SKEDMIA

ISO 9001:2026

Quality management systems — Requirements

GUIDE

Document Master List

Controlled list of every document supplied in the ISO 9001:2026 kit, with its revision status and owner.

Document control

Document number	GDE-02	**Revision**	0
Document title	Document Master List	**Issue date**	[Date]
Document owner	Management Representative	**Next review**	[Date]
Management system	Quality Management System	**Classification**	Internal
Standard clauses	7.5	**Status**	For approval

	Name	Position	Signature / Date
Prepared by	[Name]	[Position]	
Reviewed by	[Name]	[Position]	
Approved by	[Name]	[Position]	

Rev.	Date	Description of change	Author
0	[Date]	First issue.	[Name]

Before you issue this document

Replace every highlighted placeholder such as [Organisation Name] with your own details, delete any clause that does not apply to your scope, and have the document approved by the roles named above. Record the approval in the revision history — an auditor will ask who approved it and when.

This master list records every document in the ISO 9001:2026 Complete Documentation Kit. It is the register required by the standard's documented-information clause: when [Organisation Name] adopts the kit, update the revision and issue-date columns as each document is approved, and add any document created locally.

How to use this list

Keep one controlled copy. Every time a document is revised, raise its revision number here and on the document's own cover page. An auditor will normally ask for this list first and sample from it.

Doc. No	Document title	Clause(s)	Owner
00 Read Me First			
GDE-01	Kit Guide and Implementation Roadmap	4.1, 5.1, 6, 8, 9, 10.1	Quality Manager
01 Manual			
MAN-01	Quality Management System Manual	4.1, 4.2, 4.3, 4.4, 5.1, 5.2, 5.3, 6.1, 6.2, 6.3, 7.1, 7.2, 7.3, 7.4, 7.5, 8.1, 8.2, 8.3, 8.4, 8.5, 8.6, 8.7, 9.1, 9.2, 9.3, 10.1, 10.2	Quality Manager
MAN-02	Scope and Boundaries of the Quality Management System	4.3, 4.4	Quality Manager
MAN-03	Process Map, Process Interaction Matrix and Process Specifications	4.4, 8.1	Quality Manager
02 Policies			
POL-01	Quality Policy	5.2	Managing Director
POL-02	Quality Culture and Ethical Behaviour Charter	5.1.1, 5.1.2, 7.1.4, 7.3	Managing Director
03 Procedures			
PRO-01	Context of the Organisation and Interested Parties	4.1, 4.2	Quality Manager
PRO-02	Risks and Opportunities	6.1, 6.1.1, 6.1.2, 6.1.3	Quality Manager
PRO-03	Objectives and Planning to Achieve Them	6.2	Quality Manager
PRO-04	Planning of Changes to the QMS Procedure	6.3, 5.3	Quality Manager
PRO-05	Competence, Training and Awareness	7.2, 7.3	HR Manager
PRO-06	Communication	7.4	Quality Manager
PRO-07	Control of Documented Information	7.5, 7.5.1, 7.5.2, 7.5.3	Quality Manager
PRO-08	Monitoring and Measuring Resources and Calibration Procedure	7.1.5	Quality Manager
PRO-09	Operational Planning and Control	8.1	Operations Manager
PRO-10	Customer Requirements, Contract Review and Customer Communication Procedure	8.2, 5.1.2	Sales Manager

