



## Document Control Procedure Template

### 1. Purpose:

The purpose of this document control procedure is to ensure the effective management of documented information within [Organization Name], in accordance with ISO 9001:2015 requirements. This procedure establishes the processes for creating, reviewing, approving, distributing, and revising documents to maintain accuracy, consistency, and compliance with quality management standards.

### 2. Scope:

This procedure applies to all documented information generated, utilized, or maintained by [Organization Name] as part of its Quality Management System (QMS).

### 3. Responsibilities:

- **Quality Manager:** The Quality Manager is responsible for overseeing the implementation of this document control procedure and ensuring compliance with ISO 9001 requirements. Responsibilities include:
  - Establishing document control processes and procedures.
  - Providing training and guidance on document control requirements.
  - Monitoring the effectiveness of document control activities and initiating improvements as necessary.
- **Document Owners:** Document Owners are responsible for creating, updating, and maintaining documents within their area of expertise. Responsibilities include:
  - Ensuring that documents are accurate, up-to-date, and aligned with relevant standards and procedures.
  - Reviewing and approving documents before distribution.
  - Notifying the Quality Manager of any changes or updates to documents.
- **Document Controllers:** Document Controllers are responsible for managing the day-to-day activities related to document control. Responsibilities include:
  - Maintaining a central repository of all documents and records.
  - Ensuring that document numbering, naming, and version control are maintained consistently.

# Document Control Procedure Example

**William C. Culley**



## **Document Control Procedure Example:**

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

**Implementing ISO/IEC 17025:2017, Second Edition** Bob Mehta, 2019-02-21 The focus of this book is to demystify the requirements delineated within ISO IEC 17025 2017 while providing a road map for organizations wishing to receive accreditation for their laboratories AS9100 ISO 9001 2015 and ISO 13485 2016 are standards that have been created to 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 many diverse industries Similar to these recognized QMS standards ISO IEC 17025 2017 for laboratory accreditation serves a unique purpose It is not unusual for laboratories to retain dual certification in ISO 9001 2015 and ISO IEC 17025 2017 However ISO IEC 17025 2017 contains requirements specific to the laboratory environment that are not addressed by ISO 9001 2015 This book highlights those differences between ISO 9001 2015 and ISO IEC 17025 2017 while providing practical insight and tools needed for laboratories wishing to achieve or sustain accreditation to ISO IEC 17025 2017 For those currently or formerly accredited to the 2005 version of ISO IEC 17025 an appendix outlines the changes between the 2005 and 2017 versions of the standard Implementing ISO/IEC 17025:2017 Bhavan (Bob) Mehta, 2019-02-21 The focus of this book is to demystify the requirements delineated within ISO IEC 17025 2017 while providing a road map for organizations wishing to receive accreditation for their laboratories AS9100 ISO 9001 2015 and ISO 13485 2016 are standards that have been created to 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 many diverse industries Similar to these recognized QMS standards ISO IEC 17025 2017 for laboratory accreditation serves a unique purpose It is not unusual for laboratories to retain dual certification in ISO 9001 2015 and ISO IEC 17025 2017 However ISO IEC 17025 2017 contains requirements specific to the laboratory environment that are not addressed by ISO 9001 2015 This book highlights those differences between ISO 9001 2015 and ISO IEC 17025 2017 while providing practical insight and tools needed for laboratories wishing to achieve or sustain accreditation to ISO IEC 17025 2017 For those currently or formerly accredited to the 2005 version of ISO IEC 17025 an appendix outlines the changes between the 2005 and 2017 versions of the standard

**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 g

*Implementing Quality in Laboratory Policies and Processes* Donnell R. Christian Jr., Stephanie Drilling, 2009-11-24 In order to gain accreditation every laboratory must have a superior quality assurance program The keys to a successful program are the operational and technical manuals and associated documents which define the program and its various components Written by experts with global experience in setting up laboratories

