



I. Objective

To ensure that the Quality Manual, Procedures, Forms and other internally generated quality management system documents are under control. This includes creation, amendment, deletion, review, approval, maintenance and distribution.

II. Scope

This procedure defines the actions and responsibilities of the Requester, Process Owners, Document Controller and Approving Authorities in the control of new, revised and deleted documents.

III.

Responsibilities

Requester,
Process Owner

QMR

QMR

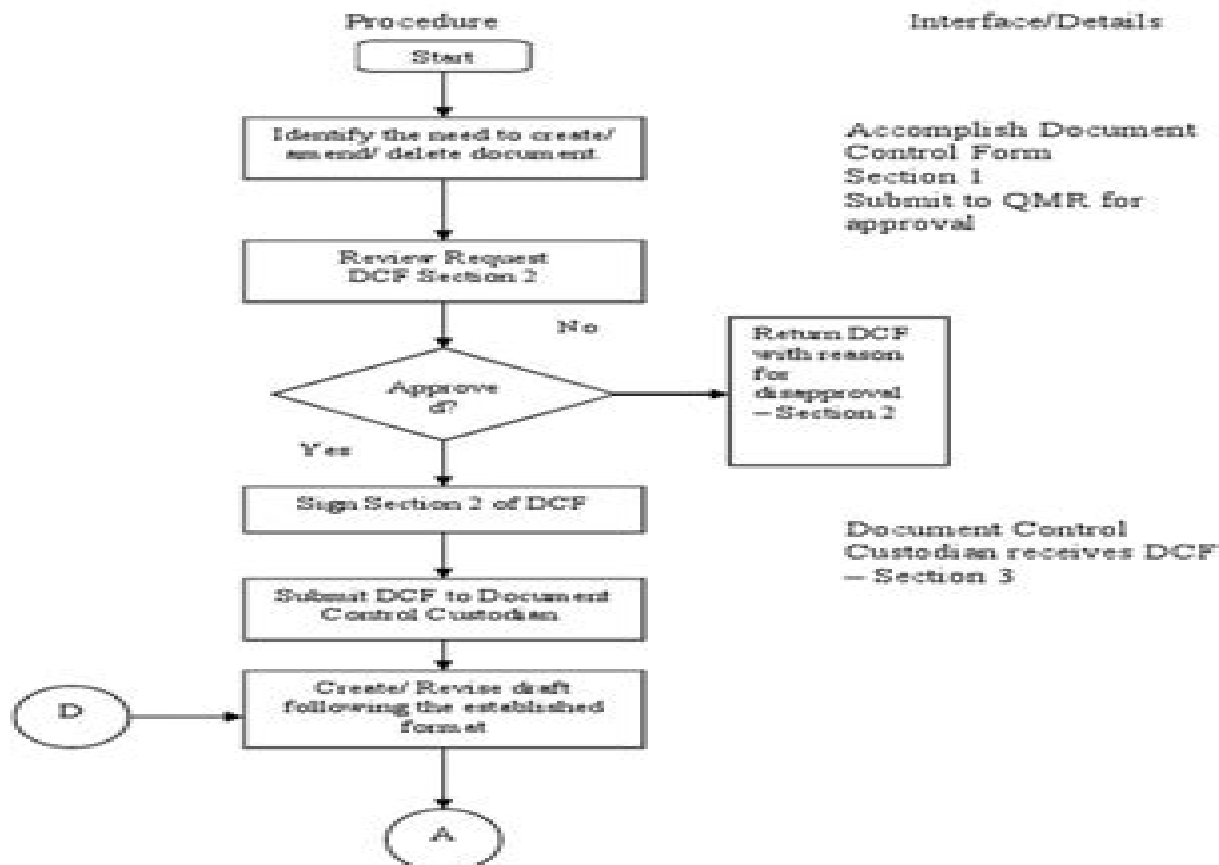
QMR

Document Control
Custodian

Process Owner

Procedure

Interface/Details



Prepared by:

Quality Management Representative

Approved by:

Senior Manager / Top Management Representatives

Sample Document Control Procedure

William C. Culley



Sample Document Control Procedure:

Document Control Denise Robitaille, 2011-09 They are supposed to be useful tools but whether they are printouts computer files flowcharts or forms documents can often give more headaches than help And yet without them most organizations couldn't function ISO 9001 and other quality management systems place great emphasis on documents and for good reason Documents aren't individual stand alone elements of the management process They are interrelated formatted in different media and controlled by various and distinct functions Keeping critical information current and in the right hands requires more than just signing off on procedures Document control is essential but where should you begin Inside you will find clear explanations about the document control process as well as practical solutions for creating organizing and maintaining documents including A discussion of different kinds of documents including electronic media and QMS requirements Identifying and defining responsibility Understanding the relationship between documents and records Tips for document writers Managing and maintaining documents Issues of accessibility Handling revisions and deviations Writing document control procedures

