

A18-2017 – CROSS REFERENCE TO LABORATORY MANAGEMENT SYSTEM

Revision 2.0

March 16, 2018

Laboratory Name: _____

Laboratory Number: _____



CALA
Trust, measured accurately

Rating Guide Item (A02)	Cross reference
4 General Requirements	
4.1 Impartiality	
<ul style="list-style-type: none"> • Structured and managed 	
<ul style="list-style-type: none"> • Management commitment 	
<ul style="list-style-type: none"> • Impartiality of lab activities 	
<ul style="list-style-type: none"> • Identification of risks 	
<ul style="list-style-type: none"> • Elimination or minimizing risk 	
4.2 Confidentiality	
<ul style="list-style-type: none"> • Management of information 	
<ul style="list-style-type: none"> • Release of confidential information 	
<ul style="list-style-type: none"> • Information from third parties 	
<ul style="list-style-type: none"> • Personnel external to the laboratory 	
5 Structural Requirements	
5.1 Legal Entity	
5.2 Management with overall responsibility	
5.3 Range of Activities	
5.4 Meeting requirements	
5.5 Organization and management structure	

Rating Guide Item (A02)	Cross reference
5 Structural Requirements	
5.6 Responsibility for management system	
5.7 Communication from lab management on:	
<ul style="list-style-type: none"> • Effectiveness of management system 	
<ul style="list-style-type: none"> • Planned changes. 	
6 Resource Requirements	
6.1 General	
6.2 Personnel	
<ul style="list-style-type: none"> • Requirements for internal or external staff 	
<ul style="list-style-type: none"> • Document competence requirements 	
<ul style="list-style-type: none"> • Competence 	
<ul style="list-style-type: none"> • Communication of duties & responsibilities 	
<ul style="list-style-type: none"> • Procedures and records 	
<ul style="list-style-type: none"> • Authorizations 	
6.3 Facilities and environmental conditions	
<ul style="list-style-type: none"> • Suitability 	
<ul style="list-style-type: none"> • Documentation of requirements 	
6 Resource Requirements	

Rating Guide Item (A02)	Cross reference
<ul style="list-style-type: none"> • Monitor, control and record conditions 	
<ul style="list-style-type: none"> • Measures to control facilities 	
6.4 Equipment	
<ul style="list-style-type: none"> • Access 	
<ul style="list-style-type: none"> • Outside the lab's permanent control 	
<ul style="list-style-type: none"> • Procedure(s) for equipment 	
<ul style="list-style-type: none"> • Conformance to requirements 	
<ul style="list-style-type: none"> • Capability to produce valid results 	
<ul style="list-style-type: none"> • Equipment is calibrated 	
<ul style="list-style-type: none"> • Calibration programme 	
<ul style="list-style-type: none"> • Labelling of Equipment 	
<ul style="list-style-type: none"> • Subject to mishandling or overloading 	
<ul style="list-style-type: none"> • Intermediate checks 	
<ul style="list-style-type: none"> • Reference values or Correction factors 	
<ul style="list-style-type: none"> • Unintended adjustments 	
<ul style="list-style-type: none"> • Records 	

Rating Guide Item (A02)	Cross reference
6 Resource Requirements	
6.5 Metrological traceability	
<ul style="list-style-type: none"> • 	
<ul style="list-style-type: none"> • Established and maintained 	
<ul style="list-style-type: none"> • Traceability to the SI 	
<ul style="list-style-type: none"> • When traceability to the SI is not possible 	
6.6 Externally provided products and services	
<ul style="list-style-type: none"> • Use of suitable products and services 	
<ul style="list-style-type: none"> • Procedures and records 	
<ul style="list-style-type: none"> • Communication of requirements 	
7 Process Requirements	
7.1 Review of requests, tenders and contracts	
<ul style="list-style-type: none"> • Procedure 	
<ul style="list-style-type: none"> • Inappropriate method requested 	
<ul style="list-style-type: none"> • Statement of conformity 	
<ul style="list-style-type: none"> • Differences between request and contract 	
<ul style="list-style-type: none"> • Customer informed of deviations 	
<ul style="list-style-type: none"> • Amendments 	
7 Process Requirements	

