

D101: DEMO OF ISO 9001:2026 DOCUMENTS KIT **Price 499 USD**

Totally Editable Documentation Package (System manual, Procedures, Forms, Process Approach, Exhibits, Policies, Audit Checklist, etc.) For ISO 9001:2026 Quality Management System Certification and Quick Process Improvement

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Chapter-1.0 Contents of ISO 9001:2026 Quality Management System Documentation Kit (188 document files)

A. This Editable Documents Kit Has 10 Main Directories as Below:

Sr. No.	List of Directory	Details of Documents
1.	QMS Manual	14 Files in MS-Word
2.	Procedures	9 Procedures in MS-Word
3.	Process Approach	11 Files in MS-Word
4.	SOP	31 SOPs in MS-Word
5.	Exhibits	12 Exhibits in MS-Word
6.	Blank Formats /Templates to retain documented information Name of departments:	69 Blank formats in MS-Word / Excel
	SYS (Quality Management System)	18 Formats in MS-Word
	Training (TRG)	06 Formats in MS-Word
	Purchase (PUR)	06 Formats in MS-Word
	Marketing (MKT)	04 Formats in MS-Word
	Production (PRD)	07 Formats in MS-Word
	Quality Control (QCD)	03 Formats in MS-Word
	Design & Development (RND)	07 Formats in MS-Word
	Maintenance (ENG)	04 Formats in MS-Word
	Dispatch (DES)	02 Formats in MS-Word
	Store (STR)	05 Formats in MS-Word
	HRD	03 Formats in MS-Word
	Installation and servicing (SER)	04 Formats in MS-Word
7.	Filled Formats /Templates to retain documented information Name of departments:	42 formats in MS-Word
	SYS (Quality Management System)	09 Formats in MS-Word
	Training (TRG)	06 Formats in MS-Word
	Purchase (PUR)	03 Formats in MS-Word
	Marketing (MKT)	03 Formats in MS-Word
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	Design & Development (RND)	04 Formats in MS-Word
	Maintenance (ENG)	05 Formats in MS-Word
	Dispatch (DES)	01 Formats in MS-Word
	Store (STR)	02 Formats in MS-Word
	HRD	02 Formats in MS-Word
	Installation and servicing (SER)	04 Formats in MS-Word
8.	ISO 9001:2026 Audit Checklists (Clause wise & Department wise)	02 Files in MS-Word (Total 400+Questions)
9.	ISO 9001:2026 Compliance Matrix (Requirements wise reference documented information)	01 File in MS-Excel
10.	Sample MRM Documents	02 Files in MS-Word

Total 188 files in editable form for Quick Download by e-delivery

B. Documented Information Package:

Our documentation kit comprises sample documents required for ISO 9001:2026 Quality Management System implementation and certification. **All documents are in editable MS-Word / Excel format.** You can easily make changes in our sample documents as per your organization's needs and **within few days your entire documents** with all necessary system requirements can be made ready to use. In the ISO 9001:2026 Standard, documented information is required at few places. But for making the system better, we have provided many document templates from which a user can select documents to make own system with minor changes, as applicable. Though ISO 9001:2026 standard does not require manual, procedures, etc., it requires maintaining and/or retaining some documented information specified in various clauses of the standard. Therefore, this kit contains two types of documented information as listed below:

1. **Maintain documented information: QMS Manual, Policy, Procedures, etc.**
2. **Retain documented information: Records (Forms)- templates**

Under the main directories of this kit, further files are provided in MS Word/Excel document as per the details given below.

1. QMS Manual (10 Chapters and 4 Annexures):

This document is a sample copy of an organization's system manual which describes clause wise details on how ISO 9001:2026 Quality Management System is implemented. It covers the context of organization, scope, policy, objectives, and organization structure as well as macro level details of implementation of each requirement from clause no. 4 to 10 of ISO 9001:2026. This sample manual will also help you prepare company profile, amendment sheet, Quality policy statement and organization chart etc.