How to Establish a Document Control System for Compliance with ISO 9001:2015, ISO 13485:2016, and FDA Requirements Stephanie L. Skipper, 2015-10-14 This book explains the requirements for compliance with FDA regulations and ISO standards 9001 13485 for documented information controls and presents a methodology for compliance The document control system DCS or documented information control system DICS is the foundation of a quality management system It is the first quality system element that must be implemented because the establishment and control of documented processes and information in a quality controlled environment is dependent on the ability to proactively manage access to documents and the movement of documents through the document life cycle A well developed document control system benefits business by Improving knowledge retention and knowledge transfer within and across business units Improving access to knowledge based information Improving employee performance by providing standardized processes and communicating clear expectations Improving customer communication and satisfaction by providing documented information from which common understanding can be achieved Providing traceability of activities and documentation throughout the organization Improving organization of and access to documents and data Sample documents are included in the appendixes of this book to help clarify explanations and a full set of formatted procedures and document templates are available for download to get you off to an even faster start This book provides a process based approach that can be used for controlling all forms of documented information that are required to be managed under the quality management system

Environmental and Quality Systems Integration William C. Culley, 2019-09-17 Global competition corporate downsizing and corporate restructuring have forced many firms to reevaluate their operating methods Today corporations must do more with less while still watching the bottom line and improving profitability ISO 14000 and ISO 9000 because of their similar management system requirements and auditing procedures are going to become ISO 14001 Step by Step Naeem Sadiq, Asif

Hayat Khan,2019-01-22 This book provides practical advice on how to achieve compliance with ISO 14001 2015 the international standard for an EMS environmental management system With an EMS certified to ISO 14001 you can improve the efficiency of your business operations and fulfil compliance obligations while reassuring your employees clients and other stakeholders that you are monitoring your environmental impact This easy to follow guide takes a step by step approach and provides many sample documents to help you understand how to record and monitor your organisation s EMS processes Ideal for compliance managers IT and general managers environmental officers auditors and trainers this book will provide you with The confidence to plan and design an EMS Detailed descriptions of the ISO 14001 2015 requirements will give you a clear understanding of the standard even if you lack specialist knowledge or previous experience Guidance to build stakeholder support for your EMS Information on why it is important for an organisation to have an environmental policy and a sample communications procedure will help you to raise awareness of the benefits of implementing an EMS andAdvice on how to become an ISO 14001 certified organisation The book takes a step by step approach to implementing an ISO 14001 compliant EMS Key features A concise summary of the ISO 14001 2015 requirements and how you can meet them An overview of the documentation needed to achieve ISO 14001 2015 accreditation Sample documents to help you understand how to record and monitor your organisation s environmental management processes New for the second edition Updated for ISO 14001 2015 including terms definitions and references Revised approach to take into account requirements to address risks and opportunities Your practical guide to implementing an EMS that complies with ISO 14001 2015 buy this book today to get the help and guidance you need *Guidelines for Risk Based Process Safety* CCPS (Center for Chemical Process Safety),2007-04-10 *Guidelines for Risk Based Process Safety* provides guidelines for industries that manufacture consume or handle chemicals by focusing on new ways to design correct or improve process safety management practices This new framework for thinking about process safety builds upon the original process safety management ideas published in the early 1990s integrates industry lessons learned over the intervening years utilizes applicable total quality principles i e plan do check act and organizes it in a way that will be useful to all organizations even those with relatively lower hazard activities throughout the life cycle of a company *Guidelines for Risk Based Process Safety* Center for Chemical Process Safety (CCPS),2010-08-26 *Guidelines for Risk Based Process Safety* provides guidelines for industries that manufacture consume or handle chemicals by focusing on new ways to design correct or improve process safety management practices This new framework for thinking about process safety builds upon the original process safety management ideas published in the early 1990s integrates industry lessons learned over the intervening years utilizes applicable total quality principles i e plan do check act and organizes it in a way that will be useful to all organizations even those with relatively lower hazard activities throughout the life cycle of a company **Guidance Manual for Collection of Samples During RCRA Inspections** Laura L. Gentile,1995 **The Laboratory Quality Assurance System** Thomas A. Ratliff,2005-02-25 Both the 17025 1999

standard and especially ANSI ISO ASQ 9001 2000 standard require that a laboratory document its procedures for obtaining reliable results The Laboratory Quality Assurance Manual details to the user how to prepare a new laboratory quality assurance manual which will be appropriate to use as a procedures manual for a particular laboratory a sales tool to attract potential customers a document that can be to answer regulatory questions and ultimately a tool to become a registered ISO 9001 2000 Lab and gain related certifications based on the standard The Laboratory Quality Assurance Manual Incorporates changes to ANSI ISO ASQ 9001 2000 pertaining to laboratories Provides blank forms used in preparing a quality manual Provides information on the interrelationship of ANSI ISO 17025 1999 and ANSI ISO ASQ 9001 2000 **Wiley**