Rating Guide Item (A02)	Cross reference
<ul style="list-style-type: none"> Clarification of requests 	
<ul style="list-style-type: none"> Records of review 	
7.2 Selection, verification and validation of methods	
<ul style="list-style-type: none"> Appropriate methods and procedures 	
<ul style="list-style-type: none"> Current and available 	
<ul style="list-style-type: none"> Latest valid version in use 	
<ul style="list-style-type: none"> Lab selection of methods 	
<ul style="list-style-type: none"> Verification of standard methods 	
<ul style="list-style-type: none"> Method development 	
<ul style="list-style-type: none"> Deviations from methods 	
<ul style="list-style-type: none"> Validation of methods 	
<ul style="list-style-type: none"> Changes to validated methods 	
<ul style="list-style-type: none"> Fit-for-purpose 	
<ul style="list-style-type: none"> Records of validation 	
7.3 Sampling	
<ul style="list-style-type: none"> Sampling plan and method 	
<ul style="list-style-type: none"> Sampling method content 	
<ul style="list-style-type: none"> Records 	
7 Process Requirements	
7.4 Handling of test or calibration items	

Rating Guide Item (A02)	Cross reference
<ul style="list-style-type: none"> • Procedure 	
<ul style="list-style-type: none"> • 	
<ul style="list-style-type: none"> • System for identification 	
<ul style="list-style-type: none"> • Deviations from conditions 	
<ul style="list-style-type: none"> • Storage conditions 	
7.5 Technical Records	
<ul style="list-style-type: none"> • Sufficient information 	
<ul style="list-style-type: none"> • Amendments to technical records 	
7.6 Evaluation of measurement uncertainty	
<ul style="list-style-type: none"> • Identification of contributions 	
<ul style="list-style-type: none"> • Evaluation of MU for all calibrations 	
<ul style="list-style-type: none"> • Evaluation of MU for quantitative results 	
<ul style="list-style-type: none"> • Estimation of MU for other types of tests 	
7.7 Ensuring the validity of results	
<ul style="list-style-type: none"> • Procedure for monitoring 	
<ul style="list-style-type: none"> • Data recorded to detect trends 	
7 Process Requirements	
<ul style="list-style-type: none"> • PT or inter-laboratory comparisons 	
<ul style="list-style-type: none"> • Analysis of data to control and improve 	
<ul style="list-style-type: none"> • Data outside pre-defined criteria 	

Rating Guide Item (A02)	Cross reference
7.8 Reporting of results	
<ul style="list-style-type: none"> Results reviewed and authorized 	
<ul style="list-style-type: none"> Results reported clearly, unambiguously 	
<ul style="list-style-type: none"> Simplified reports 	
<ul style="list-style-type: none"> Common requirements for reports 	
<ul style="list-style-type: none"> Specific requirements for testing 	
<ul style="list-style-type: none"> Calibration certificates 	
<ul style="list-style-type: none"> Reporting sampling 	
<ul style="list-style-type: none"> Statements of conformity 	
<ul style="list-style-type: none"> Opinions and interpretations 	
<ul style="list-style-type: none"> Amendments to reports 	
7.9 Complaints	
<ul style="list-style-type: none"> Documented process 	
<ul style="list-style-type: none"> Availability of process 	
7 Process Requirements	
<ul style="list-style-type: none"> Content of process 	
<ul style="list-style-type: none"> Responsibility 	
<ul style="list-style-type: none"> Acknowledgement of complaint 	
<ul style="list-style-type: none"> Impartiality of outcomes 	
<ul style="list-style-type: none"> Formal notice of end of process 	

