Gamp 5 A Quality Risk Management Approach To Computer

Data Integrity and Data Governance
Computerized Laboratory Systems
GAMP Good Practice Guide
Pharmaceutical Microbiology
Method Validation in Pharmaceutical Analysis
Quality Risk Management in the FDA-Regulated Industry
Pharmaceutical Manufacturing Handbook
International IT Regulations and Compliance
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Pharmaceutical Computer Systems Validation
A Risk-based Approach to Operation of GxP Computerized Systems
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Risk Management Applications in Pharmaceutical and Biopharmaceutical Manufacturing
EBOOK: Quality, Risk and Control in Health Care
ISPE Guide
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Guideline on General Principles of Process Validation
Biocontamination Control for Pharmaceuticals and Healthcare
Computer System Validation and GAMP 5
Factfulness
Multivariate Analysis in the Pharmaceutical Industry
Pharmaceutical Manufacturing Handbook
GAMP Good Practice Guide

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Data Integrity and Data Governance John Wiley & Sons
The purpose of this book is to help you understand how computerized systems are validated using the GAMP5framework. The information will be presented in a project life cycle format. This will give you a solid idea howComputerized System Validation projects are conducted. This book is suited for anyone new to Computer SystemsValidation. It is written in a simple manner and can serve as starter guide which includes many high-level sample templates and illustration.
Computerized Laboratory Systems John Wiley & Sons
Adopting a practical approach, the authors provide a detailed interpretation of the existing regulations (GMP, ICH), while also discussing the appropriate calculations, parameters and tests. The book thus allows readers to validate the analysis of pharmaceutical compounds while complying with both the regulations as well as the industry demands for robustness and cost effectiveness. Following an introduction to the basic parameters and tests in pharmaceutical validation, including specificity, linearity, range, precision, accuracy, detection and quantitation limits, the text focuses on a life-cycle approach to validation and the integration of validation into the whole analytical quality assurance system. The whole is rounded off with a look at future trends. With its first-hand knowledge of the industry as well as regulating bodies, this is an invaluable reference for analytical chemists, the pharmaceutical industry, pharmaceutists, QA officers, and public authorities.
GAMP Good Practice Guide Springer Nature
Standards, technologies, and requirements for computer validation have changed dramatically in recent years, and so have the interpretation of the standards and the understanding of the processes involved. International IT Regulations and Compliance brings together current thinking on the implementation of standards and regulations in relation to IT for a wide variety of industries. The book provides professionals in pharmaceutical and semiconductor industries with an updated overview of requirements for handling IT systems according to various Quality Standards and how to ?translate? these requirements in the regulations.
Pharmaceutical Microbiology John Wiley & Sons
In today's uncertain times, risk has become the biggest part of management. Risk management is central to the science of prediction and decision-making; holistic and scientific risk management creates resilient organizations, which survive and thrive by being adaptable. This book is the perfect guide for anyone interested in understanding and excelling at risk management. It begins with a focus on the foundational elements of risk management, with a thorough explanation of the basic concepts, many illustrated by real-life examples. Next, the book focuses on equipping the reader with a working knowledge of the subject from an organizational process and systems perspective. Every concept in almost every chapter is calibrated to not only ISO 9001 and ISO 31000, but several other international standards. In addition, this book presents several tools and methods for discussion. Ranging from industry standard to cutting edge, each receives a thorough analysis and description of its role in the risk management process. Finally, you'll find a detailed and practical discussion of contemporary topics in risk management, such as supply chain risk management, risk-based auditing, risk in 4.0 (digital transformation), benefit-risk analyses, risk-based design thinking, and pandemic/epidemic risk management. Jayet Moon is a Senior ASQ member and holds ASQ CQE, CSQP, and CQIA certifications. He is also a chartered quality professional in the U.K. (CQP-MCQI). He earned a master's degree in biomedical engineering from Drexel University in Philadelphia and is a Project Management Institute (PMI) Certified Risk Management Professional (PMI-RMP). He is a doctoral candidate in Systems and Engineering Management at Texas Tech University