Practitioner's Guide to GAAS 2008 Michael J. Ramos, 2007-12-04 The clearest easiest to use guide to understanding GAAS 2008 on the market fully updated The 2008 edition includes Risk Assessment Standards effective for December 31 2007 Financial Statement Audits The only reference that offers the most comprehensive coverage of generally accepted auditing standards practices and procedures Wiley Practitioner's Guide to GAAS 2008 explains and interprets the standards in clear easy to understand language Offering the accounting professional a clear accessible distillation of the official language of Statements on Standards for Attestation Engagements SSAEs and Statements on Standards for Accounting and Review Services SSARs this invaluable resource provides advice on exactly when and how to remain fully compliant with each Michael J Ramos Denver CO was an auditor with KPMG and now works as an author and consultant He is the author of How to Comply with Sarbanes Oxley Section 404 Assessing the Effectiveness of Internal Control Second Edition 0 471 74066 7 and The Sarbanes Oxley Section 404 Implementation Toolkit Practice Aids for Managers and Auditors 0 471 71225 6

Code of Federal Regulations, 1996 Special edition of the Federal Register containing a codification of documents of general applicability and future effect with ancillaries *Guidelines for the establishment and management of seed testing laboratories* Food and Agriculture Organization of the United Nations, International Seed Testing Association, 2023-06-19 This Joint ISTA and FAO Handbook provides comprehensive guidelines for the setting up and managing of seed testing laboratories including those that cater to small and medium scale seed enterprises and farmers cooperatives which operate in low input production systems The adherence to the guidelines by seed testing laboratories results in the use of uniform procedures across the board thereby facilitating inter laboratory transferability of data This harmonization enhances confidence in seed quality assurance mechanisms The publication is an update of a previous one that was also jointly published by both organizations in 1969 and revised in 1979 and 1983 Its utility is enhanced by a set of new up to date information that builds upon existing data The Handbook provides guidance on the range of seed testing equipment procedures and management systems that have become available in the several decades since the previous editions Seed testing is a critical component of the seed value chain and is essential for the production of quality seeds i.e. seeds that are alive can germinate and produce vigorous seedlings are healthy and come from lots that meet set thresholds for genetic and

analytical purity The Handbook covers all aspects of seed testing from laboratory design and equipment selection to sample preparation testing methods and data analysis Importantly the publication also addresses the challenges of testing seeds in low input systems and provides guidance on how to adapt procedures to these settings Therefore it serves as a reference material and training resource for everyone involved in seed quality assurance procedures in particular the personnel of seed testing laboratories and regulatory agencies seed producers and farmers The updated publication represents a significant improvement over previous guidelines as it incorporates advances in technology and provides guidance on a broader range of testing procedures The guidelines are designed to be adaptable to different contexts and seed types and include recommendations for quality management systems proficiency testing and inter laboratory comparisons which are essential for ensuring the accuracy and reliability of seed testing results

Best Practices for Environmental Project Teams

Stephen Massey, 2011-08-16 Best Practices for Environmental Project Teams provides project managers and their teams government managers and regulatory agencies with practical guidelines for continuously improving performance Project managers and team members can pick from a variety of chapter topics stated as Actions to address existing skill gaps with practical tools and guidelines

Integrating Business Management Processes Titus De Silva, 2020-08-18 Integrating Business Management Processes Volume 3 Harmonising Quality Food Safety and Environmental Processes 978 0 367 48547 4 Shelving Guide Business Management The backbone of any organisation is its management system It must reflect the needs of the organisation and the requirements of its customers Compliance with legal requirements and ethical environmental practices contributes towards the sustainability of the management system Whatever the state of maturity of the management this book one of three provides useful guidance to design implement maintain and improve its effectiveness and is intended to provide readers with practical how to methods for integrating quality safety and environmental management processes This volume sets out procedures and flowcharts to show how the integration of these processes can be achieved Separated into management procedures core procedures support procedures and assurance procedures and complemented by practical examples this book is an invaluable resource for complete systems development and integration This book along with its two companion volumes is a practical guide for real managers designed to help them manage their business more effectively and gain competitive advantage Titus De Silva is a consultant in management skills development pharmacy practice quality management and food safety and an advisor to the newly established National Medicines Regulatory Authority NMRA in Sri Lanka

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