the relationship to upstream?
- Building with a data and strategic thought about the product
- Data integration: connecting the data to the product
- Data integration: connecting the data to the product
- Building with a data and strategic thought about the product

11:30 AM - 12:00 PM

PLANNING

Ampho

Ampho
Head of North America Outsourcing and

LEARNING NETWORK AND EXTERNAL PLANS TO SUPPORT A GLOBAL MANUFACTURING NETWORK

- Building a learning network of suppliers and customers, internal quality and external supply chain experts
- Data integration: connecting the data to the product
- Building a learning network of suppliers and customers, internal quality and external supply chain experts
- Building a learning network of suppliers and customers, internal quality and external supply chain experts
- Building a learning network of suppliers and customers, internal quality and external supply chain experts
- Building a learning network of suppliers and customers, internal quality and external supply chain experts

American Pharmaceutical Outsourcing Validation Of

Leon Shargel, Isadore Kanfer



American Pharmaceutical Outsourcing Validation Of:

ICH Quality Guidelines Andrew Teasdale, David Elder, Raymond W. Nims, 2017-09-29 Examining the implications and practical implementation of multi disciplinary International Conference on Harmonization ICH topics this book gives an integrated view of how the guidelines inform drug development strategic planning and decision making Addresses a consistent need for interpretation training and implementation examples of ICH guidelines via case studies Offers a primary reference point for practitioners addressing the dual challenge of interpretation and practical implementation of ICH guidelines Uses case studies to help readers understand and apply ICH guidelines Provides valuable insights into guidelines development with chapters by authors involved in generating or with experience implementing the guidelines Includes coverage of stability testing analytical method validation impurities biotechnology drugs and products and good manufacturing practice GMP

Pharmaceutical Process Validation Robert A. Nash, Alfred H. Wachter, 2003-03-27 The third edition of this text contains additional chapters which cover troubleshooting procedures validation in contract manufacturing and current harmonization trends

Process Validation & cGMP (Part - 1) Mr. Rohit Manglik, 2024-03-24 Covers fundamentals of process validation documentation regulatory guidelines and GMP principles in pharmaceutical manufacturing

Ligand-Binding Assays Masood N. Khan, John W. A. Findlay, 2009-10-22 A consolidated and comprehensive reference on ligand binding assays Ligand binding assays LBAs stand as the cornerstone of support for definition of the pharmacokinetics and toxicokinetics of macromolecules an area of burgeoning interest in the pharmaceutical industry Yet outside of the Crystal City Conference proceedings little guidance has been available for LBA validation particularly for assays used to support macromolecule drug development Ligand Binding Assays Development Validation and Implementation in the Drug Development Arena answers that growing need serving as a reference text discussing critical aspects of the development validation and implementation of ligand binding assays in the drug development field Ligand Binding Assays covers essential topics related to ligand binding assays from pharmacokinetic studies the development of LBAs assay validation statistical LBA aspects and regulatory aspects to software for LBAs and robotics and other emerging methodologies for LBAs Highlights include A general discussion of challenges and proven approaches in the development of ligand binding assays More detailed examination of characteristics of these assays when applied to support of pharmacokinetic and toxicokinetic studies of compounds at different stages in the discovery or development timeline A concise but detailed discussion of validation of ligand binding assays for macromolecules A practical approach to fit for purpose validation of assays for biomarkers those molecules receiving increased attention as potentially demonstrating that the target chosen in discovery is being modulated by the candidate therapeutic both in nonclinical and clinical studies Written by a team of world recognized authorities in the field Ligand Binding Assays provides key information to a broad range of practitioners both in the pharmaceutical and allied industries and in related contract research

organizations and academic laboratories and perhaps even in the field of diagnostics and clinical chemistry