Doc. No	Document title	Clause(s)	Owner
PRO-11	Design and Development Procedure	8.3	Design and Engineering Manager
PRO-12	Control of Externally Provided Processes, Products and Services Procedure	8.4, 8.1	Procurement Manager
PRO-13	Production and Service Provision Control Procedure	8.5.1, 8.5.2, 8.5.3, 8.5.4, 8.5.5, 8.5.6	Operations Manager
PRO-14	Release of Products and Services and Control of Nonconforming Outputs Procedure	8.6, 8.7	Quality Manager
PRO-15	Monitoring, Measurement, Analysis and Evaluation	9.1, 9.1.1, 9.1.3	Quality Manager
PRO-16	Customer Satisfaction, Feedback and Complaints Procedure	9.1.2, 9.1.3, 5.1.2, 8.2.1	Quality Manager
PRO-17	Internal Audit	9.2, 9.2.1, 9.2.2	Lead Internal Auditor
PRO-18	Management Review	9.3, 9.3.1, 9.3.2, 9.3.3	Managing Director
PRO-19	Nonconformity, Corrective Action and Continual Improvement	10.2, 10.1	Quality Manager
04 Forms, Registers & Templates			
FRM-01	Context and Issues Analysis	4.1, 4.2	Quality Manager
FRM-02	Roles, Responsibilities and Authorities Matrix	5.3, 5.1	Managing Director
FRM-03	Competence Matrix and Training Record	7.2	HR Manager
FRM-04	Communication Log and Suggestion Form	7.4	Quality Manager
FRM-05	Contract and Order Review Record	8.2.3, 8.2.4	Sales Manager
FRM-06	Design and Development Record Pack	8.3.2, 8.3.3, 8.3.4, 8.3.5, 8.3.6	Design and Engineering Manager
FRM-07	Production Records: Traceability, Property and Change Control	8.5.2, 8.5.3, 8.5.6	Operations Manager
FRM-08	Release, Nonconforming Output and Concession Report	8.6, 8.7	Quality Manager
FRM-09	Internal Audit Plan and Report	9.2, 9.2.2	Lead Internal Auditor
FRM-10	Management Review Minutes and Actions	9.3.2, 9.3.3	Managing Director
FRM-11	Document Change Request	7.5	Quality Manager
04 Forms, Registers & Templates			
REG-01	Interested Parties and Requirements Register	4.2	Quality Manager
REG-02	Statutory and Regulatory Requirements Register	8.2.2, 4.2, 5.1.2	Quality Manager
REG-03	Risk Register	6.1.1, 6.1.2, 9.1.3	Quality Manager

Doc. No	Document title	Clause(s)	Owner
REG-04	Opportunity Register	6.1.1, 6.1.3, 9.1.3, 9.3.2	Quality Manager
REG-05	Objectives and Action Plan Tracker	6.2	Quality Manager
REG-06	Resources, Infrastructure, Work Environment and Organizational Knowledge Register	7.1.1, 7.1.2, 7.1.3, 7.1.4, 7.1.6	Operations Manager
REG-07	Monitoring and Measuring Equipment Register and Calibration Record	7.1.5	Quality Manager
REG-08	External Provider Evaluation and Approved Provider Register	8.4.1, 8.4.2	Procurement Manager
REG-09	Nonconformity and Corrective Action Register	10.2	Quality Manager
05 Plans & Programmes			
PLN-01	QMS Implementation Plan	4.1, 6, 8, 9	Quality Manager
PLN-02	Annual Internal Audit Programme	9.2, 9.2.2	Lead Internal Auditor
PLN-03	Competence and Training Plan	7.2, 7.3	HR Manager
PLN-04	Communication Plan	7.4	Quality Manager
06 Checklists			
CHK-01	ISO 9001:2026 Gap Analysis Checklist	—	Quality Manager
CHK-02	ISO 9001:2015 to ISO 9001:2026 Transition Gap Checklist	4.1, 4.2, 5.1.1, 5.2, 5.3, 6.1.2, 6.1.3, 6.3, 7.1.3, 7.1.4, 7.1.6, 7.3, 8.1, 9.1.2, 9.1.3, 9.2.2, 9.3.2, 9.3.3, 10.1, 10.2	Quality Manager
CHK-03	Internal Audit Checklist	9.2	Lead Internal Auditor
CHK-04	Certification Readiness Checklist	—	Quality Manager
07 Internal Auditor Training Pack			
TRN-01	Internal Auditor Participant Handbook	9.2	Quality Manager
TRN-02	Trainer's Guide	9.2	Quality Manager
TRN-03	Presentation Slides Outline	9.2	Quality Manager
TRN-04	Exercises and Case Studies	9.2	Quality Manager
TRN-05	Examination	9.2	Quality Manager
TRN-06	Examination Answer Key and Marking Scheme	9.2	Quality Manager
TRN-07	Course Forms	9.2	Quality Manager
TRN-08	Certificate of Training Template	9.2	Quality Manager

Revision status

Record the approved revision of each document as it is issued. Rows are pre-filled at revision 0 (first issue).

