WASHINGTON PROCESS GROUP

ISO 9001:2015 DIY Implementation Kit

Sample Preview

A structured DIY kit for organizations building or improving a Quality Management System aligned to ISO 9001:2015.

SAMPLE PREVIEW — LIMITED CONTENT

This is a limited sample preview. The full kit includes complete templates, trackers, checklists, and implementation support materials.

FULL KIT CONTENTS

What's Included in the Full Kit

The ISO 9001:2015 DIY Implementation Kit includes structured tools and support materials organized around the full QMS implementation process — from initial gap assessment through internal audit preparation and certification readiness. The following is a high-level overview of what the full kit contains.

ISO 9001:2015 Implementation Roadmap	A phase-by-phase implementation roadmap that walks your team through the logical sequence for building a conforming QMS from the ground up.
QMS Readiness Checklist	A structured checklist covering all ISO 9001:2015 clause requirements — organized so your team can assess and track readiness status across each area.
Clause-by-Clause Gap Assessment Support	Gap assessment tools mapped to the ISO 9001:2015 clause structure, helping your team identify what is in place, what is missing, and what needs to be developed.
Quality Manual Support Materials	Support materials and structure for developing a Quality Manual aligned to ISO 9001:2015 requirements and customized to your organization.
Policy and Procedure Templates	Core quality policies and operating procedure templates aligned to ISO 9001:2015 clause requirements — ready to customize for your organization.
Internal Audit Tools	Internal audit planning, checklist, and tracking tools to support a structured internal audit program prior to certification.
Corrective Action and Nonconformance Tools	Tools for identifying, documenting, tracking, and closing corrective actions and nonconformances in a format aligned to ISO 9001:2015 Clause 10.
Management Review Support Materials	Agenda templates, input/output tracking, and documentation support for conducting management reviews that meet ISO 9001:2015 Clause 9.3 requirements.
QMS Implementation Tracker	A master tracker for monitoring implementation progress across all clause areas, owners, and target completion dates.
Audit Readiness Support Tools	Checklists and tools to help your team prepare for external certification body audits and stage reviews.

Note: This sample preview shows the format and structure of selected kit components only. Full templates, complete trackers, checklists, and all support materials are included in the purchased kit.

SAMPLE — QMS READINESS CHECKLIST

Sample QMS Readiness Checklist

The full kit includes a structured readiness checklist covering all ISO 9001:2015 clause requirements. The sample below shows the format and column structure only — it does not represent the complete checklist.

ISO 9001 Area	Sample Requirement	Example Evidence	Status	Notes
Context of the Organization (Cl. 4)	Determine internal and external issues relevant to the QMS	Context analysis document, SWOT or stakeholder register	In Progress	
Leadership (Cl. 5)	Top management shall demonstrate leadership and commitment to the QMS	Quality policy, signed management review records	Not Started	
Planning (Cl. 6)	Identify risks and opportunities and plan actions to address them	Risk and opportunity register	In Progress	
Support (Cl. 7)	Determine and maintain competence for personnel affecting QMS performance	Job descriptions, training records, competency matrix	Complete	
Operation (Cl. 8)	Plan, implement, and control processes needed to meet requirements	Process documentation, work instructions, SOPs	Not Started	
Performance Evaluation (Cl. 9)	Monitor, measure, analyze, and evaluate QMS performance	KPI reports, customer satisfaction data, internal audit records	In Progress	
Improvement (Cl. 10)	Identify and address nonconformities and implement corrective actions	Corrective action log, nonconformance reports	Not Started	

SAMPLE ONLY — The full kit checklist covers all ISO 9001:2015 clauses and sub-clauses across the complete standard. This preview shows format and column structure only.

SAMPLE — GAP ASSESSMENT TRACKER

Sample Gap Assessment Tracker

The full kit includes a structured gap assessment tracker organized by ISO 9001:2015 clause area. The sample below shows the format and column structure only.