Specification of Drug Substances and Products Christopher M. Riley, Thomas W. Rosanske, George L. Reid, 2020-07-23 Specification of Drug Substances and Products Development and Validation of Analytical Methods Second Edition presents a comprehensive and critical analysis of the requirements and approaches to setting specifications for new pharmaceutical products with an emphasis on phase appropriate development validation of analytical methods and their application in practice This thoroughly revised second edition covers topics not covered or not substantially covered in the first edition including method development and validation in the clinical phase method transfer process analytical technology analytical life cycle management special challenges with generic drugs genotoxic impurities topical products nasal sprays and inhalation products and biotechnology products The book s authors have been carefully selected as former members of the ICH Expert Working Groups charged with developing the ICH guidelines and or subject matter experts in the industry academia and in government laboratories Presents a critical assessment of the application of ICH guidelines on method validation and specification setting Written by subject matter experts involved in the development and application of the guidelines Provides a comprehensive treatment of the analytical methodologies used in the analysis control and specification of new drug substances and products Covers the latest statistical approaches including analytical quality by design in the development of specifications method validation and shelf life prediction

Biopharmaceutical Manufacturing Gary Gilleskie, Charles Rutter, Becky McCuen, 2021-09-07 Biopharmaceuticals medicines made by or from living organisms including cells from living organisms are extremely effective in treating a broad range of diseases Their importance to human health has grown significantly over the years as more biopharmaceutical products have entered the market and now the biggest selling drugs in the world are biopharmaceuticals Biopharmaceutical Manufacturing Principles Processes and Practices provides concise comprehensive and up to date coverage of biopharmaceutical manufacturing Written in a clear and informal style the content has been influenced by the authors substantial industry experience and teaching expertise That expertise enables the authors to address the many questions posed over the years both by university students and professionals with experience in the field Consequently the book will appeal both to undergraduate or graduate students using it as a textbook and specialized industry practitioners seeking to understand the big picture of biopharmaceutical manufacturing This book

Comprehensive Toxicology, 2017-12-01 Comprehensive Toxicology Third Edition Fifteen Volume Set discusses chemical effects on biological systems with a focus on understanding the mechanisms by which chemicals induce adverse health effects Organized by organ system this comprehensive reference work addresses the toxicological effects of chemicals on the immune system the hematopoietic system cardiovascular system respiratory system hepatic toxicology renal toxicology gastrointestinal toxicology reproductive and endocrine toxicology neuro and behavioral toxicology developmental toxicology and carcinogenesis also including critical sections that cover the general principles of

toxicology cellular and molecular toxicology biotransformation and toxicology testing and evaluation Each section is examined in state of the art chapters written by domain experts providing key information to support the investigations of researchers across the medical veterinary food environment and chemical research industries and national and international regulatory agencies Thoroughly revised and expanded to 15 volumes that include the latest advances in research and uniquely organized by organ system for ease of reference and diagnosis this new edition is an essential reference for researchers of toxicology Organized to cover both the fundamental principles of toxicology and unique aspects of major organ systems Thoroughly revised to include the latest advances in the toxicological effects of chemicals on the immune system Features additional coverage throughout and a new volume on toxicology of the hematopoietic system Presents in depth comprehensive coverage from an international author base of domain experts

Outsourcing in Clinical Drug Development Roy Drucker, Graham Hughes, 2015-05-06 Sponsor companies and CROs alike will appreciate the industry wide analysis practical how to advice and helpful charts and checklists provided by Outsourcing in Clinical Drug Development A panel of experts discuss supplier identification and selection financial considerations and the ethical issues They cover contracting out laboratory analy