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(A) Table of Contents

Chapter No.	Subject	Page No.	ISO 9001:2026 Clause Reference
Section – 1			
1.	Company profile	1 – 3	-----
2.	Table of contents	1 – 2	-----
3.	Control and distribution	1 – 3	-----
Section – 2			
4.	Context of the Organization	1 – 4	4.0
	Understanding the organization and its context		4.1
	Understanding the needs and expectations of interested parties		4.2
	Determining the scope of the Quality Management System		4.3
	Quality Management System		4.4
5.	Leadership	1 – 4	5.0
	Leadership & Commitment		5.1
	Quality Policy		5.2
	Roles, responsibilities and authorities		5.3
6.	Planning	1 – 5	6.0
	Action to address risks and opportunities		6.1
	Quality objectives and planning to achieve them		6.2
	Planning of changes		6.3
7.	Support	1 – 5	7.0
	Resources		7.1
	Competence		7.2

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	Awareness		7.3
	Communication		7.4
	Documented information		7.5
8.	Operation	1 – 6	8.0
	Operational planning and control		8.1
	Requirements for products and services		8.2
	Design and development of products and services		8.3
	Control of externally provided processes, products and services		8.4
	Production and service provision		8.5
	Release of products and services		8.6
	Control of nonconforming outputs		8.7
9.	Performance evaluation	1 – 4	9.0
	Monitoring, measurement, analysis and evaluation		9.1
	Internal audit		9.2
	Management review		9.3
10.	Improvement	1 – 2	10.0
	Continual improvement		10.1
	Nonconformity and corrective action		10.2
Annexures			
ANX–I	List of Documented information	1 – 1	=====
ANX–II	Glossary of terms	1 – 1	=====
ANX–III	Company activity process flow chart	1 – 2	=====
ANX–IV	Organization structure	1 – 1	=====

2. Procedures (09 procedures):

Sample copies of necessary procedures are given as per ISO 9001:2026, covering all the details like purpose, scope, responsibility, activities to be followed as well as the list of exhibits, reference documents and formats, as applicable. The list of sample procedures provided is as below.

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List of Procedures

1. Procedure for Management of document
2. Procedure for corrective action
3. Procedure for internal quality management system audit
4. Procedure for management review meeting
5. Procedure for Risk, Biodiversity and Ecosystem Assessment
6. Procedure for Monitoring and Measuring Equipment control
7. Procedure for non-conformance Product Control
8. Procedure for QMS Training implementation in Organization
9. Procedure for Quality Culture & Ethical Behaviour

3. Process Approach (11 process flow charts):

List of Process Approach

1. Process flow chart of customer service
2. Process flow chart of dispatch
3. Process flow chart of engineering
4. Process flow chart of marketing
5. Process flow chart of production
6. Process flow chart of purchase
7. Process flow chart of quality control
8. Process flow chart of QMS coordinator processes
9. Process flow chart of research and development
10. Process flow chart of stores
11. Process flow for training activity

4. SOP (31 SOPs)

List of SOPs

1. Finished product handling
2. Awareness regarding environment, health and safety

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3. Review of country specific requirements
4. Personnel security
5. Change control
6. House keeping
7. Label control
8. Ware house operations
9. Raw material issue
10. Operation of Power Plant
11. Effluent treatment plant
12. Waste filling and transportation
13. Testing of raw materials
14. Chemical reaction
15. Filtration
16. Blending
17. Tray dryer
18. Product change over (cleaning and washing)
19. Product withdrawal
20. Process / Equipment related Change Control process
21. Research and development
22. General requirement of sampling
23. Diesel generating set
24. Steam boiler
25. Water softening plant
26. Air compressor
27. Thermic fluid heater
28. Chilling plant
29. Reverse osmosis plant
30. Hot air generator
31. Ice Plant

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5. Exhibits (12 Exhibits)

It covers Skill Requirements, Multi skill requirements, Document Identification and Codification System and Need and Expectations of interested parties, etc.

List of exhibits

1. Organization level QMS Skill Requirements
2. Department level Multi skill Identification
3. Disposal Of Non-Conforming Products
4. Operation control plan
5. Quality Plan – Incoming Inspection and Testing
6. Material Specifications
7. Exhibit for Document Identification and Codification System
8. Organization level Risk Management Sheet
9. Needs and Expectation of interested parties
10. Internal & External Communication Matrix
11. Department level Risk and Opportunity
12. Organization Knowledge Sharing Matrix

6. Blank Sample Formats For All Departments (69 sample formats)

This directory includes sample copy of blank forms that are required to maintain evidence/records as well as establish control and create a good system in the organization. The sample forms for various departments/areas can be used as a guide for the users to prepare own formats with minor changes as required by their organization. The organization is free to change the formats to suit their own requirements. A total of 62 blank formats is provided as per the list given below.