Clause Area	Requirement Summary	Current State	Gap Identified	Action Needed	Owner	Status
4.1 Context	Understand the organization and its context	No formal analysis documented	Yes	Develop context analysis document	Quality Lead	Not Started
5.2 Quality Policy	Establish, implement, and maintain a quality policy	Draft policy exists but not formally approved	Yes	Finalize, approve, and communicate quality policy	Management	In Progress
6.1 Risk & Opportunity	Address risks and opportunities in QMS planning	No risk register in place	Yes	Develop risk and opportunity register	Quality Lead	Not Started
7.2 Competence	Ensure personnel are competent based on education, training, and experience	Training records exist but not systematically maintained	Partial	Develop competency matrix and training log	HR / Quality	In Progress
8.1 Operational Planning	Plan, implement, and control processes to meet requirements	Key processes documented as SOPs	No	Review SOPs for clause alignment	Operations	Complete
9.2 Internal Audit	Conduct internal audits at planned intervals	No internal audit program established	Yes	Develop audit program, schedule, and checklist	Quality Lead	Not Started
10.2 Corrective Action	React to nonconformities and take corrective action	Informal issue tracking via email only	Yes	Implement formal corrective action log	Quality Lead	Not Started

SAMPLE ONLY — The full gap assessment tracker covers all ISO 9001:2015 clause areas and sub-requirements. This preview shows format and column structure only.

SAMPLE — INTERNAL AUDIT TRACKER

Sample Internal Audit Tracker

The full kit includes structured internal audit tools — including a planning template, audit checklist, and finding tracker. The sample below shows the format only.

Audit Area	Audit Question	Evidence Reviewed	Finding Type	Action Required	Owner	Status
Quality Policy (Cl. 5.2)	Is the quality policy documented, approved, and communicated to relevant staff?	Quality policy document, staff acknowledgment records	Minor NC	Formalize staff communication and acknowledgment process	Management	Open
Competence (Cl. 7.2)	Are training records maintained and up to date for all relevant personnel?	Training log, job descriptions	Observation	Establish annual competency review process	HR	Open
Document Control (Cl. 7.5)	Are controlled documents identified with version, date, and approval status?	Procedure documents, document register	Conformance	No action required	Quality Lead	Closed
Internal Audit (Cl. 9.2)	Is there a documented internal audit program with a defined schedule?	Audit schedule, audit records	Major NC	Develop and implement a documented audit program	Quality Lead	Open
Corrective Action (Cl. 10.2)	Are corrective actions documented with root cause analysis and verification?	Corrective action log	Minor NC	Add root cause and verification fields to CA log	Quality Lead	Open

SAMPLE ONLY — The full internal audit tracker covers all planned audit areas across the QMS. This preview shows format, column structure, and example entries only.

SAMPLE — CORRECTIVE ACTION TRACKER

Sample Corrective Action Tracker

The full kit includes a structured corrective action and nonconformance tracker aligned to ISO 9001:2015 Clause 10.2. The sample below shows the format only.

Issue / Nonconformance	Root Cause	Corrective Action	Owner	Target Date	Status
Internal audit found training records missing for 3 of 8 production staff	No standardized onboarding checklist requiring training record completion	Develop onboarding checklist; update training records for affected staff	HR Manager	2026-03-15	In Progress
Customer complaint: delivered report contained outdated version of procedure	Document control procedure not followed — document register not checked prior to delivery	Update document control procedure; conduct team refresher; add pre-delivery checklist step	Quality Lead	2026-02-28	Open
Management review not conducted in the past 12 months	No scheduled review cadence established in the QMS	Schedule quarterly management reviews; document agenda and output template	Top Management	2026-03-01	Open
Supplier did not meet delivery timeline on two consecutive orders	No formal supplier evaluation or performance criteria in place	Develop supplier evaluation form and performance criteria	Procurement	2026-04-01	Open
Nonconformance report from Stage 1 audit: no documented risk register	Risk assessment process not formalized	Develop risk and opportunity register; review at management meeting	Quality Lead	2026-02-15	Complete

SAMPLE ONLY — The full corrective action tracker includes all nonconformances identified across the QMS. This preview shows format, column structure, and example entries only.

SAMPLE — PROCEDURE TEMPLATE

Sample Template Preview: Document Control Procedure

SAMPLE PREVIEW

The following is a partial excerpt from the Document Control Procedure template included in the full kit. Actual procedure language must be customized to reflect your organization's specific processes, document types, and approval structure.