Generic Drug Product Development Leon Shargel, Isadore Kanfer, 2013-10-24 In this era of increased pharmaceutical industry competition success for generic drug companies is dependent on their ability to manufacture therapeutic equivalent drug products in an economical and timely manner while also being cognizant of patent infringement and other legal and regulatory concerns *Generic Drug Product Development Solid Oral Dosage Forms Second Edition* presents in depth discussions from more than 30 noted specialists describing the development of generic drug products from the raw materials to the development of a therapeutic equivalent drug product to regulatory approval Major topics discussed include Active pharmaceutical ingredients Experimental formulation development including a new section on Quality by Design QbD Scale up Commercial product formulation Quality control and bioequivalence Drug product performance ANDA regulatory process Post approval changes Post marketing surveillance Legislative and patent challenges This second edition also contains a new chapter on the relationship between the FDA and the United States Pharmacopeia and in Chapter 4 using specific examples the application of Quality by Design QbD during formulation development is examined The book is a thorough guide to the development of solid oral generic dosage formulations This textbook is ideal for the pharmaceutical industry graduate programs in pharmaceutical sciences and health professionals working in the area of generic drug development

23 European Symposium on Computer Aided Process Engineering Yang Yang, Suzanne S. Farid, Nina F. Thornhill, 2013-06-10 The impact of upstream and downstream process fluctuations in bioprocesses becomes more pronounced during technology transfer of pilot scale processes into large scale facilities Mismatches in equipment sizes upon scale up combined with process fluctuations can lead to discarding expensive product Understanding and predicting the root causes of product loss is critical to determining the necessary facility modifications

required to meet manufacturing targets In this work a decision tree classification method CART classification and regression tree was used to interrogate stochastic Monte Carlo simulation datasets that mimic batch fluctuations in industrial biopharmaceutical batch processes The resulting binary decision tree not only reveals the impact of process fluctuations on product mass loss but also extracts a series of if then rules on the critical combinations of factors that lead to unacceptable mass loss levels The decision tree analysis therefore allows rapid prediction of the likelihood of facility fit issues as well as the batch characteristics that result in product loss

23rd European Symposium on Computer Aided Process Engineering, 2013-06-10 Computer aided process engineering CAPE plays a key design and operations role in the process industries from the molecular scale through managing complex manufacturing sites The research interests cover a wide range of interdisciplinary problems related to the current needs of society and industry ESCAPE 23 brings together researchers and practitioners of computer aided process engineering interested in modeling simulation and optimization synthesis and design automation and control and education The proceedings present and evaluate emerging as well as established research methods and concepts as well as industrial case studies Contributions from the international community using computer based methods in process engineering Reviews the latest developments in process systems engineering Emphasis on industrial and societal challenges

Practical Pharmaceutical Engineering Gary Prager, 2018-11-28 A practical guide to all key the elements of pharmaceuticals and biotech manufacturing and design Engineers working in the pharmaceutical and biotech industries are routinely called upon to handle operational issues outside of their fields of expertise Traditionally the competencies required to fulfill those tasks were achieved piecemeal through years of self teaching and on the job experience until now Practical Pharmaceutical Engineering provides readers with the technical information and tools needed to deal with most common engineering issues that can arise in the course of day to day operations of pharmaceutical biotech research and manufacturing Engineers working in pharma biotech wear many hats They are involved in the conception design construction and operation of research facilities and manufacturing plants as well as the scale up manufacturing packaging and labeling processes They have to implement FDA regulations validation assurance quality control and Good Manufacturing Practices GMP compliance measures and to maintain a high level of personal and environmental safety This book provides readers from a range of engineering specialties with a detailed blueprint and the technical knowledge needed to tackle those critical responsibilities with confidence At minimum after reading this book readers will have the knowledge needed to constructively participate in contractor user briefings Provides pharmaceutical industry professionals with an overview of how all the parts fit together and a level of expertise that can take years of on the job experience to acquire Addresses topics not covered in university courses but which are crucial to working effectively in the pharma biotech industry Fills a gap in the literature providing important information on pharmaceutical operation issues required for meeting regulatory guidelines plant support design and project engineering Covers the basics

of HVAC systems water systems electric systems reliability maintainability and quality assurance relevant to pharmaceutical engineering Practical Pharmaceutical Engineering is an indispensable tool of the trade for chemical engineers mechanical engineers and pharmaceutical engineers employed by pharmaceutical and biotech companies engineering firms and consulting firms It also is a must read for engineering students pharmacy students chemistry students and others considering a career in pharmaceuticals