Rating Guide Item (A02)	Cross reference
7.10 Nonconforming work	
<ul style="list-style-type: none"> Responsibilities and authorities 	
<ul style="list-style-type: none"> Actions based on risk levels 	
<ul style="list-style-type: none"> Evaluation of significance 	
<ul style="list-style-type: none"> Decision on acceptability 	
<ul style="list-style-type: none"> Customer is notified or work recalled 	
<ul style="list-style-type: none"> Responsibility for resumption of work 	
<ul style="list-style-type: none"> Record retention 	
<ul style="list-style-type: none"> Need for corrective action 	
7.11 Control of data and information management	
<ul style="list-style-type: none"> Access to data and information 	
<ul style="list-style-type: none"> Validation of LIMS 	
7 Process Requirements	
<ul style="list-style-type: none"> Protected from unauthorized access 	
<ul style="list-style-type: none"> Safeguarded 	
<ul style="list-style-type: none"> Complies with specifications 	
<ul style="list-style-type: none"> Maintenance 	
<ul style="list-style-type: none"> System failures 	
<ul style="list-style-type: none"> 	

Rating Guide Item (A02)	Cross reference
<ul style="list-style-type: none"> • Off-site providers or operators 	
<ul style="list-style-type: none"> • Availability of manuals etc... 	
<ul style="list-style-type: none"> • Calculations and data transfers 	

Rating Guide Item (A02)	Cross reference
8.0 Management System Requirements	
<ul style="list-style-type: none"> Option A 	
<ul style="list-style-type: none"> Option B 	
8.2 Management System Documentation	
<ul style="list-style-type: none"> Policies and objectives 	
<ul style="list-style-type: none"> Management commitment 	
<ul style="list-style-type: none"> Supporting procedures and processes 	
<ul style="list-style-type: none"> Access by personnel 	
8.3 Control of management system documents	
<ul style="list-style-type: none"> Control of documents 	
<ul style="list-style-type: none"> Approval of documents 	
<ul style="list-style-type: none"> Periodic review 	
<ul style="list-style-type: none"> Changes identified 	
<ul style="list-style-type: none"> Distribution 	
<ul style="list-style-type: none"> Uniquely identification 	
<ul style="list-style-type: none"> Obsolete documents 	
8.4 Control of records	
<ul style="list-style-type: none"> Requirement to retain records 	
<ul style="list-style-type: none"> Record integrity 	

Rating Guide Item (A02)	Cross reference
8.0 Management System Requirements	
8.5 Risk	
<ul style="list-style-type: none"> • Consideration of risks and opportunities 	
<ul style="list-style-type: none"> • Actions to address risks and opportunities 	
<ul style="list-style-type: none"> • Actions proportional to impact 	
8.6 Improvement	
<ul style="list-style-type: none"> • Identify and Select 	
<ul style="list-style-type: none"> • Feedback 	
8.7 Corrective actions	
<ul style="list-style-type: none"> • Correction 	
<ul style="list-style-type: none"> • Evaluation of need for corrective action 	
<ul style="list-style-type: none"> • Implement corrective action 	
<ul style="list-style-type: none"> • Effectiveness 	
<ul style="list-style-type: none"> • Update risks and opportunities 	
<ul style="list-style-type: none"> • Changes to system, if necessary 	
<ul style="list-style-type: none"> • Appropriateness 	
<ul style="list-style-type: none"> • Record retention 	
8.8 Internal audits	
<ul style="list-style-type: none"> • Objectives and frequency 	

Rating Guide Item (A02)	Cross reference
8.0 Management System Requirements	
<ul style="list-style-type: none"> Audit criteria defined 	
<ul style="list-style-type: none"> Reporting of outcomes 	
<ul style="list-style-type: none"> Follow-up 	
<ul style="list-style-type: none"> Record retention 	
8.9 Management reviews	
<ul style="list-style-type: none"> Objectives and frequency 	
<ul style="list-style-type: none"> Inputs 	
<ul style="list-style-type: none"> Outputs 	