**Implementing Quality in Labora ISO 9001:2015 for Small Businesses** Ray Tricker, 2016-10-04 Small businesses face many challenges today including the increasing demand by larger companies for ISO 9001 compliance a challenging task for any organisation and in particular for a small business without quality assurance experts on its payroll Ray Tricker has already guided hundreds of businesses through to ISO accreditation and this sixth edition of his life saving ISO guide provides all you need to meet the new 2015 standards ISO 9001 2015 for Small Businesses helps you understand what the new standard is all about and how to achieve compliance in a cost effective way Covering all the major changes to the standards this book provides direct accessible and straightforward guidance This edition includes down to earth explanations to help you determine what you need to enable you to work in compliance with and or achieve certification to ISO 9001 2015 a contextual explanation of ISO 9001 within the structure of ISO 9000 family of standards a detailed description of the structure of ISO 9001 2015 and its compliance with Annex SL coverage of the new requirements for Risk Management and Risk Analysis a guide to the costs involved in implementing ISO 9001 2015 and advice on how to control costs an example of a complete generic Quality Management System consisting of a Quality Manual plus a whole host of Quality Processes Quality Procedures and Word Instructions and access to a free software copy of these generic QMS files to give you a starting point from which to develop your own documentation This book is also supported with a complete bibliography containing abbreviations and acronyms as well as a glossary of terms This comprehensive text will provide you and your

small business with a complete guide on your way to ISO compliance      **ISO 9001:2008 for Small Businesses** Ray Tricker,2014-01-21 Small businesses face many challenges today including the increasing demand by larger companies for ISO compliance Compliance is a challenging task for any organisation and can often be time consuming and costly particularly for small businesses who are unlikely to have quality assurance experts on the payroll However it is still possible to achieve compliance without the need for expensive consultancy or training that takes you out of the office Ray Tricker has already guided hundreds of businesses through the challenge and this the 5th edition of his life saving ISO guide has been rewritten and refined following 5 years field use of working with the standard The one area that an organisation particularly a small business always wants to know is how much is it going to cost to implement and operate a QMS compliant with ISO 9001 2008 and is it going to be worth the trouble Due to popular demand Edition 5 now includes a brand new chapter on the cost of implementing ISO 9001 2008 This edition provides Relevant examples that put the concepts and requirements of the standard into a real life context Down to earth explanations to help you determine what you need to work in compliance with and or achieve certification to ISO 9001 2008 An example of a complete generic Quality Management System consisting of a Quality Manual plus a whole host of Quality Processes Quality Procedures and Work Instructions Access to a free software copy of this generic QMS files available from the author to give you a starting point from which to develop your own documentation ISO 9001 2008 is the most widely followed quality management standard and the rewards can be great opening up new business opportunities as well as bringing real improvements to your processes and outputs      **Complete Guide to ISO 14000** Richard Barrett Clements,1996 This guide offers an overview of ISO 14000 a newly developed set of international standards for environmental management Their benefits and effects on a wide variety of companies are discussed      *ISO 9000 Quality Systems Development Handbook* David Hoyle,1998 Quality system developers may be managers consultants or quality assurance staff who spend a lot of time away from their offices They need a quick reference guide to the many issues they face in day to day communications which current books do not seem to provide      *Green Profits* Nicholas P Cheremisinoff,Avrom Bendavid-Val,2001-05-17 Green Profits covers two tightly connected topics environmental management systems EMS and pollution prevention P2 in a single volume Authored by an environmental engineer and an economist planner Green Profits shows how to implement an EMS especially ISO 14001 so that it leads to profitable pollution prevention innovations and how to identify and implement pollution prevention measures in a sound strategic business framework Green Profits provides the knowledge and tools for enterprise managers to achieve the benefits of both EMS and P2 and to do so in ways that fit in with existing management systems in their enterprises Environmental management systems are planned and organized ways for an enterprise to manage its interactions with the environment in particular those interactions that consume resources degrade the environment and create human health risk Part I of Green Profits provides a thorough and practical understanding of the elements of EMSs in general and ISO 14001 in particular tools