Method Validation in Pharmaceutical Analysis PharmaLogika Books
This book elevates alarm management from a fragmented collection of procedures, metrics, experiences, and trial-and-error, to the level of a technology discipline. It provides a complete treatment of best practices in alarm management. The technology and approaches found here provide the opportunity to completely understand the what, the why, and the how of successful alarm systems. No modern industrial enterprise, particularly in such areas as chemical processing, can operate without a secure and reliable infrastructure of alarms and controls-they are an integral part of all production management and control systems. Improving alarm management is an effective way to provide operators with high-value support and guidance to successfully manage industrial plant operations. Readers will find: Recommendations and guidelines are developed from fundamental concepts to provide powerful technical tools and workable approaches; Alarms are treated as indicators of abnormal situations, not simply sensor readings that might be out of position; Alarm improvement is intimately linked to infrastructure management, including the vital role of plant maintenance to alarm management, the need to manage operators' charter to continue to operate during abnormal situations vs. cease operation, and the importance of situation awareness without undue reliance upon alarms. The ability to appreciate technical issues is important, but this book requires no previous specific technical, educational, or experiential background. The style and content are very accessible to a broad industrial audience from board operator to plant manager. All critical tasks are explained with workflow processes, examples, and insight into what it all means. Alternatives are offered everywhere to enable users to tailor-make solutions to their particular sites.
Quality Risk Management in the FDA-Regulated Industry McGraw-Hill Education (UK)
This book provides practical and detailed advice on how to implement data governance and data integrity for regulated analytical laboratories working in the pharmaceutical and allied industries.
Pharmaceutical Manufacturing Handbook IChemE
Sets forth tested and proven risk management practices in drug manufacturing Risk management is essential for safe and efficient pharmaceutical and biopharmaceutical manufacturing, control, and distribution. With this book as their guide, readers involved in all facets of drug manufacturing have a single, expertly written, and organized resource to guide them through all facets of risk management and analysis. It sets forth a solid foundation in risk management concepts and then explains how these concepts are applied to drug manufacturing. Risk Management Applications in Pharmaceutical and Biopharmaceutical Manufacturing features contributions from leading international experts in risk management and drug manufacturing. These contributions reflect the latest research, practices, and industry standards as well as the authors' firsthand experience. Readers can turn to the book for: Basic foundation of risk management principles, practices, and applications Tested and proven tools and methods for managing risk in pharmaceutical and biopharmaceutical product manufacturing processes Recent FDA guidelines, EU regulations, and international standards governing the application of risk management to drug manufacturing Case studies and detailed examples demonstrating the use and results of applying risk management principles to drug product manufacturing Bibliography and extensive references leading to the literature and helpful resources in the field With its unique focus on the application of risk management to biopharmaceutical and pharmaceutical manufacturing, this book is an essential resource for pharmaceutical and process engineers as well as safety and compliance professionals involved in drug manufacturing.
International IT Regulations and Compliance Ispe Headquarters
With its coverage of Food and Drug Administration regulations, international regulations, good manufacturing practices, and process analytical technology, this handbook offers complete coverage of the regulations and quality control issues that govern pharmaceutical manufacturing. In addition, the book discusses quality assurance and validation, drug stability, and contamination control, all