List of blank formats

1.	Master List of Documents	2.	Spray Dryer Log Sheet
3.	Change Control Note	4.	Spin Flash Dryer Log Sheet
5.	Equipment Calibration Status Report	6.	Reverse Osmosis Log Sheet
7.	Master List of Records	8.	Blender / Ball Mill Log Sheet

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9.	Quality Objective Monitoring Report	10.	Process change Request form
11.	ISO 9001 IQA Plan	12.	Equipment change Request form
13.	QMS Clause wise Audit compliance Report	14.	pH Meter Calibration Report
15.	QMS Internal Audit NCR Form	16.	Test Request Slip for Incoming Sample
17.	Quality Objective Achievement Plan	18.	Test Request Slip for in process / Finish product Sample
19.	Corrective Action Report	20.	Research And Development Request Report
21.	List of Legal compliance	22.	Research And Development Plan
23.	Communication Report	24.	Design Verification Report
25.	Meeting of Management (MOM)	26.	Design and Development Monitoring Register
27.	Ecosystem impact analysis	28.	Design Review Report
29.	Bio diversity impact register	30.	Experiment Data Sheet
31.	Department Wise Quality Audit Checklist	32.	MoM for Design Review
33.	Quality Culture & Ethical Behaviour Awareness Training Record	34.	Equipment History Card
35.	Ethical Behaviour & Quality Commitment Declaration Form	36.	Equipment Annual Preventive maintenance Schedule
37.	Training Calendar	38.	Equipment Maintenance Checklist
39.	Employee Competence Record Sheet	40.	Machine Breakdown Record
41.	New Employee Induction Training Report	42.	Packing Slip
43.	Job Responsibility & Authority Sheet	44.	Drum / Bag / Carton Inspection Report
45.	Training Attendance Report	46.	Gate Pass
47.	Skill Matrix Sheet	48.	Material Issue Slip
49.	Purchase Order	50.	Preservation Assessment Report
51.	Material Inward Record	52.	Goods Receipt note
53.	List of Approved Supplier / Service Provider	54.	Vehicle inspection checklist – Loading
55.	New Supplier / Vendor Registration Form	56.	Performance Appraisal Records–Functional Heads
57.	Annual Purchase Contract Order	58.	Performance Appraisal Records–Staff

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59.	Vendor Evaluation Form	60.	Manpower Requirement form
61.	Order Form / Order Confirmation	62.	Machine Installation & Commissioning Check sheet
63.	Customer Complaint Report	64.	Service Call Report
65.	Customer Feed Back Form	66.	Machine Installation & Commissioning Report
67.	Customer Property Monitoring Register	68.	Repairing Card
69.	Non-Conforming Product Record Sheet		

7. Filled Formats (42 sample filled formats)

This directory includes sample copy of filled forms to maintain/retain records as well as establish control and create system in the organization. The given samples are just a guide to follow and the organization should modify the same according to follow-up of related processes. Filled formats can be used as templates. A total of 40 formats is provided as per the list given below.

List of filled formats

1.	Master List of Documents	22.	Customer Property Monitoring Register
2.	Change Control Note	23.	Non-Conforming Product Record Sheet
3.	Master List of Records	24.	Spray Dryer Log Sheet
4.	Quality Objective Monitoring Report	25.	Reverse Osmosis Log Sheet
5.	ISO 9001 IQA Plan	26.	Process Change Form
6.	QMS Internal Audit NCR Form	27.	pH Meter Calibration Report
7.	Quality Objective Achievement Plan	28.	Test Request Slip for in process / Finish product Sample
8.	Corrective Action Report	29.	Research And Development Plan
9.	Communication Report	30.	Design and Development Monitoring Register
10.	Training Calendar	31.	Design Review Report
11.	Employee Competence Record Sheet	32.	MoM for Design Review
12.	New Employee Induction Training Report	33.	Equipment Maintenance Checklist
13.	Job Responsibility & Authority Sheet	34.	Machine Breakdown Record
14.	Training Attendance Report	35.	Packing Slip
15.	Skill Matrix Sheet	36.	Gate Pass

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16.	Material Inward Record	37.	Preservation Assessment Report
17.	List of Approved Supplier / Service Provider	38.	Performance Appraisal Records–Functional Heads
18.	New Supplier / Vendor Registration Form	39.	Manpower Requirement form
19.	Order Form / Order Confirmation	40.	Machine Installation & Commissioning Check sheet
20.	Customer Complaint Report	41.	Service Call Report
21.	Repairing Card	42.	Machine Installation & Commissioning Report

8. ISO 9001:2026 Audit Checklists (02 Files in MS-Word (Total 400+ Questions))

This covers audit questions based on the ISO 9001:2026 requirements. It will be a very good tool for the auditors to make their own audit questionnaire for auditing a quality management system. It will bring effectiveness in auditing. A total of more than 400 questions are prepared on the basis of ISO 9001:2026 requirements and given according to the sequence of the requirements clauses of the standard. Department-wise audit questionnaire is also provided in this kit. The checklists can be logically used for auditing during internal audit for ISO 9001:2026 to establish proper audit trail.

