
Quality Procedure ISO 9001: 2008 – Control of Documents

1 Purpose

FablessSemi Inc¹ controls all documents that are required by our Quality Management System (QMS).

The purpose of this procedure is to define the process and responsibilities to ensure that internal documents, external and customer-supplied documents (standards, specifications, procedures, etc.), and data within FablessSemi Inc are controlled.

2 Scope

The requirements of this procedure apply to all drawings, procedures, work instructions, forms, etc. used within the QMS that affect the quality of products or services at FablessSemi Inc.

3 Revision History

3.1.1 Approval

This Quality Procedure has been approved for use throughout FablessSemi Inc:

Date of Initial Version: February 2nd, 2009
Name and Title of Approver: Any Name (Director)
Date of Approval: April 21st, 2009

This Quality Procedure was written and published by **Any Name** of FablessSemi Inc. Please contact any.name@fablesssemi.com with questions, comments and improvement ideas.

3.1.2 Revision History

The following revisions have been made to this procedure since its initial publication:

Revision Date	List of Changes	Author	Approval
02/02/09	First draft	PW	
10/03/09	Incorporated review comments from the QA team and up-issued	PW	
21/04/09	Issued as an example document and published to the Extranet Library at www.cognidox.com	PW	PW

4 Related Documents

All references to ISO 9001 in this procedure refer to the requirements of ISO 9001:2008.

¹ FablessSemi Inc is a fictional company. The domain www.fablesssemi.com is registered and controlled by Cognidox Limited (www.cognidox.com) as a demonstration of its document control software.

Sample Iso Document Control Procedure

Chad Kymal



Sample Iso Document Control Procedure:

How to Establish a Document Control System for Compliance with ISO 9001:2015, ISO 13485:2016, and FDA Requirements Stephanie L. Skipper, 2015-10-12 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 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 **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 and Advice 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

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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 The ISO 14000 EMS Audit Handbook Greg Johnson,2020-04-28 The ISO 14000 EMS Audit

Handbook is an innovative and cost effective approach for the Environmental Management System EMS audit to ISO 14001 The Handbook presents comprehensive strategies for conducting all phases of the EMS audit including effective assessment processes for determining improved environmental performance **Quick Reference To Quality System Procedure As**

Per ISO 15189:2022 Dr. Vinodkumar Swamy (V.K. Swamy),2025-10-29 This quick reference guide on Quality System Procedures is aligned with NABL ISO 15189 2022 providing a practical roadmap for medical laboratories seeking accreditation It details a structured Quality Management System QMS covering all laboratory processes from pre to post examination ensuring reliability accuracy and consistency Key procedures include personnel competency equipment

calibration sample security method validation internal and external quality control and participation in external quality programs The book emphasizes management roles in policy resource allocation and continuous improvement integrating risk management documentation and quality assurance principles Designed for practical implementation in the Indian context it supports laboratory professionals quality managers and auditors in achieving technical competence regulatory compliance and excellence in patient centered diagnostic services

The ITSM Process Design Guide Donna Knapp,2010-08-15 The ITSM Process Design Guide Developing Rengineering and Improving IT Service Management closes the knowledge gap by providing detailed guidance on assessing designing measuring and integrating ITSM processes The advice and techniques in this book apply unilaterally to every IT service provider and ITSM framework standard and maturity model

Digital Forensics Processing and Procedures David Lilburn Watson,Andrew Jones,2013-08-30 This is the first digital forensics book that covers the complete lifecycle of digital evidence and the chain of custody This comprehensive handbook includes international procedures best practices compliance and a companion web site with downloadable forms Written by world renowned digital forensics experts this book is a must for any digital forensics lab It provides anyone who handles digital evidence with a guide to proper procedure throughout the chain of custody from incident response through analysis in the lab A step by step guide to designing building and using a digital forensics lab A comprehensive guide for all roles in a digital forensics laboratory Based on international standards and certifications

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

The ISO/TS 16949 Auditor

Handbook Chad Kymal, 2007

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