Doc. No	Rev.	Issue date	Approved by	Next review
GDE-01	0	[Date]	[Name]	[Date]
MAN-01	0	[Date]	[Name]	[Date]
MAN-02	0	[Date]	[Name]	[Date]
MAN-03	0	[Date]	[Name]	[Date]
POL-01	0	[Date]	[Name]	[Date]
POL-02	0	[Date]	[Name]	[Date]
PRO-01	0	[Date]	[Name]	[Date]
PRO-02	0	[Date]	[Name]	[Date]
PRO-03	0	[Date]	[Name]	[Date]
PRO-04	0	[Date]	[Name]	[Date]
PRO-05	0	[Date]	[Name]	[Date]
PRO-06	0	[Date]	[Name]	[Date]
PRO-07	0	[Date]	[Name]	[Date]
PRO-08	0	[Date]	[Name]	[Date]
PRO-09	0	[Date]	[Name]	[Date]
PRO-10	0	[Date]	[Name]	[Date]
PRO-11	0	[Date]	[Name]	[Date]
PRO-12	0	[Date]	[Name]	[Date]
PRO-13	0	[Date]	[Name]	[Date]
PRO-14	0	[Date]	[Name]	[Date]
PRO-15	0	[Date]	[Name]	[Date]
PRO-16	0	[Date]	[Name]	[Date]
PRO-17	0	[Date]	[Name]	[Date]
PRO-18	0	[Date]	[Name]	[Date]
PRO-19	0	[Date]	[Name]	[Date]
FRM-01	0	[Date]	[Name]	[Date]
REG-01	0	[Date]	[Name]	[Date]
REG-02	0	[Date]	[Name]	[Date]
FRM-02	0	[Date]	[Name]	[Date]
REG-03	0	[Date]	[Name]	[Date]
REG-04	0	[Date]	[Name]	[Date]
REG-05	0	[Date]	[Name]	[Date]
REG-06	0	[Date]	[Name]	[Date]

Doc. No	Rev.	Issue date	Approved by	Next review
REG-07	0	[Date]	[Name]	[Date]
REG-08	0	[Date]	[Name]	[Date]
FRM-03	0	[Date]	[Name]	[Date]
FRM-04	0	[Date]	[Name]	[Date]
FRM-05	0	[Date]	[Name]	[Date]
FRM-06	0	[Date]	[Name]	[Date]
FRM-07	0	[Date]	[Name]	[Date]
FRM-08	0	[Date]	[Name]	[Date]
FRM-09	0	[Date]	[Name]	[Date]
FRM-10	0	[Date]	[Name]	[Date]
REG-09	0	[Date]	[Name]	[Date]
FRM-11	0	[Date]	[Name]	[Date]
PLN-01	0	[Date]	[Name]	[Date]
PLN-02	0	[Date]	[Name]	[Date]
PLN-03	0	[Date]	[Name]	[Date]
PLN-04	0	[Date]	[Name]	[Date]
CHK-01	0	[Date]	[Name]	[Date]
CHK-02	0	[Date]	[Name]	[Date]
CHK-03	0	[Date]	[Name]	[Date]
CHK-04	0	[Date]	[Name]	[Date]
TRN-01	0	[Date]	[Name]	[Date]
TRN-02	0	[Date]	[Name]	[Date]
TRN-03	0	[Date]	[Name]	[Date]
TRN-04	0	[Date]	[Name]	[Date]
TRN-05	0	[Date]	[Name]	[Date]
TRN-06	0	[Date]	[Name]	[Date]
TRN-07	0	[Date]	[Name]	[Date]
TRN-08	0	[Date]	[Name]	[Date]

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ISO 9001:2026

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Clause Cross-Reference Matrix

Shows which kit document answers each clause of ISO 9001:2026 — the table certification bodies ask for at stage 1.

Document control

Document number	GDE-03	**Revision**	0
Document title	Clause Cross-Reference Matrix	**Issue date**	[Date]
Document owner	Management Representative	**Next review**	[Date]
Management system	Quality Management System	**Classification**	Internal
Standard clauses	—	**Status**	For approval

	Name	Position	Signature / Date
Prepared by	[Name]	[Position]	
Reviewed by	[Name]	[Position]	
Approved by	[Name]	[Position]	

Rev.	Date	Description of change	Author
0	[Date]	First issue.	[Name]

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Use this matrix to demonstrate that every requirement of ISO 9001:2026 is addressed by documented information. Complete the final column with the evidence or record that proves the requirement is being met in practice, and take it to the stage 1 audit.