key aspects of pharmaceutical manufacturing that are heavily influenced by regulatory guidelines. The team of expert authors offer you advice based on their own firsthand experience in all phases of pharmaceutical manufacturing.

Good Research Practice in Non-Clinical Pharmacology and Biomedicine CRC Press

Multivariate Analysis in the Pharmaceutical Industry provides industry practitioners with guidance on multivariate data methods and their applications over the lifecycle of a pharmaceutical product, from process development, to routine manufacturing, focusing on the challenges specific to each step. It includes an overview of regulatory guidance specific to the use of these methods, along with perspectives on the applications of these methods that allow for testing, monitoring and controlling products and processes. The book seeks to put multivariate analysis into a pharmaceutical context for the benefit of pharmaceutical practitioners, potential practitioners, managers and regulators. Users will find a resources that addresses an unmet need on how pharmaceutical industry professionals can extract value from data that is routinely collected on products and processes, especially as these techniques become more widely used, and ultimately, expected by regulators.

Standards, Quality Control, and Measurement Sciences in 3D Printing and Additive Manufacturing Woodhead Publishing

The purpose of this new edition is to offer an updated view of the risk management field as it applies to medical products. Since the publication of the first edition (2012), the emphasis on risk-based processes has growth exponentially across all sectors, and risk management is now considered as significant as quality management. ISO 9001 was revised and now requires that top management promote the use of risk-based thinking. ISO 13485:2016, which specifies the requirements for a quality management system specific to the medical devices industry, also now shows a greater emphasis on risk management and risk-based decision making. In addition, the FDA Food Safety Modernization Act (FSMA) is the most important reform of U.S. food safety laws in more than 70 years. This indispensable book presents a systematic and comprehensive approach to quality risk management. It will assist medical and food product manufacturers with the integration of a risk management system or risk management principles and activities into their existing quality management system by providing practical explanations and examples. The appropriate use of quality risk management can facilitate compliance with regulatory requirements such as good manufacturing practice or good laboratory practice. All chapters have been updated and revised, and a new chapter has been added to discuss some of the most common pitfalls and misunderstandings regarding risk management, specifically those related to the use of FMEA as the only element of risk management programs. One of the appendices includes 12 case studies, and the companion CD-ROM contains dozens of U.S. FDA and European guidance documents as well as international harmonization documents (ICH and GHTF-IMDRF) related to risk management activities, as well as a 30-question exam (with answers) on the material discussed in the book.

GAMP Good Practice Guide Ispe Headquarters

“With better governance a key issue in the NHS boardroom, this book provides a comprehensive underpinning to future developments.” Roger Moore, Chief Executive, NHS Appointments Commission, UK "This book provides a much needed integration of different streams in the quality movement, examining the need and methods for control and accountability as well as the continuous improvement approach." John Ovretveit, The Karolinska Institute Medical Management Centre, Stockholm, Sweden “This excellent book is both informative and challenging...[it] helps us work our way through the contradictory and often inconsistent health maze that is bound by quality, risk, control, governance, trust, regulation, private activity, accountability, assurance and outcome.” Adam Graycar, Cabinet Office of South Australia This book explores the concepts of trust, control and risk management as key components of organisational accountability in the public sector. It explores how the concept of risk management has been introduced into the public sector and how this has impacted on the definition of governance in the National Health Service. It also addresses the concept of controls assurance by placing it in the context of developments both in local health care management and central government. Key questions that are addressed include: ·How can devolved public sector organisations be held accountable? · What is the relationship between risk, control and governance? ·How do private sector ideas about governance translate into the provision of public health services? Quality, Risk and Control in Health Care is essential reading for health policy makers, health practitioners and professionals, as well as students and academics in the fields of health policy, health services management, social policy and public policy.

ISPE Baseline® Guide Quality Press

This open access book, published under a CC BY 4.0 license in the Pubmed indexed book series Handbook of Experimental Pharmacology, provides up-to-date information on best practice to improve experimental design and quality of research in non-clinical pharmacology and biomedicine.

Risk-based Software Validation HIMSS

Author D. H. Stamatis has updated his comprehensive reference book on failure mode and effect analysis (FMEA). This is one of the most comprehensive guides to FMEA and is excellent for professionals with any level of understanding.!--nl--This book explains the process of conducting system, design, process, service, and machine FMEAs, and provides the rationale for doing so. Readers will understand what FMEA is, the different types of FMEA, how to construct an FMEA, and the linkages between FMEA and other tools. Stamatis offer a summary of tools/methodologies used in FMEA along with a glossary to explain key terms and principles. The updated edition includes information about the new ISO 9000:2000 standard, the Six Sigma approach to FMEA, a special section on automotive requirements related to ISO/TS 16949, the “robustness” concept, and TE 9000 and the requirements for reliability and maintainability. Also includes FMEA forms and samples, design review checklist, criteria for evaluation, basic reliability formulae and conversion failure factors, guidelines for RPN calculations and designing a reasonable safe product, and diagrams, and examples of FMEAs with linkages to robustness.

ISPE GAMP® RDI Good Practice Guide Flatiron Books

Biocontamination Control for Pharmaceuticals and Healthcare outlines a biocontamination strategy that tracks bio-burden control and reduction at each transition in classified areas of a facility. The first edition of the book covered many of the aspects of the strategy, but the new official guidance signals that a roadmap is required to fully comply with its requirements. Completely updated with the newest version of the EU-GPM (EN17141) the new edition expands the coverage of quality risk management and new complete examples to help professionals bridge the gap between regulation and implementation. Biocontamination Control for Pharmaceuticals and Healthcare offers professionals in pharma quality control and related areas guidance on building a complete biocontamination strategy. Includes the most current regulations Contains three new chapters, including Application of Quality Risk Management and its Application in Biocontamination Control, Designing an Environmental Monitoring Programme, and Synthesis: An Anatomy of a Contamination Control Strategy Offers practical guidance on building a complete biocontamination strategy

