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GAMP Good Practice Guide, 2005-01-01 **GMP im Labor** Nadine Frankenberg, 2022-07-15

Der perfekte Leitfaden für Neulinge und Quereinsteigerinnen zur Etablierung eines GMP konformen Qualitätssystems mit besonderem Augenmerk auf die speziellen Bedürfnisse im Labor Good Manufacturing Practice GMP oder gute Herstellungspraxis ist ein System zur Qualitätssicherung von Herstellungsprozessen und Produkten basierend auf nationalen und internationalen Gesetzen. Dabei beziehen sich die Richtlinien auf Produktionsabläufe und Umgebung in der Herstellung von Arzneimitteln und Wirkstoffen, aber auch bei Kosmetika, Lebens- und Futtermitteln. GMP im Labor beschließt sich mit der Umsetzung von GMP Anforderungen in der Laborpraxis und präsentiert eine Grundlage für die Etablierung eines GMP konformen Qualitätssystems. Besonders nützlich für Einsteigerinnen werden die speziellen Bedürfnisse im Laboralltag detailliert beleuchtet und praktische Lösungssansätze geboten. Dabei wird genauer auf die gesetzlichen Grundlagen und Abgrenzung zu anderen Systemen GLP, GCP, ISO etc. eingegangen und die verschiedenen Punkte eines Qualitätssystems wie etwa Dokumentation, Validierung und Risikomanagement beleuchtet. Des Weiteren werden die fundamentalen Prozesse in einem GMP konformen Qualitätssystem wie etwa Gerätequalifizierung, Abweichungen und Selbstinspektionen behandelt und mit praktischen Lösungssansätzen präsentiert. Dringend benannt: Das einzige Buch, das einen umfassenden und doch komprimierten Einblick in die komplexe GMP-Thematik bietet. Praxisnah. Es wird besondere Rücksicht auf die speziellen Bedürfnisse im Labor genommen und bietet praktische Lösungssansätze. Leitfaden. Behandelt die notwendigen Grundlagen und dient als Leitfaden zur Etablierung eines GMP konformen Qualitätssystems. GMP im Labor richtet sich an Studierende in Bachelor- und Masterstudiengängen, Doktorandinnen, Naturwissenschaftlerinnen sowie Mitarbeiterinnen in Laboratorien und in der Industrie.

IT-Compliance Michael Rath, Rainer Sponholz, 2009 **Practical Approaches to Method Validation and Essential Instrument Qualification** Chung Chow Chan, Herman Lam, Xue-Ming Zhang, 2011-03-01

Practical approaches to ensure that analytical methods and instruments meet GMP standards and requirements. Complementing the authors' first book *Analytical Method Validation and Instrument Performance Verification*, this new volume provides coverage of more advanced topics focusing on additional and supplemental methods, instruments and electronic systems that are used in pharmaceutical, biopharmaceutical and clinical testing. Readers will gain new and valuable insights that enable them to avoid common pitfalls in order to seamlessly conduct analytical method validation as well as instrument operation qualification and performance verification. Part 1: Method Validation begins with an overview of the book's risk-based approach to phase appropriate validation and instrument qualification; it then focuses on the strategies and requirements for early phase drug development including validation of specific techniques and functions such as process analytical technology, cleaning validation and validation of laboratory information management systems. Part 2: Instrument Performance Verification explores the underlying principles and techniques for verifying instrument performance; coverage includes analytical instruments that

are increasingly important to the pharmaceutical industry such as NIR spectrometers and particle size analyzers and offers readers a variety of alternative approaches for the successful verification of instrument performance based on the needs of their labs At the end of each chapter the authors examine important practical problems and share their solutions All the methods covered in this book follow Good Analytical Practices GAP to ensure that reliable data are generated in compliance with current Good Manufacturing Practices cGMP Analysts scientists engineers technologists and technical managers should turn to this book to ensure that analytical methods and instruments are accurate and meet GMP standards and requirements