DOCUMENT CONTROL PROCEDURE — SAMPLE PREVIEW ONLY

1. Purpose

This procedure establishes requirements for the control of documented information within the [Organization Name] Quality Management System (QMS). The purpose is to ensure that documented information is properly created, reviewed, approved, distributed, and maintained in a current and accessible state, and that obsolete information is prevented from unintended use.

2. Scope

This procedure applies to all documented information required by the ISO 9001:2015 standard and all documented information determined by [Organization Name] to be necessary for the effective operation of the QMS. This includes policies, procedures, work instructions, forms, records, and externally originated documents.

3. Responsibilities

Quality Manager: Responsible for maintaining the master document register, ensuring documents are reviewed and approved prior to use, and controlling distribution of controlled documents.

Document Authors / Process Owners: Responsible for drafting, reviewing, and updating documents within their assigned process areas.

All Staff: Responsible for using only the current approved version of controlled documents and reporting obsolete documents found in use.

4. Sample Procedure Statements

4.1 All controlled documents shall be identified with a document number, title, revision level, effective date, and approval signature prior to distribution and use.

4.2 Prior to issuing a new document or revising an existing document, the Quality Manager or designated authority shall review and approve the content for adequacy and compliance with QMS requirements.

4.3 Obsolete documents shall be removed from all points of use or clearly identified as obsolete to prevent unintended use. Retained obsolete documents shall be marked accordingly and stored separately.

[Additional procedure sections covering document review cycles, revision history, external document control, and record retention requirements are included in the full template.]

SAMPLE ONLY — This is a partial excerpt for preview purposes. The full Document Control Procedure template and all other procedure templates are included in the purchased kit.

INTENDED AUDIENCE

Who This Kit Is For

The ISO 9001:2015 DIY Implementation Kit is designed for organizations that need a structured, practical path to QMS implementation and want to manage the work internally using templates, trackers, and implementation tools.

- Small businesses building a Quality Management System for the first time
- Service-based organizations preparing for ISO 9001:2015 certification
- Government contractors needing a conforming QMS to support contract requirements
- Consulting firms looking to establish a documented quality framework
- Organizations preparing to implement ISO 9001:2015 before engaging a certification body
- Teams that need templates, trackers, and structure before working with a consultant
- Organizations that want to organize and document their QMS internally

LIMITATIONS

What This Kit Does Not Replace

This kit is a structured DIY implementation and documentation support resource. It is important to understand what it is not intended to provide.

- Does not guarantee ISO 9001:2015 certification
- Does not replace a certification body audit (Stage 1 or Stage 2)
- Does not replace legal, technical, or professional advisory services
- Is a DIY implementation and documentation support kit — not a managed consulting engagement
- Does not write your policies or procedures — your team customizes the templates to your actual processes and operations

GET THE FULL KIT

Purchase the Complete ISO 9001:2015 DIY Implementation Kit

ISO 9001:2015 DIY Implementation Kit

Washington Process Group

Full Kit Price

$997

One-time purchase · Instant digital download · Self-paced

The full kit includes the complete QMS readiness checklist, clause-by-clause gap assessment tools, Quality Manual support materials, policy and procedure templates, internal audit tools, corrective action and nonconformance tools, management review support materials, QMS implementation tracker, and audit readiness support tools.

Visit washingtonprocessgroup.com/iso-9001.html to purchase the full ISO 9001:2015 DIY Implementation Kit.

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Full Kit Includes — Complete Components

ISO 9001:2015 Implementation Roadmap	Phase-by-phase guide from gap assessment to certification readiness
QMS Readiness Checklist	All ISO 9001:2015 clause requirements — complete
Clause-by-Clause Gap Assessment Support	Structured assessment tools mapped to the full ISO 9001:2015 standard
Quality Manual Support Materials	Structure and support materials for developing your organization's Quality Manual
Policy & Procedure Templates	Core quality policies and procedures aligned to ISO 9001:2015 requirements
Internal Audit Tools	Audit planning, checklist, and finding tracker for a conforming audit program
Corrective Action & Nonconformance Tools	Full tracking and documentation tools aligned to Clause 10.2
Management Review Support Materials	Agenda, input/output tracker, and records aligned to Clause 9.3
QMS Implementation Tracker	Master implementation progress tracker across all clause areas
Audit Readiness Support Tools	Preparation checklists and tools for Stage 1 and Stage 2 audit readiness

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