Validation of Pharmaceutical Processes ASTM International

Pharmaceutical Microbiology: Essentials for Quality Assurance and Quality Control presents that latest information on protecting pharmaceutical and healthcare products from spoilage by microorganisms, and protecting patients and consumers. With both sterile and non-sterile products, the effects can range from discoloration to the potential for fatality. The book provides an overview of the function of the pharmaceutical microbiologist and what they need to know, from regulatory filing and GMP, to laboratory design and management, and compendia tests and risk assessment tools and techniques. These key aspects are discussed through a series of dedicated chapters, with topics covering auditing, validation, data analysis, bioburden, toxins, microbial identification, culture media, and contamination control. Contains the applications of pharmaceutical microbiology in sterile and non-sterile products Presents the practical aspects of pharmaceutical microbiology testing Provides contamination control risks and remediation strategies, along with rapid microbiological methods Includes bioburden, endotoxin, and specific microbial risks Highlights relevant case studies and risk assessment scenarios

All Art Is Propaganda Academic Press

Completely revised and updated to reflect the significant advances in pharmaceutical production and regulatory expectations, this third edition of Validation of Pharmaceutical Processes examines and blueprints every step of the validation process needed to remain compliant and competitive. The many chapters added to the prior compilation examine va

Good Informatics Practices (GIP) Module: Risk Management Elsevier

Thoroughly revised to include the latest industry developments, the Second Edition presents a comprehensive overview of computer validation and verification principles and how to put them into practice. To provide the current best practice and guidance on identifying and implementing improvements for computer systems, the text extensively reviews regulations of pharmaceuticals, healthcare products, blood processing, medical devices, clinical systems, and biotechnology. Ensuring that organizations transition smoothly to the new system, this guide explains how to implement the new GMP paradigm while maintaining continuity with current practices. In addition, all 24 case studies from the previous edition have been revised to reflect the new system.

Quality Risk Management in the FDA-Regulated Industry Royal Society of Chemistry

GAMP 5 provides pragmatic and practical industry guidance to achieve compliant computerized systems fit for intended use in an efficient and effective manner. This technical document describes a flexible risk-based approach to compliant GxP regulated computerized systems, based on scalable specification and verification. It points to the future of computer systems compliance by centering on principles behind major industry developments such as PQLI; ICH Q8, Q9, Q10; and ASTM E2500. This revolutionary Guide addresses the entire lifecycle of an automated system and its applicability to a wide range of information systems, lab equipment, integrated manufacturing systems, and IT infrastructures. It contains new information on outsourcing, electronic batch recording, end user applications (such as spreadsheets and small database applications), and patch management.

Foundations of Quality Risk Management John Wiley & Sons

Continuous pharmaceutical manufacturing is currently receiving much interest from industry and regulatory authorities, with the joint aim of allowing rapid access of novel therapeutics and existing medications to the public, without compromising high quality. Research groups from different academic institutions have significantly contributed to this field with an immense amount of published research addressing a variety of topics related to continuous processing. The book is structured to have individual chapters on the different continuous unit operations involved in drug substance and drug product manufacturing. A wide spectrum of topics are covered, including basic principles of continuous manufacturing, applications of continuous flow chemistry in drug synthesis, continuous crystallization, continuous drying, feeders and blenders, roll compaction and continuous wet granulation.The underlying theme for each of these chapters is to present to the reader the recent advances in modeling, experimental investigations and equipment design as they pertain to each individual unit operation. The book also includes chapters on quality by design (QbD) and process analytical technology (PAT) for continuous processing, process control strategies including new concepts of quality-by-control (QbC), real-time process management and plant optimization, business and supply chain considerations related to continuous manufacturing as well as safety guidelines related to continuous chemistry. A separate chapter is dedicated to discussing regulatory aspects of continuous manufacturing, with description of current regulatory environment quality/GMP aspects, as well as regulatory gaps and challenges. Our aim from publishing this book is to make it a valuable reference for readers interested in this topic, with a desire to gain a fundamental understanding of engineering principles and mechanistic studies utilized in understanding and developing continuous processes. In addition, our advanced readers and practitioners in this field will find that the technical content of Continuous Pharmaceutical Processing is at the forefront of recent technological advances, with coverage of future prospects and challenges for this technology.

Alarm Management for Process Control, Second Edition John Wiley & Sons

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.