> **Good practice**
> Auditors use a matrix like this to plan their sample. A complete, honest matrix shortens stage 1 considerably; an aspirational one invites a deeper audit.

Clause	Kit document(s)	Evidence / records in use
4.1	CHK-02, FRM-01, GDE-01, MAN-01, PLN-01, PRO-01	
4.2	CHK-02, FRM-01, MAN-01, PRO-01, REG-01, REG-02	
4.3	MAN-01, MAN-02	
4.4	MAN-01, MAN-02, MAN-03	
5.1	FRM-02, GDE-01, MAN-01	
5.1.1	CHK-02, POL-02	
5.1.2	POL-02, PRO-10, PRO-16, REG-02	
5.2	CHK-02, MAN-01, POL-01	
5.3	CHK-02, FRM-02, MAN-01, PRO-04	
6	GDE-01, PLN-01	
6.1	MAN-01, PRO-02	
6.1.1	PRO-02, REG-03, REG-04	
6.1.2	CHK-02, PRO-02, REG-03	
6.1.3	CHK-02, PRO-02, REG-04	
6.2	MAN-01, PRO-03, REG-05	
6.3	CHK-02, MAN-01, PRO-04	
7.1	MAN-01	
7.1.1	REG-06	
7.1.2	REG-06	
7.1.3	CHK-02, REG-06	
7.1.4	CHK-02, POL-02, REG-06	
7.1.5	PRO-08, REG-07	
7.1.6	CHK-02, REG-06	
7.2	FRM-03, MAN-01, PLN-03, PRO-05	
7.3	CHK-02, MAN-01, PLN-03, POL-02, PRO-05	
7.4	FRM-04, MAN-01, PLN-04, PRO-06	
7.5	FRM-11, MAN-01, PRO-07	

Clause	Kit document(s)	Evidence / records in use
7.5.1	PRO-07	
7.5.2	PRO-07	
7.5.3	PRO-07	
8	GDE-01, PLN-01	
8.1	CHK-02, MAN-01, MAN-03, PRO-09, PRO-12	
8.2	MAN-01, PRO-10	
8.2.1	PRO-16	
8.2.2	REG-02	
8.2.3	FRM-05	
8.2.4	FRM-05	
8.3	MAN-01, PRO-11	
8.3.2	FRM-06	
8.3.3	FRM-06	
8.3.4	FRM-06	
8.3.5	FRM-06	
8.3.6	FRM-06	
8.4	MAN-01, PRO-12	
8.4.1	REG-08	
8.4.2	REG-08	
8.5	MAN-01	
8.5.1	PRO-13	
8.5.2	FRM-07, PRO-13	
8.5.3	FRM-07, PRO-13	
8.5.4	PRO-13	
8.5.5	PRO-13	
8.5.6	FRM-07, PRO-13	
8.6	FRM-08, MAN-01, PRO-14	
8.7	FRM-08, MAN-01, PRO-14	
9	GDE-01, PLN-01	
9.1	MAN-01, PRO-15	
9.1.1	PRO-15	
9.1.2	CHK-02, PRO-16	
9.1.3	CHK-02, PRO-15, PRO-16, REG-03, REG-04	
9.2	CHK-03, FRM-09, MAN-01, PLN-02, PRO-17, TRN-01, TRN-02, TRN-03, TRN-04, TRN-05, TRN-06, TRN-07, TRN-08	

Clause	Kit document(s)	Evidence / records in use
9.2.1	PRO-17	
9.2.2	CHK-02, FRM-09, PLN-02, PRO-17	
9.3	MAN-01, PRO-18	
9.3.1	PRO-18	
9.3.2	CHK-02, FRM-10, PRO-18, REG-04	
9.3.3	CHK-02, FRM-10, PRO-18	
10.1	CHK-02, GDE-01, MAN-01, PRO-19	
10.2	CHK-02, MAN-01, PRO-19, REG-09	

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