

Master List of Key QMS Documents

Objective:

Objective of this document is to provide references to second level documents including documented procedures as required by the standard and other reference documents defined by the company. This document also provides references to “Forms” to be used to prepare records with respect to various clauses of ISO 9001:2000

Clause No.	Second Level Documents		Forms for Records	
	Document No.	Document Title	Form No.	Form Title
4.1	DFC 4.1	Description of Interaction of QMS processes		
4.2.3	P-CD	Procedure for Control of Documents	M-01	Document Approval and Revision Control register
	R-4.2.3	Control of External Origin Documents	M-02	Document Amendment Note
			M-03	Document Issue record
4.2.4	P-CR	Procedure for Control of records	---	Permission for records destruction
5.4.1	R-5.4.1	Quality Objectives		
5.5	R-5.5 A	Organization Chart		
	R-5.5 B	Responsibility and authority		
5.6	DFC 5.6	Management Review	M-04	Minutes and Action Plan: Management

Clause No.	Second Level Documents		Forms for Records	
	Document No.	Document Title	Form No.	Form Title
	R-5.6	Yearly Plan for Management Review and internal Quality Audit		Review
6.2	DFC 6.2	Human Resource Development	M-11	Employee Card
	R-6.2 A	Competence requirement for Human resources	M-12	Training Attendance Sheet
	R-6.2 B	Skill Matrix	R-6.2 B	Skill Matrix
	R-6.2 C	Questionnaire for Leadership Evaluation		
	R-6.2 D	Questionnaire for Internal Auditor Evaluation		
6.3, 6.4	DFC 6.3	Methodology for Resource Maintenance		
	R-6.3	List of Machinery with their capability	MT-01	Machine History Register
7.1	DFC 7.1	Product Realization Processes		
7.2	DFC 7.2	Review of product related requirements (contract review)	Mkt-01	Contract/ Order Review Form
7.3	SOP 7.3	Methodology for Product Design and Development	M-13	Product Design and Development Planning

Clause No.	Second Level Documents		Forms for Records	
	Document No.	Document Title	Form No.	Form Title
7.4.1	SOP 7.4.1	Methodology for Evaluation and Re-evaluation	PR-01	Evaluation and Selection of Supplier
			PR-02	List of Supplier with Grade
			PR-03	Periodic Re-evaluation of Supplier
7.4.2	SOP 7.4.2	Purchasing	PO	Purchase Order
7.5.1	DFC 7.5.1	Production Flow Chart		
	SOP 7.5.1	Control of Production Processes		
7.5.2	Exclusion			
7.5.3	SOP 7.5.3	Methodology for Product identification and Traceability	----	Product ID Tag
7.5.4	SOP 7.5.4	Control of Customer Property	----	Letter to Customer for damage to customer property
7.5.5	SOP 7.5.5	Methodology for Preservation of Product		
7.6	SOP 7.6	Methodology for control of monitoring and measurement devices		
	R-7.6	List of Monitoring and Measurement Devices		

Clause No.	Second Level Documents		Forms for Records	
	Document No.	Document Title	Form No.	Form Title
8.2.1	SOP 8.2.1	Methodology Customer Satisfaction Survey	M-05	Customer Feedback Form
8.2.2	P-IQA	Procedure for Internal Quality Audit	M-06	Internal Audit Schedule
	R-5.6	Yearly Plan for Management Review and internal Quality Audit	M-07	Internal Audit Report
			M-08	Action Request (AR)
8.2.3	----	All DFC's and SOP's	M-14	Monitoring of Processes of QMS
	R-5.4.1	Quality Objectives	R-5.4.1	Quality Objectives
8.2.4	PQP-1	Product Quality Plan – Receiving Stage		
	PQP-2	Product Quality Plan – In-process Stage		
	PQP-3	Product Quality Plan – Final Stage		
8.3	P-NCP	Procedure for Control of Non conforming products	M-08	Action Request (AR)
8.4	DFC 8.4	Analysis of Data		
8.5.2, 8.5.3	P-CPA	Procedure for Corrective and	M-08	Action Request (AR)

Clause No.	Second Level Documents		Forms for Records	
	Document No.	Document Title	Form No.	Form Title
		Preventive Actions	M-09	Supplier Corrective Action Request (SCAR)
			M-10	Corrective - Preventive Action (CPA)