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Abstract:

July Shed This document describes the procedure for control of records.

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TABLE OF CONTENTS

1.0	PURPOSE			ارمير
2.0	THEORY			10
3.0	RULES FOR CONTROL OF	RECORDS	O	•••••
Appe	endix A: Records Matrix			
			160	
			250	
			10	
			×5	
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1.0 PURPOSE

This procedure defines the requirements for the control of records within the quality management system (QMS). The scope of this procedure is to control only the records referenced in this document; other records are not controlled.

2.0 THEORY

A record is any written or electronic piece of evidence that may be needed later to provide evidence of conformity to requirements. Typically a blank "form" becomes a "record" when it is completed. Records must be controlled so that the information on them is accessible, legible and suitably maintained.

3.0 RULES FOR CONTROL OF RECORDS

- 3.1 The controls for each type of record are defined in Appendix A of this procedure.
- 3.2 The listed "controller" must ensure their assigned records
- 3.3 Records for active contracts are maintained in the quality department handling the operations. Records are
- 3.4 The Document Control Center maintains archive files for records. Records shall be maintained a minimum of
- 3.5 Records that are discarded after retention shall be
- 3.6 Hardcopy records are to be stored in suitable cabinets that
- 3.7 Records are available for review by the Customer and copies of non-proprietary records are furnished to the Customer upon request. Non-disclosure agreements are required for non-Governmental entities.
- 3.8 Records are verified for
- 3.9 The Company does not require vendors to maintain records for the Company; instead,
- 3.10 To ensure protection of records, electronic records are
- 3.11 Local computer data that is stored on company computers must
- 3.12 When making corrections to written record entries, the error is
- 3.13 Correction fluid or correction tape is not to be used on any quality records.



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Appendix A: Records Matrix

Required Record or Document Type	Company Record	Controller	Туре	Location	Minimum Retention
Calibration records	Calibration		Form		
Contract review					N
records	Contract review		Form		
Control of				eive	
Nonconforming Product	RFS		Form	7	
Corrective and	KIB		TOIM	60	
preventive actions	RFS		Form	0	
			Ca		
Design change records	Engineering order		Form		
Design input			. (1)		
records	Engineering order		Form		
Design review records	Engineering order		Form		
Design validation	Production	~ *			
records	inspection	<u> </u>	Form		
Design verification	Production		_		
records	inspection	V	Form		
First Article Inspection	First article	•	Form		
Internal audit records	Internal audit		Form		
Lost, damaged or unsuitable Customer	Ciar				
property	Customer property		Form		
Management review meeting minutes	Management review report		Form		
Record of realization	•				
process	Engineering order		Form		
Record of release of	Production				
product	inspection		Form		
Supplier evaluation	Supplier review		Form		
	Production		Б		
Traceability records	inspection		Form		
Training records	Training record		Form		

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