GAMP Good Practice Guide, 2011 This GAMP Good Practice Guide A Risk Based Approach to GxP Process Control Systems is a revision of the GAMP Good Practice Guide Validation of Process Control Systems It provides guidance and examples on the application of the principles and framework of GAMP 5 A Risk Based Approach to Compliant GxP Computerized Systems to a wide range of systems from basic instruments to large complex distributed control systems This Guide aims to achieve process control systems that are fit for intended use and compliant with applicable regulations providing recommended good practice based on a life cycle approach for the development maintenance and management of process control systems The Guide applies science based Quality Risk Management as described in ICH Q9 and GAMP 5 It describes the system life cycle from concept to retirement providing a high level overview of the approach together with guidance on how activities might be scaled based on risk to product quality system novelty and complexity as well as other project specific factors EU Annex 11 Guide to Computer Validation Compliance for the Worldwide Health Agency GMP Orlando Lopez, 2015-04-06 Good Manufacturing Practice GMP ensures medicinal products are produced consistently and controlled to the quality standards appropriate for their intended use and as required by product specifications or marketing authorization Annex 11 details the European Medicines Agency EMA GMP requirements for computer systems The purpose of Annex 11 is **Das kleine QM-Lexikon** Norbert Hochheimer, 2011 Das Qualitätsmanagement mit seinen verschiedenen Ausprägungen wie GLP GCP GMP oder ISO 9000 ist heute aus Industrie und Labor nicht mehr wegzudenken Jeder der in der Praxis damit zu tun hat muss sich mit der genauen Bedeutung der QM Fachbegriffe auseinandersetzen Obendrein wird er mit zahlreichen Abkürzungen konfrontiert Hier hilft die vollständig aktualisierte und bearbeitete Auflage des kompakten Lexikons jedem weiter der sich schnell und präzise informieren möchte Der Autor der selbst über praktische QM Erfahrung in der Industrie verfügt hat rund 1500 wichtigste Begriffe und Abkürzungen gut verständlich erklärt und erläutert Stimmen zur Voraufgabe Wer in Industrie und Laboratorium mit Qualitätsmanagement zu tun hat kann von diesem kompakten Lexikon profitieren Bibliographie Fachwörterbuch Wirtschaft Alles in allem ein sehr empfehlenswertes Nachschlagewerk das sich alsbald einen bevorzugten Platz neben dem Schreibtisch erobern sollte J. Ehreke *Materials and Corrosion* 54 2003 *GAMP Good Practice Guide*, 2010 *Quality assurance of pharmaceuticals: a compendium of guidelines and related materials. Volume 2. Good manufacturing practices and inspection* World Health Organization, 2024-01-31 The GMP

Compendium for Medical Products is a valuable resource for manufacturers regulators and other stakeholders involved in producing and distributing medical products It covers various topics from quality management systems to personnel hygiene equipment validation and complaint handling The guidance provided is based on the latest scientific and technical knowledge and considers the evolving regulatory landscape and the challenges faced by the industry

Validation of Chromatography Data Systems Robert D McDowall,2016-11-23 Guiding chromatographers working in regulated industries and helping them to validate their chromatography data systems to meet data integrity business and regulatory needs This book is a detailed look at the life cycle and documented evidence required to ensure a system is fit for purpose throughout the lifecycle Initially providing the regulatory data integrity and system life cycle requirements for computerised system validation the book then develops into a guide on planning specifying managing risk configuring and testing a chromatography data system before release This is followed by operational aspects such as training integration and IT support and finally retirement All areas are discussed in detail with case studies and practical examples provided as appropriate The book has been carefully written and is right up to date including recently released FDA data integrity guidance It provides detailed guidance on good practice and expands on the first edition making it an invaluable addition to a chromatographer s book shelf

The Computer System Risk Management and Validation Life Cycle R. Timothy Stein,2006

Handbook of Validation in Pharmaceutical Processes, Fourth Edition James Agalloco,Phil DeSantis,Anthony Grilli,Anthony Pavell,2021-10-28 Revised to reflect significant advances in pharmaceutical production and regulatory expectations Handbook of Validation in Pharmaceutical Processes Fourth Edition examines and blueprints every step of the validation process needed to remain compliant and competitive This book blends the use of theoretical knowledge with recent technological advancements to achieve applied practical solutions As the industry s leading source for validation of sterile pharmaceutical processes for more than 10 years this greatly expanded work is a comprehensive analysis of all the fundamental elements of pharmaceutical and bio pharmaceutical production processes Handbook of Validation in Pharmaceutical Processes Fourth Edition is essential for all global health care manufacturers and pharmaceutical industry professionals Key Features Provides an in depth discussion of recent advances in sterilization Identifies obstacles that may be encountered at any stage of the validation program and suggests the newest and most advanced solutions Explores distinctive and specific process steps and identifies critical process control points to reach acceptable results New chapters include disposable systems combination products nano technology rapid microbial methods contamination control in non sterile products liquid chemical sterilization and medical device manufacture

Pharmaceutical Computer Systems Validation Guy Wingate,2016-04-19 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