Outsourcing and Offshoring of Professional Services: Business Optimization in a Global Economy Gupta, Amar, 2008-03-31 This book discusses the considerations and implications surrounding the outsourcing and offshoring of professional services such as software development computer aided design and healthcare from multiple global perspectives This book offers industry professionals policymakers students and educators with a balance between a broad overview and detailed analysis of offshore outsourcing would make an invaluable addition to any reference library Provided by publisher

Pharmaceutical Stability Testing to Support Global Markets Kim Huynh-Ba, 2009-12-04 The International Conference of Harmonization ICH has worked on harmonizing the stability regulations in the US Europe and Japan since the early 1990s Even though the Stability Guidelines Q1A R2 was issued over a decade ago issues surrounding this arena continue to surface as the principles described in the guideline are applied to different technical concentrations As a result the stability community has continued to discuss concerns and find ways of harmonizing regulatory requirements streamlining practices improving processes in order to bring safe and effective medical supplies to the patients around the world In 2007 the American Association of Pharmaceutical Scientists AAPS Stability Focus Group organized two workshops the Stability Workshop and the Degradation Mechanism Workshop These meetings attracted many industry scientists as well as representatives from several regulatory agencies in the world to discuss important topics related to pharmaceutical stability practices Recognizing the importance of documenting these discussions and with the permission of AAPS I have worked with speakers to assemble a collection of 30 articles from presentations given at these two meetings mainly the Stability Workshop I trust that this book will be beneficial to all of you in providing guidance and up to date information for building quality stability programs v Freedom of our mind is Mother of all inventions

Pharmaceutical Manufacturing Handbook Shayne Cox Gad, 2008-03-11 This handbook features contributions from a team of expert authors representing the many disciplines within science engineering and technology that are involved in pharmaceutical manufacturing They provide the information and tools you need to design implement operate and troubleshoot a pharmaceutical manufacturing system The editor with more than thirty years experience working with pharmaceutical and biotechnology companies carefully reviewed all the chapters to ensure that each one is thorough accurate and clear

Handbook of Nutraceuticals Volume II Yashwant Vishnupant Pathak, 2011-05-16 Due in part to an absence of universally accepted standardization methods nutraceuticals and functional foods face regulatory ignorance marketing incompetence and ethical impunity Even though many researchers believe that there is a connection between nutraceuticals and functional foods and reduced health

care expenses as well as disease prevent **American Journal of Hospital Pharmacy** ,1994 **Pharmaceutical and Biomedical Project Management in a Changing Global Environment** Scott D. Babler,2011-01-06 Pharmaceutical and Biomedical Portfolio Management in a Changing Global Environment explores some of the critical forces at work today in the complex endeavour of pharmaceutical and medical product development Written by experienced professionals and including real world approaches and best practice examples this new title addresses three key areas small molecules large molecules and medical devices and provides hard to find consolidated information relevant to and needed by pharmaceutical biotech and medical device company managers *The Almanac of American Employers 2007* Jack W. Plunkett,2006-10 This book will help you sort through America s giant corporate employers to determine which may be the best for corporate employers to determine which may be the best for you or to see how your current employer compares to others It has reference for growth and hiring plans salaries and benefits women and minority advancement industries locations and careers and major trends affecting job seekers **Good Manufacturing Practices for Pharmaceuticals** Joseph D. Nally,2016-04-19 With global harmonization of regulatory requirements and quality standards and national and global business consolidations ongoing at a fast pace pharmaceutical manufacturers suppliers contractors and distributors are impacted by continual change Offering a wide assortment of policy and guidance document references and interpretations this Sixth Edition is significantly expanded to reflect the increase of information and changing practices in CGMP regulation and pharmaceutical manufacturing and control practices worldwide An essential companion for every pharmaceutical professional this guide is updated and expanded by a team of industry experts each member with extensive experience in industry or academic settings

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American Pharmaceutical Outsourcing Validation Of Introduction

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