9. ISO 9001:2026 Compliance Matrix (01File in Excel)

The ISO 9001:2026 requirement wise list of documented information reference of this kit is given in the compliance matrix for easy reference of user to understand how documentation system is made.

10. Sample MRM Documents (02 Files in MS Word)

The ISO 9001:2026 Sample MRM documents and records are given in the document kit for easy reference of users to understand how management review meeting is conducted and record is retained.

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Chapter-2.0 ABOUT COMPANY

Punyam Academy Pvt. Ltd. is a progressive company promoted by a group of qualified engineers and management graduates having rich experience of more than 30 years in ISO and management area consultancy and online training as well. The company serves the global customers through on-site and off-site modes of service delivery system. We offer a full range of consulting services geared towards helping all types of organizations to achieve competitiveness, certifications and compliance to international standards and regulations. So far, we have **more than 4500 clients in more than 101 countries. Our ready-made editable documents kit helps the client in making their documents with ease and helps them comply with the related ISO standard faster.**

1. Our promoters and engineers have experience in providing management training, ISO series consultancy for **more than 4500 companies** globally. We have clients **in more than 101 countries.**
2. We are a highly qualified team of 80 members (M.B.A., Degree Engineers). Our director has a rich professional experience in this field (since 1991).
3. We have 100% success rate in ISO series certifications for our clients from reputed certification bodies. We possess a branded image and are a leading name in the global market.
4. We also suggest continual improvement and cost reduction measures as well as provide highly informative training presentations and other products that give you payback within 2 months against our cost.
5. We have spent more than 60000 man-days (170 man-years) in the preparation of documents and training slides.

We are committed for:

1. Personal involvement and commitment from the day one
2. Optimum charges
3. Professional approach
4. Hard work and updating the knowledge of team members
5. Strengthening clients by system establishment and providing best training materials in any areas of management to make their house in proper manner
6. Establishing strong internal control with the help of system and use of the latest management techniques.

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Chapter-3.0 USER FUNCTION

3.1 Hardware and Software Requirements

A. Hardware

- Our documentation kit can better perform with the help of P4 and higher computers with a minimum of 10 GB hard disk space.
- For better visual impact, you may keep the setting at high color.

B. Software

- Documents are written in MS-Office 2007 and Windows XP programs. You are, therefore, required to have MS-Office 2007 or higher versions with Windows XP.

3.2 Features of Documentation kit

- The kit contains all necessary documents as listed, and complies with the requirements of system standards.
- The documents are written in easy-to-understand English language.
- This kit will save much time in typing and preparing your documents at your own.
- The kit is user-friendly to adopt and easy to learn.
- The contents of this kit are developed under the guidance of experienced experts.
- The kit provides a model of the management system that is simple and free from excessive paperwork.

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Chapter-4.0 BENEFITS OF USING OUR DOCUMENTATION KIT

1. By using these documents, you can save a lot of your precious time while preparing the ISO 9001:2026 Quality Management System documents.
2. The kit takes care of all the sections and sub-sections of ISO 9001:2026 and helps you to establish better Quality Management System.
3. This documentation kit enables you to change the contents and print as many copies as you need. The users can modify the documents as per their industry requirements and create their own ISO 9001:2026 documents for their organization.
4. It will save much cost in document preparation.
5. You will get a better control in your system due to our proven formats.
6. You will also get a better control in your system as our proven documents and templates are developed under the guidance of experts and globally proven consultants. The team has a rich experience of more than 30 years in the ISO series and other management system documentation consultancy.
7. Our products are highly sold across the globe and are used by many multinational companies. They have got total satisfaction as well as experienced value for money.
8. In the preparation of documentation kit, our team has verified and evaluated the entire content at various levels. More than 1000 hours have been spent in the preparation of this documentation kit.
9. The entire kit is prepared by a globally proven team of leading ISO 9001 consultants.

Chapter-5.0 METHOD OF ONLINE DELIVERY

On secured completion of the purchase, we provide a user name and password to download the product from our FTP server. Hence, we provide an instant online delivery of our products to the user by sending an email of user name and password.

For purchase Click Here ➡ 

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