

ISO 17025:2017 LMS - Manual-Procedures-Forms-Matrix -P-xxx Numbers

QMS Clause	Level 1 - Topic	Level 2 Procedure	Procedure Name	Level 3 - Work Instruction	Work Instruction Name	Level 4 Form #	Form Name	Flow Diagram	Flow Diagram Name	Attachment #	Attachment Name
All	Laboratory Management System Manual - LMS-001										
4 & 5	General and Structural requirements	P-500	Management responsibility			F-510-001 F-570-001	LMS-Process identification worksheet Comment and suggestion report			A-500-001 A-560-001	Quality policy Organization chart
6	Resource requirements	P-610	Resource management			F-610-001 F-610-002 F-610-003	Equipment problem report Resource maintenance record Environmental control log				
6.2	Personnel	P-620	Competence, awareness and training			F-620-001 F-620-002 F-620-003 F-620-004	Training action plan Group training record Job description Employee training summary				
6.4	Equipment	P-645	Control of monitoring and measuring equipment			F-645-001	Equipment calibration list				
6.6	Externally provided products and services	P-660	Control of external providers			F-660-001 F-660-002 F-660-003 F-660-004 F-660-005 F-660-006	Provider assessment report List of acceptable sources Provider corrective action request - PCAR Purchase requisition Purchase order Business agreement - contract				
7.1	Review of requests, tenders, and contracts	P-710	Customer related processes			F-710-001 F-710-002	Client assessment report Order notification				
7.2	Selection, verification and validation of methods	P-720	Operational planning of methods			F-720-001 F-720-002	Project planning worksheet Method routing summary				
7.3	Sampling										
7.4	Handling of test or calibration items	P-740	Handling of test and calibration items			F-740-001 F-740-002 F-740-003 F-740-004	Storage inspection report External property control log Identification tag / label Laboratory activity log				
7.5	Technical reports	P-755	LMS-Monitoring, analysis, and evaluation			F-755-001	Monitoring report				
7.6	Evaluation of measurement uncertainty										
7.7	ensuring the validity of results										
7.8	Reporting of results	P-780	Reporting of results			F-780-001	Statement of delivery / invoice			A-780-001 A-780-002 A-780-003	Test report - blank Calibration report - blank Report of sampling - blank
7.9	Complaint	P-790	Complaints and nonconforming outputs			F-790-001 F-790-002	Complaint response report Nonconformance report				
7.1	Nonconforming work										
8.2	Management system documentation	P-820	Control of documented information	WI-820-001	Document numbering system	F-820-001 F-820-002 F-820-003 F-820-004 F-820-005 F-820-006 F-820-007 F-820-008	List of documented information Records matrix Master documentation lists Quality records table Document change request form Document revision checklist Software inventory spreadsheet Revision status form				
8.3	Control of management system documentation										
8.4	Control of records										
8.5	Actions to address risks and opportunities	P-850	LMS-Risk management planning			F-850-001 F-850-002	LMS-Risk management worksheet LMS-Objectives planning record				
8.6	Improvement	P-860	Improvement			F-860-001 F-860-002	Data analysis worksheet Customer survey and analysis				
8.7	Corrective actions	P-870	Nonconformity and corrective action			F-870-001	Corrective action request - CAR				
8.8	Internal audits	P-880	Internal audit			F-880-001 F-880-002 F-880-003 F-880-004	Applicable procedure by work area Internal audit checklist Audit plan Audit report				
8.9	Management reviews	P-890	Management review			F-890-001 F-890-002	Management review agenda Management review output report				