and techniques for implementing an EMS and achieving ISO 14001 certification and help with getting the implementation process started Pollution prevention involves replacing process technologies that generate pollution with those that do not or that do so much less It focuses on improving production processes to minimize waste rather than treating effluents or emissions which add to costs Part II of Green Profits provides tools such as step by step guides to conducting a P2 audit and energy and material balances for identifying P2 opportunities in an enterprise examples of P2 practices in specific industry sectors and a set of tools for assessing potential P2 investments from a bottom line point of view With this New Handbook Bring your facility into compliance Improve your corporate image Reduce your company s environmental liabilities Identify and save millions of dollars from pollution prevention projects This New Handbook Includes A step by step approach to implementing ISO 14001 A step by step approach to implementing Pollution Prevention Contains nearly 100 useful charts and tables used by the experts in establishing environmental action plans gap analyses establishing an Environmental Management System Contains dozens of useful charts and calculation methods with examples for evaluating the costs and savings to your company in implementing Pollution Prevention Dozens of industry specific case studies that you can learn and profit from Shows you in stepwise fashion how project financing principles and environmental cost accounting methods when coupled with EMS can save your company money This New Handbook is unique because unlike other volumes that separately cover Environmental Management Systems and Pollution Prevention you have it all in one single volume written by Experts that are Practitioners

**Auditing Concepts and Methods** Douglas R. Carmichael, John J. Willingham, Carol A. Schaller, 1996 Instructor s Resource Manual

**Understanding and Implementing ISO 9000:2000** David L. Goetsch, Stanley Davis, 2002 As organizations that compete in the global marketplace seek to achieve peak performance worldclass quality and personnel improvement interest in ISO 9000 certification has gained worldwide momentum This practical resource and guide uses a step by step model to understanding the ISO 9000 2000 series of standards and explains how to use them in an organization Provides a detailed summary of similarities and differences between the earlier version and ISO 9000 2000 Features ISO INFO boxes throughout that show how material being discussed applies to an actual work setting Offers a Serialized Case Study after every chapter that applies the information from a given chapter to how it was used in a real setting in which ISO 9000 registration was pursued the story unfolds with each new chapter showing the problems strategies mistakes and achievements of a company going through the registration process Ideal as a step by step how to guide for use in organizations that are interested in ISO 9000 registration

**AICPA Audit and Accounting Manual**, 1993

**Managing Records for ISO 9000 Compliance** Eugenia K. Brumm, 1995 Brumm reviews and interprets each paragraph of the ANSI ISO ASQC Q9001 1994 standard that implies or specifically states a records requirement She explains how to plan develop and implement the various components of records management in order to help readers reach a higher first pass success rate She also shows how to reduce the number of hour

*Warehouse and Distribution Automation*

*Handbook* Nicholas D. Adams,1996 This practical guide to warehouse distribution automation techniques includes evaluation of automation packages strategic company plans system justification and equipment analysis testing training and follow up

Management Information Systems Patrick G. McKeown,Robert A. Leitch,1993      **AICPA Professional Standards** ,1996

**Environmental Management Quick and Easy** Joe Kausek,2007 The design and implementation of an ISO 14001 environmental management system EMS need not be complicated or costly and this book focuses on getting a basic yet effective EMS in place with minimal effort so that the organization can move quickly towards the environmental performance improvements that will be needed to meet the growing international demand for corporate environmental stewardship The real benefit of ISO 14001 is that it can significantly improve an organization s environmental performance while greatly improving its bottom line at the same time Unfortunately most companies that have implemented ISO 14001 have not yet moved beyond compliance and have not yet realized these dual benefits In order to support the goal of quick and easy ISO 14001 implementation dozens of tools checklists procedure templates and spreadsheets applicable to organizations of all sizes are provided on an accompanying CD ROM These tools not only speed the design and implementation of the EMS they also provide for efficient and effective ongoing maintenance of the system      **Manual on Quality Assurance for Installation and Commissioning of Instrumentation, Control and Electrical Equipment in Nuclear Power Plants** ,1989      **Codification of Statements on Auditing Standards** American Institute of Certified Public Accountants. Auditing Standards Board,1992

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