Practical Fermentation Technology Brian McNeil, Linda Harvey, 2008-04-15 A hands on book which begins by setting the context defining fermentation and the possible uses of fermenters and setting the scope for the book It then proceeds in a methodical manner to cover the equipment for research scale fermentation labs the different types of fermenters available their uses and modes of operation Once the lab is equipped the issues of fermentation media preservation strains and strain improvement strategies are documented along with the use of mathematical modelling as a method for prediction and control Broader questions such as scale up and scale down process monitoring and data logging and acquisition are discussed before separate chapters on animal cell culture systems and plant cell culture systems The final chapter documents the way forward for fermenters and how they can be used for non manufacturing purposes A glossary of terms at the back of the book along with a subject index will prove invaluable for quick reference Edited by academic consultants who have years of experience in fermentation technology each chapter is authored by experts from both industry and academia Industry authors come from GSK UK DSM Netherlands Eli Lilly USA and Broadley James UK USA

Data Integrity and Data Governance Robert D McDowall, 2018-11-06 Data integrity is the hottest topic in the pharmaceutical industry Global regulatory agencies have issued guidance after guidance after guidance in the past few years most of which does not offer practical advice on how to implement policies procedures and processes to ensure integrity These guidances state what but not how Additionally key stages of analysis that impact data integrity are omitted entirely The aim of this book is to provide practical and detailed help on how to implement data integrity and data governance for regulated analytical laboratories working in or for the pharmaceutical industry It provides clarification of the regulatory issues and trends and gives practical methods for meeting regulatory requirements and guidance Using a data integrity model as a basis the principles of data integrity and data governance are expanded into practical steps for regulated laboratories to implement The author uses case study examples to illustrate his points and provides instructions for applying the principles of data integrity and data governance to individual laboratory needs This book is a useful reference for analytical chemists and scientists management and senior management working in regulated laboratories requiring either an understanding about data integrity or help in implementing practical solutions Consultants will also benefit from the practical guidance provided

Pharmaceutical Manufacturing Handbook Shayne Cox Gad, 2008-03-11 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 *GMP-Qualifizierung und Validierung von Wirkstoffanlagen* Ralf Gengenbach, 2012-02-16 Unter Validierung bzw Qualifizierung versteht man die Beweisführung dass Verfahren Prozesse Ausrüstungsgegenstände Materialien Arbeitsgänge oder Systeme tatsächlich zu den erwarteten Ergebnissen führen Betroffen sind alle Unternehmen die Rohstoffe Halbfertig oder Fertigprodukte für medizinische Geräte Pharmazeutika Diagnostika Lebensmittel herstellen Ebenso sind Labore betroffen die Dienstleistungen anbieten deren Ergebnisse direkt in den Herstellungsprozess einfließen Dieses Buch liefert harte Fakten hinsichtlich der Durchführung How to do von praxiserprobten Qualifizierungs und Validierungsmaßnahmen ein Must have für Wirkstoff und Arzneimittelhersteller sowie deren Zulieferer Der deutsche Titel zur Validierung und Qualifizierung *Good Manufacturing Practices for Pharmaceuticals, Seventh Edition* Graham P. Bunn, 2019-02-04 This book provides insight into the world of pharmaceutical quality systems and the key elements that must be in place to change the business and organizational dynamics from task oriented procedure based cultures to truly integrated quality business systems that are self detecting and correcting Chapter flow has been changed to adopt a quality systems organization approach and supporting chapters have been updated based on current hot topics including the impact of the worldwide supply chain complexity and current regulatory trends Key Features Presents insight into the world of pharmaceutical quality systems Analyzes regulatory trends and expectations Includes approaches and practices used in the industry to comply with regulatory requirements Discusses recent worldwide supply chain issues Delivers valuable information to a worldwide audience regarding the current GMP practices in the industry **Computer Systems Validation** Guy Wingate, 2003-12-18 Both pervasive and ubiquitous computerized systems are now an integral component of every corporate strategy in pharmaceutical and healthcare companies However when technology is combined with high risk public safety projects or the production and control of life saving medicines or devices it is necessary to ensure that it is reliable quality Implementing ISO/IEC 17025:2005 Bhavan "Bob" Mehta, 2013-04-16 The purpose of this book is to demystify the requirements delineated within ISO IEC 17025 2005 while providing a road map for organizations that wish to receive maintain accreditation for their laboratories AS9100 ISO 9001 and ISO 13485 are standards that support the development and implementation of effective approaches to quality management and are recognized blueprints for the establishment of a quality management system QMS for diverse industries Although similar to these recognized QMS standards ISO IEC 17025 serves a unique purpose laboratory accreditation It is not unusual for laboratories to retain dual certification to ISO 9001 and ISO IEC 17025

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